

Page Change History
MBFP PSD Permit Application Dated December 31, 2007

Page Numbers	Revision Date	Action	Description
	4/23/08	Superseded	Updated Table of Contents, Acronyms
(1-1) 1-2	5/12/08	Superseded	Revised VOC total following revision to equipment leak emission calculation. Also noted Methanol emissions greater than major HAP threshold, rather than total facility HAPs as trigger for NESHAP applicability.
1-7 (1-8)	5/12/08	Superseded	Revised HAP emissions following revision to equipment leak emission calculation.
1-1	4/23/08	Superseded	Revised Saddleback Hills Mine coal production rate from 3.2 MMtpy to 3.25 MMtpy
1-2	4/23/08	Superseded	Updated emissions in Table 1.1 for PM10
(1-1) 1-2	2/12/08	Superseded	Updated emissions in Table 1.1
1-7 (1-8)	2/12/08	Superseded	Updated emissions in Table 1.2
2-1 to 2-2; 2-5 (2-6)	4/23/08	Superseded	Added sentence in Section 2.1, 2 nd paragraph, to explain conveyors C6-C10 will be ¾-covered, rather than fully enclosed. Resulting text carryover to page 2-5. (Note, Figures 2.1 and 2.2 are pages 2-3 and 2-4, with no changes.)
2-9 (2-10)	2/12/08	Superseded	Added sentence (bottom of page) about heating CO2 vent stream
(3-3) 3-4	7/21/08	Superseded	Updated Table 3.3 to match facility total emission rates listed in Table 1.1. This was inadvertently missed in earlier revisions.
(3-3) 3-4 to 3-6	5/12/08	Superseded	Revised VOC emissions in Tables 3.3 and 3.5 to reflect updated equipment leak values and total facility VOC emission rates and HAP emissions in Table 3.4 following revision to equipment leak calculations.
3-9 (3-10)	5/12/08	Superseded	Revised equipment leak definitions in Section 3.2.6.3 from 10,000 ppmv to 500 ppmv for valves and connectors and 2,000 ppmv for pumps following revised equipment leak emission calculations.
3-1 to 3.4; (3-5) 3-6	4/23/08	Superseded	Revised SBH Mine Section 3.1 to clarify that some conveyors will be ¾ covered, rather than fully enclosed; Revised Tables 3.1 through 3.5 by adding revised SBH Mine development and ongoing East Portal coal storage & conveying emission rates.

MEDICINE BOW
EXHIBIT MM
PRE-HRG MEMO

Page Numbers	Revision Date	Action	Description
3-3 to 3-10	2/12/08	Superseded	Revised emissions and emission-related descriptions to address operating hour and fuel simplifications requested by WDEQ *
4-3	7/31/08	Superseded	Correction to the total number of work hours per year for the black start generators. 250 hours were changed to 360 hours per generator per year.
4-30	7/31/08	Superseded	Correction to the total number of work hours per year for the black start generators. 250 hours were changed to 360 hours per generator per year.
4-1	7/21/08	Superseded	Updated Table 4.1 to match facility total emission rates listed in Table 1.1. This was inadvertently missed in earlier revisions.
4-31 (4-32)	5/29/03	Superseded	Correction to mercury emission rate, to reflect flow through two mercury guard beds instead of one.
4-27 (4-28)	5/12/08	Superseded	Revised equipment leak definitions in Section 4.7 from 10,000 ppmv to 500 ppmv for valves and connectors and 2,000 ppmv for pumps following revised equipment leak emission calculations. Also corrected the VOC and HAP emission rates from equipment leaks in this section.
4-29 (4-30)	4/23/08	Superseded	Clarification to first paragraph under Section 4.10, to state that the <u>expected</u> operating hours for the gasifier preheaters will be 500 hours per year, per preheater. Previously, this sentence stated the <u>maximum</u> would be 500 hours per year, per preheater, because PTE emission rates are based on this value. However, 500 hours per year per preheater is only an estimate of annual operating hours for the gasifier preheaters.
4-7 (4-8)	1/18/08	Superseded	Revised \$/ton NOx removed based on revised emissions. (Last two sentences of 1 st paragraph)
5-8	7/31/2008	Superseded	Correction to the total number of work hours per year for the black start generators. 250 hours were changed to 360 hours per generator per year.
5-3 to 5-10	2/12/08	Superseded	Added discussions of: <ul style="list-style-type: none"> --New 40 CFR Part 60, Subpart JJJJ regulations --Wyoming Chapter 6, Section 5 permitting requirements Revised discussion of Subpart DDDDD NESHAP

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General Note for Section 6, regarding equipment leak emission rates			Revisions were made in other sections of the application on 5/12/08 to reflect changes to equipment leak VOC and HAP emission rate calculations. However, these changes were not made in Section 6 because the HAP Modeling, discussed in Section 6.7, was not revised by the permittee. Rather, the WDEQ performed the revised HAP modeling and risk assessment using the revised equipment leak emission rates. Discussion of the revised modeling should be in the WDEQ technical analysis.
6-7 (6-8)	4/30/08	Superseded	Revised Table 6.4 to remove Source ID 'CoalStor,' and to provide clarifying footnotes based on conversations with JNall (4-30-08).
6-3	4/23/08	Superseded	Revised Table 6.1 for modeled PM ₁₀ emission rates
6-4	4/23/08	Superseded	Revised Table 6.2 for LP Flare model parameters and added table footnote.
6-5 to 6-6	4/23/08	Superseded	Carry-over text from page 6-3, due to edits on that page. Added footnote at bottom of Table 6.2. Deleted reference to year 2010 in Section 6.2.2.1, third paragraph.
6-7	4/23/08	Superseded	Revised Table 6.4 for coal mine area source modeling parameters and emission rates and added footnotes
6-8	4/23/08	Superseded	Added road haul volume sources to Table 6.5 and footnote.
6-9 (6-10)	4/23/08	Superseded	Replaced Figure 6.3 with updated version, showing road haul sources associated with the EMM and SBH Mine
6-19 to 6-22	4/23/08	Superseded	Updated Tables 6.10, 6.11, and Figures 6-7, 6-8 for revised 3-hr and 24-hr SO ₂ modeling results
(6-23) 6-24 to 6-26	4/23/08	Superseded	Updated Tables 6.12, 6.13, and Figures 6-10, 6-11 for revised PM ₁₀ modeling results
6-1 to 6-48	2/12/08	Superseded	Revised chapter to reflect new AERMOD near field modeling results and incorporated relevant portions from Appendix J
6-19 to 6-30	3/3/08	Superseded	Revised near-field modeling criteria pollutant results based on revised modeling for years 2000 and 2003
6-33 to 6-36	3/3/08	Superseded	Revised near-field modeling HAP results based on revised modeling for years 2000 and 2003

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7-1 (7-2)	1/18/08	Superseded	Removed first and last sentence of first paragraph after Note. Text removed was: <i>MBFP is proposing to construct a 13,000 barrel per day (BPD) Industrial Gasification & Liquefaction Plant near Medicine Bow, Wyoming.</i> <i>The proposed project is scheduled to start construction in the spring of 2008 with the construction being complete by December 2010.</i>
Appendix B	5/29/08	Superseded	Edits to pages B-1, B-2, and B-3 through B-11 to correct mercury emission rates.
Appendix B	4/23/08	Superseded, Addition	Replace pages B-1 and B-2 to reflect updated coal storage & processing emission rates Replace page B-29 (SBH Mine, coal storage emission calculations) with renumbered page B-29(1) and additional pages for coal mining emission calculations (pages B-29(2) through B-29(16)). Page B-30 reprinted, due to pagination detail.
Appendix B	2/12/08	Superseded	Emission revisions requested by WDEQ * and page numbering changes
Appendix F	1/4/08	Superseded	Updated coal storage BACT analysis
Appendix H	1/18/08	Addition	Added Incremental NO _x Removal Cost as Appendix H
Appendix I	2/12/08	Superseded	Revised to discuss far field modeling only (since near field modeling has been re-run)
Appendix J	2/12/08	Superseded	Moved and revised near field modeling discussions to Chapter 6; far field modeling description remains
Appendix N	1/18/08	Added	Added tabbed divider
Appendix O	2/13/08	Deleted	Delete Appendix O pages (see revised Appendix H)

* During a meeting on January 18, 2008, WDEQ requested emission changes to minimize recordkeeping and reporting requirements and simplify permit writing. For certain equipment, MBFP agreed to increase operating hours and base emission calculations on the highest-emitting fuel (natural gas) in order to streamline compliance. Consequently, potential emissions were increased. Notes reflecting actual equipment operations have been added to pertinent spreadsheets. WDEQ stated that BACT analyses would not be affected by these simplifying assumptions, and would instead be based on the actual operations of the equipment.

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**PREVENTION OF SIGNIFICANT
DETERIORATION PERMIT APPLICATION**

**Medicine Bow Fuel & Power LLC
Industrial Gasification & Liquefaction
(IGL) Plant**

Carbon County, Wyoming

December 31, 2007 Amended Permit Application

TABLE OF CONTENTS

Section 1	Introduction.....	1-1
1.1	General Facility Description.....	1-1
1.2	Facility Location.....	1-2
1.3	Prevention of Significant Deterioration Applicability.....	1-2
1.4	Standard Industrial Classification.....	1-7
Section 2	Process Description.....	2-1
2.1	Coal Mining.....	2-1
2.2	Gasoline Production.....	2-1
2.2.1	Coal Preparation (1100).....	2-1
2.2.2	Gasification (1200).....	2-2
2.2.3	Syngas Conditioning (1300).....	2-2
2.2.4	SELEXOL [®] Acid Gas Removal (2100).....	2-6
2.2.5	Methanol Synthesis (4500).....	2-7
2.2.6	Methanol to Gasoline (5500).....	2-7
2.3	CO ₂ Recovery (2200) and Production.....	2-9
2.4	Sulfur Recovery (3100) and Production.....	2-10
2.5	Ancillary Operations.....	2-11
2.5.1	Power Generation (7100).....	2-11
2.5.2	Air Separation Unit (6100).....	2-11
2.5.3	Intermediate and Product Storage (8100-8200).....	2-11
2.5.4	Slag Handling and Water Cleanup (1200).....	2-12
2.5.5	Water Treatment (1300 and 7100).....	2-12
2.5.6	Flares (8900).....	2-13
2.5.7	Other Utilities (8300).....	2-13
2.6	Startup Activities.....	2-13
Section 3	Emission Estimates.....	3-1
3.1	Saddleback Hills Mine.....	3-1
3.2	The Plant.....	3-2
3.2.1	Emission Sources.....	3-2
3.2.2	Normal Operations.....	3-3
3.2.3	Cold Start/Initial Year Operations.....	3-5
3.2.4	Malfunctions and Other Events.....	3-6
3.2.5	Emissions of PSD-Regulated Pollutants.....	3-7
3.2.6	Source-Specific Calculation Methods.....	3-7
Section 4	Best Available Control Technology.....	4-1
4.1	BACT Review Process.....	4-1
4.2	BACT Summary.....	4-3
4.3	Combustion Turbine Control Technology Review.....	4-4
4.3.1	Nitrogen Oxides BACT Analysis for the Combustion Turbines.....	4-4

TABLE OF CONTENTS

4.3.2	Sulfur Dioxide BACT Analysis for the Combustion Turbines	4-8
4.3.3	Volatile Organic Compound BACT Analysis for the Combustion Turbines.....	4-12
4.3.4	Particulate Emissions BACT Analysis for the Combustion Turbines	4-14
4.3.5	Startup Emissions BACT Analysis for the Combustion Turbines	4-16
4.4	Fired Heater And Boiler Control Technology Review	4-16
4.4.1	NO _x BACT Analysis for the Fired Heaters and Boiler.....	4-16
4.4.2	CO and VOC BACT Analysis for the Fired Heaters and Boiler.....	4-19
4.4.3	SO ₂ BACT Analysis for the Fired Heaters and Boiler	4-21
4.4.4	Particulate Emissions BACT Analysis for the Fired Heaters and Boiler.....	4-22
4.4.5	Startup Emissions BACT Analysis for the Fired Heaters and Boiler.....	4-23
4.5	Storage Tank Control Technology Review.....	4-23
4.6	Material Handling Control Technology Review.....	4-26
4.7	Process Fugitive Emissions Control Technology Review	4-27
4.8	Sulfur Recovery Unit (SRU) Control Technology Review	4-27
4.9	Carbon Dioxide Vent Stack (Startup Operations Only)	4-28
4.10	Gasifier Preheating Control Technology Review (Startup Operations Only).....	4-29
4.11	Black-Start Generator Control Technology Review (Startup Operations Only).....	4-30
4.12	Firewater Pump Control Technology Review (Backup Operations only)	4-31
4.13	Mercury Emission reduction.....	4-31
4.14	Mine Long-Term Coal Storage.....	4-32
Section 5	Regulatory Review	5-1
5.1	Wyoming Air Quality Regulations	5-1
5.1.1	Chapter 2 Ambient Standards	5-1
5.1.2	Chapter 3 General Emission Standards.....	5-2
5.1.3	Chapter 6 Permitting Requirements.....	5-4
5.1.4	Chapter 7 Monitoring Regulations.....	5-5
5.2	Federal Regulations	5-5
5.2.1	New Source Performance Standards (NSPS)	5-6
5.2.2	National Emissions Standards for Hazardous Air Pollutants (NESHAP)	5-8
5.2.3	Chemical Accident Prevention Provisions.....	5-9
Section 6	Near Field Air Quality Impact Analysis	6-1
6.1	Near Field Modeling Background	6-1

TABLE OF CONTENTS

6.2	Description of the Proposed Project	6-1
6.2.1	Site Location	6-1
6.2.2	Source Emissions and Parameters	6-3
6.2.3	Additional Emission Assumptions.....	6-6
6.3	Standards, Criteria Levels, And Basic Methodology.....	6-11
6.4	Near Field Modeling Method.....	6-12
6.4.1	Model Selection and Setup	6-12
6.4.2	Databases for Air Quality Assessment	6-14
6.4.3	Meteorological Data.....	6-14
6.4.4	Receptor Grid.....	6-17
6.5	Growth Analysis	6-18
6.6	Criteria Pollutant Modeling Results.....	6-18
6.6.1	SO ₂ Modeling Demonstration.....	6-18
6.6.2	PM/PM ₁₀ Modeling Demonstration.....	6-24
6.6.3	CO Modeling Demonstration.....	6-27
6.6.4	NO _x Modeling Demonstration	6-30
6.7	HAP Modeling.....	6-32
6.7.1	HAP Emission Sources	6-32
6.7.2	HAP Risk Assessment Procedures.....	6-32
6.7.3	HAP Modeling Results	6-33
6.7.4	HAP Conclusion	6-37
6.8	Impacts to Soil and Vegetation.....	6-38
6.8.1	Soil Impacts	6-41
6.8.2	Vegetation Impacts	6-41
6.9	References for Chapter 6	6-46
Section 7	Far Field Air Quality Impact Analysis	7-1
7.1	Background.....	7-1
7.2	Description of Proposed Project	7-2
7.2.1	Site Location	7-2
7.2.2	Source Emissions	7-2
7.2.3	Sources Included in CALPUFF Modeling.....	7-2
7.2.4	Reference Reports.....	7-7
7.3	Long-Range Transport Modeling Method.....	7-9
7.3.1	Long-Range Transport Modeling	7-9
7.3.2	Model Selection and Setup	7-9
7.3.3	Domain.....	7-10
7.3.4	LULC and TERREL Processing.....	7-11
7.3.5	Hourly Surface and Precipitation Data	7-12
7.3.6	Upper Air Sounding Data	7-12
7.3.7	MM5 Data.....	7-12
7.3.8	CALMET	7-12
7.3.9	PSD Class I Increment Significance Analysis	7-17
7.3.10	Class I Area Visibility Reduction Analysis	7-18
7.3.11	Total Nitrogen and Sulfur Deposition Analyses.....	7-18
7.4	Modeling Results	7-19

TABLE OF CONTENTS

7.4.1	CALPUFF Modeling Results.....	7-19
7.4.2	Soil and Vegetation Analysis.....	7-19
7.5	Conclusion	7-23

List of Tables

Table 1.1	Annual Criteria Pollutant Emissions (tpy)
Table 1.2	Annual HAPs Emissions (tpy)
Table 3.1	Mine Development Particulate Emissions
Table 3.2	Emission Units and Fugitive Sources
Table 3.3	Annual Criteria Pollutant Emissions Resulting from Normal Operations
Table 3.4	Annual HAP Emissions Resulting from Normal Operations
Table 3.5	Annual Criteria Pollutant Emissions Resulting from Cold Startup
Table 3.6	Criteria Pollutant Emissions Resulting from Malfunctions and Other Events
Table 3.7	Typical Fuel Gas Mixture Composition
Table 4.1	BACT Applicability
Table 4.2	Summary of BACT Applied to the MBFP Project
Table 4.3	Storage Tanks Summary
Table 4.4	Gasifier Preheater BACT Analysis Summary
Table 4.5	Black Start Generator BACT Analysis Summary
Table 4.6	Emergency Firewater Pump BACT Analysis Summary
Table 5.1	Ambient Air Quality Standards
Table 5.2	Subpart Kb Tanks List
Table 6.1	Maximum Combined Modeled Short-Term Emission Rates for All Sources in the Analysis
Table 6.2	Modeled Plant Point Source Parameters
Table 6.3	Modeled Cumulative (Nearby) Point Source Parameters
Table 6.4	Area Source Modeling Parameters
Table 6.5	Volume Source Modeling Parameters
Table 6.6	SILs, NAAQS, WAAQS, and PSD Class II Increments
Table 6.7	Site-Specific Elmo Meteorological Data Completeness
Table 6.8	Data Completeness Evaluation, Rawlins and Laramie NWS Hourly Surface Meteorological Data
Table 6.9	Frequency Distribution of Wind Speed and Direction of the Elmo, Rawlins, and Laramie Hourly Surface Meteorological Data (2000, 2001, 2003–2005)

TABLE OF CONTENTS

Table 6.10	Predicted SO ₂ Concentrations Compared to NAAQS / WAAQS
Table 6.11	Predicted SO ₂ Concentrations Compared to PSD Increments
Table 6.12	Predicted PM/PM ₁₀ Concentrations Compared to NAAQS / WAAQS
Table 6.13	Predicted PM/PM ₁₀ Concentrations Compared to PSD Increments
Table 6.14	Predicted CO Concentrations Compared to the NAAQS / WAAQS
Table 6.15	Predicted NO _x Concentrations Compared to the PSD Increment, NAAQS, and WAAQS
Table 6.16	Source HAP Short-Term (Maximum) Emission Rates
Table 6.17	Source HAP Emission Rates
Table 6.18	Annually Averaged Modeled Concentrations
Table 6.19	Cancer Risk Assessment Values
Table 7.1	Class I Areas and Sensitive Class II Areas Within 300 km
Table 7.2	Maximum Emission Rate from All Sources
Table 7.3	24 hour Averaged Emission Inventory for CALPUFF (3-hour and 24-hour SO ₂ , and 24-hour PM ₁₀ and Visibility)
Table 7.4	Annual Averaged Emission Inventory for CALPUFF (Annual NO _x , SO ₂ , and PM ₁₀ and Deposition)
Table 7.5	Size Distribution of Secondary Organic Aerosols (SOA)
Table 7.6	Source Location and Parameter
Table 7.7	CALMET Model Options
Table 7.8	CALPUFF Model Options
Table 7.9	2001 CALPUFF Modeling Results
Table 7-10	2002 CALPUFF Modeling Results
Table 7-11	2003 CALPUFF Modeling Results

List of Figures

Figure 1.1	Site Location Map
Figure 1.2	Mine South Portal Site Layout
Figure 1.3	Mine East Portal / Plant Layout
Figure 1.4	Process Equipment Layout
Figure 2.1	Plant Process Flow Diagram
Figure 2.2	Coal Mine Process Flow Diagram
Figure 6.1	Plant Site Area, View from South Side
Figure 6.2	Plant Site Area, View Over Coal Hills Toward Elk Mountain

TABLE OF CONTENTS

Figure 6.3	Plant and Nearby Mining Area Sources
Figure 6.4	Plant Location Relative to the WDEQ Provided Emissions Inventory
Figure 6.5	GEP Stack Height Assessment Building and Source Location Depiction
Figure 6.6	Wind Rose of AERMOD Input, Five-Year Period
Figure 6.7	2003 Maximum SO ₂ 3-Hour Impacts (PSD)
Figure 6.8	2001 Maximum SO ₂ 24-Hour Impacts (PSD)
Figure 6.9	2003 Maximum SO ₂ Annual Impacts (PSD)
Figure 6.10	2000 Maximum PM ₁₀ 24-Hour Impacts (PSD)
Figure 6.11	2005 Maximum PM ₁₀ Annual Impacts (PSD)
Figure 6.12	2001 Second High CO 1-Hour Impacts (NAAQS)
Figure 6.13	2001 Second High CO 8-Hour Impacts (NAAQS)
Figure 6.14	2001 Maximum Annual NO _x Impacts (NAAQS and PSD)
Figure 6.15	MLE Receptors for Benzene
Figure 6.16	MEI Receptors for Benzene
Figure 6.17	Aerial View of Land Use Immediately Surrounding the Plant
Figure 6.18	Land Use and Vegetation Cover near the Plant Site
Figure 7.1	Relative Location of Modeling Domain, MM5 Domain, Class I and Sensitive Class II areas, and Source based on UTM Coordinates
Figure 7.2	Modeling Domain with Receptors of Class I and Sensitive Class II Areas, Precipitation Data Monitoring Station, Ozone Monitoring Station, Surface Meteorological Data Monitoring Station, and Project Location

List of Appendices

Appendix A	Wyoming Permit Application Form
Appendix B	Emission Calculations
Appendix C	Manufacturer Specifications
Appendix D	Major Equipment List and SCCs
Appendix E	BACT Review of Recent NO _x Limits for Combined Cycle Combustion Turbines Fueled With Other Gaseous Fuels
Appendix F	Coal Storage BACT Analysis
Appendix G	Mercury Removal Cost Analysis
Appendix H	Incremental NO _x Removal Cost for SCR
Appendix I	Analysis of Criteria Pollutant Far Field Modeling Sufficiency
Appendix J	Responses to WDEQ July 17, 2007 Far Field Modeling Comments

TABLE OF CONTENTS

Appendix K	NRCS Irrigated and Nonirrigated Yields by Map Unit for Carbon County, Wyoming
Appendix L	NRCS Acreage and Proportionate Extent of the Soils for Carbon County, Wyoming
Appendix M	NRCS Rangeland Productivity and Plant Composition
Appendix N	Mesoscale Model Simulations in Quasi-Forecast Mode of the Great Western Storm of 16-20 March 2003

Acronyms

agl	Above grade level
AGR	Acid gas removal
AP-42	EPA AP-42 Emission Factors
AQRV	Air Quality Related Value
ASU	Air Separation Unit
AVO	Audio/visual/olfactory
BACT	Best Available Control Technology
BOL	Beginning of Life
BPD	Barrels per day
bpip	Building Profile Input Program
Btu	British thermal unit
CAA	Clean Air Act
CaCO ₃	Calcium carbonate
CAM	Compliance Assurance Monitoring
CDPHE	Colorado Department of Public Health and Environment
CFR	Code of Federal Regulations
Cl ₂	Chlorine
CO	Carbon monoxide
CO ₂	Carbon dioxide
COS	Carbonyl sulfide
CS ₂	Carbon disulfide
DAT	Deposition Analysis Thresholds
DEM	Digital Elevation Model
DLN	Dry Low NO _x
DME	Dimethyl ether
dscf	Dry standard cubic feet
EC	Elemental carbon
EFR	External floating roof
EMM	Elk Mountain Mine
EOL	End of life
EPA	U.S. Environmental Protection Agency
ESP	Electrostatic precipitator
°F	Degrees Fahrenheit
F	Fluorine
FGD	Flue gas desulfurization
FGR	Flue gas recirculation
FLAG	Federal Land Managers Air Quality Related Values Working Group
ft	Feet
g	Gram
gal	Gallons
GE	General Electric Co.
GEP	Good Engineering Practice
GPM	Gallons per minute
H ₂	Hydrogen
H ₂ S	Hydrogen sulfide
HAP	Hazardous air pollutant

Acronyms

HGT	Heavy gasoline treatment
HHV	Higher heating value
HNO ₃	Nitric acid
HP	High pressure
hp	Horsepower
hr	Hour
hr/yr	Hours per year
HRSG	Heat recovery steam generator
IDLH	Immediately Dangerous to Life or Health
IFR	Internal floating roof
IGCC	Integrated gasification combined cycle
IGL	Industrial Gasification and Liquefaction
in	Inch
IWAQM	Interagency Working Group on Air Quality Modeling
km	kilometer
LAC	Level of acceptable extinction change
LAER	Lowest Achievable Emission Rate
lb	Pound
lb/yr	Pounds per year
LDAR	Leak Detection and Repair
LHV	Lower heating value
LP	Low pressure
LPG	Liquefied petroleum gas
LTGC	Low-temperature gas cleanup
LULC	Land Use Land Cover
m	Meter
µg/m ³	Micrograms per cubic meter
m ³	Cubic meters
MACT	Maximum Achievable Control Technology
MDEA	Methyldiethanolamine
MEI	Maximally exposed individual
min	Minute
MLE	Most likely exposure
MMBtu	Million British thermal units
MMscf	Million standard cubic feet
MMscfd	Million standard cubic foot per day
MMtpy	Million tons per year
mol.	Molecular
MP	Medium pressure
MBFP	Medicine Bow Fuel and Power LLC
Mscf	Thousand standard cubic feet
MTBE	Methyl tertiary butyl ether
MTG	Methanol to gasoline
MW	Megawatts
MWh	Megawatt-hours
NAAQS	National Ambient Air Quality Standards

Acronyms

NCDC	National Climate Data Center
neg.	Negligible
NESHAPs	National Emission Standards for Hazardous Air Pollutants
NH ₃	Ammonia
NH ₄ NO ₃	Ammonium nitrate
(NH ₄) ₂ SO ₄	Ammonium sulfate
NIOSH	National Institute for Occupational Safety and Health
NO ₂	Nitrogen dioxide
NO ₃	Nitrate
NO _x	Nitrogen oxides
NRCS	Natural Resources Conservation Service
NSCR	Non-selective catalyst reduction
NSPS	New Source Performance Standard
NSR	New Source Review
NWS	National Weather Service
ODEQ	Oregon Department of Environmental Quality
PBL	Planetary boundary layer
PM	Particulate matter
PM ₁₀	Particulate matter, less than 10 microns
ppmv	Parts per million by volume
ppmw	Parts per million by weight
PSD	Prevention of Significant Deterioration
psi	Pounds per square inch
psig	Pounds per square inch gauge
PTE	Potential to Emit
REL	Reference Exposure Level
RACT	Reasonably Available Control Technology
RBLC	RACT/BACT/LAER Clearinghouse
RfC	Reference Concentrations for Chronic Inhalation
RH	Relative humidity
RICE	Reciprocating internal combustion engine
RMP	Risk Management Plan
RVP	Reid vapor pressure
SBH	Saddleback Hills (Mine)
SCCs	Source Classification Codes
scf	Standard cubic feet
SCFH	Standard cubic foot per hour
scm	Standard cubic meters
SCR	Selective Catalytic Reduction
SIC	Standard Industrial Classification
SILs	Significant Impact Levels
SIP	State Implementation Plan
SNCR	Selective Non-Catalytic Reduction
SO ₂	Sulfur dioxide
SO ₄	Sulfate
SOA	Secondary Organic Aerosol

Acronyms

SOCMI	Synthetic Organic Chemical Manufacturing Industry
SO _x	Sulfur oxides
SRU	Sulfur Recovery Unit
SSM	Startup, shutdown, or malfunction
TANKS	U.S. Environmental Protection Agency Tanks Version 4.0
TBD	To be determined
TPD	Tons per day
tpy	Tons per year
UOP	UOP, LLC
URF	Unit risk factor
USDA	US Department of Agriculture
USGS	U.S. Geological Survey
USNPS	US National Park Service
UTM	Universal Transverse Mercator
VOC	Volatile organic compound
vol%	Volume percent
WAQS&R	Wyoming Air Quality Standards and Regulations
WDEQ	Wyoming Department of Environmental Quality
WRAP	Western Regional Air Partnership
wt%	Weight percent
yr	Year

Acronyms

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1.1 GENERAL FACILITY DESCRIPTION

Medicine Bow Fuel & Power LLC (MBFP) is proposing to construct an underground coal mine (Mine) and industrial gasification & liquefaction (IGL) plant (Plant) that will produce transportation fuels and other products near Medicine Bow, Wyoming in Carbon County. The Mine will process approximately 8,000 tons per day (TPD) of coal (on a dry basis) to produce a variety of liquid and gaseous fuels. The Mine will be a 3.25 million ton per year (MMtpy) adjacent underground coal mine known as the Saddleback Hills Mine that will supply the coal needed for the Plant.

The Plant will utilize coal, which will be gasified to produce synthesis gas (syngas) and produce various products. In order to achieve this outcome, the Plant will use several different technologies, including: General Electric's (GE) gasification technology for the quench gasification process, UOP LLC's (UOP) SELEXOL[®] acid gas removal process, and Davy Process Technology's (Davy) methanol synthesis process followed by the Exxon-Mobil methanol-to-gasoline (MTG) process.

Saleable products produced at the Plant during normal operation are anticipated to include approximately:

- 18,500 barrels per day (BPD) of regular gasoline to be transferred via pipeline to a nearby refinery
- 42 TPD of sulfur
- 198 million standard cubic feet per day (MMscfd) of carbon dioxide (CO₂)
- 712 TPD of coarse slag

In addition to the salable products listed above, Plant operation will result in the production of the following fuels to be used onsite for power generation and process heating:

- Approximately 253 million British thermal units (MMBtu/hr) of fuel gas
- Approximately 400 to 500 MMBtu/hr of liquefied petroleum gas (LPG)

Efficient use of these fuels will provide much of the energy input needed to fuel an electric generation plant that will produce approximately 400 megawatts (MW) of electricity. The Plant will either import natural gas or divert syngas as necessary to support plant power needs not met by fuel gas, LPG, and process steam and is not expected to export power to the electrical grid. Three combustion turbines will be equipped with the best available pollution control technologies, which include low-NO_x burners, diluent injection, selective catalytic reduction (SCR), and oxidation catalyst to keep criteria pollutant emissions low.

Emission reduction technologies will be incorporated throughout the Plant. These controls are discussed in more detail in Sections 2 and 4. In addition, all roads and parking areas within the Plant fence will be either gravel or paved to control fugitive dust emissions.

This amended Prevention of Significant Deterioration (PSD) permit application contains fully updated information based on replacement of the previously planned Fischer-Tropsch and UOP upgrading processes with the Davy methanol synthesis unit and Exxon-Mobil MTG processes. This process change affects many process streams and emission calculations. Consequently, a

complete amended permit application is being submitted. This permit application contains information describing the Mine and Plant, facility emissions, applicable regulations, best available control technology (BACT) determinations, and air quality impact analyses. Wyoming Air Quality Permit Application Forms are included in Appendix A.

1.2 FACILITY LOCATION

The Mine and Plant (collectively, the MBFP Facility) will be located approximately 7.5 miles north of Interstate 80, exit 260 (Elk Mountain) on County Road #3 in Section 29 of Township 21 north and Range 79 west in Carbon County, south-central Wyoming. Figure 1.1 shows the general location of the facility. The MBFP Facility encompasses two separate areas. The Mine’s South Portal is shown in Figure 1.2. The Mine’s East Portal, near where the Plant will be located, is shown in Figure 1.3. Figure 1.4 shows the Plant process equipment layout.

1.3 PREVENTION OF SIGNIFICANT DETERIORATION APPLICABILITY

The Clean Air Act (CAA) defines 28 major source categories that have a 100 ton per year (tpy) threshold for determining prevention of significant deterioration (PSD) major source status. This facility falls within the major source category of “Fuel Conversion Plant,” and therefore is subject to the 100 tpy major source threshold. Annual emissions of criteria pollutant emissions are shown in Table 1.1 for normal operations without startup, shutdown, and malfunction (SSM) events. Estimates of the following pollutants are included: NO_x (nitrogen oxides, including nitrogen dioxide [NO₂]), carbon monoxide (CO), volatile organic compounds (VOC), and particulate matter with a diameter of less than 10 microns (PM₁₀). Emission calculation methods are summarized in Section 3 and detailed emission calculations are included in Appendix B.

Table 1.1 – Annual Criteria Pollutant Emissions (tpy)

NO _x	CO	VOC	SO ₂	PM ₁₀
251.63	176.75	188.49	32.65	195.84

Based on criteria pollutant emissions, this facility is considered to be a major source for the PSD Program (40 CFR §51.165) and the Title V Operating Permit Program (40 CFR Part 70).

Annual emissions of hazardous air pollutant (HAP) emissions from normal operations are shown in Table 1.2. HAPs with emissions greater than 0.01 tpy are included in the table. Because potential emissions of Methanol exceed 10 tpy (although total HAPs are less than 25 tpy), the facility is a major source of HAPs and is subject to some National Emission Standards for Hazardous Air Pollutants (NESHAP) in 40 CFR Parts 61 and 63.

Insert Figure 1.1 - Site Location Map

Insert Figure 1.2 – Mine South Portal Site Layout

Insert Figure 1.3 – Mine East Portal / Plant Layout

Insert Figure 1.4 –Process Equipment Layout

Table 1.2 – Annual HAP Emissions (tpy)

Pollutant	Emissions (tpy)
Acetaldehyde	0.38
Acrolein	0.06
Benzene	8.54
Carbonyl Sulfide	0.23
Ethyl Benzene	0.34
Formaldehyde	0.71
Hexane	10.26
Methanol	12.79
Naphthalene	0.01
PAH	0.02
Propylene Oxide	0.28
Toluene	1.81
Xylene	0.77
Other HAPs*	0.01
Total HAPs	24.71

*Other individual HAPs are less than 0.01 tpy each.

1.4 STANDARD INDUSTRIAL CLASSIFICATION

Two Standard Industrial Classification (SIC) Codes describe the activities associated with the MBFP Facility. These include:

1. 1222 Bituminous Coal Underground Mining
2. 1311 Crude Petroleum and Natural Gas (production of gas and hydrocarbon liquids through gasification)

Because the primary purpose, and source of revenue of the facility is to produce gasoline fuel, the main SIC code will be 1311.

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This section describes the coal mining and industrial production processes. Because coal mining is common in the area, the coal mining description is relatively short. Due to its relative newness and complexity, the Plant is described in much more detail; Figure 2.1 illustrates the process.

2.1 COAL MINING

The Mine will produce approximately 3.25 MMtpy of coal using underground continuous and longwall mining techniques. Longwall mining machines consist of multiple coal shearers mounted on a series of self-advancing hydraulic ceiling supports. Longwall mining machines are about 800 feet in width and 5 to 10 feet tall. Longwall miners extract "panels", rectangular blocks of coal, as wide as the mining machinery and as long as 12,000 feet. The shearers cut coal from a wall face, which falls onto a conveyor belt for removal. As a longwall miner advances along a panel, the roof behind the miner's path is allowed to collapse.

The mined coal will exit the mine via the East Portal. The coal will be conveyed and stored in a 300,000-ton live storage area before being conveyed to the Plant. Coal handling conveyors C1 through C5 will be fully enclosed, and conveyors C6 through C10 will be $\frac{3}{4}$ -covered (not fully enclosed). All transfer points along all conveyors (C1 through C10) will be fogged to reduce emissions. An additional 300,000-ton emergency coal stockpile will be constructed. This emergency coal stockpile is considered dead storage and will not be added to or used unless the coal supply for the live storage is interrupted. Once the emergency stockpile is constructed, it will be compacted and sealed to prevent wind erosion and spontaneous combustion.

Figure 2.2 shows the above-ground coal handling process for stacking the coal and transferring it to the Plant.

2.2 GASOLINE PRODUCTION

Figure 2.1 contains a block flow diagram illustrating the Plant production process and associated support activities. Major processes required to produce gasoline are described in this section. Additional production steps for removing CO₂ and sulfur products are described in Sections 2.3 and 2.4, respectively. Ancillary operations, such as power generation, wastewater treatment, and other activities are described in Section 2.5.

2.2.1 Coal Preparation (1100)

The Plant process begins with coal feed preparation, shown on the left side of the process block flow diagram in Figure 2.2. Raw feed coal (run of mine) from the coal storage area is routed via an enclosed conveyor to the coal crusher. The crushed coal is screened to a maximum size of 1 inch, with oversized coal recycled back to the crusher. All transfer points are fogged to reduce emissions. The crushed and screened coal is conveyed and stored in three bins and is gravity flowed to the coal-grinding mill.

The coal is crushed with water and an additive to create a slurry, which will be pumped into the gasifier under high pressure. The coal preparation process is divided into three separate trains, each with the capacity to supply 40% of the total plant requirements. The slurry produced by any of the trains can be pumped to any of the five (5) downstream gasification trains. The coal

preparation section provides a total of 8,700 tons per day (TPD) of coal to the gasifiers (wet basis); this is equivalent to 8,000 TPD of coal on a dry basis.

Drainage, wash down, and leaks in the grinding area are collected in a below-grade concrete sump. An agitator keeps the solids in suspension for pumping. Any accumulated water/solids mixture is pumped to the slurry tank.

2.2.2 Gasification (1200)

The Plant will utilize five (5) gasifier trains. Each gasifier train will be sized to handle one-fourth of the Plant's total capacity. In normal operation, four gasifier trains will be in operation with the fifth in hot standby. The gasifiers are fueled by a coal/water slurry, calcium carbonate (CaCO_3), and 98 percent pure oxygen from the air separation unit (ASU).

The gasification reaction is conducted at a pressure of 1,000 psig and generates a temperature of approximately 2,500 degrees Fahrenheit ($^{\circ}\text{F}$). The combustion chamber is lined with refractory bricks, which maintain the outer shell of the gasifier in a temperature range of 545 $^{\circ}\text{F}$ to 600 $^{\circ}\text{F}$. Each gasifier is equipped with a dedicated preheater (Gasifier Preheaters 1 through 5). During the initial gasifier startup, and during any subsequent startup following refractory replacement, the gasifier preheater combusts natural gas and slowly heats the refractory to achieve the minimum temperature needed for combustion chamber operation. Each preheater has a firing rate of 21 MMBtu/hr and is fueled with natural gas.

Combustion products of the gasification reaction consist of raw syngas, together with small amounts of a number of impurities (including chlorides, sulfides, nitrogen, argon, and methane), liquid slag, and fine solid particles. These combustion products exit the combustion chamber and flow to a quench chamber where the combustion products are cooled and most of the particle fines are removed from the syngas. The molten slag solidifies and settles to the bottom of the chamber. If necessary, calcium carbonate can be added to the coal slurry as a fluxant to facilitate free flow of the molten slag in the gasifier. Solidified coarse slag is removed from the gasifier through a lock hopper system connected to the bottom of the quench chamber, and this stream sweeps the solidified slag through a slag crusher. The crushed slag is then recycled and reused or disposed. Approximately 980 TPD of slag will be produced and approximately 712 TPD of slag will be available for sale; the remainder is recycled to the slurry because of its Btu content. The syngas exits the gasifier through a side connection.

During any startup, shutdown, or malfunction (SSM) event, the syngas will be sent to the high-pressure flare. The syngas feed to the flare is expected to have a heat rate of approximately 2,000 Btu/lb.

2.2.3 Syngas Conditioning (1300)

Syngas conditioning includes two main treatment processes:

- Scrubbing to remove particulate from the syngas
- Low-temperature gas cleanup (LTGC)

Insert Figure 2.1 - Plant Process Flow Diagram

Insert Figure 2.2 – Coal Mine Process Flow Diagram

2.2.3.1 Syngas Scrubbing

The Plant includes five (5) syngas conditioning trains, each sized for one-fourth of plant capacity. Each syngas conditioning train is integrated with a specific gasifier, with four (4) such trains operating and the fifth acting as a spare during normal operations. This description refers to one syngas conditioning train only.

Raw syngas leaves the gasifier and is mixed with process condensate in the process line to prevent the buildup of solids and thoroughly wet the entrained solids to facilitate their removal in the syngas scrubber.

The syngas scrubber is a tower that contains a water sump in the bottom and four trays in the top. Wet syngas enters the scrubber below the first tray and flows downward into the water sump, which removes most of the solids in the gas, and then flows upward through the four trays. Process condensate is supplied to the top tray and flows downward, counter-currently washing the remaining solids from the syngas. From the scrubber trays, a de-mister removes any entrained water droplets, such that an essentially particulate-free syngas exits from the top of the syngas scrubber.

2.2.3.2 Low-Temperature Gas Cleanup

The low-temperature gas cleanup (LTGC) Unit is a single system sized for 100 percent of plant capacity. The two main purposes of this system are to:

- Cool the raw syngas while producing steam; and
- Provide other gas cleanup functions, including carbonyl sulfide (COS) hydrolysis and water gas shift.

The LTGC unit receives syngas from the four (4) operating syngas scrubber trains. The syngas is then cooled in a series of two exchangers [the Syngas Interchanger against reheating treated syngas from the SELEXOL[®] unit and the low pressure (LP) steam generator which produces LP steam]. The resulting partly condensed syngas is separated, and the condensate is pumped into the return condensate stream.

After the separation, the syngas is heated to 400°F with medium pressure (MP) steam and split into two streams. The syngas either enters a water shift reactor which converts CO and H₂O to CO₂ and H₂ and hydrolyzes COS or enters a reactor where COS is hydrolyzed to hydrogen sulfide (H₂S) and CO₂. The flows are balanced to adjust the H₂ to CO ratio of the syngas for optimal methanol synthesis. The two streams are then cooled in a series of two exchangers before entering knock-out drums. Syngas in the overhead vapor streams is routed to the SELEXOL[®] Acid Gas Removal Unit as a shifted and unshifted syngas stream.

The condensate from the LTGC area flows to a stripper, which also receives the condensate streams from the gasification system. The stripper removes almost all of the ammonia (NH₃), H₂S, and COS from the condensate, along with some dissolved hydrogen (H₂) and CO. The stripper overhead gas is blended with sour flash gases from the flash separators and compressed before going to the SELEXOL[®] Unit, so that the H₂ and CO can be recovered from the sour gas. The stripper bottoms water is returned to the syngas scrubber.

2.2.4 SELEXOL® Acid Gas Removal (2100)

The SELEXOL® process, licensed by UOP, has been selected as the acid gas removal technology. Two SELEXOL® process trains will provide the following functions for the shifted and unshifted streams:

- Removal of sulfur compounds (H₂S and COS) from the syngas to a level acceptable to the downstream Methanol Synthesis Unit,
- Recovery of most of the CO₂ in the syngas for further purification, and
- Recovery of a concentrated H₂S/COS stream to be sent to the Sulfur Recovery Unit (SRU).

The quenched sour syngas from the Syngas Conditioning Unit enters a mercury removal bed, and then is mixed with recycled stripped gas and flows to the SELEXOL® Feed/Product Exchanger to cool the feed gas against treated syngas and enhance the efficiency of absorption. The cooled feed gas flows through two successive absorbers; the first absorber removes H₂S and the second absorber removes CO₂. In each absorber, the syngas enters at the bottom of a packed bed and flows upward through the bed where it contacts cool solvent entering the top of the tower. In these absorbers, H₂S, COS, CO₂, and other gases such as H₂, are transferred from the gas phase to the liquid phase. The treated gas passes through de-entrainment devices at the top of the absorbers, as well as three water wash trays to minimize solvent carry-over. The treated syngas exits the top of the CO₂ absorber and is sent to the downstream Methanol Synthesis Unit.

Treated syngas leaving the SELEXOL® Unit is expected to contain less than 0.1 parts per million by volume (ppmv) total sulfur. Further sulfur reduction through the use of sulfur beds is required to protect the catalyst in the downstream Methanol Synthesis Unit from poisoning and the risk of sulfur spikes that could be caused by SELEXOL® Unit upsets. Each of the parallel beds is sized for full plant capacity. For best performance, the syngas is heated to 400°F before entering the guard bed.

The syngas from the guard beds is then sent to a compressor, where the syngas pressure is increased to the levels required in the Methanol Synthesis Unit. The syngas is then sent to the Methanol Synthesis Unit.

The SELEXOL® solvent from the H₂S Absorber is regenerated by stripping out less soluble gases, such as CO₂, H₂, and CO. The partially regenerated SELEXOL® solvent then flows to an H₂S stripper, where the remaining H₂S, CO₂, N₂, and other compounds are transferred from the liquid phase to the gas phase by contact with steam. The steam and liberated gases exit the stripper, and then flow upward through a demister and into the trayed section of the column. In the trayed section, the rising gas is contacted with counter-current flowing reflux water to cool and partially condense the hot overhead vapor, as well as reduce solvent entrainment. The overhead stream passes through a de-entrainment device and exits the top of the column. The overhead gas then passes through a condenser in order to condense and recover a portion of the overhead steam. The liquid and vapor phases are separated; the H₂S-rich acid gas exits the unit battery limits and is sent to the SRU, and the liquid is returned to the trayed section of the H₂S stripper.

2.2.5 Methanol Synthesis (4500)

Methanol is produced from synthesis gas using a highly selective copper-based catalyst. These reactions are exothermic and occur at a temperature suitable for generating medium pressure steam. Efficient use of waste heat from the methanol synthesis process is important for overall plant economics.

The Plant will use the licensed Davy Process Technologies methanol synthesis process. Major components of this process include:

- Syngas compression
- Syngas purification
- Methanol conversion

Particulate- and acid gas-free syngas is compressed and preheated before entering the Syngas Purification Vessel, which removes any remaining low levels of impurities that could potentially poison the methanol synthesis catalyst.

Feed gas from the Syngas Purification Vessel enters the first Methanol Converters, where it flows over methanol synthesis catalyst. On leaving the reactor, the gas mixture is cooled and methanol and water condense out. The remaining gas is compressed and mixed with incoming compressed syngas and recycled through the methanol converters. A small purge is taken from recirculated gas to control the level of inerts in the loop. Part or all of this gas undergoes hydrogen recovery, while the remainder is used as high-pressure fuel gas. The crude methanol is reduced in pressure to flash off the dissolved gases, mainly CO₂. The off gases are sent to the power block as fuel gas. During normal operation, the crude methanol flows to the MTG unit. However, if the MTG unit is offline, methanol production can continue and be sent to intermediate storage.

2.2.6 Methanol to Gasoline (5500)

The Exxon-Mobil MTG process will convert methanol exiting the Methanol Synthesis Unit to approximately 18,500 BPD of high-octane gasoline. Hydrocarbons produced during the process are mainly in the gasoline boiling range (C₅+ to 412°F) with a lesser amount in the C₁-C₄ range. The process also produces a small amount of carbon oxides, a very small amount of oxygenates and coke, and a very large quantity of water. The following discussion summarizes the MTG process.

The chemistry of methanol conversion is complex. First, methanol is partially dehydrated using an alumina catalyst to an equilibrium mixture of methanol, dimethyl ether (DME), and water. Then, methanol and DME undergo a series of dehydration reactions in the MTG reactors forming light alkenes. Light alkenes oligomerize (i.e., undergoing chain growth by joining two or more alkene molecules together) and cyclise to give the final products.

One hydrocarbon produced of particular note is durene (1, 2, 4, 5-tetramethyl benzene), which is produced in greater amounts than is suitable for gasoline (unless the high-durene gasoline is blended with gasoline containing lower durene concentrations). The MTG process contains a step (Heavy Gasoline Treatment) to reduce the durene to suitable levels.

The MTG catalyst deactivates slowly due to coke deposits. Coke must be removed periodically by in situ combustion with air to restore catalyst activity. For this reason, five (5) parallel MTG reactors are provided. At any given time, one reactor will be off-line (either in regeneration or on stand-by) and the other reactors will be on-line (converting DME reactor effluent to hydrocarbons and water.)

The effluent from the MTG reactors is combined, cooled, and separated into three phases: gas, liquid water, and liquid hydrocarbon.

- Gas Phase: Most of the gas phase is recycled to the MTG reactor inlet. The remaining gas is purged to the plant's fuel gas system.
- Liquid Water Phase: The large volume of liquid water produced by the reactions contains about 0.1 weight-percent (wt%) oxygenates (alcohols, ketones, and acids).
- Liquid Hydrocarbon Phase: The liquid hydrocarbon phase from the MTG reactor is called raw MTG gasoline.

Raw MTG gasoline contains 3-6 wt% durene (1, 2, 4, 5-tetramethyl benzene) while commercial gasoline specifications typically require less than 2.0 wt% durene. A Heavy Gasoline Treatment (HGT) unit is provided to reduce the durene content to 2.0 wt%. The HGT unit fractionates raw MTG gasoline into two parts. One part is a small volume, heavy fraction with a high durene concentration; the other part is a large volume, light fraction.

The heavy fraction is heated using the HGT Reactor Charge Heater and hydrotreated in a fixed-bed reactor (the HGT reactor) to reduce its durene concentration. The hydrotreated heavy fraction is combined with the untreated light fraction to produce finished MTG gasoline meeting the durene specification.

2.2.6.1 MTG Regeneration System

During the conversion reaction in an MTG Reactor, coke forms slowly on the catalyst and reduces its activity. To restore catalyst activity, coke is periodically removed from the catalyst by controlled combustion with air, one reactor at a time.

For catalyst regeneration, one MTG Reactor is taken out of oil service and is isolated from the other reactors and hydrocarbons. After isolation, the reactor is depressurized to the HP flare. Hydrocarbon vapors are then removed from the reactor and are replaced with nitrogen. Regenerator gas consisting primarily of nitrogen is recycled and mixed with a controlled quantity of air. The hot gas flows to the MTG Reactor where coke on the catalyst is removed by controlled combustion. Regeneration flue gas leaves the reactor and is cooled and separated.

Following coke combustion, the reactor is again evacuated, purged with nitrogen, and filled with recycle gas. The reactor is brought back on-line by flowing recycle gas through the Reactivation Heater and then starting DME reactor effluent feed when the bed temperature is high enough to sustain reaction.

At an appropriate time, another MTG reactor is taken out of service for regeneration.

2.2.6.2 MTG Water Treatment Unit

The MTG water is processed to remove most organics and oxygenates so that it will meet GE specifications for process water recycle to the gasification unit.

The water from the MTG Unit is heated against hot stripped water in the Feed/Product Exchanger before entering the MTG Water Stripper. There, most of the oxygenates and any residual hydrocarbons are driven overhead as vapor. The stripper overhead is condensed by the air-cooled Stripper Overhead Condenser and the condensate is recovered in the Receiver. LP steam is used to drive the Stripper Reboiler. The aqueous stripper condensate, containing most of the oxygenates, is pumped from to the Power Block where it will be vaporized into one of the power plant fuel streams. Any insoluble organics are decanted in the Receiver and pumped to the slops system. Any trace non-condensables are sent to flare.

Because acetic acid and any heavier acids cannot be completely stripped from the water, provision is made for caustic injection into the stripper sump to neutralize the acids to ensure that the pH is above 5.5. The stripped, neutralized water from the bottom of the stripper is pumped by the Stripper Bottoms Pump, cooled in the Stripper Overhead Condenser against the feed water, and routed to one of the Gasification Units.

2.2.6.3 LPG Processing Unit

The MTG Process produces a significant LPG byproduct stream consisting of approximately 60 percent olefin and 40 percent paraffin materials. LPG average production is expected to be 27,171 lb/hr, which is approximately 3,380 BPD.

In the Plant's geographic area, LPG has no significant market value. Therefore, LPG will be used as in-plant fuel or a blending stock for RVP control. The RVP pressure specification changes month to month. Any LPG not used for RVP control will be used as fuel and can provide approximately 500 MMBtu/hr to the plant in summer. LPG fuel usage will reduce the quantity of natural gas or syngas used by the Plant.

2.3 CO₂ RECOVERY (2200) AND PRODUCTION

Under normal operations, a CO₂-rich stream exits the SELEXOL[®] Unit. At this point in the process, the CO₂ contains less than 10 parts per million (ppm) total sulfur. The CO₂ flows into the CO₂ Recovery Unit, where it is compressed in one of three parallel four-stage centrifugal compressor trains and dried in a drying unit installed upstream of the third stage compressor suction. Some of the CO₂ is then refrigerated to provide liquid coolant to the Methanol Synthesis and SELEXOL[®] Units. The remaining CO₂ is ready for sale.

During startup, shutdown, and malfunction (SSM) events at the site, the CO₂ exiting the SELEXOL[®] Unit may be vented either because the CO₂ does not meet downstream specifications or because the site does not have sufficient power to start the CO₂ compression trains. This venting will occur through the CO₂ Vent Stack until the gas meets specifications and the compressors have been started, at which point no further emissions will occur from this stack. When venting occurs, the vent stream will be heated to 75°F by heat exchangers using steam from the existing processes (no new fired heater is required).

2.4 SULFUR RECOVERY (3100) AND PRODUCTION

In the Sulfur Recovery Area, the H₂S and COS in the acid gas from the SELEXOL[®] Unit is converted to elemental sulfur. After recovery of the sulfur, the non-sulfur portions of the Claus gas are treated to remove residual sulfur species.

The acid gas feed to the Sulfur Recovery Unit (SRU) is first washed with stripped sour water. The washed acid gas is then injected into a reaction furnace, where it is partially combusted with oxygen from the Air Separation Unit. The combustion products, which include sulfur, H₂S, SO₂, and CO₂, are cooled in the waste heat boiler to produce MP steam, and then further cooled in a condenser, where elemental sulfur is condensed.

Since the reaction of H₂S and SO₂ to produce sulfur is limited by equilibrium, the vapors from the first sulfur condenser are reheated against MP steam and reacted to form more sulfur over a special catalyst. These reaction products are once again cooled to condense more sulfur. To maximize the conversion of the sulfur species to elemental sulfur, two more subsequent stages of reheat, reaction and sulfur condensation are included. This is a three-stage Claus process, and about 42 TPD of sulfur will be produced and sold.

The raw sulfur recovered from the condensers flows as a liquid to a below-ground concrete pit. Since the raw sulfur contains dissolved H₂S and other volatile sulfur species, a sulfur degassing system, including transfer pump, reaction vessel, and ejector is used to remove the volatiles. The purified sulfur is then pumped to liquid sulfur storage before being shipped as a liquid to the customer.

The unconverted gas from the last sulfur conversion stage (SRU tail gas) still contains about 5% of the sulfur in the feed acid gas, mostly COS and CS₂ that are difficult to convert to sulfur. To remove these sulfur species, the SRU tail gas passes through a hydrogenation reactor that reduces them to H₂S. The reducing gas (hydrogen and CO) is produced by partially combusting fuel gas in the Reducing Gas Generator. The effluent from the reducing gas generator is cooled by generating LP steam, and then washed with water before proceeding to tail gas treatment.

The SRU tail gas is compressed and injected at the inlet of the SELEXOL H₂S Stripper where it is combined with the SELEXOL H₂S flash gas. During normal operation, the SRU tail gas will be recycled back to the SELEXOL[®] Unit. However, SRU tail gas will be routed to one of the flares in the event of a SELEXOL[®] or Claus unit upset. There are no continuous or intermittent purge gas streams from the SELEXOL[®] Unit.

When tail gas from the Claus units is routed to the SELEXOL[®] Unit, there are no vapor emissions to atmosphere from the SELEXOL[®] Unit. The following three vapor streams originate in the SELEXOL[®] Unit and flow to other plant areas:

- CO₂ product stream — The CO₂ product stream is compressed and sent to a pipeline customer. In an emergency or shutdown this stream may be vented; however, the stream is vented from the CO₂ recovery area, not from the SELEXOL[®] Unit.
- Claus gas stream — The Claus Gas is reacted to produce elemental sulfur, with any residual gas recycled to the SELEXOL[®] Unit. In an emergency or shutdown situation, the stack gas is vented from the sulfur plant area, not from the SELEXOL[®] Unit.
- Treated syngas — The treated syngas stream flows to the methanol synthesis area.

2.5 ANCILLARY OPERATIONS

2.5.1 Power Generation (7100)

The Power Block will consist of three parallel GE 7EA gas turbines normally fueled by a mixture of fuel gas, LPG, syngas, and natural gas that will produce approximately 185 MW in simple cycle mode at 100% firing rates at average normal operating annual ambient conditions. In addition, a heat recovery system on the gas turbine exhaust will superheat the medium pressure (MP) steam from the Methanol Synthesis area and the low pressure (LP) steam from the Syngas Conditioning area, and also produce and superheat HP steam. The superheated HP steam, MP steam and LP steam will then flow to a single, three-stage steam turbine, thereby producing approximately 215 MW of additional power, for a total nominal 400 MW.

If one of the three gas turbines is off-line, the two operating gas turbines with the heat recovery system would be capable of producing enough power to maintain the facility at full operating rates. Duct firing may be required in this scenario during summer operations. This operating flexibility is expected to considerably improve the overall availability of the Plant.

During the initial facility startup, power will be supplied by three, 1.6 MW Black Start Generators (Gen 1, 2 and 3). These generators will fire natural gas and will be operated until the Power Block can supply sufficient power.

2.5.2 Air Separation Unit (6100)

Two (2) identical air separation trains are provided, each of which will produce 3,700 short tons per day of 98 percent by volume (vol%) oxygen.

Atmospheric air is compressed to approximately 100 pounds per square inch absolute (psia) using an electric-driven compressor, treated to remove condensables, and fed to the air separation unit (ASU) where oxygen is separated cryogenically from atmospheric air. Following separation, the oxygen product with a purity of 98 vol% is pumped to high pressure as a cryogenic liquid and vaporized against a stream of condensing high pressure air within the ASU main heat exchanger. Almost all the gaseous oxygen product at 1,250 pounds per square inch gauge (psig) is fed as oxidant to the gasifiers. A small portion of the oxygen is let down in pressure and routed to the SRU, where it is used for sulfur production.

Since water is at a premium in the facility, ASU compressor intercooling and aftercooling is provided by a closed-loop, 66,000 gallons per minute (GPM) circulating glycol system, with heat rejection to the atmosphere by air-coolers.

A quantity of nitrogen is taken from the ASU and compressed for general plant usage, such as purging and tank inerting.

2.5.3 Intermediate and Product Storage (8100-8200)

Twelve (12) intermediate and product storage tanks will store large quantities of volatile materials. The largest of these storage tanks will include ten 150 ft diameter, 48 ft high, fully enclosed internal floating roof tanks. Two of these 150 ft diameter tanks will store methanol intermediate to provide some process buffering. The remaining eight of the largest tanks will store gasoline product, providing 60 days of product storage. An additional 130 ft diameter, 48

ft high tank will store heavy gasoline intermediate and a 7,000 gallon tank will store slops containing some volatile components.

Fifteen (15) additional small vessels will store the materials listed below.

- Liquid sulfur product
- Process water
- Additive
- Coolant
- Filtrate
- Glycol
- Liquid nitrogen
- Liquid oxygen
- LPG

2.5.4 Slag Handling and Water Cleanup (1200)

Slag slurry and black water from the Gasification Area enter the Slag Handling and Water Cleanup Area. The slag is dewatered using a flash system with hot water blowdown streams from the Gasifiers and Syngas Scrubber. The slag is conveyed to a stockpile where it will be loaded into trucks for offsite uses by others. There may be some slagscreening performed, as determined by customer demand. The slag is a vitreous (glass-like), high-density material and is not expected to become airborne. However, the stockpile will be kept wet as needed to prevent particulate emissions.

Gray water from the Water Cleanup system is routed to the Sour Water Stripper.

2.5.5 Water Treatment (1300 and 7100)

The Plant uses water for processing and as a heating and cooling medium in both liquid and steam phases. Raw water enters the Plant and is pumped to the Raw Water Tank located within the Power Block. From there, the raw water is filtered and processed by reverse osmosis (RO) and/or demineralizer units to produce the boiler feed water and the process water requirements of the overall facility.

The Plant is designed to be a zero-liquid process discharge facility. Water is re-used as much as possible and only a small portion of the total water with a high concentration of dissolved minerals flows to one of two evaporation ponds.

The brine concentrate from the RO system, along with gasification purge water, contain high concentrations of dissolved minerals such as sodium chloride. The combined reject water streams are sent to the steam-assisted evaporation pond within the Power Block, in which LP steam and solar energy are used to evaporate the residual water. The minerals are deposited in a layer at the bottom of the evaporation ponds, from which they may be eventually removed for off-site disposal.

Aqueous effluents (including gasification quench blowdown and steam generation blowdown) that cannot be recycled within the process areas will be sent to the Raw Water Processing Unit within the Power Block. If possible, this water will be re-used as substitute raw water feed, otherwise it will be sent to the Water Treatment Area for evaporation. The evaporation pond is sized to handle facility effluents and plant storm water runoff that has been through oil/water separation. Biological treatment of process water is not expected to be required.

2.5.6 Flares (8900)

Two continuous pilot flare systems will be operated at the facility: a HP flare and a LP flare. The large HP flare will be designed to handle the largest flare loads, such as, for example, the total syngas flow from the gasifiers in the event that they must be isolated from the downstream units. The HP system will operate at a positive pressure to minimize the cost of piping and equipment. The smaller LP flare system will operate at close to atmospheric pressure and will handle smaller flare loads such as the MTG stripper vent emergency releases. Sections 3 (Emission Estimates) and 4 (BACT) include detailed information about the flares.

2.5.7 Other Utilities (8300)

2.5.7.1 Instrument Air / Plant Air

Instrument air and plant air will be supplied by four (4) 50% capacity packaged units, one of which is powered by a generator in case of plant-wide power failure. No nitrogen backup for plant air is included. Each unit will supply 18,700 standard cubic feet per hour (SCFH) instrument air and 5,600 SCFH plant air. This system is included within the Power Block.

2.5.7.2 Nitrogen

Plant nitrogen for purging, tank inerting, and general plant purposes, as well as process nitrogen will be supplied from the ASU at 125 psig. A 10,000 gallon liquid nitrogen storage tank, with ambient air vaporizer, will be provided for backup supply and for startup service.

2.5.7.3 Cooling

All ambient temperature cooling is done, directly or indirectly, with air coolers.

2.5.7.4 Natural Gas / Plant Fuel Gas

Natural gas will be used for startup and as part of the fuel mix on an as-needed basis for the power generation system and process heaters.

2.6 STARTUP ACTIVITIES

The first step in the startup process is to obtain the power required for energizing the critical control and safety systems. Power for initial startup of the gas turbines is provided by the three “black start” natural gas electric generators (Gen 1, Gen 2, and Gen 3), which will be used to provide power for approximately 1 week or less. Other key utility systems such as instrument

air, water supply and purification, fire water, and nitrogen will be made operational as required to start the first gas turbine. It is especially important that the flare system be ready for service before any flammable gas is present.

Once critical utilities are in service, one of the three gas turbines (Turbine 1-3) is started on pipeline quality natural gas. This will produce enough power to displace all of the black start generators, start the circulating glycol cooling system, start the auxiliary boiler circulation and gasification quench water system, and begin the startup of one of the ASUs. During normal operations, the turbine fuel will be a combination of natural gas, fuel gas, and LPG.

One of the two ASUs can be started up once adequate electric power is available. The circulating glycol cooling system must be in service before the ASU compressors can operate. From an initial warm condition, the ASU startup can take several days for cool down of the cold box equipment. When online, the ASU will initially produce enough oxygen to begin operation of two of the four (4) coal gasifiers needed for full-capacity operation. At this time, a second gas turbine is started up, also on natural gas, to provide enough power for full capacity operation of one ASU.

Before each gasifier can be started, the refractory in that gasifier must be heated. Refractory heating is accomplished using the natural gas-fired preheaters (Preheater 1-5) and takes approximately 500 hours per gasifier. Multiple gasifiers may be preheated simultaneously. In addition to completing the refractory heating, the plant quench water circulation must be in service, along with the sour water stripper and low temperature syngas cooling system before the startup of any gasifier. To start the first gasifier, the natural gas fired preheat burner is shut down, removed and replaced with the coal slurry feed injector. Coal slurry and oxygen are then fed to the injector to initiate the gasification of the coal. A second gasifier is then started up in the same manner as the first. By this time, the single ASU is operating at full rates and is producing enough oxygen to feed two (2) gasifiers. The initial raw syngas product is flared until the syngas conditioning unit is on-line, which is anticipated to take approximately 1 week during the initial startup.

Circulation of SELEXOL[®] solvent through the Acid Gas Removal System is commenced at this time. The refrigeration package must also be in operation to chill the solvent to operating temperature. Once the SELEXOL[®] unit is ready, and when the two gasifiers are in service at full operating pressure and temperature, the syngas is allowed to enter the SELEXOL[®] unit. The CO₂ recovered by the SELEXOL[®] unit is initially vented (CO₂ Stack) until the CO₂ meets pipeline specifications, which may take some days. The desulfurized syngas from the SELEXOL[®] unit is flared until the methanol synthesis unit is ready to receive feed. During the cold start there will be a brief period (anticipated to be approximately 10 hours) where off-spec gas may be flared.

After the SELEXOL[®] unit is in service, the gasifier system operation is adjusted if necessary to make syngas of the proper composition so that, after acid gas removal, the syngas is an acceptable feed for the Methanol Synthesis Unit.

The SRU can be started up once a sufficient flow of sulfur-rich acid gas (Claus Gas) is available from the SELEXOL[®] unit. Once desulfurized syngas that meets the Methanol Synthesis Unit specifications is available, the methanol synthesis unit can be started up to produce methanol which is routed to an intermediate storage tank. Once methanol of sufficient quantity is available to assure startup of the methanol to gasoline (MTG) unit, the MTG unit will convert methanol to

hydrocarbons (primarily gasoline) and water in fixed-bed reactors. Methanol is then converted to an equilibrium mixture in the DME reactor. The effluent from the DME reactor is then combined with recycled gas and converted to gasoline and water through the MTG reactors. The MTG reactor effluent is collected and separated into three phases. (1) A portion of the gas phase is recycled with the remaining gas being sent to the plant fuel gas system. (2) The liquid water phase produces water which is recycled into the gasifier unit, and (3) The liquid hydrocarbon phase becomes raw MTG gasoline. Following hydrotreating, the facility produces finished gasoline, LPG and fuel gas of high quality.

When MP steam is available in adequate quantity from the syngas cooldown and methanol unit, the MP steam is routed through the gas turbine superheat coils, permitting the steam turbine to be started up to produce additional power. The flow to the steam turbine is augmented by LP steam from gasification low temperature syngas cooling.

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3.1 SADDLEBACK HILLS MINE

Originally Arch of Wyoming LLC (subsidiary of Arch Coal, Inc.) permitted the Mine (underground) and the Elk Mountain (surface) Mines together under one air quality permit (Permit # CT-4136). The combined facilities were known as the Carbon Basin Mines. Arch Coal has entered into an option agreement to sell the underground coal reserve and surface real property to MBFP. Once MBFP exercises this option, Arch Coal has retained the rights to operate the Elk Mountain Mine and market the surface coal. As a result of this agreement, a determination was made by the Wyoming Department of Environmental Quality (WDEQ)/Air Quality Division (AQD) that the Saddleback Hills Mine was considered a support activity under the definition of a facility and should be included in the MBFP PSD application.

During the underground mine’s development phase, approximately 2.5 million tons of coal will need to be mined over a 3-year period. The development phase constructs the underground infrastructure required to support the longwall mining system which will commence operations at approximately the time when the Plant achieves full capacity. During the development or construction phase of the mine, coal will be conveyed from the South Portal where it will be stored in a small stockpile. It is anticipated that this production will either be loaded into trucks at the South portal and hauled to the Seminoe II train loadout in Hanna, Wyoming, or placed in the designated long term storage stockpile.

During the MBFP construction phase, development will also occur at the East Portal. The following activities will occur at the East Portal:

- Construction of the East Portal entry areas that will consist of a reinforced concrete retaining wall;
- Installation of conveyors from the portal face to the coal storage facilities (some conveyors will be fully enclosed, some will be ¾-covered);
- Construction of the coal storage facilities;
- Construction of a ¾-covered overland conveyor system from the coal storage facilities to the Plant;
- Construction of the Mine’s office, maintenance shop, and warehouse facilities.

Emission sources associated with the Mine during the development phase are shown in Table 3.1.

Table 3.1 – Mine Development Particulate Emissions

Development Year	Coal Conveying and Loading PM ₁₀ (tpy)	Coal to Seminoe II PM ₁₀ (tpy)
1	3.04	26.8
2	5.17	109.3
3	4.20	71.6

Only particulate emissions associated with the Mine are included in the table above. Emissions from mine area fuel combustion (on-site machinery) are based on calculations provided in Permit Application AP 2989 for the Carbon Basin Mines.

Detailed Mine Development emission calculation spreadsheets are included in Appendix B.

3.2 THE PLANT

3.2.1 Emission Sources

Emissions associated with this Plant include both point source and fugitive emission sources. The three combustion turbines account for the majority of NO_x, CO, SO₂, and PM₁₀ emissions, while storage tanks and equipment leaks emit the most VOCs and HAPs. Table 3.2 shows significant point and fugitive sources of emission.

Manufacturer specifications for the turbines and certain other equipment are included in Appendix C. With regard to the combustion turbines, a General Electric (GE) specification sheet has been included in Appendix C; this specification does not constitute a vendor guarantee from GE. Equipment-specific guarantees could not be obtained from vendors at this time. Guarantees for some equipment will be obtained at the time purchase contracts are signed.

Due to the long lead-time needed to design this Plant, specific manufacturers and models have not yet been identified for many equipment items, and manufacturer specifications are not yet available.

A list of other major equipment is included in Appendix D, along with a list of source classification codes (SCCs) for point source equipment.

Table 3.2 – Emission Units and Fugitive Sources

Description	Identification	Size	Use
<i>Normally Operating Equipment and Fugitive Sources</i>			
Combustion Turbine 1	CT-1	66 MW	Electrical and steam generation
Combustion Turbine 2	CT-2	66 MW	Electrical and steam generation
Combustion Turbine 3	CT-3	66 MW	Electrical and steam generation
Auxiliary Boiler	AB	66 MMBtu/hr	Steam generation (normal service is standby at 25% load to prevent freeze ups if there is a Plant shutdown)
Catalyst Regenerator*	B-1	21.53 MMBtu/hr	Catalyst regeneration (only during catalyst regeneration; average continuous rate is approximately 9 MMBtu/hr)
Reactivation Heater*	B-2	12.45 MMBtu/hr	Reactivation heating
HGT Reactor Charge Heater	B-3	2.22 MMBtu/hr	Reactor charge heating
HP Flare (pilot only)	FL-1	0.82 MMBtu/hr	For safety and VOC control
LP Flare (pilot only)	FL-2	0.20 MMBtu/hr	For safety and VOC control
Equipment Leaks	EL	N/A	N/A
Storage Tanks	Tanks	Various	Primarily methanol and gasoline storage
Coal Storage & Processing	CS	N/A	Coal conveyance & feedstock storage
<i>SSM Equipment</i>			
Gasifier Preheater 1*	GP-1	21 MMBtu/hr	Gasifier refractory preheating
Gasifier Preheater 2*	GP-2	21 MMBtu/hr	Gasifier refractory preheating
Gasifier Preheater 3*	GP-3	21 MMBtu/hr	Gasifier refractory preheating
Gasifier Preheater 4*	GP-4	21 MMBtu/hr	Gasifier refractory preheating
Gasifier Preheater 5*	GP-5	21 MMBtu/hr	Gasifier refractory preheating
Black-Start Generator 1*	Gen-1	2889 hp	Electrical generation
Black-Start Generator 2*	Gen-2	2889 hp	Electrical generation
Black-Start Generator 3*	Gen-3	2889 hp	Electrical generation
Firewater Pump Engine*	FW-Pump	575 hp	Supplies emergency firewater
CO ₂ Vent Stack*	CO ₂ VS	N/A	For malfunctions

* These emission units operate less than 8,760 hr/yr.

3.2.2 Normal Operations

Plant emissions are broken down into three categories (normal operation, cold startup/initial year emissions, and malfunctions). Annual emissions resulting from normal operations include emissions from equipment that operates continuously (8,760 hours per year) and equipment that operates on a regular basis. For example, the firewater pump engine may operate up to 500 hours in a typical year. Consequently, firewater pump engine emissions are included in the normal operation annual emission summary and are based on 500 hr/yr rather than 8,760 hr/yr. Note that the Auxiliary Boiler normally operates at only 25 percent load, on a hot standby basis.

However, emissions are based on 8,760 hr/yr operation at full load. Table 3.3 shows emissions resulting from normal operations and the maximum number of hours of operation per year. Detailed emission calculations are included in Appendix B.

Table 3.3 – Annual Criteria Pollutant Emissions Resulting from Normal Operations

Source ID	Description	Operating Hours (hr)	Potential Emissions (tpy)				
			NO _x	CO	VOC	SO ₂	PM ₁₀
CT-1	Power Generation	8,760	75.86	46.19	6.59	10.79	43.80
CT-2	Power Generation	8,760	75.86	46.19	6.59	10.79	43.80
CT-3	Power Generation	8,760	75.86	46.19	6.59	10.79	43.80
AB	Steam Generation ¹	8,760	14.17	23.81	1.56	0.17	2.15
B-1	Catalyst Regeneration	8,760 ²	4.62	7.77	0.51	0.06	0.70
B-2	Reactivation Heater	8,760 ²	2.67	4.49	0.29	0.03	0.41
B-3	HGT Reactor Charge Heater	8,760	0.48	0.80	0.05	0.01	0.07
Tanks	Product Storage	8,760	---	---	102.62	---	---
EL	Equipment Leaks	8,760	---	---	59.63	---	---
CS	Coal Storage & Processing	8,760	---	---	---	---	61.08
FW-Pump	Firewater Pump Engine ³	500	1.51	0.09	0.34	0.00	0.02
FL-1	HP Flare	8,760 ⁴	0.49	0.98	2.97	0.00	---
FL-2	LP Flare	8,760 ⁴	0.12	0.25	0.74	0.00	---
Total Emissions			251.63	176.75	188.49	32.65	195.84

1. Boiler will normally operate at 25% load, but potential emissions are based on continuous full load operation.
2. The catalyst regeneration heater and reactivation heaters will operate less than 8,760 hr/yr, but potential emissions are based on 8,760 hr/yr of operation.
3. The Firewater Pump combusts diesel fuel.
4. Based on continuous natural gas pilot for flares.

Table 3.4 shows annual HAP emissions resulting from normal operations. The largest HAP emission sources at the Plant are listed in the following table.

Table 3.4 – Annual HAP Emissions Resulting from Normal Operations

Pollutant	Facility-Wide Potential Emissions (tpy)	Largest Emission Source at Facility
Benzene	8.54	Equipment Leaks
Formaldehyde	0.71	Turbines
Hexane	1.29	Auxiliary Boiler ¹
Methanol	10.26	Equipment Leaks
Toluene	1.81	Turbines
Other HAPs	2.11	N/A
Total Emissions	24.71	

1. Note that HAP PTE emissions from the auxiliary boiler are calculated at continuous, full load operation. However, the boiler will normally operate at only 25% load but within compliance with its emission commitment (lb/MMBtu basis). The second-largest emission source contributing to hexane emissions at the facility will be storage tanks.

3.2.3 Cold Start/Initial Year Operations

Annual emissions have also been calculated for the initial year of operations (plant cold start). The complete Plant startup period may last as long as 180 days, and will involve bringing equipment online in a particular order. Emissions during the cold startup period will differ from those during a normal operating year. Certain equipment, such as Black-Start Generators and Gasifier Preheaters, will operate during cold startup. Individual emission units will have much shorter startup time periods; these unit-specific time periods are shown in Appendix B in the cold startup emission summary spreadsheet. Since the Plant will not have produced adequate in-plant fuels and power generation will ramp up slowly, most combustion equipment will initially burn only natural gas fuel, rather than the fuel mixture of fuel gas, LPG, and natural gas. Table 3.5 shows the annual emissions resulting from Cold Startup.

Table 3.5 – Annual Criteria Pollutant Emissions Resulting from Cold Startup

Source ID	Description	Operating Hours Fuel Gas Mixture/NG ¹	Potential Emissions (tpy)				
			NO _x	CO	VOC	SO ₂	PM ₁₀
CT-1	Power Generation	7760 / 1000	76.68	46.61	6.64	10.90	43.80
CT-2	Power Generation	7760 / 1000	76.68	46.61	6.64	10.90	43.80
CT-3	Power Generation	7760 / 1000	76.68	46.61	6.64	10.90	43.80
Gen-1	Black-Start Generator 1	0 / 360	1.15	2.79	1.03	0.00	0.00
Gen-2	Black-Start Generator 2	0 / 360	1.15	2.79	1.03	0.00	0.00
Gen-3	Black-Start Generator 3	0 / 360	1.15	2.79	1.03	0.00	0.00
AB	Steam Generation	8000 / 760	14.17	23.81	1.56	0.17	2.15
B-1	Catalyst Regeneration	8760 / 0	4.62	7.77	0.51	0.06	0.70
B-2	Reactivation Heater	8000 / 760	2.67	4.49	0.29	0.03	0.41
B-3	HGT Reactor Charge Heater	8000 / 760	0.48	0.80	0.05	0.01	0.07
GP-1	Gasifier Preheater 1	0 / 500	0.26	0.43	0.03	0.00	0.04
GP-2	Gasifier Preheater 2	0 / 500	0.26	0.43	0.03	0.00	0.04
GP-3	Gasifier Preheater 3	0 / 500	0.26	0.43	0.03	0.00	0.04
GP-4	Gasifier Preheater 4	0 / 500	0.26	0.43	0.03	0.00	0.04
GP-5	Gasifier Preheater 5	0 / 500	0.26	0.43	0.03	0.00	0.04
Tanks	Product Storage	8760	---	---	102.62	---	---
EL	Equipment Leaks	8760	---	---	59.63	---	---
CS	Coal Storage & Processing	8760	---	---	---	---	61.08
FW-Pump	Firewater Pump Engine	500 ²	1.51	0.09	0.34	0.00	0.02
CO ₂ VS	CO ₂ Vent Stack	8760	---	314.89	0.84	---	---
FL-1	HP Flare	8760 ³	10.28	81.86	3.11	187.70	0.00
FL-2	LP Flare	8,760 ⁴	0.15	0.44	0.74	36.01	0.00
Total Emissions			268.64	584.48	192.87	256.69	196.04

1. Operating hours shown for firing fuel gas mixture and natural gas (NG) are based on expected operations. However, emissions are conservatively calculated based on firing natural gas, which is the higher emitting fuel.

2. The Firewater Pump combusts diesel fuel.

3. Based on continuous natural gas pilot for flare; cold startup includes 50 hr/yr of vents to HP Flare.

3. Based on continuous natural gas pilot for flare; no vents to LP Flare are expected during cold startup.

3.2.4 Malfunctions and Other Events

Malfunctions and other events can cause unusual emissions during short periods of time.

Table 3.6 includes four types of malfunctions. Detailed emission calculations for malfunction events are included in Appendix B.

Table 3.6 – Criteria Pollutant Emissions Resulting from Malfunctions and Other Events

Source ID	Description	Operating Hours ¹	Potential Emissions (tons)				
			NO _x	CO	VOC	SO ₂	PM ₁₀
CO ₂ VS	CO ₂ Vent Stack	50	---	83.97	0.23	---	---
FL-1	HP Flare	40 ²	7.83	64.99	0.12	150.16	---
FL-2	LP Flare	8 ²	0.01	0.00	0.00	14.40	---
GP-1	Gasifier Preheater	500 ³	0.26	0.43	0.03	0.00	0.04

1. The hours shown are estimates of annual operating hours due to malfunctions.
2. Each flare is expected to combust vented gases for the number of hours shown; pilot operation will occur throughout the year.
3. During a non-cold startup year, only one of the five Gasifier Preheaters is expected to operate for up to 500 hours.

3.2.5 Emissions of PSD-Regulated Pollutants

The MTG process requires the syngas to be relatively pure in order to prevent the poisoning of the methanol synthesis catalyst. The clean syngas that is used in the MTG process is the same syngas used as fuel throughout the Plant. This cleaning is achieved by running the raw syngas from the gasifiers through a wet scrubber, which cools the raw gas and removes any particulates that are entrained in the gas stream. The raw (sour) gas then flows through the mercury vapor guard beds (mercury removal) and then through the Low Temperature Gas Cleanup process (SELEXOL[®] technology) where the raw syngas is further cleaned and where NH₃, H₂S, and COS are removed from the raw syngas. After the SELEXOL[®] process, the gas flows through a final sulfur guard bed to ensure the highest level of sulfur removal (<0.1 ppmv total sulfur).

Trace amounts of some contaminants may be emitted in very small quantities. During the feasibility study, certain trace contaminants were estimated and are shown below.

Contaminant	Concentration	Potential to Emit
Halogens (Cl ₂ and F)	<0.01 ppmv	0.001 tpy
Sulfur as H ₂ S	<0.09 ppmv	0.009 tpy

At least 90 percent of the lead in the tail gas will be removed by the activated carbon beds that remove mercury. Based on 3 million tons (8,000 TPD) of coal gasified and lead content within the coal averaging 1.93 ppmw (determined by testing), total lead exiting the gasifiers would be 5.79 tpy. Based on a conservative estimate of 90 percent removal, lead emissions from the facility are estimated to be 0.579 tpy.

3.2.6 Source-Specific Calculation Methods

The following sections provide additional detail about calculation methods used to estimate emissions from certain types of sources.

3.2.6.1 Combustion Source Methods

Most Plant combustion sources can be fueled with either a fuel gas mixture or with natural gas. The fuel gas mixture includes fuel gas and LPG that are produced within the Plant and supplementary natural gas. Mixing of the fuel gas components occurs prior to the combustion chamber of the source. The fuel gas mixture will vary between seasons and due to catalyst efficiency. Methanol production is high when the catalyst is at its beginning of life (BOL), compared to end of life (EOL). Typical molar fractions of fuel gas mixture components are shown in Table 3.7.

Table 3.7 — Typical Fuel Gas Mixture Composition¹

Fuel Component	Winter BOL	Winter EOL	Summer BOL	Summer EOL
Natural Gas	70.30%	63.01%	58.69%	50.82%
LPG	2.99%	2.75%	7.97%	7.19%
MTG Fuel Gas	4.76%	4.37%	5.94%	5.36%
Davy PSA Purge	16.87%	25.19%	21.05%	30.89%
Davy Fuel Gas 1	2.44%	2.13%	3.05%	2.61%
Davy Fuel Gas 2	2.65%	2.55%	3.30%	3.13%
Total	100.00%	100.00%	100.00%	100.00%

1. Molar percentages are given. Based on three turbines operating.

Since the fuel gas mixture is plant-specific, emission factors are not available for the fuel gas mixture. However, since the fuel has a significant methane component and also includes large quantities of C3 and C4 fuels, use of natural gas emission factors is a reasonable approximation. Consequently, emission calculations for non-diesel combustion sources are based on natural gas emission factors. Even so, the differences in heating values between natural gas and the fuel gas mixture causes emissions to differ.

In some circumstances, combustion of the fuel gas mixture is impractical. This is particularly true during initial startup when the plant has not yet produced sufficient quantities of syngas and LPG. Detailed emission calculation spreadsheets (Appendix B) for the combustion turbines, auxiliary boiler, and heaters clearly indicate the number of hours during which natural gas or the fuel gas mixture is being fired. (KAW question – any revision needed here? The boiler and heater sheets make mention of it, but is it enough to say it's 'clearly' indicated?)

3.2.6.2 Storage Tanks

Storage tank emissions were calculated using the EPA TANKS Program, version 4.09.d, based on use of internal floating roof tanks. TANKS reports for each type of tank having significant emissions are included in Appendix B.

The RVP of product gasoline stored at the site will vary depending on the time of year. Month-to-month vapor pressure variability was accounted for in the calculations. Tanks containing no volatile organic components and those with insignificant emissions are listed on the Tanks detailed calculation page within Appendix B.

3.2.6.3 Equipment Leaks

Equipment leak estimates were calculated using the average emission factor approach described in EPA's "Protocol for Equipment Leak Emission Estimates" (EPA-453/R-95-017). EPA-approved Synthetic Organic Chemical Manufacturing Industry (SOCMI) factors were used for the calculations. Although use of the Refinery emission factors was considered, use of the Refinery factors was deemed inappropriate for the following reasons.

- The Plant process is a chemical synthesis process rather than a refinery process.
- SOCMI factors are recommended for use in all industries, except refineries.
- Even within refineries, SOCMI factors are recommended for chemical processes, such as production of methyl tertiary butyl ether (MTBE).
- The refinery emission factor equation usage guidelines specifically disallow corrections for methane concentrations exceeding 10 wt% and some process streams at the Plant will contain more than 10 wt% methane.

Process streams within the Plant were grouped according to composition and service type (gas, light liquid, heavy liquid) and the number of potential equipment leak components was estimated for each process stream group. All streams were assumed to contain fluids for 8,760 hr/yr. Within Appendix B, detailed equipment leak calculations show controlled and uncontrolled emissions. Controlled emissions were calculated using control effectiveness factors for valves in gas or light liquid service and pump seals in light liquid service. The control effectiveness factors are based on implementation of a monthly Leak Detection and Repair (LDAR) program and assume a leak definition of 500 ppm for valves and connectors in VOC service and 2,000 ppm for pumps in VOC service. As discussed in the BACT analysis, the Plant will implement an LDAR program.

3.2.6.4 Flares

Flaring emission calculations are based on procedures included in "TCEQ Guidance Document for Flares and Vapor Oxidizers" (RG-109, October 2000). This document provides emission factors for NO_x and CO and advises use of 98% destruction efficiency for VOCs / HAPs and H₂S.

The HP and LP Flares will be operated with continuous pilots. Consequently, normal operations include combustion emissions based on the design heat input for each flare and assume natural gas firing. Emissions from normal operation at both flares represent pilot gas combustion only, because no process streams will be routinely directed to either flare.

Emissions from large malfunction events were estimated for the HP and LP Flares, due to the possible significant nature of a malfunction event affecting these flares. Malfunction-related emissions from the HP Flare are based on directing all syngas to the flare, which is the largest stream, by volume, that could potentially be directed to the HP Flare. Malfunction-related events affecting the LP Flare for a potential worst-case (high flow rate, high H₂S content) vent stream that could be directed to the LP Flare.

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The proposed Plant is one of the 28 named source categories in 40 CFR §52.21(b)(1) and is classified as a new major source of regulated emissions under the PSD New Source Review (NSR) program. An analysis of the Best Available Control Technology (BACT) is required for sources with potential emissions greater than the PSD established significance thresholds. The BACT analysis evaluates the technical feasibility and cost-effectiveness of emission control options to determine the applicable control technology and emission limits.

BACT is determined on a case-by-case basis taking into consideration technical practicability and economic reasonableness. For PSD BACT requirements, energy and environmental impacts should also be considered. Control technology alternatives are identified for each new or modified source of pollutants based on knowledge of the applicant’s particular industry and previous regulatory decisions for other identical or similar sources.

The proposed Plant will be located in Carbon County, Wyoming. Carbon County is currently designated attainment or unclassifiable for all national ambient air quality standards. Table 4.1 evaluates the applicability of BACT requirements.

Table 4.1 – BACT Applicability

Pollutant	Significance Threshold (tpy)	Estimated Facility Potential to Emit (tpy)	BACT Applicable
CO	100	176.75	Yes
NO _x	100	251.63	Yes
SO ₂	100	32.65	No ¹
PM ₁₀	100	195.84	Yes
VOC	100	188.49	Yes

1. Although federal PSD regulations do not require BACT for sources with less than 100 tpy of potential emissions, WDEQ requires BACT reviews for minor sources.

4.1 BACT REVIEW PROCESS

In a December 1, 1987 memorandum from the EPA Assistant Administrator for Air and Radiation, the agency provided guidance on the “top-down” methodology for determining BACT. The “top-down” process involves the identification of all potentially applicable emission control technologies according to control effectiveness. Evaluation begins with the top or most stringent emission control alternative. If the most stringent control technology is shown to be technically or economically infeasible, or if environmental impacts are severe enough to preclude its use, then it is eliminated from consideration and the next most stringent control technology is similarly evaluated. This process continues until the BACT option under consideration cannot be eliminated. The top control alternative not eliminated is determined to be BACT. This process involves the following five steps from “New Source Review Workshop Manual,” DRAFT October 1990, EPA Office of Air Quality Planning and Standards.

- Step 1: Identify all available control technologies with practical potential for application to the specific emission unit for the regulated pollutant under evaluation;
- Step 2: Eliminate all technically infeasible control technologies;
- Step 3: Rank remaining control technologies by control effectiveness and tabulate a control hierarchy;
- Step 4: Evaluate most effective controls and document results; and
- Step 5: Select BACT, which will be the most effective practical option not rejected based on economic, environmental, and/or energy impacts.

Formal use of these steps is not always necessary. However, the BACT requirements have consistently been interpreted to contain two core components that must be met in any determination. First, the BACT analysis must consider the most stringent available technologies (those with the potential to provide the maximum reductions). Second, a determination to use a technology with a lesser potential control efficiency must be supported by an objective analysis of the associated energy, environmental, and economic impacts. Additionally, the minimum control efficiency evaluated in the BACT analysis must at least achieve emission rates equivalent to applicable New Source Performance Standards (NSPS) or other applicable state or federal rules.

The process of identifying potential control technologies involves researching many resources, including a review of existing and historical technologies that have been proposed or implemented for other projects and a survey of available literature. Evaluating the applicability of each control option entails an assessment of feasibility and cost-effectiveness. This process determines the potential applicability of a control technology by considering its commercial availability (as evidenced by past or expected near-term deployment on the same or similar types of emission units). An available technology is one that is deemed commercially available because it has progressed through the following development steps: concept stage; research and patenting; bench scale/laboratory testing; pilot scale testing; licensing and commercial demonstration; and commercial sales.

The evaluation process also considers the project specific physical and chemical characteristics of the gas stream to be controlled. A control method applicable to one emission unit may not be applicable to a similar unit because of differences in the physical and chemical characteristics of gas streams to be controlled.

The following BACT analysis for the proposed Plant was conducted in a manner consistent with the top-down approach. As part of this analysis, control options for potential reductions were identified by researching the EPA Reasonably Available Control Technology (RACT)/BACT/Lowest Achievable Emission Rate (LAER) Clearinghouse (RBLC) database and by drawing upon engineering, integrated gasification combined cycle (IGCC) process, and industrial gasification permitting experience, and by surveying available literature. IGCC facilities employ several processes similar to the proposed Plant. Potential controls identified were then evaluated as necessary on a technical, economic, environmental, and energy basis.

4.2 BACT SUMMARY

Table 4.2 summarizes BACT proposed for this project:

Table 4.2 - Summary of BACT Applied to the Plant

Source	Proposed BACT Method
Combustion Turbine/HRSG/Steam Turbine Combined Cycle Trains (3x3x1)	NO _x SCR with NO _x control to 6 ppmvd NO _x (corrected to 15% O ₂) in the HRSG exhaust when firing fuel gas mixture or natural gas CO: Catalytic Oxidation control to 6 ppmvd CO (corrected to 15% O ₂) in the HRSG exhaust when firing fuel gas mixture or natural gas VOC: Collateral control from Catalytic Oxidation control to 1.4 ppmvw CO (corrected to 15% O ₂) in the HRSG exhaust when firing fuel gas mixture or natural gas PM/PM10: Good combustion practices SO ₂ : SRU system designed to reduce fuel sulfur concentrations to 0.1 ppmvd and combustion of low sulfur natural gas as supplementary fuel
Auxiliary Boiler and Process Heaters	NO _x : Low NO _x burners CO, VOC, PM/PM10: Good combustion practices SO ₂ : SRU system designed to reduce fuel sulfur concentrations to 0.1 ppmvd and combustion of low sulfur natural gas as supplementary fuel
Storage Tanks	Gasoline, Methanol, Heavy Gasoline, and Slop Storage tanks will have internal floating roofs; all other tanks will have fixed roofs
Coal Handling	Dust suppression (fogging) used in combination with fully enclosed conveyors and passive engineering design at transfer points
Equipment Fugitives	VOC: Leak Detection and Repair (LDAR) program
Sulfur Recovery Unit (flare and thermal oxidizer)	Re-route tail gas to upstream point in SELEXOL [®] Unit
Carbon Dioxide Vent	Startup, shutdown, upset conditions only (<50 hours/year), optimized process design
Gasifier Preheaters	Low sulfur fuel (natural gas), good combustion practices, restricted operation (initial startup and new refractory only, < 500 hours/year per gasifier)
Black-Start Generators	Low sulfur fuel (natural gas), good combustion practices, restricted operation (initial startup only, <360 hours/year)
Firewater Pump	Restricted operation (<500 hours/year), ultra-low-sulfur diesel fuel (15 ppm sulfur), good combustion practices

4.3 COMBUSTION TURBINE CONTROL TECHNOLOGY REVIEW

The following is the BACT analysis for the proposed combustion turbines. Each of the three proposed combustion turbines will be a GE 7EA model turbine with a nominal capacity of 66 MW at average ambient conditions. Each combustion turbine will have a heat recovery steam generator (HRSG), and all three will utilize one steam turbine generator, in a 3 x 3 x 1, combined cycle configuration. The primary fuel will be a fuel gas mixture comprised of imported natural gas plus process generated fuels including: LPG from the MTG process, and fuel gas from both the Davy and MTG synthesis processes. By volume, the combustible portion of this natural gas based fuel mixture will consist primarily of methane (61.4%), hydrogen (15.3%), and butane (5.1%). Each combustion turbine will also be capable of firing natural gas, for startup, fuel enrichment, and backup purposes. Finally, under certain market conditions, each combustion turbine may also be fired with a syngas-based fuel mixture. By volume, the combustible portion of this syngas-based fuel mixture will consist primarily of hydrogen (46.1%) and CO (44.5%) with a small amount of hydrocarbons.

4.3.1 Nitrogen Oxides BACT Analysis for the Combustion Turbines

NO_x is formed during combustion primarily by the reaction of combustion air nitrogen and oxygen within the high temperature combustion zone (thermal NO_x), or by the oxidation of nitrogen in the fuel (fuel NO_x). Because the tail gas contains negligible amounts of fuel-bound nitrogen, essentially all combustion turbine NO_x emissions originate as thermal NO_x.

The rate of thermal NO_x formation in the combustion turbines is primarily a function of the fuel residence time, availability of oxygen, and peak flame temperature. Several NO_x control technologies are available to reduce the impacts of these variables during the combustion process, including diluent injection and dry low NO_x burner technology. Post-combustion control technologies have also been used in some processes to remove NO_x from the exhaust gas stream.

Identify Control Technologies

The following NO_x control technologies were evaluated for the proposed combustion turbines:

Combustion Process Controls

- Diluent Injection
- Dry Low NO_x Burners
- Low NO_x Burners
- Flue Gas Recirculation

Post-Combustion Controls

- EMxTM
- Selective Non-Catalytic Reduction (SNCR)
- Selective Catalytic Reduction (SCR)

*Evaluate Technical Feasibility***Diluent Injection**

Higher combustion temperatures may increase thermodynamic efficiency, but may also increase the formation of thermal NO_x. A diluent, such as water, steam, or nitrogen can be added to the fuel gas mixture to effectively reduce the combustion temperature and formation of thermal NO_x. The fuel gas mixture combusted in the combustion turbines contains small amounts of N₂ and CO₂, both of which act as a diluent. However, additional dilution is would be necessary to achieve meaningful NO_x reductions. Diluent injection is a technically feasible control technology for the proposed combustion turbines while firing the fuel gas mixture. N₂ produced in the ASU could be introduced to the turbine burners in this instance to reduce combustion temperatures. In addition, when the turbines are firing natural gas only, nitrogen from the ASU could be introduced as a diluent also. There may be brief periods of time when the turbines are first started (on natural gas) when no diluent from the ASU is available. This is expected to be a very short time period as the ASU is one of the first units started during the startup sequence.

Dry Low NO_x Burners

Dry Low NO_x (DLN) burner technology has successfully been demonstrated to reduce thermal NO_x formation from combustion turbines firing natural gas. This technology utilizes a burner design that controls the stoichiometry and temperature of combustion by regulating the distribution and pre-mixing of fuel and air, which minimizes localized fuel-rich pockets that produce elevated combustion temperatures and higher NO_x emissions.

Available DLN burner technologies for combustion turbines are designed for natural gas (methane-based) fuels, but are not applicable to combustion turbines utilizing a fuel gas mixture, which has a different heating value, gas composition, and flammability characteristics. Research is ongoing to develop DLN technologies for tail gas (or fuel gas mixtures) and syngas-fueled combustion turbines, but no designs are currently available. In particular, the turbine vendor has stated that DLN is not feasible for fuels that contain less than 85% by volume methane or that contain substantial amounts of hydrogen. The fuel gas mixture that will be utilized in the turbines contains too little methane (61.4%) and too much hydrogen (15.3%). Therefore, DLN burner technology is not technically feasible for the Plant turbines due to potential explosion hazards in the combustion section associated with the high content of hydrogen in the fuel gas mixture.

Low NO_x Burners

Low NO_x burners are widely used to reduce NO_x emissions. A conventional low NO_x burner is designed to control fuel and air mixing at each burner in order to create larger and more branched flames. This reduces peak flame temperature and results in less NO_x formation. In addition, the improved flame structure reduces the amount of oxygen available in the hottest part of the flame and improves burner efficiency. In contrast to DLN burners, low NO_x burners can be used with a variety of gaseous fuels. Low NO_x burner technology is technically feasible for Plant turbines.

Flue Gas Recirculation

Flue gas recirculation is being researched by combustion turbine manufactures, but is not currently an available control technology. While the technology may be a future option to

reduce NO_x emissions, significant development work is required to complete maturation and integration of the concept into a power plant system, including validating all emissions characteristics and overall plant performance and operability. Additionally, current research efforts have focused on pre-mixed natural gas combustion, and results would need to be expanded to assess fuel gas mixture applications. Thus, flue gas recirculation is not technically feasible for the proposed combustion turbines.

EMx™

EMx™ (formerly known as SCONO_x) is a control technology that utilizes a single catalyst to minimize CO, VOC, and NO_x emissions. All installations of the technology have been on small natural gas facilities. EMx™ has not been applied to large-scale fuel gas mixture/syngas combustion turbines, which creates concerns regarding the timing, feasibility of scaling up to a larger unit and use of different fuel, cost-effectiveness of necessary design improvements, and potential catalyst fouling. Therefore, EMx™ is not technically feasible.

Selective Non-Catalytic Reduction (SNCR)

SNCR is a post-combustion NO_x control technology in which a reagent (ammonia or urea) is injected in the exhaust gas to react with NO_x to form nitrogen and water without the use of a catalyst. The success of this process in reducing NO_x emissions is highly dependent on the ability to uniformly mix the reagent into the flue gas, which must occur in a very narrow high temperature range. The consequences of operating outside the optimum temperature range are severe. Above the upper end of the temperature range, the reagent will be converted to NO_x. Below the lower end of the temperature range, the reagent will not react with the NO_x, resulting in excess ammonia emissions. SNCR technology is occasionally used in conventional coal-fired heaters or boilers, but it has never been applied to natural gas combined cycle or syngas/fuel gas mixture units because no locations exist in the heat recovery steam generator with the optimal temperature and residence time that are necessary to accommodate the technology. Therefore, SNCR is not technically feasible.

Selective Catalytic Reduction (SCR)

SCR technology has never been attempted at an IGCC plant using coal-derived syngas. BACT analyses for previously permitted IGCC plants have determined SCR is not technically feasible due to concerns regarding a back pressure energy penalty, catalyst performance, and potential operational impacts to downstream equipment from the sulfur content in the fuel. Several analyses noted the unavailability of meaningful performance guarantees from SCR suppliers. In other cases, the application of SCR to the IGCC process was not deemed cost effective due to increased operation and maintenance costs and the costs associated with reducing syngas sulfur to levels that are assumed to be adequate to minimize operational impacts.

MBFP's initial evaluation of the application of SCR to the Plant indicates that due to the extremely high sulfur removal necessary for the MTG process, catalyst fouling and other operational concerns due to sulfur in the fuel would be alleviated. The gas fed to the Methanol Synthesis Unit requires less than 30 ppb sulfur. All fuel gas used throughout the plant is first desulfurized in the acid gas removal (AGR) unit and sulfur beds, and therefore contains less than 30 ppb sulfur (expressed as H₂S). In summary, under the proposed fuel gas mixture-firing scenario, SCR is believed to be technically feasible.

During most startup operations, the combustion turbines will be fired with fuel gas mixture. However, for the initial startup and some cold startup scenarios, natural gas will be used to fire the combustion turbines. SCR is not technically feasible during the initial startup operations due to the low temperature where the SCR would be applied. Whether firing natural gas or the fuel gas mixture, the SCR will be utilized as soon as the exhaust temperature reaches the operational range of the SCR.

Rank Control Technologies

Low NO_x burners, SCR, and diluent injection are the NO_x control technologies that are technically feasible for the proposed combustion turbines during normal operations when firing either the fuel gas mixture or natural gas.

Evaluate Control Options

The use of low NO_x burners and SCR was identified as the only technically feasible NO_x control technology for the proposed combustion turbines during normal operations. The low NO_x burners are expected to achieve 25 ppm NO_x in turbine exhaust. The use of SCR will further reduce NO_x emissions to 6 ppmvd (at 15% O₂) when firing syngas (fuel gas mixture). The nominal gross output for the 3 x 3 x 1 generator/HRSG/ steam turbine configuration is 400 MW. Therefore, the equivalent potential NO_x emission rate is approximately 0.135 lb/MWh, significantly lower than the applicable NSPS Subpart Da or KKKK limit of 1.0 and 3.6 lb/MWh respectively.

The use of low NO_x burners and diluent injection combined with SCR was identified as the only technically feasible combination of NO_x control technologies for the proposed combustion turbines during natural gas firing operations. These combined technologies will reduce NO_x emissions to 6 ppmvd (at 15% O₂).

With one exception, the proposed NO_x BACT limit of 6 ppmvd (corrected to 15% O₂) is well below emission limits found on the RACT/BACT/LAER Clearinghouse for similar turbines firing either syngas or tail gas. Appendix E provides a summary of emission control determinations for these turbines. For completeness, all RACT/BACT/LAER emission control determinations for process type 15.250 (explained in Appendix E) are included. The most stringent NO_x BACT limit for a combined cycle combustion turbine firing syngas or tail gas is 1.9 ppmvd (corrected to 15% O₂ and based on an annual average) for the Bayport Energy Facility. However, this facility utilizes DLN technology to achieve this level of NO_x emissions. For reasons described above, DLN is not technically feasible for the Plant. The next most stringent NO_x BACT limit is 8 ppmvd (corrected to 15% O₂ and based on a 30-day rolling average) for the Exxon Mobil Shute Creek facility. The Exxon-Mobil facility uses a proprietary mix of gas that includes syngas as one component. All other fueled combustion turbines shown in Appendix E have NO_x emission limits of 15 ppmvd (corrected to 15% O₂) or more.

As the first implementer of SCR technology on this type of turbine/fuel combination, the 6 ppmvd NO_x emission limit reflects a level of control within the accepted range of SCR control efficiencies (70-90 percent control efficiency). Specifically, a reduction from 25 ppmvd to 6 ppmvd is estimated, representing a long-term 76 percent reduction in NO_x from 80 percent SCR performance when the system is new and clean. Technical issues such as pressure loss in

the combustion turbine and ammonia slip argue against expecting the highest level of control efficiency for this innovative installation of SCR.

Moreover, the additional cost of reducing NO_x emissions to below 6 ppm has been estimated, although MBFP believes that achieving NO_x emissions less than 6 ppmvd (corrected to 15% O₂) is a technical feasibility issue rather than a cost issue. Variability in plant-generated fuel could potentially increase NO_x emissions and prevent burner optimization. Consequently, exhaust from the turbines may be somewhat higher than expected. With a 6-ppm NO_x limit, the facility will have some ability to compensate for high NO_x concentrations entering the SCR system by increasing NO_x removal efficiency beyond the 76 percent that would be achieved assuming 25 ppm NO_x concentration in the turbine exhaust. Based on equipment and operating costs provided by SNC Lavalin, the incremental cost of reducing NO_x emissions from 6 ppm to 4 ppm, is estimated to be \$2,455/ton removed. This cost estimate is included as Appendix H.

Select NO_x Control Technology

The use of SCR with diluent injection is proposed as BACT for the proposed combustion turbines during normal operations to reduce NO_x emissions to 6 ppm when firing fuel gas mixture. The use of SCR with diluent injection is also proposed for natural gas combustion during start up operations. The proposed BACT NO_x limits are presented below for each combustion turbine.

Proposed NO_x BACT Limit when burning fuel gas mixture: 6 ppmvd (corrected to 15% O₂)

Proposed NO_x BACT Limit when burning natural gas: 6 ppmvd (corrected to 15% O₂)

The NO_x BACT limits expressed for each combustion turbine are for normal operations. During startup and shutdown operations, NO_x emissions may be greater for certain periods due to unstable combustion associated with lower combustion turbine efficiencies and transitional periods between fuels. Potential emissions for startup and shutdown operations are provided in the Emissions Inventory and are evaluated as part of the air dispersion modeling analysis. See Section 4.3.5 for more information regarding startup operations.

4.3.2 Sulfur Dioxide BACT Analysis for the Combustion Turbines

The combustion turbines oxidize sulfur compounds in fuel primarily into sulfur dioxide (SO₂). Emissions can be controlled by limiting the fuel sulfur content or by removing SO₂ from the exhaust gas.

Identify Control Technologies

The following SO₂ control technologies were evaluated for the proposed Plant combustion turbines.

Pre-Combustion Process Controls

Chemical Absorption Acid Gas Removal

Physical Absorption Acid Gas Removal

Low Sulfur Fuel

Post-Combustion Controls

Flue Gas Desulfurization

*Evaluation Technical Feasibility***Chemical and Physical Acid Gas Removal Systems**

During the gasification process, sulfur in the feedstock converts primarily into H₂S, and will also convert into minor quantities of other sulfur species, such as COS. Commercially available AGR systems are capable of removing greater than 99% of the sulfur compounds from syngas/tail gas. AGR systems are commonly used for gas sweetening processes of refinery fuel gas or tail gas treatment systems, and are typically coupled with processes that produce useful sulfur byproducts.

AGR systems can employ either chemical or physical absorption methods. Chemical absorption methods are amine-based systems that utilize solvents, such as methyldiethanolamine (MDEA), to bond with the H₂S in the tail gas. A stripper column is then used to regenerate the solvent and produce an acid gas stream containing H₂S that can be processed into useful sulfur by-products. An MDEA AGR system has been determined as BACT for all operating and permitted IGCC facilities. The two operating IGCC facilities in the United States both use amine (MDEA) systems to reduce the syngas total sulfur concentration to 100 to 400 ppm. The process involves taking the gas out of the AGR removal process and passing it through a methanol synthesis process, and the gases coming out of the methanol and MTG processes (fuel gas mixture) are used as fuel in the combustion turbines. In order for the methanol process to function properly the sulfur content in the gas must be less than 0.1 ppm sulfur. Therefore, chemical absorption methods, even with the use of sulfur beds, are not technically feasible for the Plant's process.

Other types of AGR systems utilize physical absorption methods that employ a physical solvent to remove sulfur from gas streams, such as mixtures of dimethyl ethers of polyethylene glycol (SELEXOL[®]) or methanol (Rectisol). These systems operate by absorbing H₂S under pressure into the solvent. Dissolved acid gases are removed resulting in a regenerated solvent for reuse and the production of an acid gas stream containing H₂S that can be processed into useful sulfur by-products. Physical absorption methods have historically been used to purify gas streams in the chemical processing and natural gas industries, and can achieve sulfur removal to the level required by methanol process of less than 0.1 ppm sulfur. This sulfur concentration can feasibly be reduced to the sulfur content required by the methanol unit through the use of sulfur removal beds. Physical acid gas removal systems are a technically feasible control technology.

Low Sulfur Fuel

Providing low sulfur fuel to the turbines is another pre-combustion emission control method. The AGR system described above removes sulfur from the fuel gas streams in order to provide

low sulfur fuel gas to the combustion turbines. However, additional fuel is needed for the turbines. Natural gas is a low sulfur fuel that can be used to supplement fuel gases produced at the Plant. The combustion turbines' burners are compatible with Plant-produced fuel gases, natural gas, and a combination of both types of fuels. When firing natural gas exclusively, SO₂ emissions are conservatively estimated to be 0.0034 lb per MMBtu. Use of natural gas as the supplementary fuel is a technically feasible option.

Flue Gas Desulfurization

Flue gas desulfurization (FGD) is a post-combustion SO₂ control technology that reacts an alkaline compound with SO₂ in the exhaust gas. FGD systems are most commonly used by conventional pulverized coal units and can typically achieve greater than 95% removal efficiency on new facilities. The FGD process results in a solid by-product that requires the installation of a significant number of ancillary support systems to accommodate treatment, handling, and disposal. FGD is more readily applied to high SO₂ concentration gas streams, such as those present with direct combustion coal units. No examples were identified where an FGD system has been applied to a tail gas/syngas fired combustion turbine facility or similar process, such as a natural gas fired unit. Therefore, FGD is not technically feasible for the proposed combustion turbines. Even if feasible to the tail gas fired processes, FGD could not achieve the high removal efficiencies associated with AGR systems and would not provide appreciable SO₂ removal.

Rank Control Technologies

The use of physical acid gas removal for process fuels and use of low sulfur natural gas fuel were identified as the only technically feasible SO₂ and acid gas emissions control technologies applicable to the proposed combustion turbines.

Evaluate Control Options

With regard to Plant-produced fuels, physical acid gas removal is the only feasible control technology identified, and is proposed as BACT for this project. Sulfur removal will occur prior to the methanol catalyst and will reduce the sulfur content to less than 0.1 ppmvd.

The AGR design reduces syngas sulfur concentrations by greater than 99%, and produces a secondary gas stream that can be processed into potentially useful sulfur byproducts. The solvent used by the AGR system will be regenerated and reused. Any related water streams will be treated, as the facility will be a zero water discharge facility. Overall, no collateral environmental issues have been identified that would preclude the AGR design option from consideration as BACT for the proposed project.

With regard to supplementary fuels, use of natural gas is the only feasible control method.

Select SO₂ Control Technology

A physical absorption AGR system designed to reduce tail gas sulfur concentrations to 0.1 ppm (expressed as H₂S) is proposed as BACT for SO₂ emissions from the proposed combustion turbines. The proposed AGR system will reduce fuel gas mixture sulfur content by greater than 99%. The gas fed to the Methanol Synthesis Unit requires less than 0.1 ppm, and therefore sulfur guard beds will be used to reach less than 0.1 ppm of sulfur. All fuel gas used throughout

the plant is first desulfurized in the AGR units and sulfur beds, and therefore contains less than 0.1 ppm sulfur (expressed as H₂S).

Although the fuel gas has very low sulfur content, the turbines burn a large proportion of natural gas as part of the fuel gas mixture (see Table 3.6 for fuel gas mixture components).

Consequently, the proposed BACT limits associated with combustion of the fuel gas mixture, as well as natural gas, are based on AP-42 factors of 0.0034 lb/MMBtu.

Proposed SO₂ BACT Limit when burning fuel gas mixture: 0.0034 lb/MMBtu

Proposed SO₂ BACT Limit when burning natural gas: 0.0034 lb/MMBtu

Carbon Monoxide BACT Analysis for the Combustion Turbines

CO emissions are a result of incomplete combustion. Providing adequate fuel residence time and higher temperatures in the combustion zone to ensure complete combustion can reduce CO emissions. However, these same control factors can increase NO_x emissions. Conversely, reduce NO_x emission rates achieved through flame temperature control (by diluent injection) can increase CO emissions. The design strategy is to optimize the flame temperature to reduce potential NO_x emissions, while minimizing the impact to potential CO emissions. The combustion turbines for the proposed project will be a GE 7EA model, which is designed to optimally consume fuel gas mixture. Post-combustion control technologies have also been used to reduce CO emissions in some processes.

Identify Control Technologies

The following CO control technologies were evaluated for the proposed combustion turbines.

Combustion Process Controls

Good Combustion Practices

Post-Combustion Controls

EMxTM

Oxidation Catalyst

Evaluate Technical Feasibility

Good Combustion Practices

Good combustion practices include the use of operational and design elements that optimize the amount and distribution of excess air in the combustion zone to ensure complete combustion. This technology has been determined to be BACT for CO emissions for combustion turbines, which use syngas/fuel gas mixture fired combustion turbines.

EMxTM

The EMxTM system was evaluated in the NO_x BACT analysis, and determined to not be technically feasible.

Oxidation Catalysts

Catalytic oxidation is a post-combustion control technology that utilizes a catalyst to oxidize CO into CO₂. Due to the significant portion of natural gas in the fuel gas mixture, oxidation catalyst is technically feasible for the Plant's turbines.

Rank Control Technologies

Good combustion practice and catalytic oxidation are the only technically feasible CO control technology identified.

Evaluate Control Options

Good combustion practice and catalytic oxidation are the only feasible control technology identified, and has been determined to be BACT for CO emissions for combustion turbines.

Select CO Control Technology

Good combustion practice and catalytic oxidation are proposed as BACT for CO emissions from the proposed combustion turbines. The use of good combustion practices is expected to achieve CO emissions of 6 ppmvd (at 15% O₂).

Proposed CO BACT Limit when burning fuel gas mixture: 6 ppmvd (corrected to 15% O₂)

Proposed CO BACT Limit when burning natural gas: 6 ppmvd (corrected to 15% O₂)

The CO BACT limits expressed for each combustion turbine are for normal operations. During startup and shutdown operations, CO emissions may be greater for certain periods due to unstable combustion associated with lower combustion turbine efficiencies and transitional periods between fuels. Potential emissions for startup and shutdown operations are provided in the Emissions Inventory and are evaluated as part of the air dispersion modeling analysis. See Section 4.3.5 for more information regarding startup operations.

4.3.3 Volatile Organic Compound BACT Analysis for the Combustion Turbines

VOC emissions are a product of incomplete combustion. Providing adequate fuel residence times and higher temperatures in the combustion zone to ensure complete combustion can reduce VOC emissions. The design strategy is to optimize the flame temperature to reduce potential NO_x emissions, while minimizing the impact to potential VOC emissions. The combustion turbines for the proposed project will be a GE 7EA model, designed to optimally consume fuel gas mixture. Post-combustion control technologies have also been used to reduce VOC emissions in some processes.

Identify Control Technologies

The following VOC technologies were evaluated for the proposed combustion turbines.

Combustion Process Controls

Good Combustion Practices

Post-Combustion Controls

EMx™

Catalytic Oxidation

Evaluate Technical Feasibility

Good Combustion Practices

Good combustion practices include the use of operational and design elements that optimize the amount and distribution of excess air in the combustion zone to ensure complete combustion. This technology has been determined to be BACT for VOC emissions from syngas fired combustion turbines in IGCC permits nationwide.

EMx™

The EMx™ system was evaluated in the NO_x BACT analysis, and determined to not be technically feasible.

Catalytic Oxidation

Catalytic oxidation, primarily a CO control device with limited VOC control, was evaluated in the CO BACT analysis, and determined to be technically feasible.

Rank Control Technologies

Good combustion practice and catalytic oxidation are the only technically feasible VOC control technology identified.

Evaluate Control Options

Good combustion practice and catalytic oxidation are the only feasible control technology identified, and has been selected as BACT for syngas fired combustion turbines.

Select VOC Control Technology

Good combustion practice and catalytic oxidation are proposed as BACT for VOC emissions from the proposed combustion turbines. The BACT emission limit is proposed below.

Proposed VOC BACT Limit when burning fuel gas mixture: 1.4 ppmvw (corrected to 15% O₂)

Proposed VOC BACT Limit when burning natural gas: 1.4 ppmvw (corrected to 15% O₂)

The VOC BACT limit expressed for each combustion turbine is for normal operations. During startup and shutdown operations, VOC emissions may be greater for certain periods due to unstable combustion associated with lower combustion turbine efficiencies and transitional periods between fuels. Potential emissions for startup and shutdown operations are provided in the Emissions Inventory and are evaluated as part of the air dispersion modeling analysis. See Section 4.3.5 for more information regarding startup operations.

4.3.4 Particulate Emissions BACT Analysis for the Combustion Turbines

Fuel quality and combustion efficiency are key drivers impacting the quantity and disposition of potential particulate emissions. In some processes, post-combustion control technologies can also be used to reduce particulates.

Identify Particulate Emission Control Technologies

The following particulate emission control technologies were evaluated for the proposed combustion turbines.

Combustion Process Controls

- Clean Fuels with Low Potential Particulate Emissions
- Good Combustion Practices

Post-Combustion Controls

- Electrostatic Precipitation
- Baghouse

Evaluate Technical Feasibility

Clean Fuels with Low Potential Particulate Emissions

Higher ash content fuels have the potential to produce greater particulate emissions. In addition, fuels containing sulfur have the potential to produce sulfur compounds that may form condensable particulate emissions. Combustion turbine operations require fuels that contain negligible amounts of fuel bound particulate in order to minimize performance impacts. The Plant's process inherently produces a fuel gas mixture containing minimal amounts of particulate. The control of fuel gas mixture sulfur compounds as discussed in the SO₂ BACT analysis will reduce potential condensable particulates. Therefore, the use of clean fuels is a technically feasible control technology.

Good Combustion Practices

The use of good combustion practices is a technically feasible control technology that minimizes particulate emissions resulting from incomplete combustion, and was proposed as BACT for CO and VOC emissions.

Electrostatic Precipitation

Electrostatic precipitation (ESP) is a post-combustion particulate control technology most commonly applied to large volume gas streams containing high particulate concentrations, such

as with direct combustion coal units. An ESP has not been applied to syngas/fuel gas mixture combustion turbine operations due to the low particulate concentrations of the associated exhaust gas streams. The use of ESP is not technically feasible based on the particulate matter present in the exhaust gas at the Plant. The particulate matter content will be less than 0.003 grains of PM/dscf.

An ESP can consistently provide PM emission reductions down to 0.002 to 0.015 grains of PM/dscf (from “Controlling Stack Emissions in the Wood Products Industry,” Gerry Graham). Therefore, an ESP would not provide additional control. Operation of an ESP is not considered technically feasible for the proposed combustion turbines.

Baghouse

A baghouse is a post-combustion control technology that uses a fine mesh filter to remove particulate emissions from gas streams, and is most commonly applied to industries producing large volume gas streams with high particulate concentrations. A baghouse has not been applied to syngas/fuel gas mixture combustion turbine operations due to the reduced volume and minimal particulate concentration of the associated exhaust gas streams. Use of a baghouse is not technically feasible based on the particulate matter present in the exhaust gas at the Plant. The particulate matter content is less than 0.003 grains of PM/dscf. A baghouse can consistently provide PM emission reductions down to 0.02 grains of PM/dscf. More stringent control can be achieved, but not greater than 0.003 grains of PM/dscf (per The Tenant Company, Griffin Filters, Farr Air Pollution Control). Therefore, a baghouse would not provide additional control and is not considered technically feasible for the proposed combustion turbines.

Rank Control Technologies

The use of clean fuels with low potential particulate emissions and good combustion practices were identified as the only technically feasible particulate emissions control technologies applicable to the proposed combustion turbines.

Evaluate Control Technologies

The use of clean fuels with low potential particulate emissions and good combustion practices were identified as the only technically feasible particulate emissions control technologies applicable to the proposed combustion turbines. These technologies have been determined to be BACT for syngas fired combustion turbines.

Select Particulate Emissions Control Technology

The use of clean fuels with low potential particulate emissions and good combustion practices are proposed as BACT for particulate emissions from the proposed combustion turbines. The following emission limit resulting from the implementation of these technologies is proposed for each combustion turbine.

Proposed Particulate Emissions (PM₁₀ – filterable) BACT limit when burning fuel gas mixture: 0.013 lb/MMBtu. Based on the Lower Heating value (LHV).

Proposed Particulate Emissions (PM₁₀ – filterable) BACT limit when burning natural gas: 0.013 lb/MMBtu. Based on the LHV.

The potential particulate combustion turbine emission rates during startup and shutdown operations are less than or equal to the aforementioned BACT limits for normal operations while firing fuel gas mixture. Potential emissions for startup and shutdown operations are provided in the Emissions Inventory and are evaluated as part of the air dispersion modeling analysis. See Section 4.3.5 for more information regarding startup operations.

4.3.5 Startup Emissions BACT Analysis for the Combustion Turbines

Turbine startup emissions are quantified separately from normal operating emissions. The SCR system used on the turbine/HRSG units does not initially reduce NO_x emissions since the system must heat up to achieve the operating temperature conducive for proper pollution control operation. When the temperature range is achieved during fuel gas mixture and natural gas startup operations, the SCR system will be engaged and the catalyst will begin to minimize NO_x emissions.

To satisfy BACT during the startup mode of the turbines, the duration of the startups will be minimized to the best extent possible for each turbine unit.

4.4 FIRED HEATER AND BOILER CONTROL TECHNOLOGY REVIEW

The BACT analysis for the proposed fired heaters and auxiliary boiler applies to three heaters with firing capacity of 21.5 MMBtu/hr to 2.2 MMBtu/hr and a 66 MMBtu/hr boiler. The fuel gas mixture, comprised primarily of methane and hydrogen, will fuel the fired heaters and auxiliary boiler during normal operations. Backup fuel for the heaters and boiler will be natural gas for startup and upset conditions, and is discussed in Section 4.4.5.

4.4.1 NO_x BACT Analysis for the Fired Heaters and Boiler

NO_x is formed during combustion primarily by the reaction of combustion air nitrogen and oxygen in the high temperature combustion zone (thermal NO_x), or by the oxidation of nitrogen in the fuel (fuel NO_x). The rate of NO_x formation is a function of fuel residence time, oxygen availability, and temperature in the combustion zone. Primary fired heater and auxiliary boiler NO_x control technologies focus on combustion process controls.

Identify All Control Technologies

The following potential NO_x control technologies were evaluated for the proposed auxiliary boiler and fired heaters.

Combustion Process NO_x Controls

Low NO_x Burners

Low NO_x Burners with Flue Gas Recirculation

Post-Combustion NO_x Controls

Selective Catalytic Reduction (SCR)

Selective Non-Catalytic Reduction (SNCR)

Non-Selective Catalytic Reduction (NSCR)

EMxTM

Evaluate Technical Feasibility**Low NO_x Burners**

Low NO_x burners reduce the formation of thermal NO_x by incorporating a burner design that controls the stoichiometry and temperature of combustion by regulating the distribution and mixing of fuel and air. As a result, fuel-rich pockets in the combustion zone that produce elevated temperatures and higher potential NO_x emissions are minimized. Historically, low NO_x burners have been selected as BACT for syngas/tail gas-fired heaters and boilers. Therefore, low NO_x burner technology is technically feasible for the proposed auxiliary boiler and fired heaters.

Low NO_x Burners with Flue Gas Recirculation

Flue gas recirculation (FGR) is used to reduce NO_x emissions in some processes by recirculating a portion of the flue gas into the main combustion chamber. This process reduces the peak combustion temperature and oxygen in the combustion air/flue gas mixture, which reduces the formation of thermal NO_x. FGR has the potential to reduce combustion efficiency and cause greater carbon monoxide emissions. A RBLC search was performed over the previous 10-year period for other gaseous fuels and gaseous fuel mixtures in boilers and process heaters less than 100 MMBtu/hr (Process Type 13.390). The search encompassed 24 facilities and 110 processes. Application of FGR was not identified for process heaters less than 100 MMBtu/hr in this search. All the process heaters and the auxiliary boiler at the facility will be less than 100 MMBtu/hr and will emit relatively small quantities of NO_x. Therefore, FGR has not been previously demonstrated for the intended operation of the fired heaters.

Selective Catalytic Reduction (SCR)

SCR is a post-combustion technology that reduces NO_x emissions by reacting NO_x with ammonia in the presence of a catalyst. SCR technology has been most commonly applied to pulverized coal generating units and to natural gas fired combustions turbines. A RBLC search was performed over the previous 10-year period for other gaseous fuels and gaseous fuel mixtures in boilers and process heaters less than 100 MMBtu/hr (Process Type 13.390). The search encompassed 24 facilities and 110 processes. Application of SCR was identified at two out of the 24 facilities. Therefore, SCR is technically feasible for the intended operation of the fired heaters. However, at one of the facilities that employed SCR, the RBLC stated that the project was "...to meet the new NO_x requirements dictated by the SIP." The other facility that employed SCR is located in an area regulated by the same SIP, and fired a fuel comprised primarily of hydrocarbons. Both of the facilities are located in an ozone nonattainment area and SCR was implemented to comply with the state NO_x rules (SIP). The Plant is not located in a nonattainment area and is therefore not subject to the same stringent NO_x rules as these two facilities with SCR. Additionally, based on the difference in fuels, the uncontrolled NO_x emissions would be higher from the hydrocarbon-fired heater as compared to the fuel gas fired heaters proposed by MBFP. Therefore, the NO_x reductions for the auxiliary boiler and fired heaters at the MBFP facility would receive comparatively less NO_x reduction benefit with the application of SCR, and the cost would not be warranted.

Selective Non-Catalytic Reduction (SNCR)

SNCR is a post-combustion NO_x control technology where ammonia or urea is injected into the exhaust to react with NO_x to form nitrogen and water without the use of a catalyst. Use of this technology requires uniform mixing of the reagent and exhaust gas within a narrow high

temperature range (1,600°F–1,900°F). Operations outside of this temperature range will significantly reduce removal efficiencies and may result in ammonia emissions or increased NO_x emissions. The auxiliary boiler and fired process heaters exhaust temperatures range from approximately 700°F to 900°F. Thus, SNCR is not technically feasible for the proposed auxiliary boiler or fired process heaters.

Non-Selective Catalytic Reduction (NSCR)

NSCR is a post-combustion control technology that utilizes a catalyst to reduce NO_x emissions under fuel-rich conditions. The technology has been utilized in the automobile industry and for reciprocating engines. A RBLC search was performed over the previous 10-year period for other gaseous fuels and gaseous fuel mixtures in boilers and process heaters less than 100 MMBtu/hr (Process Type 13.390). The search encompassed 24 facilities and 110 processes. Application of NSCR was not identified for process heaters or boilers less than 100 MMBtu/hr in this search. NSCR technology requires a fuel-rich environment for NO_x reduction, which will not be available in the proposed auxiliary boiler or fired heaters. Therefore, NSCR is not a technically feasible for the proposed auxiliary boiler or fired heaters.

EMx™

EMx™ is a post-combustion control technology that utilizes a single catalyst to minimize CO, VOC, and NO_x emissions. Installations of the technology have been limited to small natural gas combustion turbine applications. Recent analyses by state agencies have determined that the technology is currently not feasible for syngas/tail gas fired process heater applications. For example, the Oregon Department of Environmental Quality (ODEQ) concurred that EMx™ was not technically feasible for a proposed 140 MMBtu/hr auxiliary boiler project. ODEQ also noted that a small boiler (4.2 MMBtu/hr) project in California installed an EMx™ system, but the South Coast Air Quality Management District determined application of the technology could not demonstrate the necessary emission reductions. Based on these determinations and the limited scope of commercial installations, EMx™ is not technically feasible for the proposed auxiliary boiler or fired heaters.

Rank Control Technologies

SCR and the use of low NO_x burner technology were the only technically feasible control options identified for reducing NO_x emissions. The only applications of SCR identified by the RBLC search were located in an area where the SIP influenced the NO_x reductions which were more stringent than BACT. The total potential NO_x emissions proposed at the MBFP facility during normal operations for all heaters and the auxiliary boiler combined are 4.11 tpy. The use of SCR is not warranted at the Plant based on the relatively small amount of aggregate NO_x emissions from all of the fired process heaters.

Evaluate Control Options

Low NO_x burner technology has historically been selected as BACT for syngas/tail gas fired process heaters and provide good NO_x control through prevention of NO_x formation. As discussed earlier in this section, SCR is not warranted for these process heaters due to the small amount of NO_x emissions from the heaters.

Select NO_x Best Available Control Technology

The use of low NO_x burner technology is proposed as BACT for NO_x emissions from the proposed auxiliary boiler and fired process heaters. The proposed BACT emission limits for each unit are presented below for operation on both fuel gas mixture and natural gas.

Proposed NO_x BACT Limits:

Auxiliary Boiler: 0.036 lb/MMBtu (fuel gas mixture)

50.0 lb/MMscf (natural gas)

Catalyst Regen Heater: 30 lb/MMscf (fuel gas mixture)

Reactivation Heater: 30 lb/MMscf (fuel gas mixture)

50 lb/MMscf (natural gas)

HGT Reactor Charge Heater: 30 lb/MMscf (fuel gas mixture)

50 lb/MMscf (natural gas)

4.4.2 CO and VOC BACT Analysis for the Fired Heaters and Boiler

Potential CO and VOC emissions are due to incomplete combustion that is typically a result of inadequate air and fuel mixing, a lack of available oxygen, or low temperatures in the combustion zone. Fuel quality and good combustion practices can limit CO and VOC emissions. Good combustion practice has commonly been determined as BACT for syngas/tail gas fired heaters. Post-combustion control technologies using catalytic oxidation have also been used in some processes to reduce CO and VOC emissions.

Identify Control Technologies

The following CO and VOC control technologies were evaluated for the proposed fired heaters.

Combustion Process Controls

Good Combustion Practices

Post-Combustion Controls

Oxidation Catalyst

EM_xTM

Evaluate Technical Feasibility

Good Combustion Practices

Good combustion practices include the use of operational and design elements that optimize the amount and distribution of excess air in the combustion zone to ensure complete combustion. Good combustion practice has historically been determined as BACT for CO and VOC emissions from syngas-fired process heaters and is a technically feasible control strategy for the proposed auxiliary boiler and fired heaters.

Oxidation Catalyst

Catalytic oxidation is a post-combustion control technology that utilizes a catalyst to oxidize CO and VOC into CO₂ or H₂O. The technology has most commonly been applied to natural gas fired combustion turbines. No examples were identified where oxidation catalyst technology has been applied to a syngas-fired process heater. Because of the low potential CO and VOC emissions without an oxidation catalyst during normal operations (less than 6.92 tpy CO and less than 0.57 tpy VOC from the auxiliary boiler and all heaters combined), the use of catalytic oxidation technology is determined not to be warranted due to the small emission reduction potential.

EM_xTM

EM_xTM technology is discussed in the NO_x BACT analysis and determined not to be technically feasible.

Rank Control Technologies

Good combustion practice is the only feasible control strategy identified, and has historically been selected as BACT for CO and VOC emissions from syngas/tail gas fired process heaters.

Evaluate Control Options

Good combustion practice is the only feasible control strategy identified, and has historically been selected as BACT for CO and VOC emissions from syngas/tail gas fired process heaters.

Select CO and VOC Control Technology

The use of good combustion practices is proposed as BACT for potential CO and VOC emissions from the auxiliary boiler and proposed process heaters. The BACT limits for CO and VOC emissions are proposed below.

Proposed CO BACT Limit:

- Auxiliary Boiler: 0.037 lb/MMBtu (fuel gas mixture)
84.0 lb/MMscf (natural gas)
- Catalyst Regen Heater: 84.0 lb/MMscf (fuel gas mixture)
- Reactivation Heater: 84.0 lb/MMscf (fuel gas mixture)
84.0 lb/MMscf (natural gas)
- HGT Reactor Charge Heater: 84.0 lb/MMscf (fuel gas mixture)
84.0 lb/MMscf (natural gas)

Proposed VOC BACT Limit:

- Auxiliary Boiler: 0.004 lb/MMBtu (fuel gas mixture)
5.50 lb/MMscf (natural gas)
- Catalyst Regen Heater: 5.50 lb/MMscf (fuel gas mixture)
- Reactivation Heater: 5.50 lb/MMscf (fuel gas mixture)
5.50 lb/MMscf (natural gas)
- HGT Reactor Charge Heater: 5.50 lb/MMscf (fuel gas mixture)
5.50 lb/MMscf (natural gas)

4.4.3 SO₂ BACT Analysis for the Fired Heaters and Boiler

The auxiliary boiler and fired heaters oxidize any residual sulfur compounds present in the fuel gas mixture into SO₂. The control of SO₂ emissions is most directly associated with low-sulfur fuel.

Identify SO₂ Control Technologies

The following SO₂ control technologies were evaluated for the proposed process heaters.

Pre-Combustion Control

Lower Sulfur Fuels

Post-Combustion Control

Flue Gas Desulfurization

Evaluate Technical Feasibility

Low Sulfur Fuels

Potential SO₂ emissions are directly related to the sulfur content of fuels. The gas fed to the Methanol Unit requires less than 0.1 ppmvd, and therefore the SELEXOL[®] process in the AGR unit and sulfur beds will be used to achieve this low sulfur level. All fuel gas used throughout the plant is first desulfurized in the AGR unit, and therefore contains less than 0.1 ppmvd sulfur (expressed as H₂S). The concentration in the exhaust of each fired heater will be less than 0.2 ppmvd. Minimizing fuel sulfur content through the use of natural gas (startup only) or low sulfur fuel gas has been determined to be BACT for many combustion processes, including fired process heaters. Therefore, using low-sulfur-fuel is a technically feasible control technology.

Flue Gas Desulfurization

FGD is a post-combustion SO₂ control technology that reacts an alkaline solution with SO₂ in the exhaust gas. FGD systems are more readily applied to high SO₂ concentration gas streams, such as with a pulverized coal unit. FGD has not been applied to small process heaters due to the low SO₂ concentrations of exhaust streams associated with tail gas combustion. Therefore, FGD technology is not technically feasible for the proposed fired heaters.

Rank Control Technologies

The use of low-sulfur fuels is the only technically feasible SO₂ control technology identified for the proposed fired heaters.

Select SO₂ Best Available Control Technology

The use of low sulfur fuels (tail gas) is proposed as BACT for SO₂ emissions from the proposed auxiliary boilers and fired heaters. As emissions of SO₂ are negligible, BACT limits are not proposed for the auxiliary boiler and fired heaters.

4.4.4 Particulate Emissions BACT Analysis for the Fired Heaters and Boiler

Fuel quality and combustion efficiency are key drivers affecting the quantity and disposition of potential particulate emissions. In some processes, post-combustion control technologies can also be used to reduce particulate.

Identify Control Technologies

The following particulate emissions control technologies were evaluated for the proposed auxiliary boiler and fired process heaters.

Pre-Combustion Control

Clean Fuels

Good Combustion Practices

Post-Combustion Control

Electrostatic Precipitation

Baghouse

Evaluate Technical Feasibility**Clean Fuels**

Fuels containing ash have the potential to produce particulate matter emissions. Additionally, fuels containing sulfur have the potential to produce sulfur compounds that may form condensable particulate matter emissions. The fuel gas mixture consumed by the proposed auxiliary boilers and fired heaters will contain negligible amounts of particulate matter and is considered a low sulfur fuel. Therefore, the use of clean fuels is a technically feasible control technology for the process heaters.

Good Combustion Practice

The use of good combustion practice is a technically feasible technology that can minimize the potential particulate emissions associated with incomplete combustion.

Electrostatic Precipitation

ESP is a post-combustion particulate emissions control most readily applied to large volume gas streams containing high particulate concentrations. No examples have been found where an ESP has been applied to a syngas/tail gas fired process heater due to the reduced volume and minimal particulate concentration of the associated exhaust gas stream. Therefore, ESP is not technically feasible for the auxiliary boiler and proposed process heaters.

Baghouse

A baghouse is a post-combustion control technology that utilizes a fine mesh filter to remove particulate emissions primarily from large volume gas streams containing high particulate concentrations. No examples have been found where a baghouse has been applied to a syngas/tail gas fired process heater due to the reduced volume and minimal particulate

concentration of the associated exhaust gas stream. Therefore, baghouse technology is not technically feasible for the auxiliary boiler and proposed process heaters.

Rank Control Technologies

The use of clean fuels and good combustion practices are the only technically feasible control technologies identified.

Select Particulate Emissions Control Technology

The use of clean fuels and good combustion practices has been proposed as BACT. The proposed PM BACT limit is presented below.

Proposed PM BACT Limit:

Auxiliary Boiler: 0.005 lb/MMBtu (fuel gas mixture)

7.60 lb/MMscf (natural gas)

Catalyst Regen Heater: 7.60 lb/MMscf (fuel gas mixture)

Reactivation Heater: 7.60 lb/MMscf (fuel gas mixture)

7.60 lb/MMscf (natural gas)

HGT Reactor Charge Heater: 7.60 lb/MMscf (fuel gas mixture)

7.60 lb/MMscf (natural gas)

Please note that these emission limits were all calculated with emission factors from EPA's AP-42 "Compilation of Air Pollutant Emission Factors" document. AP-42 particulate emissions from fuel gas firing have been demonstrated to underestimate actual emissions in some cases. At this time, it cannot be determined if the particulate emissions presented here are underestimated for these process heaters based on the use of AP-42 factors. All heater particulate emission limits should be verified through stack testing, and the construction permit should be modified to reflect the more accurate emission factors obtained through testing.

4.4.5 Startup Emissions BACT Analysis for the Fired Heaters and Boiler

Fired heater startup emissions are quantified separately from normal operating emissions. During startup and upset conditions, natural gas may be used, although the fuel gas mix will still be used when available. To satisfy BACT during startup and upset operating conditions, the auxiliary boiler and fired heaters will be limited to 1,000 hours per year of natural gas firing for all startup operations including initial startup and other startup modes. The duration of the startups will also be minimized to the best extent possible for each unit. Alternatively, natural gas may be used as a backup fuel that will not increase the emissions over using fuel gas firing.

4.5 STORAGE TANK CONTROL TECHNOLOGY REVIEW

Eight gasoline product tanks are proposed for the facility, along with two methanol storage tanks, one "heavy gasoline" intermediate product tank, and one slop tank. Additionally, several smaller storage tanks and LPG storage bullet tanks are proposed. Table 4.3 lists all proposed storage tanks for the facility. VOC and HAP emissions from the storage tanks, with the exception of the closed-system LPG bullets, will occur as a result of headspace vapor displacement during filling

operations (working losses) and from diurnal temperature variations and solar heating cycles (breathing losses).

The proposed gasoline product, methanol, heavy gasoline, and slop storage tanks will be designed with internal floating roofs (IFRs), submerged fill, white exterior surfaces, and will meet NSPS Subpart Kb (Standards of Performance for Volatile Organic Liquid Storage Vessels (including Petroleum Liquid Storage Vessels) for Which Construction, Reconstruction, or Modification Commenced After July 23, 1984) requirements. The proposed smaller tanks will store water and low vapor-pressure chemicals and will be fixed roof design, with no IFRs. Because emissions from these smaller tanks will be insignificant, they are not addressed in this BACT analysis. Similarly, since the LPG bullets will be constructed as a closed system with no vents to atmosphere, they are not addressed in this analysis.

Identify VOC and HAP Control Technologies

The following VOC and HAP control technologies were evaluated for the proposed methanol, gasoline, and slop storage tanks.

1. Operate tanks under pressure, as closed systems.
2. Construct tanks with a fixed or dome roof, with vapor collection routed to fuel gas system or process system.
3. Construct tanks with a fixed or dome roof, with vapor collection routed to a control device.
4. Construct tanks with an external floating roof (EFR).
5. Construct tanks with an internal floating roof (IFR) in combination with a fixed roof.

Evaluate Technical Feasibility

Operate Storage Tank Under Pressure

Operating the storage tanks under pressure as closed systems is an inherently less-polluting process configuration because it eliminates working and breathing losses. However, this option is suitable only for materials that are gases at atmospheric pressure and temperature such as propane and butane. (Note, the proposed LPG storage tanks for the facility will be pressurized bullets, operating as closed systems.) Therefore, this option is not technically feasible for the liquid storage tanks under review.

Fixed or Dome Roof with Vapors Routed to Fuel Gas System or Process System

This option can also be considered to be an inherently less-polluting process configuration. An inert gas ‘blanket’ would be required for this option in order to ensure the tank vapor space remains outside of explosive limits. Design and operation of the gas blanket could present considerable engineering challenges, as the system and tanks must be designed and operated to prevent any under-pressure or over-pressure scenarios that could result in catastrophic tank failure. Generally, the practice of operating large storage tanks such as these (storing volatile liquid product) with a vapor space is not common due to the potential safety issues and the chance for an explosive atmosphere to be created at some point in the vapor system. The industry standard, from a safe operating perspective, for large gasoline and other volatile liquids is a floating roof.

For this control option, the vapor stream must be directed to a compatible fuel gas system or process system in order to protect plant operations and system integrity. Due to the inert gas blanket required as part of this option's design, no compatible fuel gas or process gas streams are available in the proposed facility to receive the vent stream. Based on this, in addition to the potential safety issues associated with operating a vapor system in these storage tanks, the option is considered technically infeasible.

Fixed or Dome Roof with Vapors Routed to a Control Device

This control option is very similar to the previous one, except that the vent stream would be routed to a control device, such as a thermal oxidizer, instead of a fuel gas or other process system. Similar safety issues are presented with this option as with the previous option, with regard to the vapor space in the tank and design/operation of the vapor system. However, this option is considered technically feasible, because a final destination for the vent stream is presented and available.

A certain amount of product would be "lost" to the vapor space with this option, as with the previous option. With a control device such as a thermal oxidizer, the "lost" product would not be recoverable. An advantage to the previous option is that "lost" product can be recovered through re-routing to a fuel gas or process system. Non-recoverable, lost product could present a significant economic disadvantage for this control option.

External Floating Roof (EFR) or Internal Floating Roof (IFR)

Floating roof technology is the prevalent emission control technology for large tanks storing volatile liquids. Both EFR and IFR technology provide for minimal product loss (i.e., emission prevention) as well as improved safety over fixed roof tanks. This option is technically feasible for the proposed storage tanks.

Rank Control Technologies

The three technically feasible control options are ranked as follows.

1. IFR, in combination with a fixed roof
2. Fixed or dome roof with vapors routed to a control device
3. EFR

All three technically feasible options will meet NSPS Subpart Kb requirements for VOC control. However, of the three technically feasible options, the EFR is considered to be the least effective for VOC and HAP emission control. An IFR, in combination with a fixed roof, provides better emission control for volatile liquids and is generally preferred over EFRs in similar applications.

Constructing the storage tanks with a fixed roof and a vapor collection system with the vent stream routed to a control device would also provide high control efficiency, but the option has a significant disadvantage in that operation of a thermal oxidizer will result in additional emissions from the combustion process (NO_x and CO). Based on this negative environmental impact, in addition to the safety concerns discussed earlier, this option is ranked second, below the IFR option.

Therefore, the option to construct the tanks with IFRs in combination with fixed roofs is considered the most effective control option.

Select Best Available Control Technology

An internal floating roof (IFR), in combination with a fixed roof, is proposed as BACT for the gasoline product, methanol, heavy gasoline, and slop product storage tanks. Table 4.3 presents detailed capacity and product data for each of the proposed storage tanks.

Table 4.3 – Storage Tanks Summary

Tank Name	Tank No.	No. of Tanks	Liquid Capacity per Tank (Gallons)	Roof Type	NSPS Kb Applicable
Methanol Tanks	<i>TBD</i>	2	6,341,984	IFR	Yes
Gasoline Product Tanks	<i>TBD</i>	8	6,341,984	IFR	Yes
Heavy Gasoline Tank ¹	<i>TBD</i>	1	4,763,841	IFR	Yes
Off-Spec Gasoline Tank	<i>TBD</i>	1	5,000	IFR	N/A (size)
Off-Spec Methanol Tank	<i>TBD</i>	1	5,000	IFR	N/A (size)
Slop Tank	<i>TBD</i>	1	7,000	IFR	N/A (size)
<i>Tanks with Insignificant Emission Rates</i>					
Gray Water Tank	03T-002	1	<i>TBD</i>	FR	No
Slurry Additive Tank	03T-003	1	<i>TBD</i>	FR	No
Mill Discharge Tank	01T-104	1	<i>TBD</i>	FR	No
Slurry Tank	01T-105	1	<i>TBD</i>	FR	No
Injector Coolant Tank	02T-001	1	<i>TBD</i>	FR	No
Settler	03T-001	1	<i>TBD</i>	FR	No
Filter Feed Tank	03T-004	1	<i>TBD</i>	FR	No
Filtrate Tank	03T-005	1	<i>TBD</i>	FR	No
Glycol Storage Tank	<i>TBD</i>	1	4,000	FR	No
Sulfur Storage	<i>TBD</i>	2	5,000	FR	No

1. “Heavy” gasoline is estimated to have RVP of 3-5 psia.

4.6 MATERIAL HANDLING CONTROL TECHNOLOGY REVIEW

The material handling conveyer will be fully enclosed to prevent wind blown fugitive dust. Transfer points will be controlled with fogger and passive engineering design at transfer points. This technology has been successfully used in other coal applications in Wyoming. On the MBFP Facility site there will be covered coal storage for approximately 8 hours of use.

Additionally, the coal handling operations will be subject to and will comply with the NSPS for Coal Preparation Plants (Subpart Y), as applicable.

4.7 PROCESS FUGITIVE EMISSIONS CONTROL TECHNOLOGY REVIEW

Fugitive VOC, HAP, and hydrogen sulfide (H₂S) emissions will be generated from potential leaking process equipment, primarily downstream of the coal preparation and gasification portions of the facility (SELEXOL acid gas removal, CO₂ recovery, sulfur recovery, methanol synthesis, gasoline synthesis, etc.). Additionally, fugitive ammonia emissions will be generated from potential equipment leaks in the ammonia storage and feed equipment used for the proposed SCR system (turbine NO_x control). Note that the number of piping components in ammonia service will be very small in comparison to the number of other potential leaking components at the proposed facility.

VOC and HAP emissions from equipment leaks were estimated using fugitive leak emission factors from EPA Document No. EPA-453/R-95-017, November 1995 (“Protocol for Equipment Leak Emission Estimates”). Control efficiencies reflecting a monthly leak detection program were used in the calculation, assuming a leak definition value of 500 ppmv for valves and connectors in VOC service and 2,000 ppmv for pumps in VOC service. Total facility estimated potential VOC emissions from equipment leaks are 60 tons per year, and total facility estimated potential HAP emissions from equipment leaks are 16 tons per year.

Identify VOC and HAP Control Technologies

The only available control technology for comprehensively addressing equipment leak fugitive emissions is a structured Leak Detection and Repair (LDAR) program in which certain piping components and equipment are routinely inspected for leaks, and components found to be leaking in excess of stated thresholds are repaired in a timely manner. The effect of a well-implemented LDAR program is reduced VOC and HAP emission rates due to improved maintenance and repair. LDAR programs are established as BACT in many recent RBLC determinations.

Select Best Available Control Technology

A formal, structured LDAR program is proposed as BACT for components in VOC service. Records will be maintained for all leak inspections and necessary repair work. Additionally, audio/visual/olfactory (AVO) detection is proposed for equipment potentially leaking hydrogen sulfide or ammonia. Both chemicals have low odor thresholds, and plant personnel should be able to easily detect any leaking components under routine plant operations. Leaking equipment discovered through AVO detection will be repaired in an expeditious manner in order to reduce emissions and remove potential safety issues.

4.8 SULFUR RECOVERY UNIT (SRU) CONTROL TECHNOLOGY REVIEW

The Sulfur Recovery Unit (SRU) is designed to process acid gas streams from the SELEXOL[®] acid gas removal system and Plant process into an elemental sulfur product. SRU tail gas is typically directed to a tail gas treatment unit designed to remove SO₂ from the tail gas before the tail gas is vented to atmosphere. Typical SRU design also incorporates a thermal oxidizer, also called a tail gas incinerator, to provide efficient destruction of the tail gas stream after it exits the tail gas treatment unit. In the event of a malfunction with the SRU or tail gas incinerator, or during times of cold startup, the tail gas stream may be temporarily diverted to a flare in lieu of

the tail gas incinerator. The pollutant of concern for SRUs is SO₂, although emissions of other criteria pollutants may result from the combustion process.

Identify SO₂ Emission Control Technologies

Potential control technologies for the SRU tail gas stream during times of normal operation include the following:

1. LP Flare
2. Thermal Oxidizer (Tail Gas Incinerator)
3. Re-routing Tail Gas to Process

Evaluate Technical Feasibility

The LP Flare is proposed as a low-pressure flare for the facility and will intermittently receive vent streams from various processes throughout the facility, in addition to any vents from the SRU. Control efficiency for the flare is estimated at 98%.

As mentioned earlier, a tail gas incinerator is a typical control device for SRUs and would be dedicated to the SRU tail gas, with a supplemental fuel gas or natural gas. Control efficiency is estimated between 98-99%.

Re-routing the tail gas back to the process would involve routing the tail gas to a point upstream of the H₂S absorption tower in the SELEXOL[®] acid gas removal process and would allow the stream to be reprocessed rather than being combusted and destroyed. This option results in no emissions during normal operation since nothing is emitted to the atmosphere, and therefore it has 100% control efficiency.

For the proposed Plant, all three possible control options are technically feasible during times of normal operation. However, during times of startup, shutdown, or malfunction (SSM), neither the thermal oxidizer nor re-routing the tail gas stream are considered technically feasible options, due to the variability of gas stream flowrate and composition during these times. The LP Flare is the only technically feasible option for SSM conditions.

Select Best Available Control Technology

Of the three technically feasible control options, re-routing the tail gas back into the process at an upstream point provides 100% control, and is therefore ranked higher than the LP Flare or tail gas incinerator options. BACT is chosen to be re-routing the tail gas stream during times of normal operation, with the LP Flare employed only as needed during times of SSM operations.

4.9 CARBON DIOXIDE VENT STACK (STARTUP OPERATIONS ONLY)

During initial startup operations and subsequent warm start operations, off-specification CO₂ will be vented to the atmosphere. This exhaust will contain some small amount of CO and VOC (primarily COS). Elements have been incorporated in the design and operating procedures to minimize the frequency and duration of venting this gas stream to the atmosphere. The facility is being designed so that this venting will not occur during load transitions during normal

operations. Another factor is that this carbon dioxide stream is a product. Design elements that maximize the reliability of the carbon dioxide stream and minimize startup, shutdown, and malfunction periods will reduce the frequency and duration of venting events. The venting is only anticipated for a few days during initial startup (approximately 250 hrs/yr for the first year). Since the plant will be started up at reduced load, the venting will be at a reduced rate (approximately 25% of the normal process stream flow rate). Venting is anticipated for only a few hours for subsequent warm starts, not to exceed 50 hrs/yr. Again, the venting would be at a reduced load (approximately 50% of the normal process stream flow rate).

Catalytic oxidation is not technically feasible based on the low temperature of the vent stream, approximately 100°F. Based on the temperature and large flow rate, an extremely large amount of energy would be necessary to oxidize the CO with a thermal oxidizer, and may not be possible due to the size of the stream, low temperature, and high concentration of CO₂ in the stream. RBLC ID WY-0042 contained a process identified as “Vent, CO₂ Product” where incineration was not feasible due to CO₂ concentration in the gas. RBLC ID WY-0056 contained a process identified as “CO₂ Product Vent, Train III” that also vented uncontrolled.

The total annual proposed CO emissions to be permitted from the CO₂ stack are 275 tpy for the initial year of operation. Subsequent years will be limited to 74 tpy of CO. The proposed VOC emissions are 0.02 tpy for the first year and 0.01 tpy for subsequent years. Based on the limited operating time and resultant emissions, further controls are not warranted. Thus, an optimized process design is considered BACT for this process vent.

4.10 GASIFIER PREHEATING CONTROL TECHNOLOGY REVIEW (STARTUP OPERATIONS ONLY)

During the initial startup operations, or if new refractory is in place in a gasifier, a designated 21 MMBtu/hr natural gas burner is used to preheat the refractory lining prior to commencing tail gas production. Potential emissions from the natural gas combustion in the gasifiers is exhausted from a preheat vent located on each gasifier. The primary potential emissions from the gasifier preheat vents are NO_x and CO. Each gasifier preheat vent has a potential to emit less than 1 ton per year of NO_x and CO as discussed in the emission inventory. Emissions of VOC and particulate will also be relatively small based on the short operating time, approximately one week for each gasifier, for initial startup (and refractory replacement) only. Subsequent startup operations will be warm starts and will not include this step. The expected operating hours for the gasifier preheaters are 500 hours per year per heater, for a total of 2,500 hours per year. Good combustion controls that optimize burner efficiency will minimize potential NO_x, CO, VOC and particulate emissions. Because a low-sulfur-fuel (natural gas) is being used for preheating, the potential emissions of SO₂ will also be small.

The use of a low-sulfur-fuel, restricted operating conditions, and good combustion practices are proposed as BACT for each of the five (5) gasifier preheat burners. Table 4.4 shows the proposed BACT emission rates for each gasifier preheater.

Table 4.4 – Gasifier Preheater BACT Analysis Summary

Pollutant	Proposed BACT	Proposed BACT Emission Limits (emission limits are per gasifier preheater)
NO _x	Low Sulfur Fuel Good Combustion Practices Restricted Operation (startup only)	NO _x Limit: 0.26 tpy
SO ₂		SO ₂ Limit: <0.01 tpy
CO		CO Limit: 0.43 tpy
VOC		VOC Limit: 0.03 tpy
PM		Particulate Limit: 0.04 tpy (PM ₁₀ - filterable)

4.11 BLACK-START GENERATOR CONTROL TECHNOLOGY REVIEW (STARTUP OPERATIONS ONLY)

The proposed Plant will include three (3) 1.6 MW natural gas fired generators for use during startup. The generators will be used for commissioning and initial startup. Key utility systems such as instrument air, water supply and purification, firewater, and nitrogen will be made operational prior to initiating the startup sequence for the process. It is especially important that the flare system be ready for service before any flammable gas is present. Once critical utilities are in service, one of the three gas turbines is started on natural gas. This will produce enough power to displace the Black-Start generators. The primary potential emissions from the Black-Start generators are NO_x and CO. Emissions of VOC and particulate will also be relatively small based on the short operating time and infrequent use (only initial startup and commissioning and upset conditions). The maximum hours per year proposed for the Black-Start generators are 360. Subsequent startup operations will be warm starts and are not anticipated to require firing of the Black-Start generators. Good combustion controls that optimize combustion efficiency will minimize potential NO_x, CO, VOC and particulate emissions. Because natural gas is being used, the potential emissions of SO₂ will also be small. Additionally, these natural gas fired generators will also be subject to and will comply with the NSPS for Stationary Compression Ignition Combustion Engines (Subpart IIII), as applicable.

The use of a natural gas, restricted operating conditions, and good combustion practices are proposed as BACT for the three Black-Start generators. Table 4.5 shows the proposed BACT emission rates for each Black-Start generator.

Table 4.5 – Black-Start Generator BACT Analysis Summary

Pollutant	Proposed BACT	Proposed BACT Emission Limits (emission limits are per generator)
NO _x	Natural Gas Fired Good Combustion Practices Restricted Operation (initial startup only)	NO _x Limit: 0.80 tpy
SO ₂		SO ₂ Limit: <0.01 tpy
CO		CO Limit: 1.93 tpy
VOC		VOC Limit: 0.72 tpy
PM		Particulate Limit: 0.0002 tpy (PM ₁₀ - filterable)

4.12 FIREWATER PUMP CONTROL TECHNOLOGY REVIEW (BACKUP OPERATIONS ONLY)

The Firewater Pump is used to support emergency operations at the proposed facility. Potential emissions from the Firewater Pump are controlled by restricting the hours of operation, using good combustion practices, and using ultra-low-sulfur-fuel. Operation of the emergency Firewater Pump will be limited to emergency operating scenarios or required testing by the manufacturer. The Firewater Pump will operate no more than 500 hours per year. The design will incorporate manufacturer specifications that maximize the combustion efficiency and minimize potential emissions. Based on the limited operating time and resultant emissions, further controls are not warranted. This diesel-fired pump will also be subject to and will comply with the NSPS for Stationary Compression Ignition Combustion Engines (Subpart IIII), as applicable. Assuming a displacement of <30 liters per cylinder, if model year is 2009 or after NSPS IIII would apply.

Additionally, ultra-low-sulfur diesel fuel containing less than or equal to 15 ppm sulfur will be used. Good combustion practices, restricted annual operations, and ultra-low-sulfur fuel are proposed as BACT. Table 4.6 shows the proposed BACT emission rates for the emergency Firewater Pump.

Table 4.6 – Emergency Firewater Pump BACT Analysis Summary

Pollutant	Proposed BACT	Proposed BACT Emission Limits
NO _x	Restricted Operation (<500 hr/yr) Low Sulfur Fuel Good Combustion Practices	NO _x Limit: 1.51 tpy
SO ₂		SO ₂ Limit: <0.01 tpy
CO		CO Limit: 0.09 tpy
VOC		VOC Limit: 0.34 tpy
PM		Particulate Limit: 0.02 tpy (PM ₁₀ -filterable)

4.13 MERCURY EMISSION REDUCTION

Syngas exiting the gasifiers contains some mercury. This mercury must be removed before the syngas enters the Methanol Synthesis Unit. Two mercury guard beds will be operated at the Plant and are expected to achieve 99.98% removal of mercury. The cost of the planned mercury removal system is estimated to be \$235,164 per ton of mercury removed, as shown in Appendix G.

MBFP requests a mercury emission rate of 0.02 µg/Nm³, which results in total facility mercury emissions of no more than 1.3×10⁻⁴ tpy (0.26 lb/yr). At an electrical generation rate of 66 MW per turbine (design), this results in emissions of 1.5×10⁻⁷ lb/MWh per turbine, which is significantly less than NSPS requirements in 40 CFR Part 60, Subpart Da mandating a mercury emission limit of 20×10⁻⁶ lb/MWh for affected facilities.

4.14 MINE LONG-TERM COAL STORAGE

The Mine will have two coal storage areas. The first is a 300,000-ton dead storage (emergency stockpile) and the second is a 300,000 ton active storage area. The emergency stockpile will be compacted and sealed to prevent wind erosion and spontaneous combustion. Since there will be no particulate emissions associated with this stockpile once it is constructed, it has not been included in this analysis.

Three scenarios were evaluated for the active coal storage. There are:

1. Stacking tubes located on the surface
2. Stacking tubes located in the pit excavated
3. Covered slot storage

The BACT analysis for the active storage for performed by IML Air Science (Sheridan, WY). The complete analysis is in Appendix F.

Identify Particulate Emission Control Technologies

The first two scenarios differ in the placement of the stacking tubes. Scenario 2 places the stacking tube on the pit floor on the previously mined surface coal, with the excavated spoils placed in a large berm on the west and north sides of the pit. This configuration is intended to reduce storage pile erosion and resulting PM₁₀ emissions, by sheltering the pile from prevailing winds.

The third scenario would be to construct a covered storage area (slot storage or coal barn).

Evaluate Technical Feasibility

The control strategies described above as Scenarios 2 and 3 have been implemented in Wyoming and in other parts of the country. Therefore, both are considered technically feasible.

Rank Control Technologies

The covered storage (Scenario 3) would result is zero particulate emissions (100% control effectiveness). The sheltered stacking tubes have an estimated 23% control effectiveness on the particulate emissions resulting in annual emissions of 60 tpy (Scenario 1 was estimated to be approximately 78 tpy).

An economic analysis was conducted on the incremental control cost between Scenarios 2 and 3. The incremental control cost between the two scenarios is \$6,902 per ton removed.

Evaluate Control Technologies

Although the covered storage has a greater control effectiveness, the economic analysis shows the cost for the scenario is not financially viable.

Select Particulate Emissions Control Technology

Due to the negative economic impact of the covered storage, the next most effective control option (sheltered stacking tubes) was selected.

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This section analyzes the state and federal air quality regulations that are potentially applicable to the Plant and Mine. This regulatory summary is not intended to provide a detailed explanation of all compliance requirements associated with applicable regulations.

5.1 WYOMING AIR QUALITY REGULATIONS

This section discusses the relevant Wyoming Air Quality Standards and Regulations (WAQS&R). MBFP will comply with all applicable requirements within WAQS&R.

5.1.1 Chapter 2 Ambient Standards

The Wyoming Ambient Standards set limits deemed necessary to protect public health and welfare. Table 5.1 compares the Wyoming Ambient Standards to the National Ambient Air Quality Standards (NAAQS). For many pollutants, Wyoming's ambient air quality standards are identical to national standards. However, the state has set standards for some additional pollutants.

With regard to the NAAQS, the Plant would be located within an area that is designated as attainment (or unclassifiable) for each criteria pollutant.

Table 5.1 – Ambient Air Quality Standards

Pollutant	Averaging Time	WAQS&R ($\mu\text{g}/\text{m}^3$)	NAAQS ($\mu\text{g}/\text{m}^3$)
PM ₁₀	24-hour	150 ^b	150 ^a
	Annual	50 ^f	---
PM _{2.5}	24-hour	65 ^d	35 ^c
	Annual	15 ^f	15 ^e
NO ₂	Annual	100 ^f	100 ^f
SO ₂	3-hour	1,300 ^b	---
	24-hour	260 ^b	365 ^b
	Annual	60 ^f	80 ^b
CO	1-hour	40,000 ^b	40,000 ^b
	8-hour	10,000 ^b	10,000 ^b
Ozone	1-hour	---	235 ^h
	8-hour	157 ^g	157 ^g
H ₂ S	1/2-hour	70 ⁱ (40 ^j)	70
SO ₃ (Suspended sulfates)	30-day	250 mg/100 cm ² /day	---
	Annual	500 mg/100 cm ² /day	---

Table 5.1 – Ambient Air Quality Standards

Pollutant	Averaging Time	WAQS&R ($\mu\text{g}/\text{m}^3$)	NAAQS ($\mu\text{g}/\text{m}^3$)
Fluorides	12-hour	3.0	---
	24-hour	1.8	---
	7-day	0.5	---
	30-day	0.4	---
Lead	Quarterly	1.5	1.5

^a Not to be exceeded more than once per year on average over 3 years.

^b Not more than one exceedance per year.

^c Not to exceed the 3-year average of the 98th percentile of 24-hour concentrations.

^d Not to exceed the 98th percentile of 24-hour concentrations.

^e Not to exceed the 3-year average of the weighted annual mean.

^f Not to exceed the annual mean.

^g Not to exceed the 3-year average of the fourth-highest daily maximum 8-hour average ozone concentrations.

^h Applies only to limited areas (not applicable to this project).

ⁱ Not to be exceeded more than 2 times per year.

^j Not to be exceeded more than 2 times per year in any 5 consecutive days.

5.1.2 Chapter 3 General Emission Standards

WAQS&R emission standards within Chapter 3 set forth requirements that are generally applicable to a wide variety of facilities. Applicable standards are summarized below.

5.1.2.1 Section 2 Particulate Matter

Opacity and fugitive dust are regulated under WAQS&R Chapter 3, Section 2. As a new facility, each new stationary source at the Plant and Mine may not exceed 20 percent opacity [WAQS&R Chapter 3, §2(a)]. However, brief exceedances of the 20 percent opacity limit are allowed in certain cases. An opacity of up to 40 percent is allowed for a period or periods aggregating to not more than 6 minutes in any hour [WAQS&R Chapter 3, §2(e)].

The firewater pump diesel engine would be subject to a 30 percent opacity limit except during periods not exceeding 10 consecutive seconds. This limit generally does not apply to a reasonable period of warmup following a cold start or when undergoing repairs and adjustment following a malfunction [WAQS&R Chapter 3, §2(d)].

Particulate emissions from process sources are limited by WAQS&R Chapter 3, §2(g). Coal handling, primarily movement of coal from the coal storage area, will be subject to this standard, which allows emissions up to the limit calculated by the following equation:

$$E = 17.31 P^{(0.16)}$$

Where:

E = Emissions (lb/hr)

P = Process weight (ton/hr)

Based on 8,000 TPD (333.3 ton/hr) of dry coal feed, the emission limit would be 43.84 lb/hr. Particulate emissions from coal handling will be far less than this due to the fogging system.

Fugitive dust from coal handling and storage at the Mine will be controlled by using a fogging system in order to comply with emission standards for material handling and storage at WAQS&R Chapter 3, §2(f)(ii). The IGL Plant will have about 8 hours of covered onsite storage for coal.

During construction of the Facility and associated portal areas, steps to minimize fugitive dust must be taken [WAQS&R Chapter 3, §2(f)(i)]. MBFP will require construction contractors to use control measures, such as frequent watering and/or chemical stabilization, on an as-needed basis to reduce fugitive dust emissions. In addition, contractors will be instructed to promptly remove mud or dirt that is tracked onto paved roadways [WAQS&R Chapter 3, §2(f)(i)].

5.1.2.2 Section 3 Nitrogen Oxides

The Plant will construct and operate several new gas fired fuel burning sources, such as the combustion turbines, boiler, and heaters. Under WAQS&R Chapter 3, §3(a)(i), NO_x emissions from new gas fired fuel-burning equipment calculated as nitrogen dioxide (NO₂) may not exceed 0.20 lb/MMBtu of heat input.

NO_x emissions (calculated as NO₂) from the fuel-oil burning Firewater Pump engine will be limited to 0.30 lb/MMBtu because it will have a heat input greater than 1.0 MMBtu/hr [WAQS&R Chapter 3, §3].

Internal combustion engines having a heat input of less than 200 MMBtu/hr are exempt from the NO_x emission limits given above.

5.1.2.3 Section 4 Sulfur Oxides

Sulfur oxides (SO_x) emission limits apply only to fuel burning equipment that is fueled with coal or oil. Consequently, the Firewater Pump is the only equipment subject to these standards. The Firewater Pump will be required to meet a 3-hour limit of 0.8 lb/MMBtu and a 30-day average of 0.8 lb/MMBtu [WAQS&R Chapter 3, §4(b)].

5.1.2.4 Section 5 Carbon Monoxide

Wyoming's air quality regulations do not include specific CO emission limits for stationary sources. There is, however, a general duty to prevent any exceedance of CO ambient standards [WAQS&R Chapter 3, §5]. Modeling results provided in Section 6 demonstrate that the Plant will meet this requirement.

5.1.2.5 Section 6 Volatile Organic Compounds

VOC emissions shall be limited through the application of BACT [WAQS&R Chapter 3, §6(b)]. In some cases, WDEQ regulates VOC emissions by mandating use of a flare. When a flare is required to control of VOC emissions from vapor blowdown, emergency relief systems, or VOC emissions generated from storage or processing operations, the flare shall not exceed a 20%

opacity emission standard [WAQS&R Chapter 3, §6(b)]. In addition, the flare must be a smokeless flare and must have either an automatic igniter or a continuous pilot.

5.1.2.6 Section 7 Hydrogen Sulfide

Some Plant process streams contain H₂S and will be subject to WAQS&R Chapter 3, §7. Any exit process gas stream containing H₂S that is discharged to the atmosphere must be vented, incinerated, flared or otherwise disposed of such that ambient SO₂ and H₂S standards are not exceeded. Process streams containing H₂S are treated within the Plant process to remove the sulfur. However, in the event of a malfunction, a stream containing H₂S could be vented to a flare.

5.1.2.7 Section 8 Asbestos Activities

As a new facility, the Plant will minimize use of asbestos during facility construction. Furthermore, facility personnel are unlikely to remove asbestos-containing materials from the premises in the near future. However, activities that disturb asbestos would likely be subject to extensive compliance requirements found in WAQS&R Chapter 3, §8.

5.1.3 Chapter 6 Permitting Requirements

Section 2. Best Available Control Technology (BACT)

Per the WAQS&R, Chapter 6, §2(c)(v), no permit to construct will be issued until it is demonstrated that BACT will be utilized, with consideration of the technical practicability and economic reasonableness of reducing or eliminating the proposed facility's emissions. In accordance with this requirement, and those imposed by the PSD Program discussed below, BACT analyses for all emission sources are presented in Section Four of this application.

Section 3. Operating Permits

Potential emissions from the Plant and Mine exceed the 100-tpy threshold for triggering operating permit requirements under Chapter 6, Section 3. These regulations implement the Title V Operating Permit Program required by federal law. Per the timeline established in the WAQS&R, Chapter 6, §3(c), an application for an operating permit will be submitted within twelve months of facility startup.

Section 4. Prevention of Significant Deterioration

Potential emissions from the Plant and Mine exceed the 100-tpy threshold for triggering PSD permitting. Therefore, extensive provisions within WAQS&R Chapter 6, Section 4 will apply to the facility. This permit application process, associated modeling, and installation and operation of BACT will satisfy PSD compliance requirements applicable to construction and initial operation of the facility. When facility or operational modifications are planned, PSD review may be required.

Section 5. NESHAP Source Permits

Potential emissions from the Plant and Mine exceed the 25-tpy aggregate HAP threshold for triggering major source status under the NESHAP program, and the Plant is subject to several NESHAP standards including the 40 CFR Part 63, Subpart ZZZZ Reciprocating Internal Combustion Engine NESHAP and the Subpart DDDDD Industrial-Commercial-Institutional Steam Generating Unit NESHAP. Consequently, MBFP is also subject to WDEQ's permitting requirements for construction and modification of NESHAP sources, which are codified in WAQS&R, Chapter 6, Section 5. These regulations specify requirements for submitting pre-construction permit applications and providing notifications to the WDEQ, including a notification of compliance status.

This permit application satisfies the preconstruction permitting requirements of WAQS&R, Chapter 6, Section 5. In addition to other information submitted in this application, the following construction and operation schedule information specifically requested in Chapter 6, §5(a)(iii)(A)(II)(5-7) is provided below.

- Expected construction commencement date: As soon as air quality permit is issued.
- Expected construction completion date: July 1, 2012
- Expected initial startup date: July 1, 2012

5.1.4 Chapter 7 Monitoring Regulations

Some emission units at the Plant will be subject to Compliance Assurance Monitoring (CAM) requirements in WAQS&R Chapter 7, Section 3. These regulations are based on the USEPA 40 CFR Part 64 CAM regulations. CAM requirements generally apply to each emission unit that meets all of the following criteria (with some exceptions).

- The emission unit is located at a facility that is subject to the Title V operating permit program.
- The emission unit uses a control device to achieve compliance with an emission limit and whose pre-controlled emission levels exceed major source thresholds under the Title V operating permit program.
- The unit is not subject to a New Source Performance Standard (NSPS) or a National Emissions Standard for Hazardous Air Pollutants (NESHAP) standard that was promulgated after November 15, 1990.

If the facility is subject to CAM, the affected emission units will be subject to additional monitoring, recordkeeping, and reporting requirements. In addition, the facility must prepare a CAM Plan for each affected unit. A thorough CAM applicability review and proposed CAM Plans will be submitted with the initial operating permit application.

5.2 FEDERAL REGULATIONS

The following discussion summarizes federal air quality regulations that are potentially applicable to the Plant. Due to the unique processes used by this facility, it does not fall into an

industry-specific NSPS or NESHAP. However, some equipment at the facility will be subject to NSPS or NESHAP standards.

5.2.1 New Source Performance Standards (NSPS)

Subpart A: NSPS General Provisions

Subpart A identifies a number of monitoring, recordkeeping, and notification requirements that generally apply to all NSPS Subparts. Additionally, Subpart A specifies that performance (source) tests must be conducted within 60 days of achieving the maximum production rate at which the source will be operated, but not later than 180 days after initial startup. Subpart A will apply in conjunction with any other applicable NSPS Subpart, unless otherwise noted in the specific NSPS.

Subpart Da Electric Utility Steam Generating Unit NSPS

The combustion turbines and HRSGs will not be subject to the Electric Utility Steam Generating Unit NSPS because the facility will not export power for sale. The facility is not an “electric steam generating unit,” as defined in §60.41Da, which is the key applicability criteria for 40 CFR Part 60 Subpart Da.

Subpart Db Industrial-Commercial-Institutional Steam Generating Unit NSPS

The Auxiliary Boiler, which has a heat input of 66 MMBtu/hr, will be subject to Subpart Db emission limits for NO_x and PM.

Subpart J Petroleum Refinery NSPS

As mentioned in Section One, the Plant is classified as a Crude Petroleum and Natural Gas facility (1311) that produces gas and hydrocarbon liquids through gasification. The minor or support activity is underground mining of bituminous coal (1222).

Although the facility produces gasoline, it does not do so using a refining process. Therefore, it is not subject to the Petroleum Refinery NSPS (40 CFR Part 60, Subpart J). The Plant does not meet the regulatory definition of a “petroleum refinery” because it does not engage in “. . . producing gasoline, kerosene, distillate fuel oils, residual fuel oils, lubricants, or other products through distillation of petroleum or through redistillation, cracking or reforming of unfinished petroleum derivatives [§60.2].”

Subpart Kb Storage Vessels for Petroleum Liquids NSPS

Eleven tanks, listed in Table 5.2, at the Plant are expected to be subject to the petroleum storage vessel NSPS due to their large size and volatile contents. Subpart Kb regulations set tank design and operation requirements, as well as ongoing inspection requirements. The planned IFR tank design will meet Subpart Kb requirements. Plant personnel will comply with tank inspection, repair, and recordkeeping and recording requirements.

Table 5.2 – Subpart Kb Tanks List

Tank Name	Tank	Number of Tanks	Operating Temperature (°F)	Vapor Pressure At Operational Temperature (psia)	Liquid Capacity (Gallons)	Roof Type
Methanol Tanks	TBD	2	45	0.96	6,341,984	IFR
Gasoline Product Tanks	TBD	8	45	4.14	6,341,984	IFR
Heavy Gasoline Tank ¹	TBD	1	45	2.25	4,763,841	IFR

1. “Heavy” gasoline is estimated to have RVP of 3-5 psia.

Subpart Y Coal Preparation Plant NSPS

Under 40 CFR Part 60, Subpart Y, coal transfer, crushing, and drying activities are subject to particulate matter emission limits. Specifically, emissions from coal conveying equipment may not exceed 20 percent opacity. Use of fully covered conveyors and fogging of transfer points at the Plant should maintain compliance with Subpart Y particulate emission limits and opacity standards.

Subpart VV Equipment Leaks in the SOCOMI Industry NSPS

The Plant does not meet the definition of a facility that is part of the Synthetic Organic Chemical Manufacturing Industry (SOCMI). Consequently, the Plant is not subject to this regulation.

Subpart IIII Stationary Compression Ignition Internal Combustion Engine NSPS

The diesel Firewater Pump will be subject to the compression ignition (diesel) engine NSPS. Compliance with this regulation is relatively simple for engine owners who purchase an engine that is certified by the engine manufacturer to meet new engine standards. MBFP will likely purchase a 2008 or later model year engine and will comply with this rule.

Subpart JJJJ Stationary Spark Ignition Internal Combustion Engine NSPS

The three Black-Start Generators will be subject to the spark ignition engine NSPS. In addition to purchasing engines that are certified by the engine manufacturer to meet the required new engine standards, MBFP will comply with performance testing, maintenance, and recordkeeping requirements and operate the engines in accordance with good air pollution control practices to minimize emissions. MBFP will conduct initial performance tests and, due to the limited usage of these units, will repeat performance tests every three years.

Subpart KKKK Stationary Combustion Turbines NSPS

The combustion turbines will be subject to NSPS codified in 40 CFR Part 60, Subpart KKKK. Affected units will include the three combustion turbines because they each have a heat input at peak load of more than 10 MMBtu/hr and will commence construction after February 18, 2005 [§60.4305(a)].

The combustion turbines will burn a mixture of fuel gas, LPG, and natural gas. Since more than 50 percent of the mixture will be natural gas, the turbines will be deemed to be firing natural gas [§60.4325]. Therefore, the NO_x emission limit will be based on a new turbine with a heat input of between 50 and 850 MMBtu/hr firing natural gas fuel. The applicable NO_x limit is 25 ppm (corrected to 15 percent oxygen) or 1.2 lb/MWh [40 CFR Part 60, Subpart KKKK, Table 1]. The turbines can meet the SO₂ compliance requirements by burning fuels with potential emissions of less than 0.060 lb SO₂/MMBtu [§60.4330(a)(2)]. Extensive monitoring, recordkeeping, and reporting are required by the rule. Because the combustion turbines will be subject to this recent NSPS, they will not be subject to CAM requirements.

5.2.2 National Emissions Standards for Hazardous Air Pollutants (NESHAP)

The Plant will be a major source of HAPs. Consequently, it may be subject to a variety of NESHAP regulations. The following discussion identifies NESHAPs that are potentially applicable to the facility.

Subpart ZZZZ Reciprocating Internal Combustion Engine NESHAP

Subpart ZZZZ within 40 CFR Part 63, will apply to all reciprocating internal combustion engines (RICE) at the Plant that have a site rating of more than 500 brake horsepower. The three Black-Start Generators, each nominally rated at 2,889 horsepower, will be subject to rule. However, many of the compliance requirements within Subpart ZZZZ may not apply to these units, depending on their use. They may qualify as “emergency use RICE” or as “limited use RICE,” especially if they are used less than the amount of time assumed for emission estimation purposes in this permit application (360 hr/yr, each).

Subpart DDDDD Industrial-Commercial-Institutional Steam Generating Unit NESHAP

The Industrial-Commercial-Institutional Steam Generating NESHAP (40 CFR Part 63, Subpart DDDDD) is currently being implemented by the WDEQ via provisions in WAQS&R, Chapter 3, §3(b). Although federal implementation of this NESHAP has been vacated by a federal court decision, the WDEQ continues to enforce this NESHAP.

Regulatory requirements depend on the classification of each boiler and process heater at the Plant. Proposed equipment at the Plant will likely be classified as follows.

- New small gaseous fuel equipment: HGT Reactor Charge Heater
- New large gaseous fuel equipment: Auxiliary Boiler, Catalyst Regenerator, and Reactivation Heater

Based on these classifications, the HGT Reactor Charge Heater will be subject only to initial notification requirements. In contrast, the large gaseous fuel equipment will be subject to a CO emission limit of 400 ppmv (dry basis, corrected to 3 percent oxygen). Because each of the large gaseous fuel emission units at the Plant has a maximum heat input rate of less than 100 MMBtu/hr, installation of a continuous emission monitoring system (CEMS) will not be required. MBFP will comply with all applicable Subpart DDDDD notification, performance testing, recordkeeping, and reporting requirements.

5.2.3 Chemical Accident Prevention Provisions

The Chemical Accident Prevention Provisions in 40 CFR Part 68 set forth requirements concerning the prevention of accidental releases. All facilities with extremely hazardous substances have a “general duty” to prevent accidental releases. Consequently, the Plant must design and maintain a safe facility, including taking steps to prevent releases and minimizing the consequences of any releases that do occur.

In addition, a facility that has more than a threshold quantity of a regulated substance listed in §68.130 may be subject to a variety of compliance requirements in Part 68. Guidance on how to determine if a threshold quantity exists and exceptions for certain types of facilities, processes, and materials are provided in §68.115. For example, regulated substances in gasoline need not be considered when determining if a threshold quantity exists in a process. Thus, the gasoline in the MTG process and product storage tanks will not be included in the applicability determination. The proposed methanol tanks also will not be considered in the applicability determination because methanol is not on the list of regulated sources.

With the exception of H₂S, the proposed facility will not store or use any ammonia, chlorine, methyl mercaptan, or other chemicals included as “toxic substances” in §68.130. However, several processes will contain a mixture of H₂S and/or substances listed as “flammable substances” at §68.130 (methane, ethane, propane, etc.) with concentrations high enough to possibly qualify the entire process stream, per §68.115(b)(1) and (2). As a result, this regulation may apply to some processes at the Plant if the process in question (as defined at §68.3) contains more than a threshold quantity of the listed substance. Prior to beginning operation, MBFP will determine whether it is subject to Part 68 regulations and, if necessary, prepare a Risk Management Plan for the Plant.

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6.1 NEAR FIELD MODELING BACKGROUND

To assess likely near field air quality impacts, a dispersion modeling analysis was completed for areas within 10 km (near field) of the proposed facility. The analysis was completed in accordance with a protocol approved by WDEQ (05 March 2007). The air quality dispersion modeling analysis used the USEPA-approved AERMOD suite of programs including AERMOD (version 07026), AERMAP (version 06341) and AERMET (version 06341).

The analysis included:

- 1 Determination of emission inventory source characteristics;
- 2 Development of an appropriate receptor grid, beginning at the ambient air boundary, with digital elevation model (DEM) supplied terrain heights calculated using AERMAP;
- 3 Determination of applicable direction-specific downwash parameters using the Building Profile Input Program (BPIP) PRIME (BPIPPRIME) for the many tanks and other structures associated with the project sources;
- 4 Processing of local and representative surface and upper air meteorological data to form a five-year model ready data set in AERMET;
- 5 Modeling of proposed Plant emissions in AERMOD and comparison with threshold levels; and
- 6 Modeling of project and associated coal mining feedstock operations for comparison with ambient air quality levels.

Details of these steps are provided in following subsections.

Two modeling scenarios were performed: a cumulative NAAQS / WAAQS analysis that evaluates impacts due to sources directly related to the proposed Plant as well as the nearby Elk Mountain Mine operations, and nearby sources included in a WDEQ-supplied emission inventory, and a PSD increment analysis to evaluate impacts due to the proposed Plant and onsite mining.

6.2 DESCRIPTION OF THE PROPOSED PROJECT

6.2.1 Site Location

The facility will be located approximately 7.5 miles north of Interstate 80, exit 260 (Elk Mountain) on County Road #3 in Section 29 of Township 21 north and Range 79 west in Carbon County, Wyoming as shown in Figure 1.1. The UTM coordinate (NAD27) of the center of Section 29 is 390634 meters E and 4624013 meters N. A topographic map of the facility area indicating Section 29 is shown in Figure 1.1. Photographs of the proposed site area are shown in Figure 6.1 and Figure 6.2, depicting the varying terrain.

The proposed project is classified as a 'Fuel Conversion Plant', which is one of the 28 major stationary sources for which the major source PSD threshold is 100 tpy for each criteria pollutant. As shown in Table 1.1 and Table 3.3, the estimated emissions from the facility exceed these levels for NO_x, CO, VOCs, and PM₁₀. Therefore, the project is subject to PSD review.

The project site is located in an area that is designated as attainment for all NAAQS.

Figure 6.1 – Plant Site Area, View from South Side



Figure 6.2 – Plant Site Area, View Over Coal Hills Toward Elk Mountain



6.2.2 Source Emissions and Parameters

Modeled Plant emission rates were based on the activity levels and applied control technologies described in Sections 3 and 4 of this document. Conservative emission estimates were used to predict the maximum likely impacts for each modeled pollutant. Where practicable, combinations of operations were developed to allow operational flexibility for future Plant activities. For example, cold startup and operations after cold startup, and normal operations scenarios were evaluated to determine annual emissions for modeling.

Of the emitted criteria pollutants, VOC emissions, which are precursors to ozone, were not explicitly modeled. Modeling of VOC impacts is not performed for two reasons. First, no NAAQS are established for VOCs. Second, AERMOD does not have the capability to model the chemical reactions that form ozone in the atmosphere from VOCs. Given the relatively low ambient ozone concentrations in the area surrounding the Plant and the lack of significant industrial NO_x and VOC emissions nearby, no ozone analysis was performed.

Emissions of criteria pollutants NO_x, CO, SO₂, and PM₁₀ were explicitly modeled and the maximum total short-term emission rates for all sources are shown below in Table 6.1.

Table 6.1 – Maximum Combined Modeled Short-Term Emission Rates for All Sources in the Analysis

Total NO _x (g/sec) Modeled	Total CO (g/sec) Modeled	Total SO ₂ (g/sec) Modeled	Total PM ₁₀ (g/sec) Modeled ¹
14.691	853.108	1400.80	11.42/4.21

1. Emission rate modeled with long-term analysis, including all mining-related point and fugitive sources/emission rate modeled in short-term analysis, representing only mining-related point sources (no fugitives).

Specific source model emission rates and input parameters are shown in Table 6.2. Pollutants with short-term averaging periods (CO, SO₂, and PM₁₀) were modeled at maximum short-term rates for all operating scenarios. Note that for the LP Flare, a cold startup will not occur for a full day, but during those startup hours, the expected emissions from the LP Flare may substantially exceed its normal operation short-term emission rates. The short-term modeling analysis includes these higher short-term, startup-related, emissions from the LP Flare. Modeled pollutant emissions for the long-term (annual) NO_x, SO₂, and PM₁₀ analyses were based on additive operations across the highest emitting scenarios (7760 hr/yr of normal operations after startup plus 1,000 hr/yr of cold startup conditions).

Stack input parameters such as height, diameter, velocity, and temperature, are based on vendor information or established values for similar unit operations. Effective heights and diameters for the HP and LP flares during startup and normal operations were calculated and modeled per established modeling guidance documentation.

The full cumulative modeling analysis includes a nearby (35-km) source inventory, supplied by the WDEQ, for NO_x and CO sources. Although the relative spatial distances are large, the point

Table 6.2 – Modeled Plant Point Source Parameters

Emission Unit	Emission Unit / Model ID	Location UTM			Modeled Exhaust Parameters				Modeled Emission Rates (g/s)			
		X (m)	Y (m)	Z (m)	Height (m)	Temp (K)	Velocity (m/s)	Diameter (m)	NO _x	CO	SO ₂	PM ₁₀
Turbine and HRSG Train 1	CTG1	391190.18	4624309.74	2133	45.73	366.49	7.65	5.79	2.206	1.434	0.336	1.26
Turbine and HRSG Train 2	CTG2	391190.18	4624231.74	2133	45.73	366.49	7.65	5.79	2.206	1.434	0.336	1.26
Turbine and HRSG Train 3	CTG3	391190.18	4624179.74	2133	45.73	366.49	7.65	5.79	2.206	1.434	0.336	1.26
Gasifier Preheater 1	GHEAT1	390998.86	4624266.35	2133	25.91	422.05	7.45	0.41	0.0074	0.218	0.0015	0.0197
Gasifier Preheater 2	GHEAT2	390998.46	4624253.85	2133	25.91	422.05	7.45	0.41	0.0074	0.218	0.0015	0.0197
Gasifier Preheater 3	GHEAT3	390998.18	4624241.85	2133	25.91	422.05	7.45	0.41	0.0074	0.218	0.0015	0.0197
Gasifier Preheater 4	GHEAT4	390997.86	4624229.85	2133	25.91	422.05	7.45	0.41	0.0074	0.218	0.0015	0.0197
Gasifier Preheater 5	GHEAT5	390997.46	4624217.35	2133	25.91	422.05	7.45	0.41	0.0074	0.218	0.0015	0.0197
HP Flare	Z8901	390824.94	4624353.31	2133.9	46.0 / 86.55 *	1273	20	0.152 / 13.64*	0.2956	409.4	946.02	0.0
Black-Start Generator 1	BSG1	391102.68	4623970.7	2133	30	767.6	1.96	0.41	0.033	1.95	0.0014	0.00019
Black-Start Generator 2	BSG2	391107.68	4623970.7	2133	30	767.6	1.96	0.41	0.033	1.95	0.0014	0.00019
Firewater Pump	FIREPUMP	391247.38	4624293.74	2133	6.1	739.27	45	0.15	0.0433	0.046	0.00076 ₄	0.0096
Auxiliary Boiler	AB	391085.81	4624005.5	2133	15.24	422.05	1.6	0.91	0.4076	0.685	0.005	0.062
Catalyst Regenerator	REGH	391329.29	4624467.64	2133	15.24	422.05	1.6	0.91	0.133	0.223	0.0016	0.0202
Reactivation Heater	REAH	391329.5505	4624486.43	2133	15.24	422.05	1.6	0.91	0.077	0.129	0.00092	0.0117
HGT Reactor Charge Heater	HGT	391329.29	4624447.64	2133	15.24	422.05	1.6	0.91	0.077	0.023	0.00016	0.002
LP Flare	Z8902	390856.48	4624591.43	2133.6	70.4 / 74.9*	1273	20	--- / 4.43*	0.00437	2.44	453.75	0.0
Black-Start Generator 3	BSG3	391112.68	4623970.7	2133	30	767.6	1.96	0.41	0.033	1.95	0.0014	0.00019

Table 6.2 – Modeled Plant Point Source Parameters

Emission Unit	Emission Unit / Model ID	Location UTM			Modeled Exhaust Parameters				Modeled Emission Rates (g/s)			
		X (m)	Y (m)	Z (m)	Height (m)	Temp (K)	Velocity (m/s)	Diameter (m)	NO _x	CO	SO ₂	PM ₁₀
CO ₂ Stack Vent	CO2V	390957.03	4624580.2	2133	30.48	296.88	6.99	1.83	0.0	423.21	0.0	0.0

* The second number indicates the flare’s effective stack height or effective diameter. Maximum modeled LP Flare (Source ID Z8902) height was 75 m (231 ft) for model year 2004 in the short-term NAAQS/WAAQS analysis, therefore, this will be the required height for the LP Flare.

Table 6.3 – Modeled Cumulative (Nearby) Point Source Parameters

Emission Unit / Model ID	Location UTM			Modeled Exhaust Parameters				Modeled Emission Rates (g/s)			
	X (m)	Y (m)	Z (m)	Height (m)	Temp (K)	Velocity (m/s)	Diameter (m)	NO _x	CO	SO ₂	PM ₁₀
SRC36454	421705	4587401	2225.9	13.87	672.04	12.19	1.07	15.09	-	-	-
SRC36455	421705	4587401	2225.9	13.87	672.04	12.19	0.91	6.13	2.83	-	-
SRC36456	421705	4587401	2225.9	13.87	672.04	12.19	1.07	15.09	-	-	-
SRC36457	421705	4587401	2225.9	13.87	672.04	12.19	1.07	10.38	1.32	-	-
SRC36458	421705	4587401	2225.9	8.23	842.04	78.64	0.24	3.26	0.377	-	-
SRC36459	421705	4587401	2225.9	8.23	842.04	78.64	0.24	3.26	0.377	-	-
SRC36462	421705	4587401	2225.9	12.19	685.93	41.76	1.04	0.618	0.662	-	-
SRC36463	421705	4587401	2225.9	6.4	449.82	6.12	0.46	0.154	-	-	-
SRC37392	395304.8	4649701	2023.84	7.92	596.48	24.05	0.43	0.975	0.106	-	-
SRC37393	395304.8	4649701	2023.84	7.92	596.48	24.05	0.43	0.975	0.106	-	-
SRC37771	399740	4606350	2332.8	10.97	922.04	50.51	1.01	0.710	0.518	-	-
SRC36900	375778.9	4651513	2011	11.0	730.4	71.6	0.25	0.503	0.164	-	-
SRC36901	375778.9	4651524	2011	11.0	730.4	71.6	0.25	0.503	0.164	-	-
SRC36902	375778.9	4651536	2011	11.0	762.0	38.6	0.25	0.319	0.642	-	-
SRC36903	375778.9	4651547	2011	11.0	762.0	38.6	0.25	0.319	0.642	-	-

sources included in this nearby inventory have significant emission rates. Table 6.3 details the nearby point sources used for cumulative modeling.

6.2.2.1 Coal Mine Fugitive Emission Area Sources

Existing surface and planned underground coal mining operations are located within the facility's "ambient" boundary. MBFP has an option to purchase the coal it needs from Arch Coal of Wyoming, LLC (Arch). Arch operates the existing surface mine, The Elk Mountain Mine, under permit CT – 4136 (Wyoming), which includes the projected future annual emissions and locations of its aboveground mining operations. A copy of that permit was obtained from the WDEQ.

Emission factors from the Arch surface mine permit were used to calculate future emissions from the aboveground operation locations to be constructed to support the proposed underground Saddleback Hills Mine. Area sources were created to the west of the facility for these potential future emissions.

Table 6.4 shows the area source modeling parameters for the Plant's mining operations as well as the aboveground mining operations associated with the Elk Mountain and Saddleback Hills Mine for this analysis.

6.2.2.2 Industrial Gasification & Liquefaction Plant Volume Sources

Volume sources were used to represent HAP emissions associated with storage tanks and equipment leaks. Table 6.5 shows the modeling parameters for the volume sources and Figure 6.3 shows the complete layout of all sources related to the facility (including the Elk Mountain Mine operations).

Figure 6.4 shows the locations of the Plant and the nearby sources included in the inventory sent by the WDEQ.

6.2.3 Additional Emission Assumptions

The following conservative assumptions were used when conducting this modeling analysis.

- Normal operations at the facility will not include the Black-Start Generator emissions. Therefore, simultaneous / concurrent emissions that were modeled for the Black-Start generators and turbines are not likely to occur. In other words, several emission units / sources are not likely to emit concurrently with other sources.
- Vehicle tailpipe NO_x emissions associated with the nearby mining operations (Elk Mountain Mine) were included in the PSD increment and NAAQS analysis.
- Vehicle tailpipe, surface mining, and vehicle traffic (associated with haul roads) PM₁₀, SO₂, and CO emissions (Elk Mountain Mine) were included in the NAAQS analyses to determine cumulative impacts for each pollutant.
- Surface mining emissions are below ground level or surrounded by high walls that could prevent the release of PM/PM10 into the ambient domain; the area sources for the surface mining for this modeling analysis are above ground level.

Table 6.4 – Area Source Modeling Parameters

Source ID (in model)	Source Type	Source Description	Corner Easting (X) (m)	Corner Northing (Y) (m)	Base Elevation (m)	Release Height (m)	Sigma z (initial dimension) (m)	Modeled Emission Rates (g/sec/m2)			
								NOx	CO	SO2	PM10 ³
MineA_SP ¹	Area	Mine Area / South Portal	384525.3	4622056.4	2252	12.0	13.95	0.0	0.0	0.0	0.0
MineA_EP ²	Area	Mine Area / East Portal	389721.7	4623411.5	2134	12.0	13.95	0.000004	0.0000034	0.00000007	0.0000082/0.0000001
MineA_S1 ²	Area	Mine Area / Surface Mining (On-Site)	389673.8	4623406.6	2134	12.0	NA	0.0000137	0.0000115	0.00000023	0.000007/0.00000057 ^{3,4}
MineA_S2 ²	Area	Mine Area / Surface Mining (Off-Site)	388229.3	4622116.0	2189	12.0	NA	0.0000137	0.0000115	0.00000023	0.000007/0.00000057 ^{3,4,5}

Notes

1. The analysis reflects development year 4 operations, where normal plant operations have begun, and all coal produced at the Saddleback Hills Mine is brought out through the mine’s East Portal (Source ID MineA_EP). Mine development emissions from the Saddleback Hills Mine South Portal (Source ID MineA_SP) will begin to decline in development year 3 and will cease in year 4.
2. Estimated sizes for these areas are as follows:
 MineA_EP: 215,578 m² (2,320,462 ft², or 53.3 acres)
 MineA_S1: 144,821.2 m² (1,558,842 ft², or 35.8 acres)
 MineA_S2: 224,918.7 m² (2,421,005 ft², or 55.6 acres)
3. Where two values are shown, the PM₁₀ modeled emissions rates represent total point source and fugitive source emissions included in the long-term (annual) analysis, and the point source emissions included in the short-term (24-hr) analyses.
4. Modeled emission rates for Arch Coal’s Elk Mountain Mine (EMM) are based on emissions calculated for construction permit CT-4136, issued to Arch of Wyoming, LLC. Emissions related to coal production from the Saddleback Hills Mine (now considered as a support facility to the proposed IGL Plant) have been subtracted from previously calculated EMM emission rates in order to correctly represent emissions from the EMM surface mine only. Emissions from the Saddleback Hills Mine are represented by area sources ‘MineA_SP’ and ‘MineA_EP.’
5. Modeled emissions at the EMM area source ID ‘MineA_S2’ are conservatively assumed to be the same as those at the EMM area source ID ‘MineA_S1.’ Estimated emissions in [tons/year] for these two mine area sources are assumed to be equal at 35.5 tpy each; however, the mine areas are differing sizes, with MineA_S2 at a larger size than MineA_S1. As a result, the modeled emissions in [g/s/m²] for MineA_S2 are calculated at 0.00000454 g/s/m². Rather than use the smaller emission rate for modeling, the higher emission rate for MineA_S1 was conservatively assumed in the model for MineA_S2 and is listed here.

Table 6.5 – Volume Source Modeling Parameters

Source ID (in model)	Source Type	Source Description	Easting (X) (m)	Northing (Y) (m)	Base Elevation (m)	Release Height (m)	Sigma y (initial dimension) (m)	Sigma z (initial dimension) (m)	Modeled Emission Rates (g/sec)			
									NOx	CO	SO2	PM10
T_A	Volume	Gasoline Tank	390966.4	4624652	2133.2	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_B	Volume	Gasoline Tank	391021.3	4624652	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_C	Volume	Gasoline Tank	391109.2	4624652	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_D	Volume	Gasoline Tank	391175.2	4624652	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_E	Volume	Gasoline Tank	390966.4	4624712	2133.2	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_F	Volume	Gasoline Tank	391021.3	4624712	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_G	Volume	Gasoline Tank	391109.2	4624712	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_H	Volume	Gasoline Tank	391175.2	4624712	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_I	Volume	Methanol Tank	390966.4	4624822	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_J	Volume	Methanol Tank	391021.3	4624822	2133	14.6304	10.6325581	2.32	0.0	0.0	0.0	0.0
T_K	Volume	Heavy Gas Tank	391173.8	4624840	2133	14.6304	9.21488372	2.32	0.0	0.0	0.0	0.0
V1	Volume	Equipment Leaks	391224.369	4624457.507	2133	2.0	61.12	4.65	0.0	0.0	0.0	0.0
V_1 through V_112	Volume	Haul Roads	varying	varying	varying	2.0	1.63	2.79	0.0	0.0	0.0	0.0284/0.0 ¹

Notes

1. Haul road PM₁₀ emissions are fugitives, and per WDEQ policy, are not included in short-term (24-hr) modeling analyses.

Figure 6.3 – Plant and Nearby Mining Area Sources

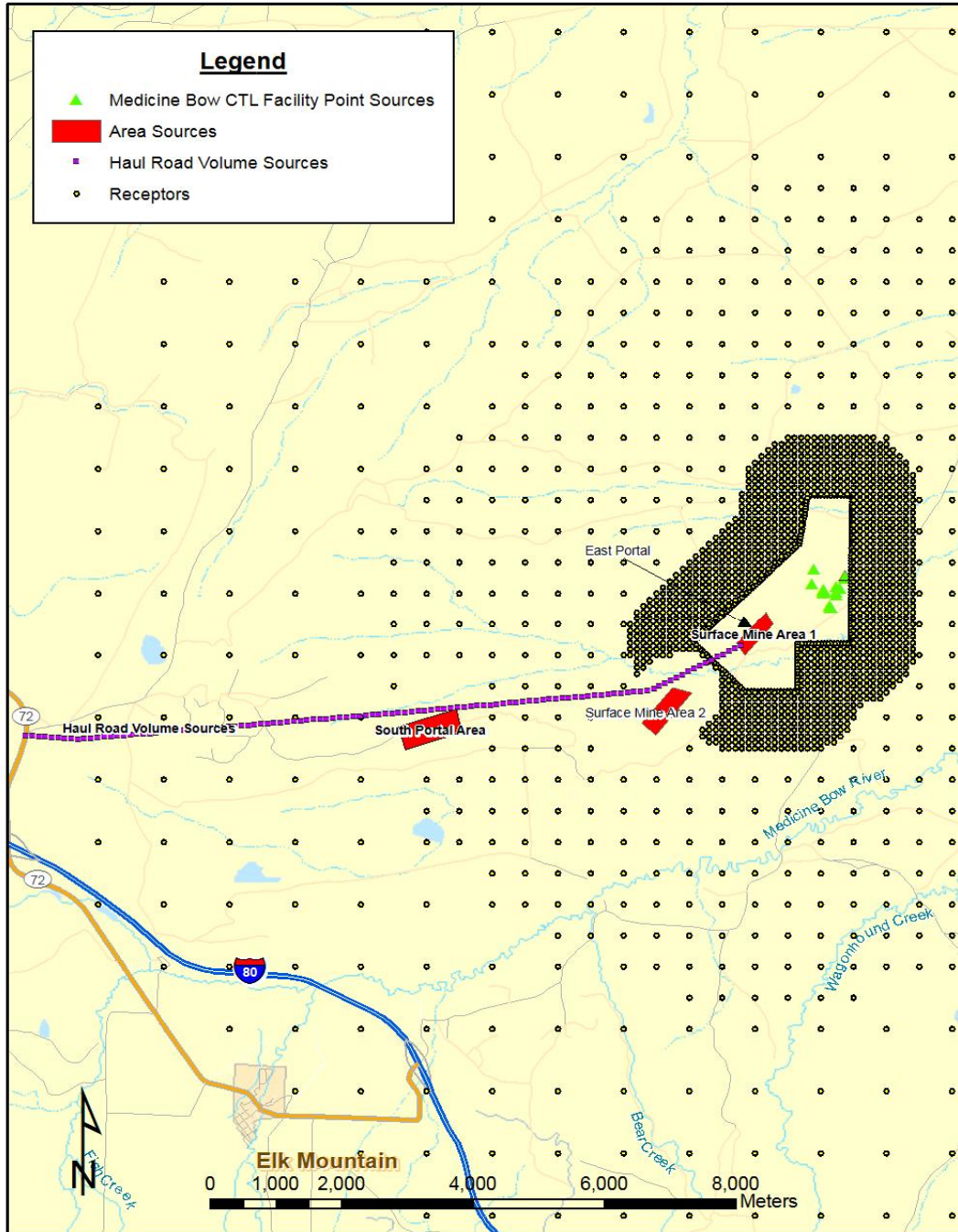
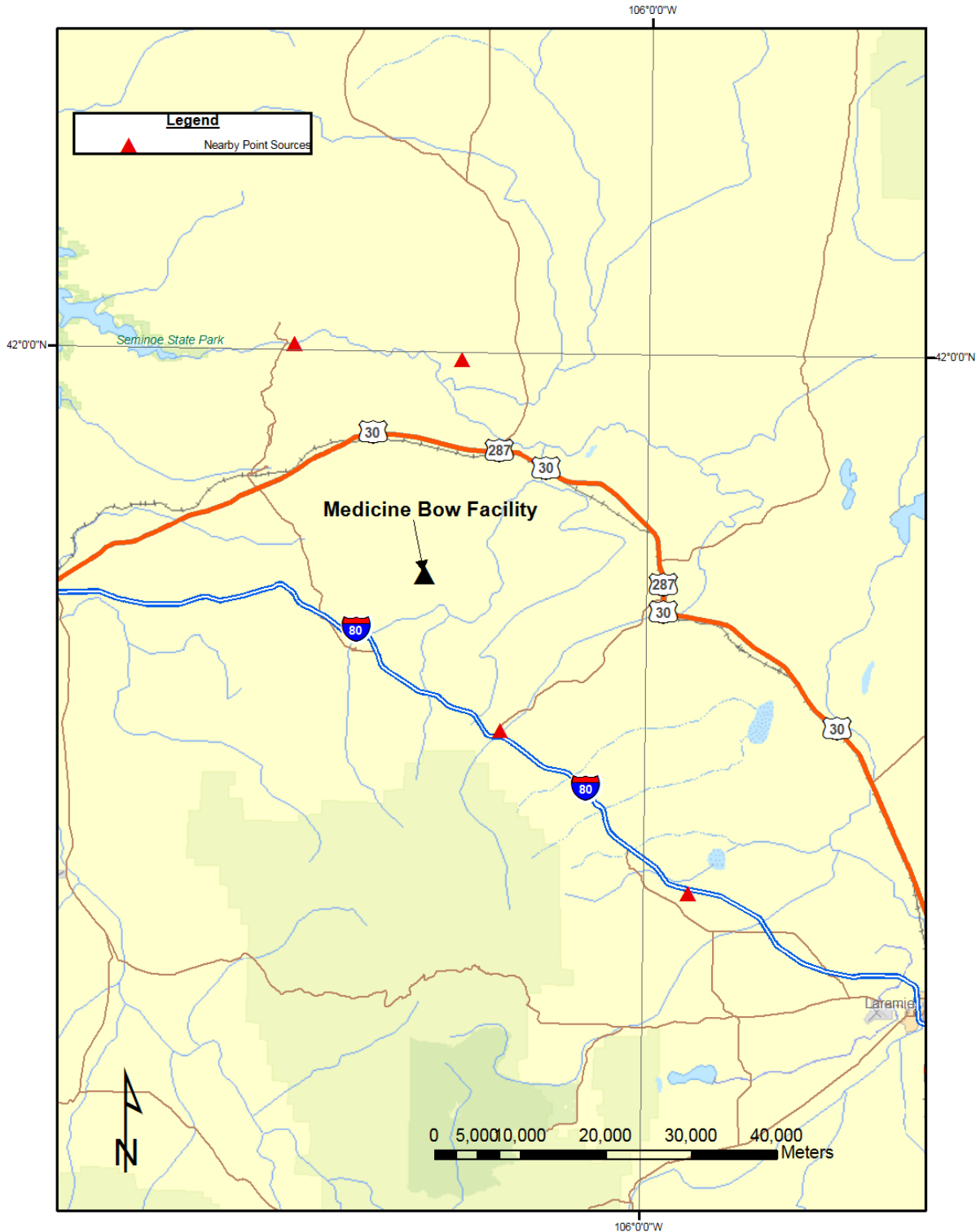


Figure 6.4 – Plant Location Relative to the WDEQ Provided Emission Inventory



6.3 STANDARDS, CRITERIA LEVELS, AND BASIC METHODOLOGY

The results of the air quality dispersion modeling analysis are compared with various ambient levels to assess potential impacts to local air quality resulting from the proposed Plant. The proposed Plant’s source emissions must not cause an exceedance of any national or Wyoming ambient air quality standards, and the increase in ambient air concentrations must not exceed the allowable PSD increments.

The dispersion modeling analysis typically involves a two-step approach. The first step looks at the proposed facility’s emission sources and is referred to as the significant impact analysis (SIA). Only the proposed facility is considered in the SIA analysis; nearby sources and background ambient air quality concentrations are not considered. The highest predicted off-site concentration for each pollutant and each averaging period is compared to the modeling significant impact levels (SILs) listed in Table 6.6. If the estimated concentration levels are below the applicable SIL, no further analysis is required and the source is considered to have an insignificant impact. For the proposed Plant, SIA modeling results indicated exceedance of the SILs for each of the pollutants shown in Table 6.6.

The next phase is more robust and includes the NAAQS / WAAQS and the PSD increment analyses, which require modeling the proposed Plant emission sources as well as nearby sources and taking the background air quality concentration into account. The NAAQS and WAAQS are maximum concentration “ceilings” measured in terms of the total concentration of a pollutant in the atmosphere. The proposed Plant’s source emissions cannot cause a NAAQS or WAAQS exceedance. A PSD increment is the maximum increase in ambient concentration that is allowed to occur above a baseline concentration for a pollutant. Significant deterioration is said to occur when the amount of new pollution would exceed the applicable PSD increment. The NAAQS, WAAQS, and Class II PSD Increments are listed in Table 6.6. Because the proposed Plant emissions resulted in SIL exceedances for all modeled criteria pollutants, full NAAQS / WAAQS and PSD increment analyses were performed.

Table 6.6 – SILs, NAAQS, WAAQS, and PSD Class II Increments

Pollutant	Averaging Period	SIL (µg/m³)	NAAQS ¹ / WAAQS (µg/m³)	Allowable Increment (µg/m³)
Nitrogen Dioxide	Annual	1	100	25
Sulfur Dioxide	3-hour	25	1,300	512
	24-hour	5	365 / 260	91
	Annual	1	80 / 60	20
Particulate Matter <10 µm [PM ₁₀]	24-hour	5	150	30
	Annual	1	Revoked / 50	17
Carbon Monoxide	1-hour	2,000	40,000	N/A
	8-hour	500	10,000	N/A

1. Primary NAAQS are noted in this table. Secondary NAAQS are addressed in Section 6.8 (Impacts to Soil and Vegetation).

For a new source, compliance with any NAAQS is based upon the total estimated air quality, which is the sum of the background concentration and the estimated ambient impacts of the Plant's proposed emissions. A complete PSD increment "consumption" and NAAQS comparison evaluation was completed for this modeling analysis.

6.4 NEAR FIELD MODELING METHOD

Near field impact analysis modeling was conducted for Plant sources of NO_x, CO, SO₂ and PM₁₀ emissions using the methodology outlined in the previous section. This section includes a detailed description of the modeling approach and data requirements for assessing air quality impacts due to the proposed Plant.

6.4.1 Model Selection and Setup

The air quality impacts were modeled at near field receptors using the latest version of the EPA regulatory model (AERMOD) (Version 07026). The AERMOD model is designed to predict ground-level pollutant concentrations from a wide variety of sources associated with industrial facility source types. AERMOD contains algorithms for: (1) dispersion in both the convective and stable boundary layers; (2) plume rise and buoyancy; (3) plume penetration into elevated inversions; (4) computation of vertical profiles of wind, turbulence, and temperature; (5) urban nighttime boundary layer; (6) treatment of receptors on all types of terrain from the surface up to and above the plume height; (7) treatment of building wake effects; (8) improved approaches for characterizing the fundamental boundary layer parameters, and (9) treatment of plume meander. The AERMOD modeling system consists of two pre-processors: AERMET which provides AERMOD with the meteorological information it needs to characterize the planetary boundary layer (PBL); and AERMAP, which characterizes the terrain, and generates receptor grids for AERMOD.

Pursuant to WDEQ modeling guidelines (2006a and 2006b), the regulatory default options were used, including building and stack tip downwash, default wind speed profiles, exclusion of deposition and gravitational settling, consideration of buoyant plume rise, and complex terrain.

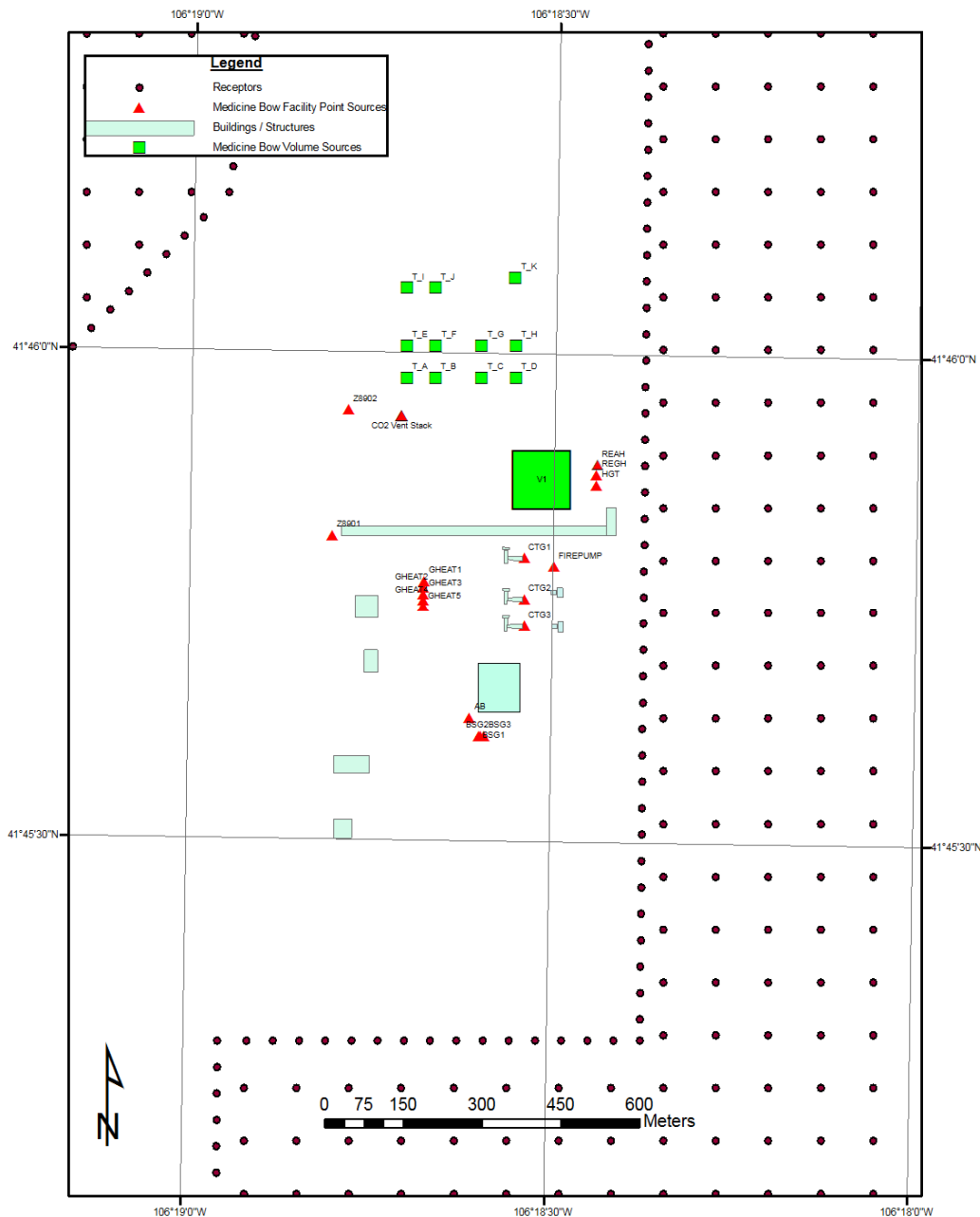
Emission sources at the Plant will be influenced by aerodynamic downwash. Since downwash is a function of projected building width and height, it is necessary to account for the changes in building projection as they relate to changes in wind direction. Once these projected dimensions are determined, they can be used as input to the AERMOD model. The USEPA Building Profile Input Program (BPIP version 04274), enhanced to include the PRIME algorithms as applicable to AERMOD, was used to conduct the good engineering practice (GEP) stack height analysis and to determine wind direction-specific building/structure dimensions.

The BPIP-PRIME program builds a mathematical representation of each building or structure to determine projected building dimensions and its potential zone of influence. These calculations are performed for 36 different wind directions (at 10-degree intervals). If the BPIP-PRIME program determines that a source is under the influence of several potential building wakes, the structure or combination of structures which has the greatest influence ($h_b + 1.5 l_b$) is selected for input to the model. Conversely, if no building wake effects are predicted to occur for a source for a particular wind direction, or if the worst-case building dimensions for that direction yield a wake region height less than the source's physical stack height, building parameters are set equal

to zero for that wind direction. For this case, wake effect algorithms are not exercised when the model is run. The building wake criteria influence zone is 5 l_b downwind, 2 l_b upwind, and 0.5 l_b crosswind. These criteria are based on recommendations by USEPA. The PRIME algorithm addresses the entire structure of the wake, from the cavity immediately downwind of the building, to the far wake.

Input to the BPIP/PRIME program consisted of the location of Plant emission units and the coordinates and heights of the buildings and structures. The structures used in the analysis are shown in Figure 6.5 along with the source locations.

Figure 6.5 – GEP Stack Height Assessment Building and Source Location Depiction



6.4.2 Databases for Air Quality Assessment

The databases required for the air quality impact assessment included emissions inventory, meteorological data, receptor points, and terrain data. The emissions inventory was described in Section 6.2.2 and presented in Tables 6.2 through 6.5. The following sections describe the meteorological data, receptor points, and terrain data databases required to perform the air quality impact assessment.

6.4.3 Meteorological Data

Nearby sources of meteorological data (three surface sites and one upper air site) were identified, and six years of recent (2000–2005) meteorological data were obtained, reviewed for completeness, and the valid years were processed in AERMET. The surface sites included a nearby meteorological tower installation with automatic recording instrumentation located outside of Elmo, WY, about 24 km northwest of the Plant site, and two National Weather Service (NWS) ASOS sites, one located at the Rawlins Municipal Airport approximately 70 km west of the Plant location and one located at the Laramie Gen. Brees Airport approximately 73 km southeast of the Plant location.

Inter-Mountain Labs (IML) operated the Elmo meteorological station in accordance with *Meteorological Monitoring Guidance for Regulatory Modeling Applications* (EPA-454/R-99-005). IML performed semi-annual quality assurance audits on the station and the IML staff conducted quality control procedures on the data. IML submitted quarterly reports (including semiannual quality assurance audits) to Dennis Wuertz at Seminoe (Arch of Wyoming, LLC), who then submitted the reports to Bob Schick at the WDEQ. Cara Keslar in the Air Quality Monitoring Division may be contacted with regard to this data. The UTM coordinates (Zone 13, NAD27) of this station are 372052 meters E, 4638122 meters N.

In order to meet the completeness criteria for PSD-quality meteorological data, only 10 percent of the data in any given year can be missing. The Elmo, WY data was reviewed for completeness and the results are shown in Table 6.7. The Elmo data collected during 2002 does not satisfy the completeness criteria because only 64%, 40%, and 81% of the data were available during the 2nd, 3rd, and 4th quarters of the year. Therefore, 2000, 2001, 2003, 2004, and 2005 on-site data were used for the AERMET processing and AERMOD modeling.

Therefore, a five year meteorological data set was developed for the years 2000, 2001, and 2003–2005 with the Elmo site noted as the “on-site” location and the Laramie and Rawlins sites as the NWS surface locations, respectively. The Rawlins NWS site meteorology set for years 2000 and 2003 lacked sufficient cloud cover data necessary to establish completeness. Consequently, Rawlins NWS surface meteorology data was used only for the years 2001, 2004, and 2005. Because the Laramie NWS had complete cloud cover data for the two years for which the Rawlins data was incomplete, Laramie NWS surface meteorology was used for the years 2000 and 2003. The full five-year data set was processed in AERMET into model-ready format.

Table 6.7 – Site-Specific Elmo Meteorological Data Completeness

Months	Year	1st Quarter	2nd Quarter	3rd Quarter	4th Quarter
		January-March	April-June	July-September	October-December
Total Hours per Quarter		2184 or 2160	2184	2208	2208
Number of Missing Hours	2000	0	193	0	1
	2001	0	2	0	1
	2002	159	787	1316	420
	2003	0	1	1	2
	2004	2	0	1	50
	2005	2	50	1	0
Percent Completed (%)	2000	100.0	91.2	100.0	100.0
	2001	100.0	99.9	100.0	100.0
	2002	92.6	64.0	40.4	81.0
	2003	100.0	100.0	100.0	99.9
	2004	99.9	100.0	100.0	97.7
	2005	99.9	97.7	100.0	100.0

Three years of hourly surface observations (2001, 2004, and 2005) from the Rawlins Municipal Airport, WY were obtained from the National Climatic Data Center (NCDC) in AERMET-compatible TD3505 format. The Rawlins NWS site is located approximately 70 km west of the proposed facility at UTM coordinates (NAD27) 317221 meters E and 4629697 meters N.

Two years of hourly surface observations (2000 and 2003) from the Laramie Gen. Brees Airport, WY were obtained from the NCDC in AERMET-compatible TD3505 format. The Laramie NWS site is located approximately 73 km southeast of the proposed facility at UTM coordinates (NAD83 Zone 13 North) 443640.9 meters E and 4573759.8 meters N.

The Rawlins and Laramie hourly surface meteorology data sets were reviewed to establish completeness. The result of the completeness review of the Rawlins and Laramie data is shown in Table 6.8. The frequency distribution of wind speed and direction for the Elmo, Rawlins, and Laramie combined / AERMET processed surface data is shown in Table 6.9.

Table 6.8 – Data Completeness Evaluation, Rawlins and Laramie NWS Hourly Surface Meteorological Data

Year	NWS Site	Number of Missing Hours	Percent Complete (%)
2000	Laramie	328	96.3
2001	Rawlins	504	94.2
2003	Laramie	151	98.3
2004	Rawlins	447	94.9
2005	Rawlins	514	94.1

Table 6.9 – Frequency Distribution of Wind Speed and Direction of the Elmo, Rawlins, and Laramie Hourly Surface Meteorological Data (2000, 2001, 2003–2005)

Wind Direction	Wind Speed						Total
	0.5 - 2.1	2.1 - 3.6	3.6 - 5.7	5.7 - 8.8	8.8 - 11.1	>= 11.1	
348.75 - 11.25	43	75	79	58	46	102	403
11.25 - 33.75	60	150	120	90	66	520	1006
33.75 - 56.25	75	245	260	223	292	3793	4888
56.25 - 78.75	64	428	608	752	965	10043	12860
78.75 - 101.25	47	539	1070	1280	1188	5847	9971
101.25 - 123.75	54	310	482	466	398	1537	3247
123.75 - 146.25	45	152	101	149	177	609	1233
146.25 - 168.75	33	126	98	70	76	191	594
168.75 - 191.25	64	129	148	108	83	200	732
191.25 - 213.75	37	248	464	491	324	393	1957
213.75 - 236.25	58	286	432	381	297	564	2018
236.25 - 258.75	43	205	212	218	244	1055	1977
258.75 - 281.25	44	185	161	149	178	930	1647
281.25 - 303.75	50	152	111	62	49	226	650
303.75 - 326.25	68	111	77	45	29	66	396
326.25 - 348.75	34	68	61	26	19	47	255
Sub-Total:	819	3409	4484	4568	4431	26123	43834
Calms:							0
Missing/Incomplete:							14
Total:							43848

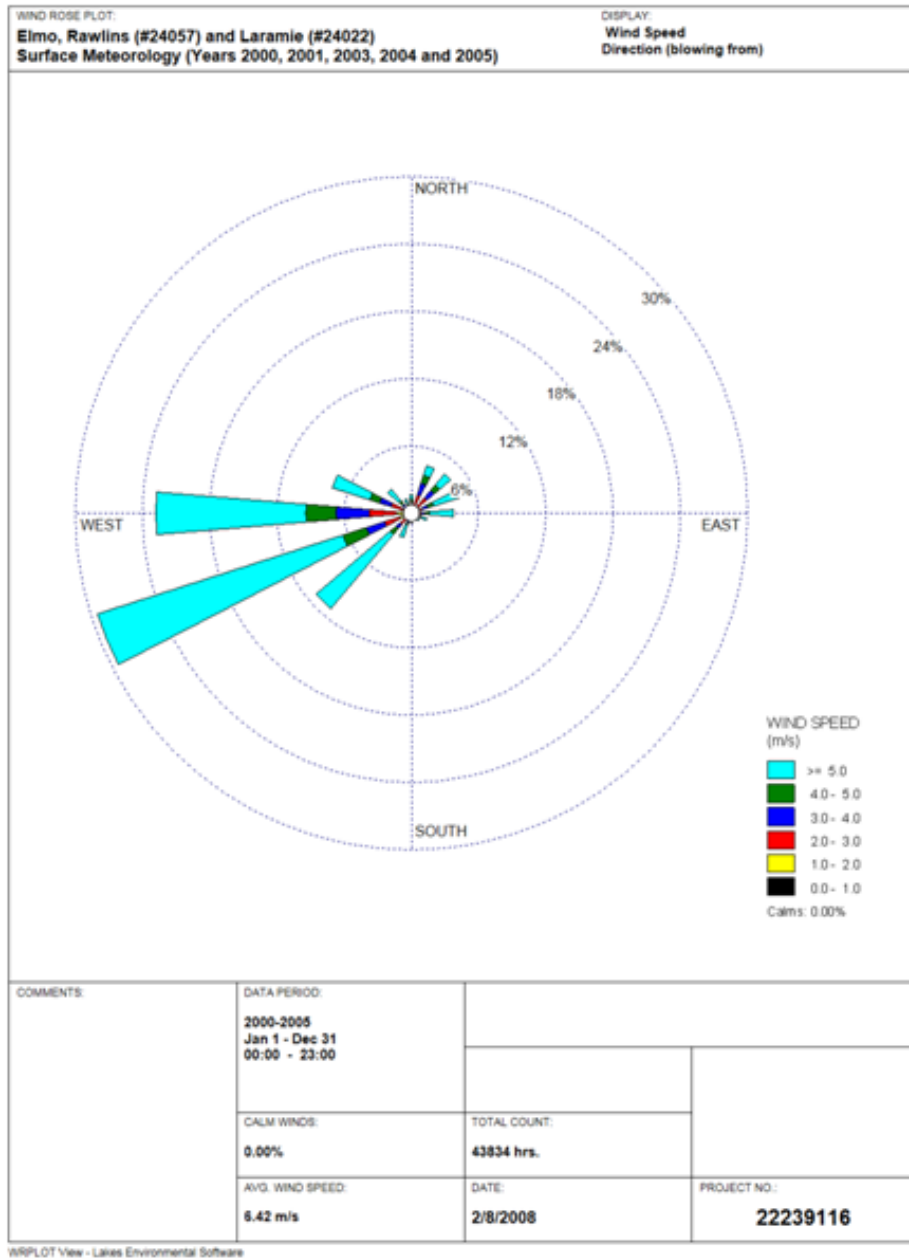
Upper air data are needed to estimate hourly mixing heights, which are required inputs to the AERMOD dispersion model. The most suitable NWS station to the project site that routinely performs upper air soundings is the NWS station in Riverton, WY (WBAN 24061), which is located approximately 250 km northwest of the proposed project site. The UTM coordinates (NAD27) of the Riverton NWS station are 217421 meters E and 4773109 meters N. Twice-daily upper air sounding data was obtained from the National Oceanic & Atmospheric Administration (NOAA), <http://raob.fsl.noaa.gov/>.

As discussed with WDEQ, the same five years (2000, 2001, 2003, 2004, and 2005) were used for both the NWS surface and upper air data in the AERMET processing so that the upper air data coincided with the surface data. Five parameters for each hour were collected at the Elmo, WY monitoring site, including wind direction (degree), wind speed (meters per seconds), sigma theta (degrees), temperature (Celsius), and precipitation (millimeters). Sensor elevations are 10 meters above grade level (agl) for wind speed and direction, 2 meters (agl) for temperature, and approximately 1 meter (agl) for precipitation.

An average of the desert scrubland and grassland surface characteristics values for albedo, the Bowen Ratio and surface roughness length were applied to AERMET Stage 3.

The windrose of the processed AERMET data based on the site-specific Elmo, Laramie, and Rawlins hourly surface meteorological data is shown in Figure 6.6.

Figure 6.6 – Wind Rose of AERMOD Input, Five-Year Period



6.4.4 Receptor Grid

The receptor grid used in the modeling analysis was designed to identify the maximum air quality impact due to the proposed project. The receptor grid began at the ambient air boundary and extended outward 10 km into ambient air. The following receptor spacing was used:

- 1 50 m spacing along the Plant's ambient air boundary;
- 2 100 m spacing from the boundary to 1 km;
- 3 500 m spacing from 1 km out from the proposed project to 5 km;
- 4 1 km spacing from 5 km to 10 km from the proposed project; and
- 5 500 meter from nearby mining area sources to closest receptor.

Receptor elevations were included for all receptor points and were obtained from digital elevation 7.5 minute topographic maps (<http://data.geocomm.com>). The DEM domain was extended to approximately 25 km from the proposed Plant to include the potential terrain of 10 percent slope or greater for complex terrain modeling. Source elevations were also obtained from the same data using AERMAP. The receptor grid is shown in Figure 6.3.

6.5 GROWTH ANALYSIS

During normal operations, the Plant is expected to employ 300 to 400 people with various trades. Many of these trades are commonly found in the coal mining industry. These employees are expected to live in the existing communities, such as Elk Mountain, Medicine Bow, Hanna, Saratoga, Rawlins, and Laramie. Carbon County has historically been a coal mining area with mining activity from the turn of the century through 2005. Population in the county has been declining since the 1990s (approximately 1,300) possibly resulting from the declining coal industry. The commercial support industries are already in place in Hanna and along the I-80 corridor.

6.6 CRITERIA POLLUTANT MODELING RESULTS

The following sections describe the results of the ambient air quality impact analysis. Modeling files are included in a CD-ROM provided along with this application. The README file included on the CD-ROM explains modeling file organization.

6.6.1 SO₂ Modeling Demonstration

Table 6.10 presents the maximum predicted 3-hour, 24-hour, and annual average SO₂ concentrations due to all cumulative source emissions. The second-highest concentration for each year is presented for the 3-hour and 24-hour averaging periods, while the maximum value for each annual average is presented. The total concentration (cumulative predicted concentration plus background) is compared to the NAAQS and WAAQS. As shown in the table, all predicted total concentrations are well below the respective NAAQS and WAAQS values.

Table 6.11 presents the maximum predicted 3-hour, 24-hour, and annual average concentrations due to project-specific source emissions and compares these values to the PSD increment. These emissions include on-site mining operations and emissions from Plant equipment. For the PSD increment comparison, each year's maximum concentration is selected for the 3-hour, 24-hour, and annual average time period. The predicted concentration is compared to the PSD increment directly, without including the background concentration. As shown in the table, all predicted concentrations are below the respective PSD increments.

Figures 6.7, 6.8, and 6.9 illustrate maximum PSD increment impacts for 3-hour, 24-hour, and annual averaging times.

Table 6.10 – Predicted SO₂ Concentrations Compared to NAAQS / WAAQS

Averaging Period	Data Period			Receptor Location (m)		Predicted Cumulative Concentration (ug/m3)	Background Concentration (ug/m3)	Total Concentration (Cumulative + Background) (ug/m3)	NAAQS / WAAQS (ug/m3)
	Year	Month/Day	Hour	East	North				
3 Hour ^{1,2}	2000	8/6	03	388955.38	4627705	1108.55	31.4	1139.95	N/A / 1300
	2001	6/22	24	381955.38	4628205	1097.85	31.4	1129.25	N/A / 1300
	2003	7/26	06	389455.38	4628205	1008.50	31.4	1039.90	N/A / 1300
	2004	5/22	03	395455.38	4624205	1033.64	31.4	1065.04	N/A / 1300
	2005	8/1	06	381955.38	4628205	1034.36	31.4	1065.76	N/A / 1300
24 Hour ^{1,2}	2000	5/30	24	389972.38	4624361	190.70	7.84	198.54	365 / 260
	2001	10/19	24	392055.38	4625005	197.92	7.84	205.76	365 / 260
	2003	12/27	24	391955.38	4625205	201.96	7.84	209.80	365 / 260
	2004	12/26	24	395455.38	4624205	241.39	7.84	249.23	365 / 260
	2005	9/15	24	395455.38	4624205	205.98	7.84	213.82	365 / 260
Annual	2000	N/A	N/A	391421.4	4624635	4.25	2.62	6.87	80 / 60
	2001	N/A	N/A	391421.4	4624585	4.51	2.62	7.13	80 / 60
	2003	N/A	N/A	391422.4	4624685	4.43	2.62	7.05	80 / 60
	2004	N/A	N/A	391420.4	4624485	4.01	2.62	6.63	80 / 60
	2005	N/A	N/A	391420.4	4624435	4.09	2.62	6.71	80 / 60

1. Based on the second-highest maximum.
2. Short-term analyses based on actual LP Flare (Source ID Z8902) maximum height of 70 m (231 ft) for model year 2004; all other model years, flare maximum height was set less than this height.

Table 6.11 – Predicted SO₂ Concentrations Compared to PSD Increments

Averaging Period	Data Period			Receptor Location (m)		Predicted Concentration (ug/m3)	PSD Increment (ug/m3)
	Year	Month/Day	Hour	East	North		
3 Hour ¹	2000	12/8	06	389410.38	4623014	7.04	512
	2001	9/24	03	389410.38	4623014	11.28	512
	2003	5/7	03	389410.38	4623014	8.95	512
	2004	8/27	06	389445.38	4622979	10.24	512
	2005	8/21	06	389410.38	4623014	8.23	512
24 Hour ¹	2000	2/14	24	391455.38	4624505	1.58	91
	2001	9/24	24	389410.38	4623014	2.46	91
	2003	12/13	24	391455.38	4624505	1.97	91
	2004	1/7	24	391555.38	4624505	1.62	91
	2005	8/21	24	389516.38	4622908	2.02	91
Annual	2000	N/A	N/A	391421.4	4624635	4.25	20
	2001	N/A	N/A	391421.4	4624585	4.51	20
	2003	N/A	N/A	391422.4	4624685	4.43	20
	2004	N/A	N/A	391420.4	4624485	4.01	20
	2005	N/A	N/A	391420.4	4624435	4.09	20

1. Short-term analyses does not include LP Flare (Source ID Z8902).

Figure 6.7 – 2003 Maximum SO₂ 3-Hour Impacts (PSD)

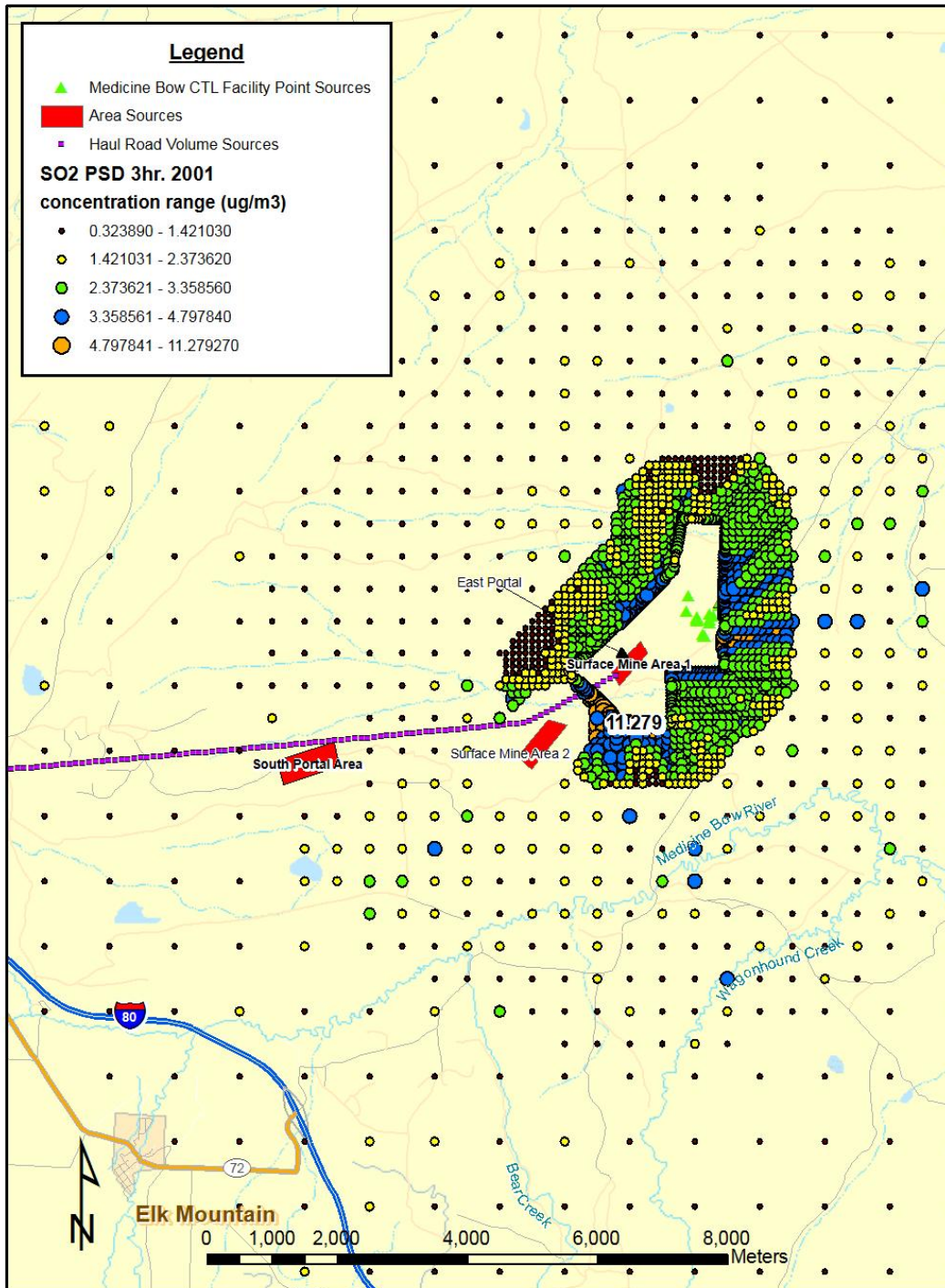


Figure 6.8 – 2000 Maximum SO₂ 24-Hour Impacts (PSD)

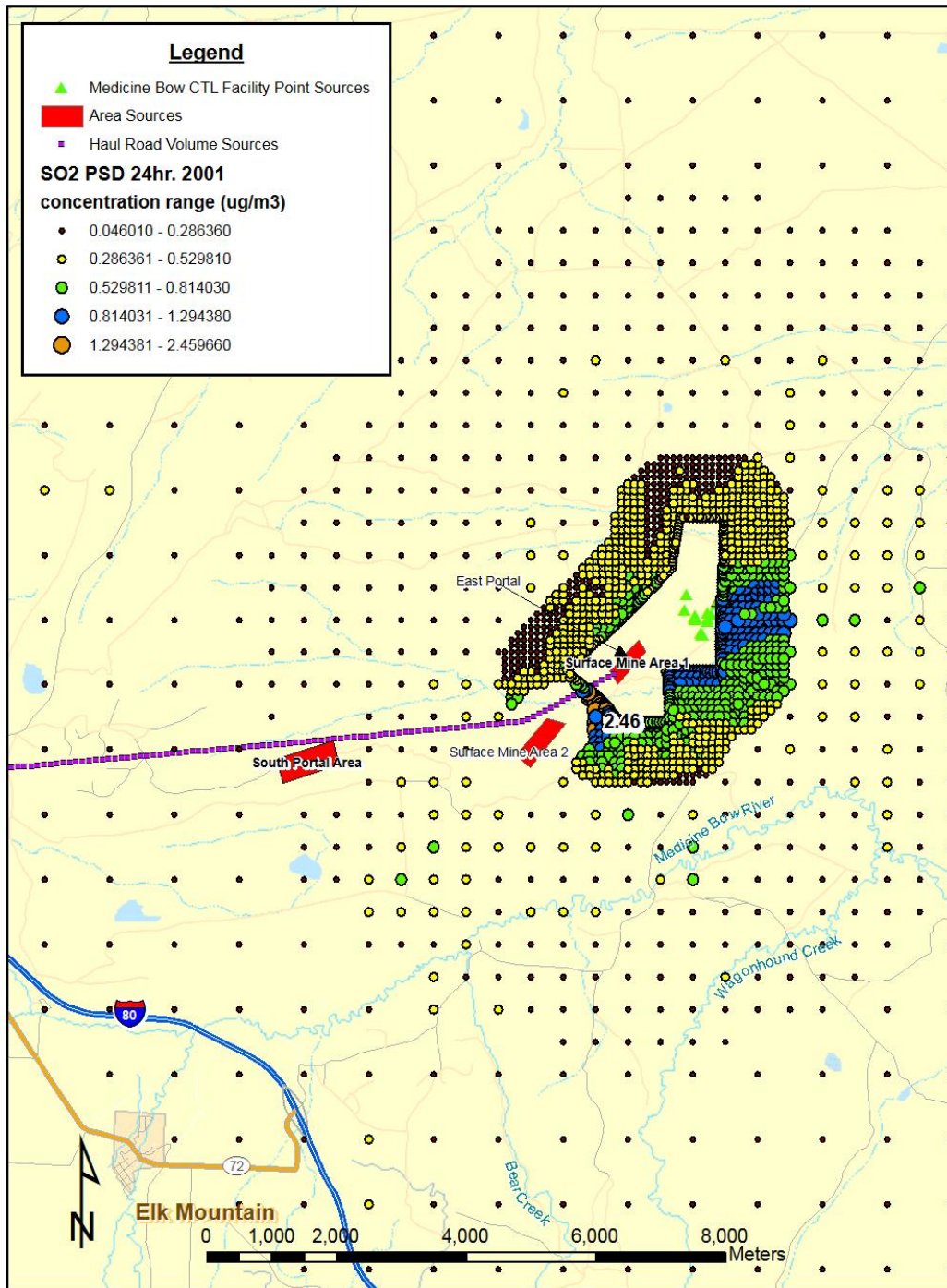
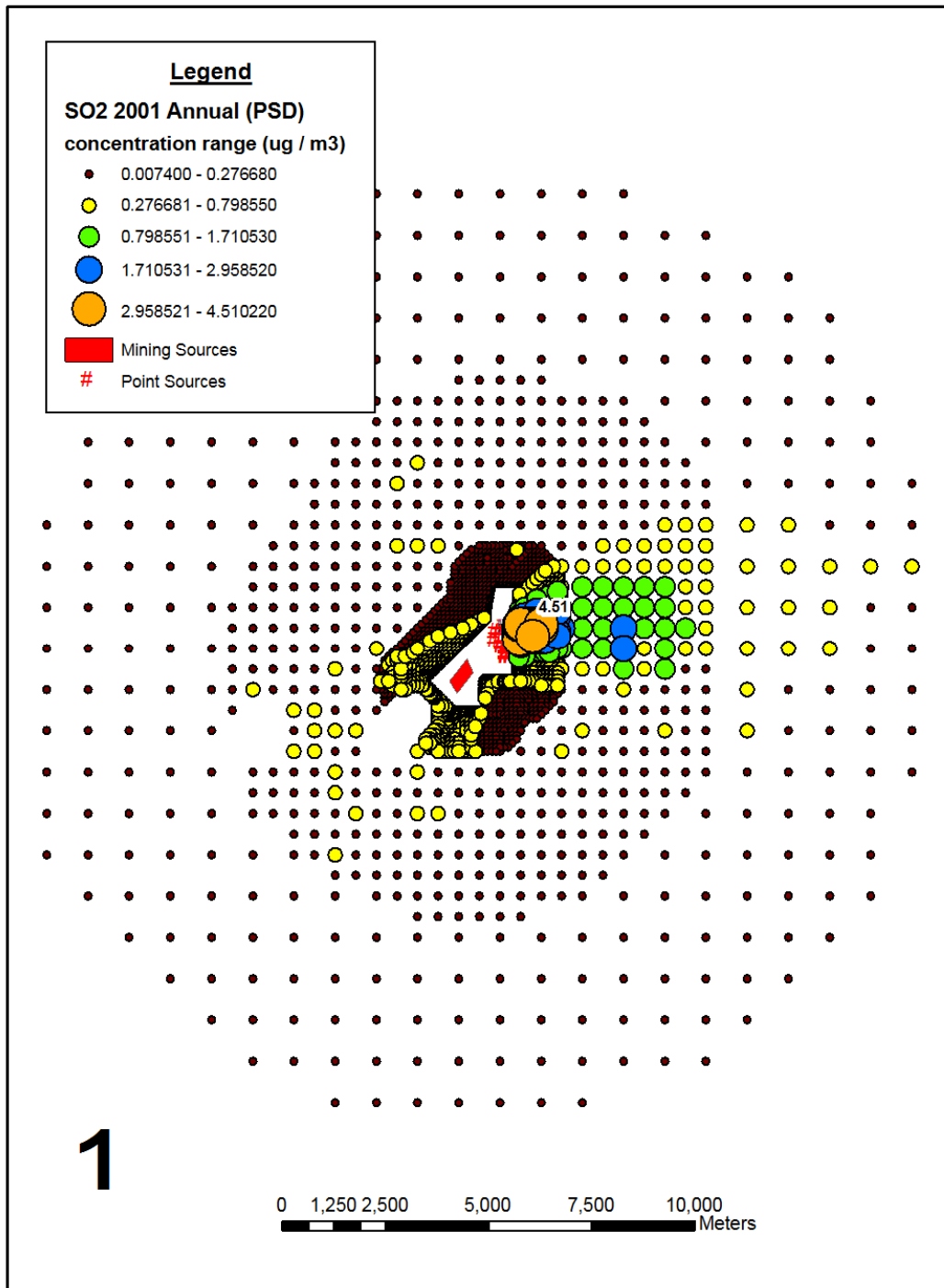


Figure 6.9 – 2001 Maximum SO₂ Annual Impacts (PSD)



6.6.2 PM/PM₁₀ Modeling Demonstration

Table 6.12 presents the maximum predicted 24-hour and annual average cumulative PM/PM₁₀ concentrations compared to the NAAQS and WAAQS. Emissions in this analysis include nearby mining operations and vehicle traffic and the proposed Plant. The predicted second-highest 24-hr value is presented, along with the highest predicted maximum annual value. Each of these values is added to the respective 24-hr and annual background concentration for comparison to the NAAQS and WAAQS. As shown in the table, all predicted total concentrations are below the respective NAAQS and WAAQS values.

Table 6.13 presents the results of the PM/PM₁₀ PSD increment analysis. The maximum predicted 24-hr and annual PM/PM₁₀ values are compared to the respective PSD increment. As shown in the table, all predicted concentrations are below the applicable PSD increment value.

Table 6.12 – Predicted PM/PM₁₀ Concentrations Compared to NAAQS / WAAQS

Averaging Period	Data Period			Receptor Location (m)		Predicted Cumulative Concentration (ug/m3)	Background Concentration (ug/m3)	Total Concentration (Cumulative + Background) (ug/m3)	NAAQS / WAAQS (ug/m3)
	Year	Month / Day	Hour	East	North				
Annual	2000	N/A	N/A	389355.4	4623005	14.33	26	40.33	NA / 50
	2001	N/A	N/A	389355.4	4623005	14.45	26	40.45	NA / 50
	2003	N/A	N/A	389355.4	4623005	13.62	26	39.62	NA / 50
	2004	N/A	N/A	389410.4	4623014	15.03	26	41.03	NA / 50
	2005	N/A	N/A	389355.4	4623005	14.80	26	40.80	NA / 50
24 Hour (H2H)	2000	12/14	24	391455.38	4624505	5.76	56	61.76	150 / 150
	2001	12/13	24	391655.38	4624505	5.20	56	61.20	150 / 150
	2003	12/30	24	391455.38	4624505	6.81	56	62.81	150 / 150
	2004	12/31	24	391455.38	4624605	5.28	56	61.28	150 / 150
	2005	11/2	24	391555.38	4624505	5.12	56	61.12	150 / 150

Table 6.13 – Predicted PM/PM₁₀ Concentrations Compared to PSD Increments

Averaging Period	Data Period			Receptor Location (m)		Predicted Concentration (ug/m3)	PSD Increment (ug/m3)
	Year	Month / Day	Hour	East	North		
Annual	2000	N/A	N/A	389355.4	4623005	14.33	17
	2001	N/A	N/A	389355.4	4623005	14.45	17
	2003	N/A	N/A	389355.4	4623005	13.62	17
	2004	N/A	N/A	389410.4	4623014	15.03	17
	2005	N/A	N/A	389355.4	4623005	14.80	17
24 Hour (H1H)	2000	1/16	24	391455.4	4624605	6.33	30
	2001	11/29	24	391555.4	4624505	5.49	30
	2003	12/13	24	391555.4	4624505	7.72	30
	2004	1/7	24	391555.4	4624505	6.30	30
	2005	1/8	24	391455.4	4624605	6.00	30

Figures 6.10 and 6.11 illustrate maximum PSD increment impacts for 24-hour and annual averaging times.

Figure 6.10 – 2005 Maximum PM₁₀ 24-Hour Impacts (PSD)

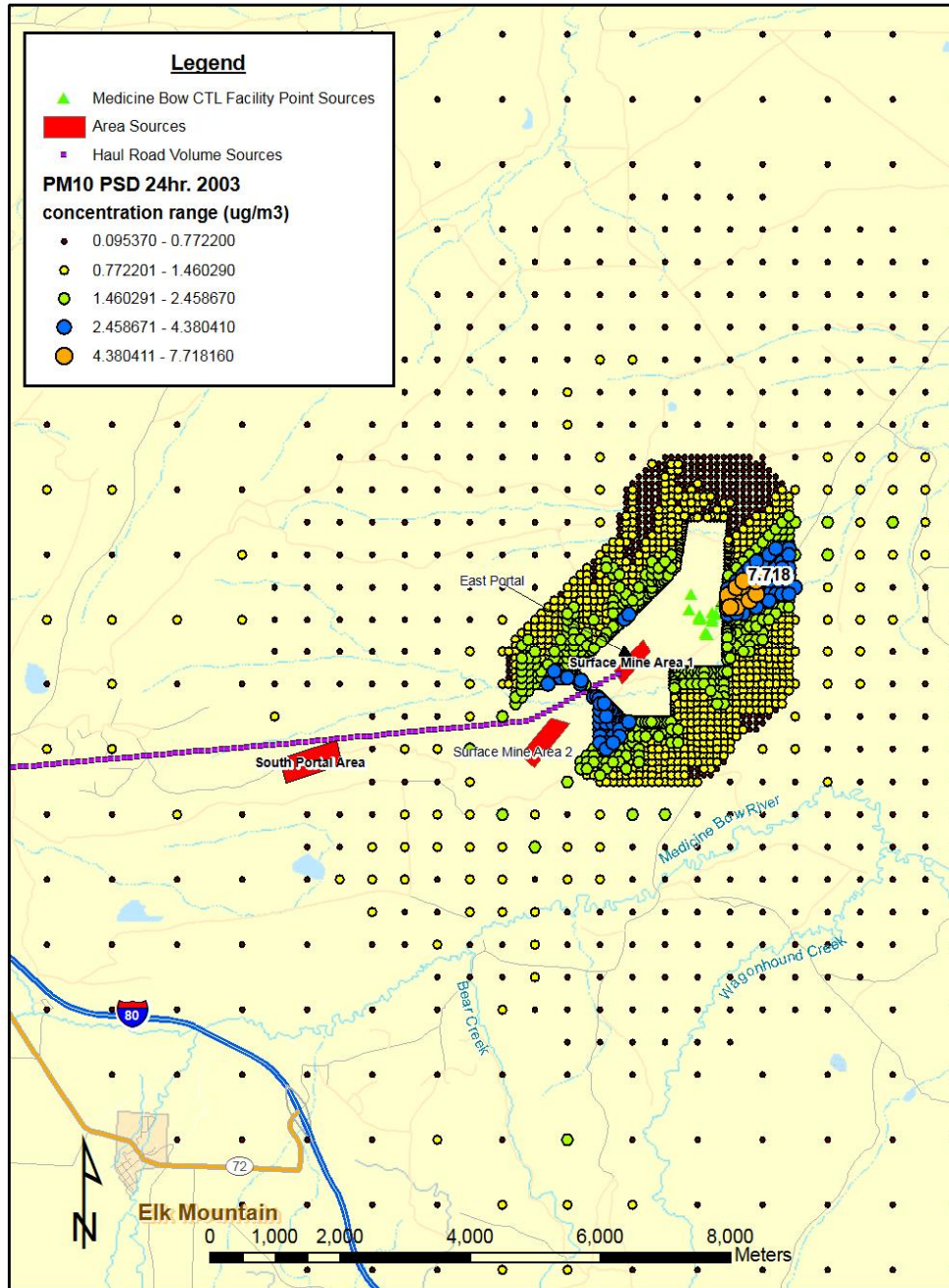
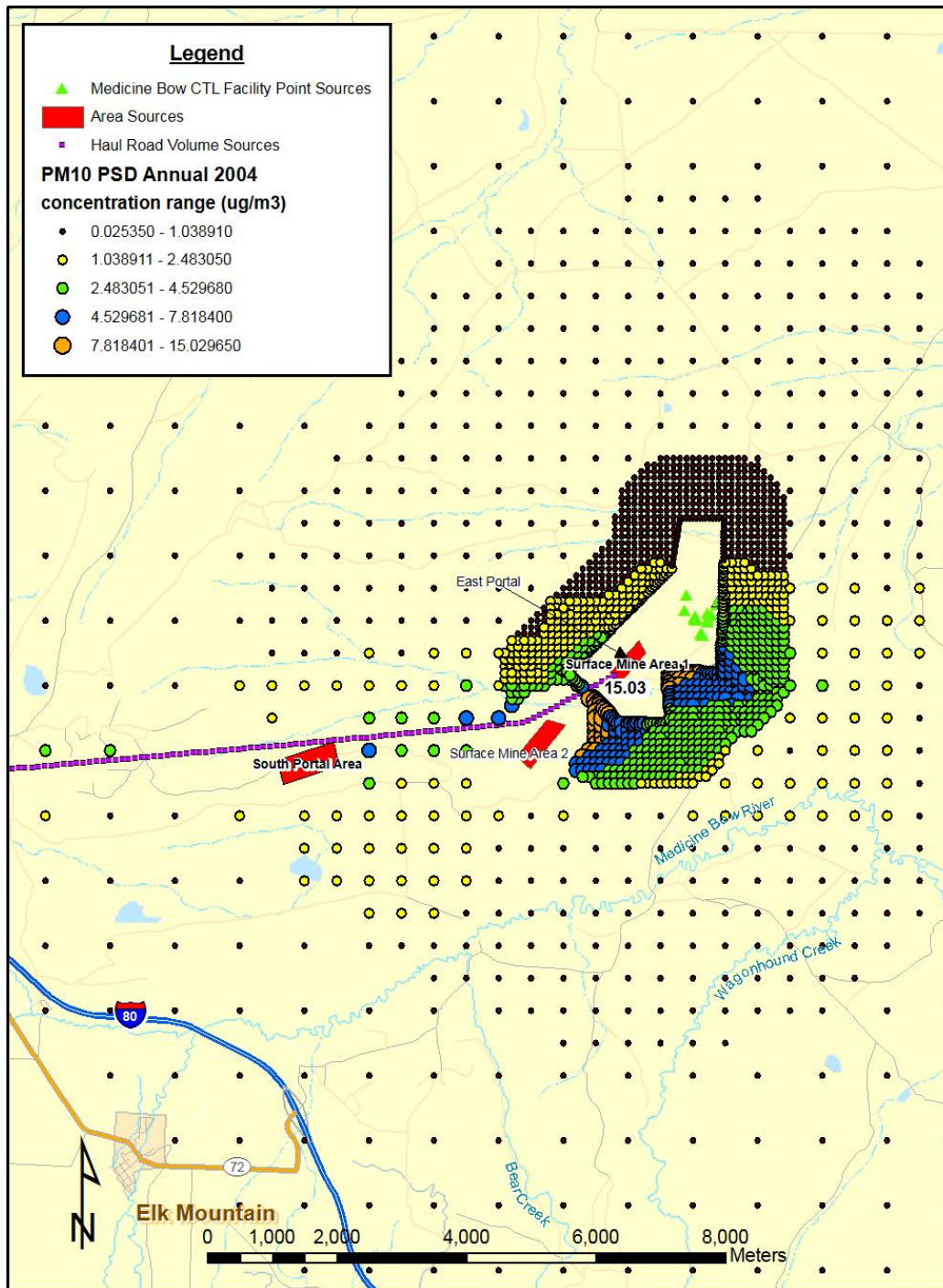


Figure 6.11 – 2005 Maximum PM₁₀ Annual Impacts (PSD)



6.6.3 CO Modeling Demonstration

Table 6.14 presents the maximum predicted 1-hour and 8-hour average cumulative CO concentrations compared to the NAAQS and WAAQS. Emissions in this analysis include nearby point sources (from WDEQ emission inventory data), nearby mining operations and vehicle traffic, and the proposed Plant. The maximum predicted second-high values are presented and added to the respective 1-hour and 8-hour background concentrations for comparison to the NAAQS and WAAQS. As shown in the table, all predicted total concentrations are below the respective NAAQS and WAAQS values. No PSD increment analysis was conducted for CO, as no PSD increments exist for CO.

Table 6.14 – Predicted CO Concentrations Compared to the NAAQS / WAAQS

Averaging Period	Data Period			Receptor Location (m)		Predicted Concentration (ug/m3)	Background Concentration (ug/m3)	Total Concentration (Predicted + Background) (ug/m3)	NAAQS & WAAQS (ug/m3)
	Year	Month / Day	Hour	East	North				
8 Hour	2000	10/3	24	392955.4	4622205	3366.56	916	4,282.56	10,000
	2001	8/3	08	392455.4	4622705	4321.5	916	5,237.5	10,000
	2003	4/11	08	390255.4	4621705	3674.5	916	4,590.5	10,000
	2004	7/26	08	392955.4	4622205	3098.76	916	4,014.76	10,000
	2005	8/8	08	392455.4	4622705	3443.05	916	4,359.05	10,000
1 Hour	2000	10/23	24	392955.4	4622205	26917.83	1946	28,863.83	40,000
	2001	8/3	05	392455.4	4622705	33584.77	1946	35,530.77	40,000
	2003	7/15	04	390355.4	4621705	27086.87	1946	29,032.87	40,000
	2004	5/10	01	392455.4	4621705	21204.38	1946	23,150.38	40,000
	2005	8/8	04	392455.4	4622705	27289.27	1946	29,235.27	40,000

Figures 6.12 and 6.13 illustrate the second high CO 1-hour impacts with respect to the NAAQS and WAAQS.

Figure 6.12 – 2001 Second High CO 1-Hour Impacts (NAAQS)

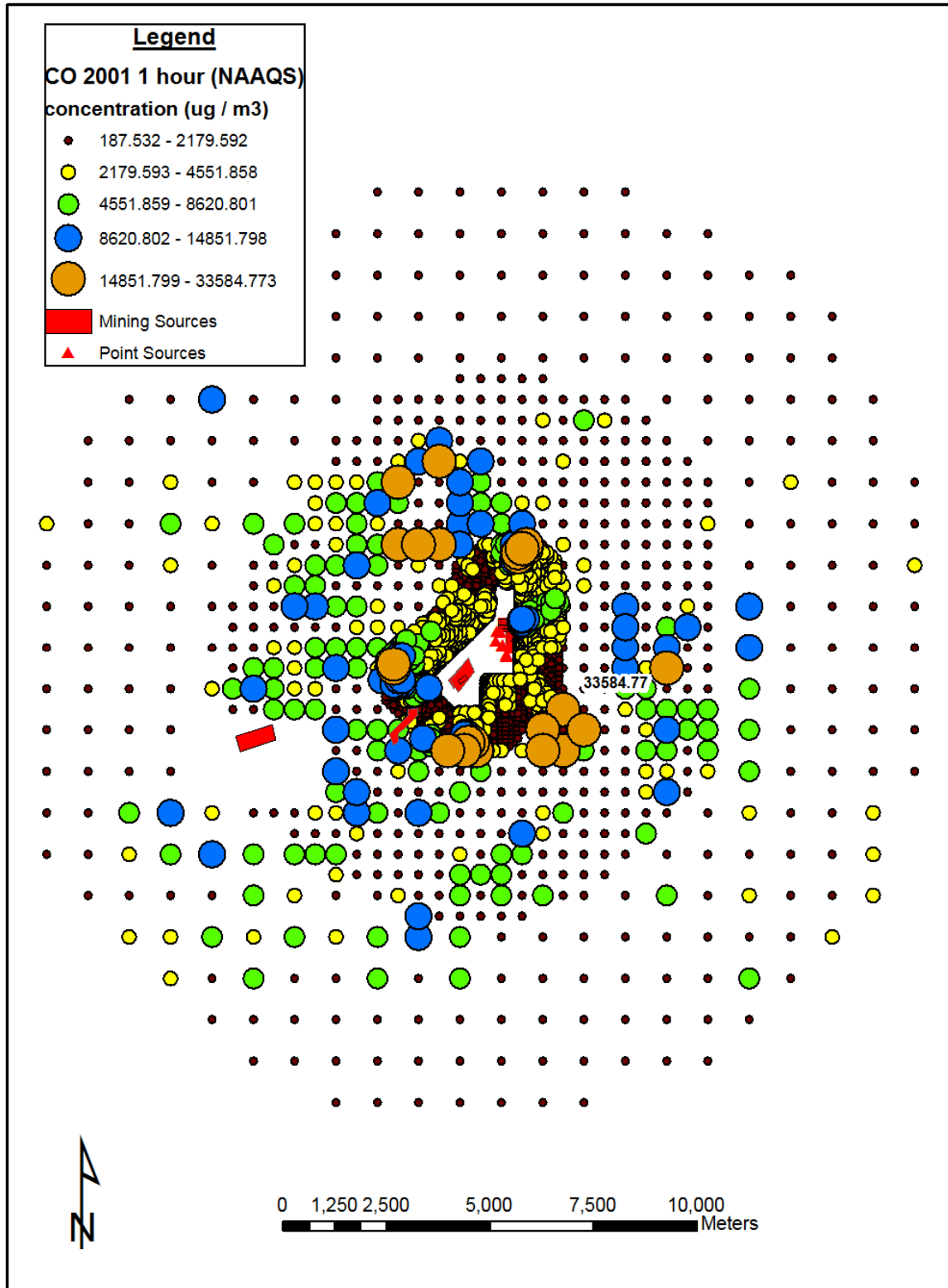
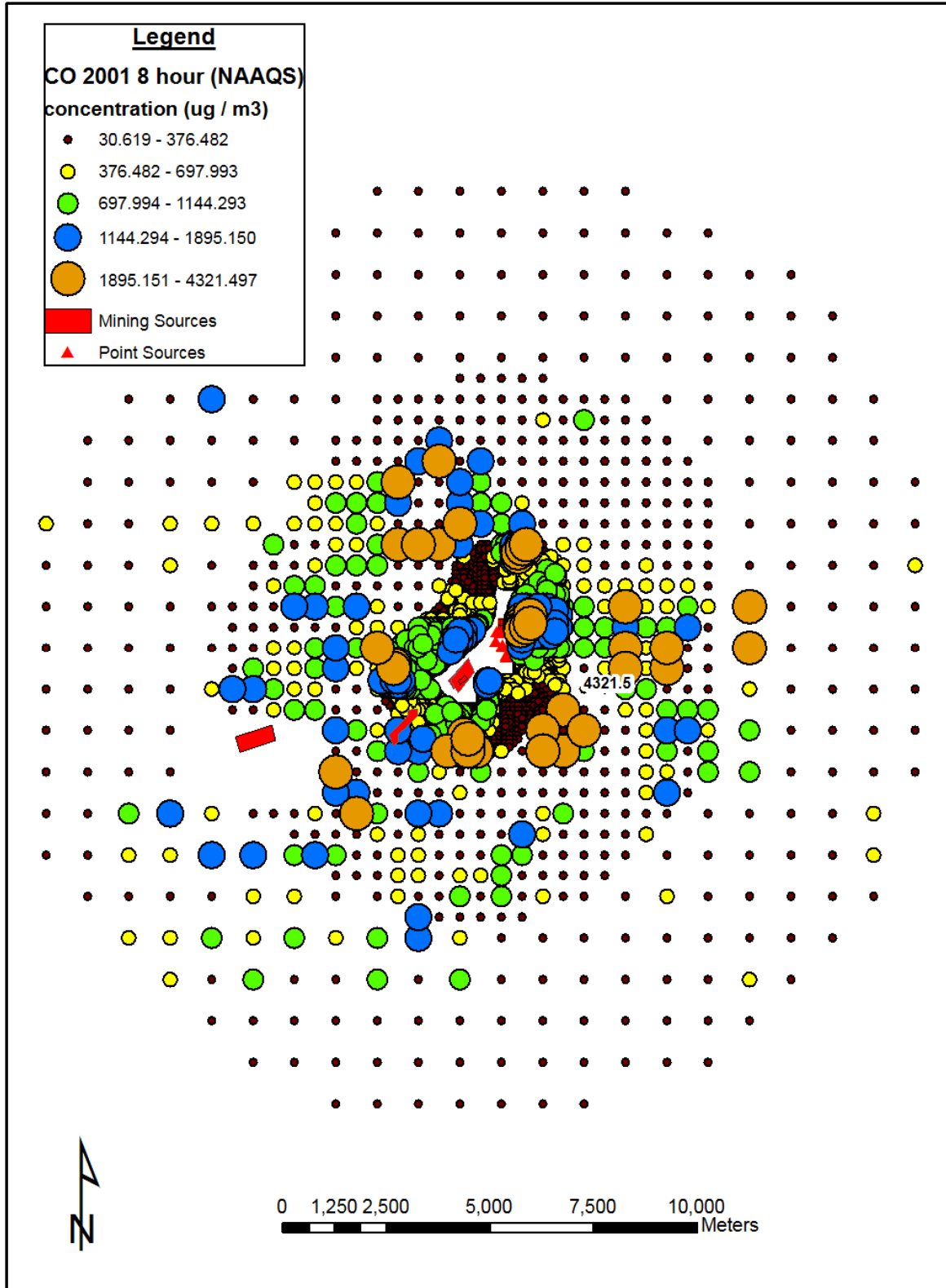


Figure 6.13 – 2001 Second High CO 8-Hour Impacts (NAAQS)



6.6.4 NO_x Modeling Demonstration

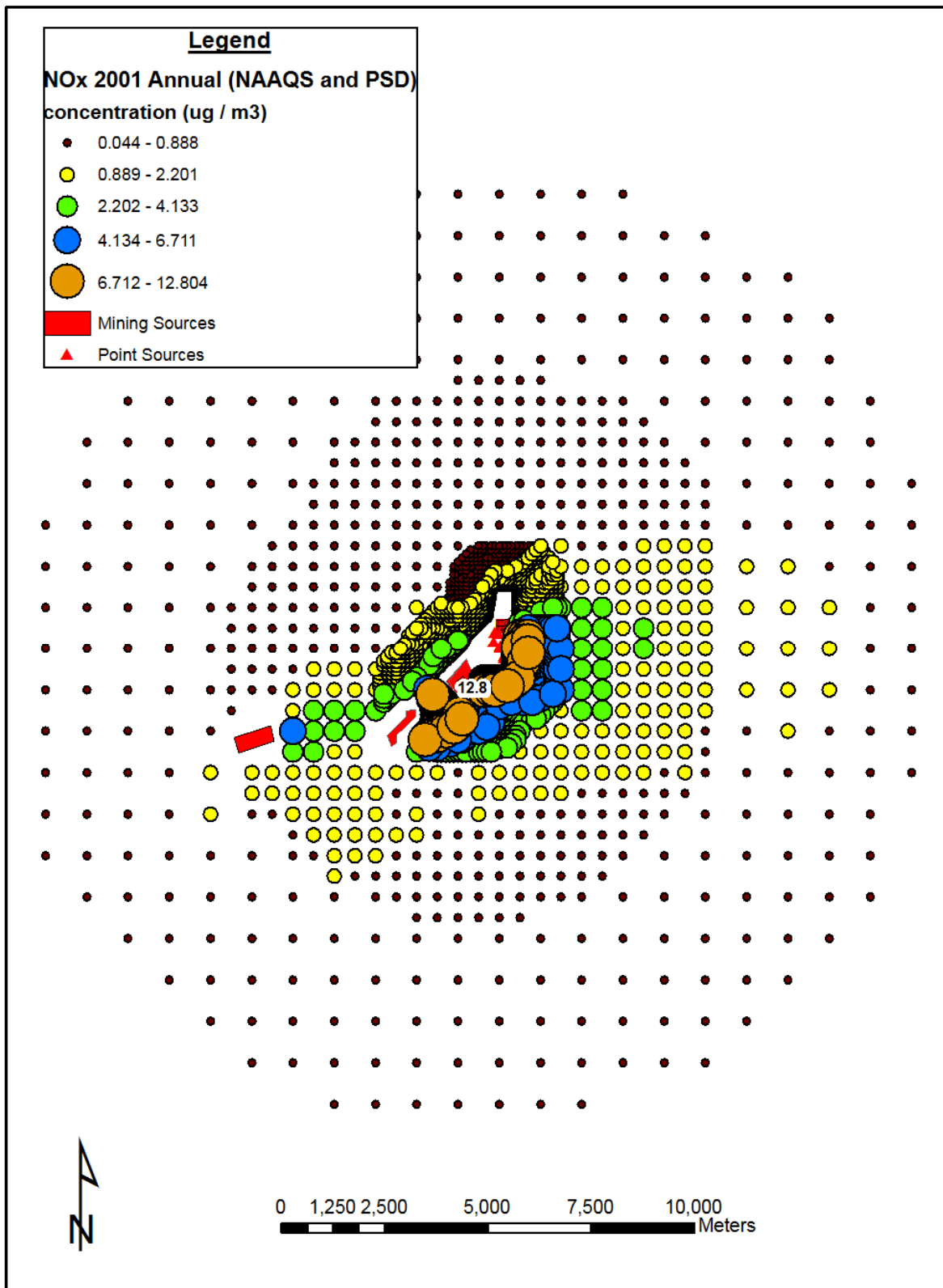
Table 6.15 presents the maximum predicted annual average NO_x concentrations compared to the NAAQS, WAAQS, and the NO_x PSD increment value. Emissions in this analysis include nearby point sources (from WDEQ emission inventory data), nearby mining operations and vehicle traffic, and the proposed Plant. The maximum predicted annual average concentrations are presented and added to the background concentration for comparison to the NAAQS and WAAQS. As shown in the table, all predicted total concentrations are well below the respective PSD increments, and the total concentrations fall well below the NAAQS and WAAQS.

Table 6.15 – Predicted NO_x Concentrations Compared to the PSD Increment, NAAQS, and WAAQS

Averaging Period	Data Period	Receptor Location (m)		Predicted Concentration (ug/m3)	PSD Increment (ug/m3)	Background Concentration (ug/m3)	Total Concentration (Predicted + Background) (ug/m3)	NAAQS & WAAQS (ug/m3)
		East	North					
Annual	2000	389455.4	4622605	12.68	25	9.43	22.11	100
	2001	389455.4	4622605	12.80	25	9.43	22.23	100
	2003	389455.4	4622605	11.48	25	9.43	20.91	100
	2004	390604.4	4623395	11.60	25	9.43	21.03	100
	2005	390604.4	4623395	12.16	25	9.43	21.59	100

Figure 6.14 illustrates the maximum annual NO_x impacts.

Figure 6.14 – 2001 Maximum Annual NO_x Impacts (NAAQS and PSD)



6.7 HAP MODELING

HAP modeling was conducted using AERMOD and the five years of meteorological data described previously for the criteria pollutant modeling.

6.7.1 HAP Emission Sources

During normal operations, the largest HAP emission sources are fugitive emissions from equipment leaks and tanks. A ground-based volume source was modeled to represent fugitive HAP emissions associated with process equipment leaks. This fugitive HAP volume source is geographically located in the synthesis process areas of the Plant and was given a release height of 2 meters. Total equipment leak emissions for each pollutant were emitted from this allocated volume source. Table 6.4 has a complete listing of the volume sources for this modeling analysis.

For tank emissions, eleven volume sources were created for the storage tank emissions. Eight tanks will contain gasoline, two will store methanol, and the remaining tank is a heavy gasoline tank. Total emissions for each pollutant were divided equally among the eleven tank volume sources. Each tank volume source release height was set equal to the tank's height.

6.7.2 HAP Risk Assessment Procedures

HAP emissions were modeled and compared to the appropriate corresponding USEPA thresholds in order to evaluate the potential health risks due to short-term and long-term exposures. Benzene, formaldehyde, xylene, toluene, and methanol maximum 1-hour (short-term) averaged concentrations were compared to the Reference Exposure Levels (RELs) obtained from the EPA Air Toxics Database, Table 2 (EPA, 2005a). An REL is defined as the concentration level at or below which no adverse health effects are anticipated for specified exposure duration. The REL is designed to protect the most sensitive individuals in the population. Exceeding the REL does not automatically indicate an adverse health impact.

No RELs are available for ethyl benzene and n-hexane. Instead, the available Immediately Dangerous to Life or Health values divided by 100 (IDLH/100) were used. Dividing by 100 is a very conservative approach to reduce a pollutant's concentration threshold of concern to only 1 percent of the level that is considered to be "immediately dangerous." IDLH values are determined by the National Institute for Occupational Safety and Health (NIOSH) and were obtained from the EPA's Air Toxic Database (EPA, 2005a). The maximum of the two short-term (g/sec) emission rates due to cold startup and normal operations for each pollutant and source were modeled and are shown in Table 6.16. For example, for a particular pollutant, several sources' emissions will be highest during startup (generators) and other sources' emissions are highest during normal operations (tank operations at full plant production). For each type of source, the highest emission rates (from startup, malfunction or normal operations) were modeled simultaneously to conservatively estimate air quality impacts.

Table 6.16 – Source HAP Short-Term (Maximum) Emission Rates

Source ID ¹ (in model)	Formaldehyde (g/sec)	Benzene (g/sec)	Methanol (g/sec)	n - Hexane (g/sec)	Toluene (g/sec)	Ethyl benzene (g/sec)	Xylene (g/sec)
CTG1	0.007	0.0012	0.0	0.0	0.013	0.00317	0.0063
CTG2	0.007	0.0012	0.0	0.0	0.013	0.00317	0.0063
CTG3	0.007	0.0012	0.0	0.0	0.013	0.00317	0.0063
GHEAT1	0.000195	0.0000054	0.0	0.00467	0.0000088	0.0	0.0
GHEAT2	0.000195	0.0000054	0.0	0.00467	0.0000088	0.0	0.0
GHEAT3	0.000195	0.0000054	0.0	0.00467	0.0000088	0.0	0.0
GHEAT4	0.000195	0.0000054	0.0	0.00467	0.0000088	0.0	0.0
GHEAT5	0.000195	0.0000054	0.0	0.00467	0.0000088	0.0	0.0
Z8901	0.0	0.0	0.0	0.0	0.0	0.0	0.0
BSG1	0.1297	0.00052	0.0	0.000273	0.001	0.0	0.00045
BSG2	0.1297	0.00052	0.0	0.000273	0.001	0.0	0.00045
FIREPUMP	0.00057	0.00045	0.0	0.0	0.0002	0.0	0.000128
AB	0.00061	0.000017	0.0	0.0147	0.000028	0.0	0.0
REGH	0.0002	0.0000056	0.0	0.0048	0.000009	0.0	0.0
REAH	0.000115	0.0000032	0.0	0.00277	0.0000052	0.0	0.0
HGT	0.000021	0.0000006	0.0	0.0005	0.0000009	0.0	0.0
Z8902	0.0	0.0	0.0	0.0	0.0	0.0	0.0
BSG3	0.1297	0.00052	0.0	0.000273	0.001	0.0	0.00045
CO2V	0.0	0.0	0.0	0.0	0.0	0.0	0.0
T_A	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_B	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_C	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_D	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_E	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_F	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_G	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_H	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_J	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_J	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
T_K	0.0	0.00138	0.00624	0.0013	0.0015	0.000099	0.00042
V1	0.0	0.3	0.3	0.0	0.0	0.0	0.0

1. Tanks are shown as sources T_A through T_K. V1 is the equipment leak volume source.

6.7.3 HAP Modeling Results

6.7.3.1 Maximum 1-Hour HAP Concentrations

Table 6.17 shows the highest short-term (1-hour) averaged concentrations using worst-case assumptions and the corresponding RELs. Each of the seven modeled HAPs has a predicted maximum 1-hour concentration well below the applicable REL.

Table 6.17 – Source HAP Emission Rates

HAP	Maximum 1-hour Averaged Modeled Concentrations ($\mu\text{g}/\text{m}^3$)	Reference Exposure Levels (RELs) ($\mu\text{g}/\text{m}^3$)
Benzene ¹	311.5	1,300
Toluene ¹	5.73	37,000
Ethylbenzene ²	0.38	35,000
Xylene ¹	1.61	22,000
n-Hexane ²	4.98	39,000
Formaldehyde ¹	80.37	94
Methanol ¹	311.5	28,000

1. EPA Air Toxics Database, Table 2 (EPA, 2005b).

2. No REL available for these HAPs. Values shown are from (IDLH/100) EPA Air Toxics Database, Table 2 (EPA, 2005b).

6.7.3.2 Maximum Annual HAP Concentrations

Annually averaged modeled HAP concentrations due to normal operations were compared to the Reference Concentrations for Chronic Inhalation (RfCs). An RfC is defined by the EPA as the daily inhalation concentration (maximum annually averaged for this analysis) at which no long-term adverse health effects are expected. RfCs exist for both non-carcinogenic and carcinogenic effects on human health (EPA, 2005b). Annually averaged modeled benzene, methanol, toluene, ethylbenzene, xylene, n-hexane, and formaldehyde concentrations were compared to the non-carcinogenic RfCs shown in Table 6.18. Maximum annual predicted concentrations are well below the applicable RfCs for each pollutant.

Table 6.18 – Annually Averaged Modeled Concentrations

HAP	Maximum Annually Averaged Modeled Concentrations ($\mu\text{g}/\text{m}^3$)	Non-Carcinogenic (RfCs) ($\mu\text{g}/\text{m}^3$) ¹
Benzene	13.56	30
Toluene	0.101	400
Ethyl benzene	0.007	1000
Xylene	0.028	100
n-Hexane	0.117	200
Formaldehyde	0.042	9.8
Methanol	13.61	4000

1. EPA Air Toxics Database, Table 1 (EPA, 2005c).

6.7.3.3 Carcinogen Analysis

RfCs for suspected carcinogens benzene and formaldehyde are expressed as unit risk factors (URS) and accepted methods for risk assessment are used to evaluate the incremental cancer risk for these pollutants. Since the closest residence, viewed in aerial photographs, is 3.3 km to the

south of the Plant, the maximum annually averaged modeled concentration predicted at a distance of 3 km and beyond for Benzene and the maximum annually averaged modeled concentration for Formaldehyde are multiplied by EPA’s URFs (based on 70-year exposure), and then multiplied by an adjustment factor which represents the ratio of projected exposure time to 70 years.

The adjustment factors represent two scenarios: a most likely exposure (MLE) scenario and one reflective of the maximally exposed individual (MEI). The MLE duration is assumed to be 9 years, which corresponds to the mean duration that a family remains at a residence (EPA, 1993). This duration corresponds to an adjustment factor of $9/70 = 0.13$. The duration of exposure for the MEI is assumed to be 70 years and the corresponding adjustment factor is 1.0.

A second adjustment is made for time spent at home versus time spent elsewhere. For the MLE scenario, the at-home time fraction is 0.64 (EPA, 1993), and it is assumed that during the rest of the day the individual will remain in an area where annually averaged HAP concentrations would be one-quarter as large as the maximum annual average concentration. Therefore, the MLE adjustment factor is calculated as follows.

$$\text{MLE Adjustment Factor} = (0.13) \times [(0.64 \times 1.0) + (0.36 \times 0.25)] = 0.095.$$

The MEI scenario assumes that the individual is at home 100 percent of the time, for the final adjustment factor of $(1.0 \times 1.0) = 1.0$.

The values for the cancer risk assessment from benzene and formaldehyde emissions from the proposed Plant are shown in Table 6.19.

Table 6.19 – Cancer Risk Assessment Values

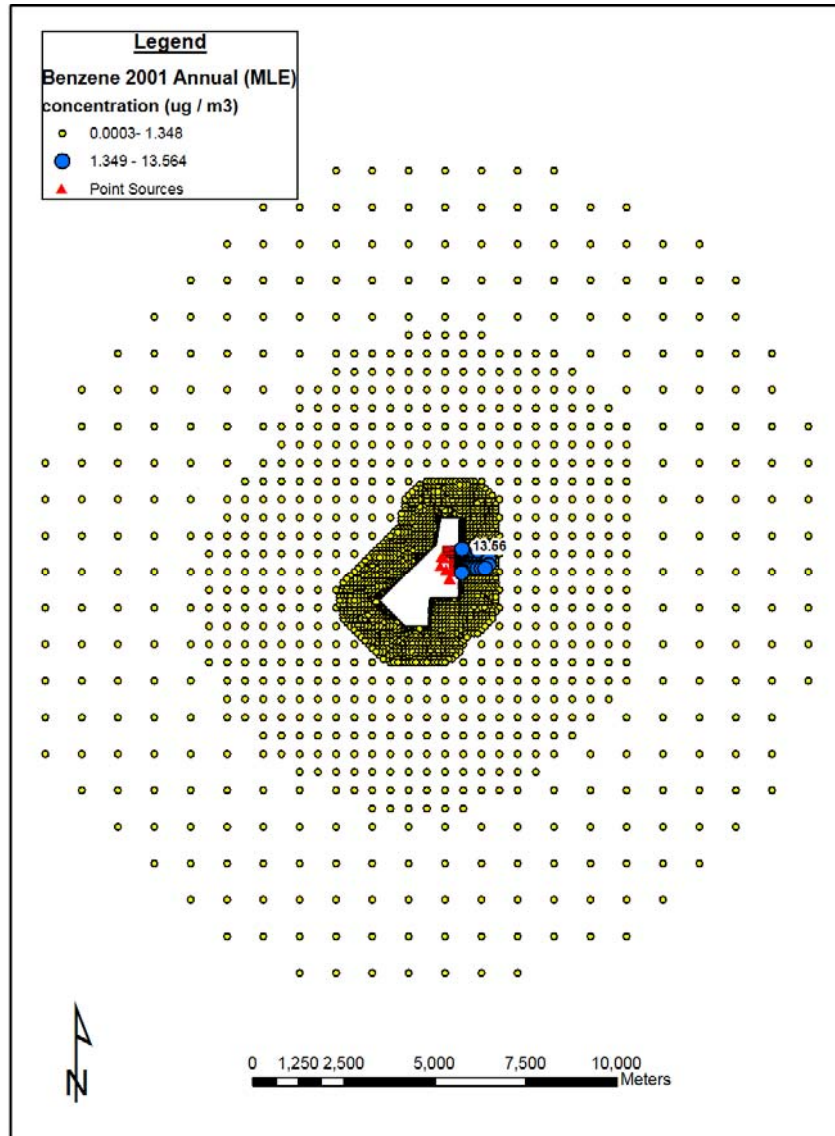
Analysis ¹	HAP	Carcinogenic RfC (Risk Factor) ² (1/μg/m ³)	Exposure Adjustment Factor	Maximum Annually Averaged Modeled Concentrations (μg/m ³)	Estimated Long-Term Exposure Risk
MLE	Benzene	7.80E-06	0.095	0.21611	1.60E-07
MLE	Formaldehyde	5.50E-09	0.095	0.00615	3.21E-12
MEI	Benzene	7.80E-06	1	0.21611	1.69E-06
MEI	Formaldehyde	5.50E-09	1	0.00615	3.38E-11

1. EPA Air Toxics Database, Table 1 (EPA, 2005c).

Figures 6.15 and 6.16 show the receptor locations with respect to the Plant including the maximum annually averaged concentrations for benzene for each receptor. Concentration ranges are colored based on the incremental cancer risk analysis. Figure 6.15 corresponds to the MLE and Figure 6.16 corresponds to the MEI. Each blue dot represents receptors that have concentrations that are at a 1×10^{-6} (1-in-a-million) risk or greater of developing cancer. Yellow receptors indicate a lower risk of developing cancer. Formaldehyde concentrations do not translate to the 1×10^{-6} risk threshold and therefore are not shown graphically.

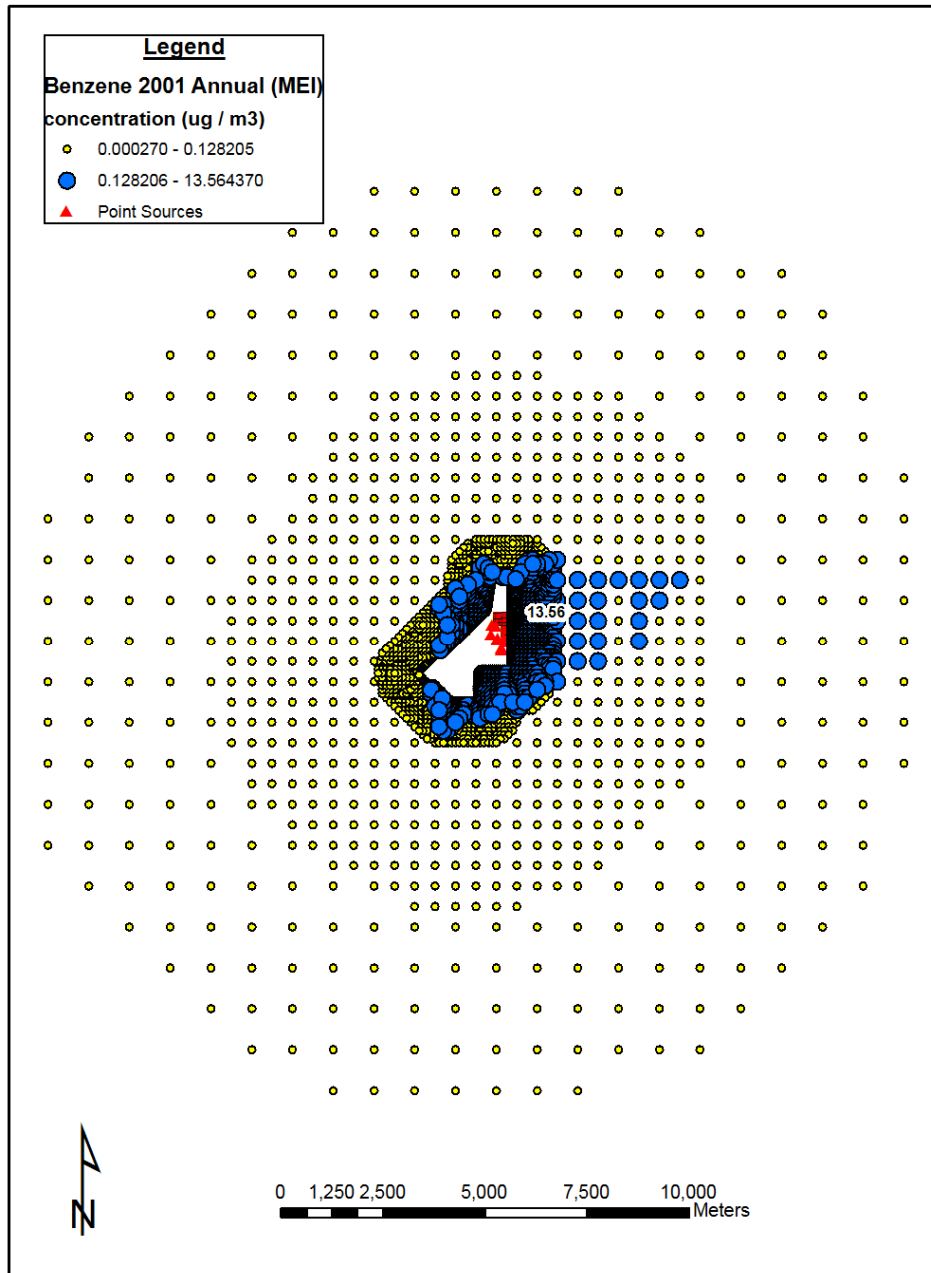
For the MLE analysis; a concentration of $1.349528 \mu\text{g}/\text{m}^3$ corresponds to a 1×10^{-6} risk of developing cancer due to benzene exposure from Plant emissions.

Figure 6.15 – MLE Receptors for Benzene



For the MEI exposure analysis; a concentration of $0.128205 \mu\text{g}/\text{m}^3$ corresponds to 1×10^{-6} risk.

Figure 6.16 – MEI Receptors for Benzene



6.7.4 HAP Conclusion

All maximum 1-hour and maximum annual predicted HAP concentrations are below the applicable RELs and RfCs, respectively. Based on these recognized EPA thresholds, short-term HAP exposure resulting from Plant emissions meets applicable criteria.

With regard to carcinogenic pollutants, predicted formaldehyde concentrations do not exceed a 1×10^{-6} risk at any modeled receptor. In contrast, benzene concentrations do exceed this risk

threshold at some locations. MLE greater than 1×10^{-6} risk occurs only along the east side of the Plant, while MEI exposure greater than 1×10^{-6} risk occurs along the south, east, and north Plant boundaries. The 1×10^{-6} MEI risk begins to fade away at 500 meters from the south and north Plant boundaries. To the east, MEI exposures greater than 1×10^{-6} risk extend beyond 500 meters out to 5 km.

As mentioned earlier, the closest residence is 3.3 km to the south of the Plant. Consequently, occupants of this residence would have a less than 1×10^{-6} risk of developing cancer due to exposure to Plant emissions of benzene or formaldehyde. As shown in the wind rose in Figure 6.6, prevailing winds blow from the west or west-southwest more than 52 percent of the time. Winds blowing from the north are rare.

6.8 IMPACTS TO SOIL AND VEGETATION

Areas surrounding the proposed Plant are of limited agricultural and commercial value and are shown in Figure 6.17 (the facility source location is indicated by coordinates). The terrain in the immediate Plant vicinity is generally rolling with a fairly uniform land cover. Views of the area presented in Figures 6.1 and 6.2 and comparison with Figure 6.17 suggests the general lack of commercial or recreational use in the project vicinity.

The potential to emit from the Plant includes four criteria pollutants (CO, NO_x, SO₂, and PM/PM₁₀) that will be emitted in excess of PSD significant impact levels. The impacts of each of these pollutant emissions from the project are below the primary and secondary NAAQS shown. Secondary NAAQS standards are expressly designed to protect public welfare, including protection of soils, vegetation, and other environmental and man-made attributes.

Figure 6.17 - Aerial View of Land Use Immediately Surrounding the Plant



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6.8.1 Soil Impacts

The US Department of Agriculture (USDA) Natural Resources Conservation Service (NRCS) has compiled a detailed list of agricultural yields and soil types for portions of Carbon County. Of the over 540,000 acres surveyed, land capability is classified as Class 3 or worse (no soils are designated as Class 5). Soil within the surveyed areas of the county is classified as follows:

- Class 3: Soils have severe limitations that reduce the choice of plants or that require special conservation practices, or both.
- Class 4: Soils have very severe limitations that reduce the choice of plants or that require very careful management, or both.
- Class 6: Soils have severe limitations that make them generally unsuitable for cultivation. Rangeland or forestry improvements can be applied.
- Class 7: Soils have very severe limitations that make them unsuitable for cultivation. They can be used for forestry or grazing, but rangeland improvements are impractical.
- Class 8: Soils and miscellaneous areas have limitations that nearly preclude their use for commercial crop production.

Only 1 percent of the surveyed land produces alfalfa or hay without using irrigation. With regard to irrigated land (accounting for a small portion of the county), the most productive land produces up to 5 tons of alfalfa per acre. Assuming a value of \$130/ton of alfalfa, maximum cropland production value is \$650/acre on the best-producing land included in the NRCS survey of Carbon County. Based on this information, most Carbon County land does not have significant commercial value. NRCS crop yields are provided in Appendix K. The NRCS soil survey is provided in Appendix L.

Little information on direct gaseous air pollutant effects on soil is available in the current literature. While certain soils can be an effective sink for gaseous pollutants such as NO₂ and some studies have been done, accurate methods for routinely quantifying the effects of NO₂ and other pollutants on soil in the field do not exist. The rate of adsorption is dependent on the distance from the source, concentrations in the air, soil properties, vegetative cover, and the prevailing hydrological and meteorological conditions. No significant impacts on soils from exposures to acidic gases such as NO₂ occur unless the soils experience a large decrease in buffering capacity and the pH of precipitation drops dramatically (Smith, 1981). Because NO_x ambient concentration increases attributable to the Plant and surrounding sources represent less than 13 percent of the secondary NAAQS for this pollutant, soil impacts are expected to be low.

6.8.2 Vegetation Impacts

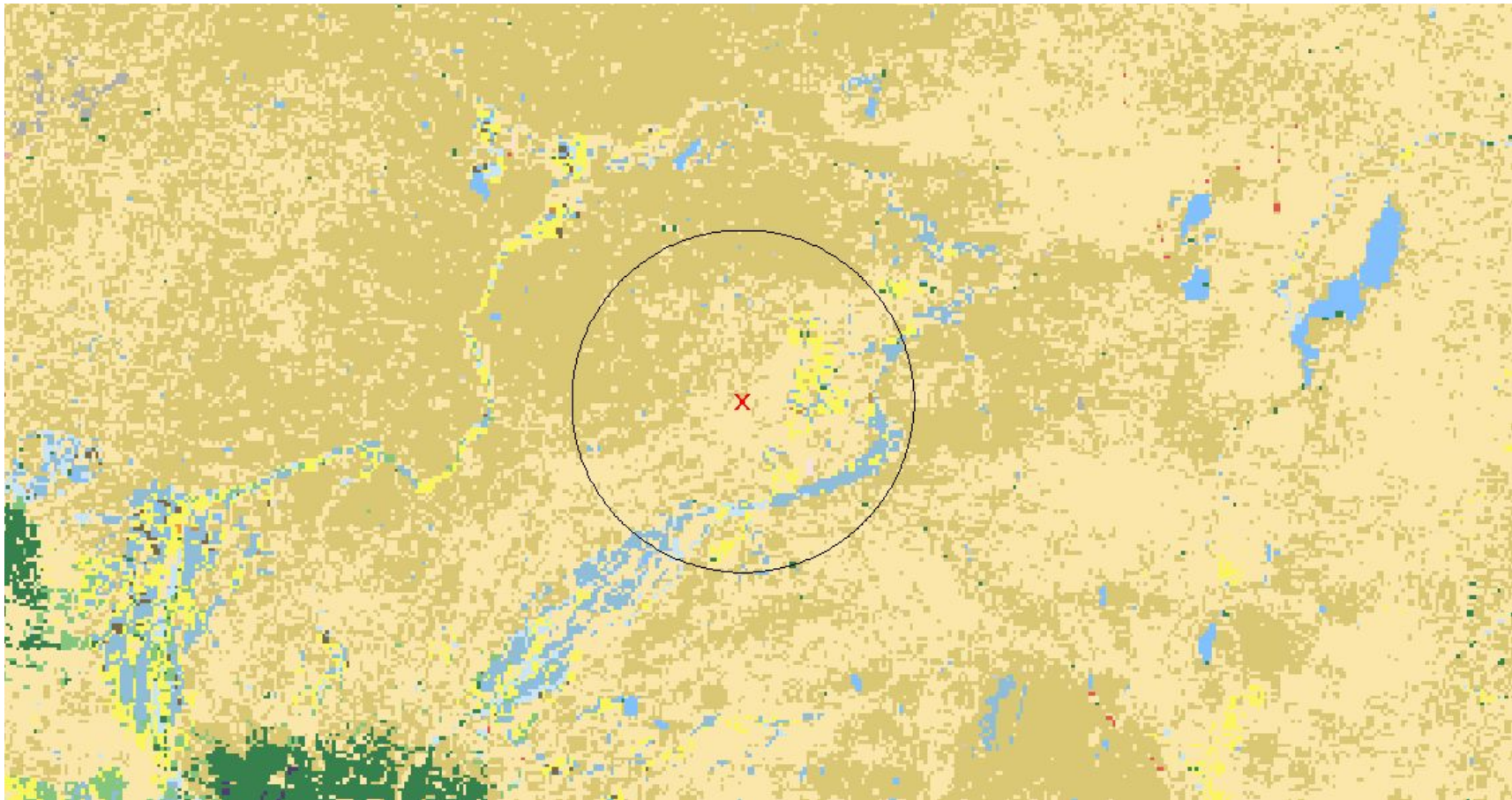
The Plant is located within a gently rolling landscape. The commercial productivity of the lands around the immediate Medicine Bow area is very low. There are some areas with limited agriculture within 10 km of the site. The closest cropland is approximately 2.3 km from the Plant. Primary land use and vegetation cover is depicted in Figure 6.18, which shows that the predominant land use is fallow or shrubland. Only a small percentage of the land surrounding the facility is cropland. A review of the Wyoming Department of Agriculture and livestock census suggests that Carbon County lands are generally low in productivity (see Appendix M).

Damage or injury to plants from air pollutants is caused primarily through foliage injury and not by absorption through the plant roots. As a result, ambient air concentrations of pollutants are the primary indicators of potential impact. The concentration of a pollutant and the duration of the exposure period are collectively referred to as the dose; the lowest dose that produces an effect is called the threshold dose. However, because of the relationship between concentration and time, there is no single threshold dose for an effect.

Reduction in yield, whether quantitative or qualitative, is also of prime importance but is difficult to measure. Foliar damage to root crops, for example, may bear no relationship to the amount of economic damage incurred. If injury occurs near harvest time, there may be no detectable yield loss (Capron and Mansfield, 1976).

Figure 6.18 - Land Use and Vegetation Cover near the Plant Site

(Light yellow denotes cropland, darker green is forest, blue is water, light tan is fallow, and dark tan is shrubland)



Distance Ring Radius is 10 km

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6.8.2.1 Effects of NO_x

The direct effects of NO_x on vegetation are usually associated with and confined to areas near specific industrial sources. For example, vegetation injury from exposure to high NO₂ concentrations has been observed near nitric acid factories and arsenals, but there is little published information regarding vegetation injury in the field due to NO or other NO_x (U.S. EPA, 1982a).

Many reports, however, have substantiated NO_x effects on vegetation grown in laboratory conditions (Hill and Bennett, 1970; Capron and Mansfield, 1976; Czeh and Nothdruff, 1951; Taylor et al., 1975; Kress, 1982). A threshold value of 191 µg/m³ for long-term (10,000-hour) laboratory exposures of crops and trees has been widely used (U.S. EPA, 1982a). The maximum modeled NO_x increase from the proposed Plant and surrounding sources is low (12.80 µg/m³ based on annual averaging) and well below the threshold value (191 µg/m³). Therefore, no detrimental effects on vegetation in the project area will likely result from NO_x emissions from the Plant.

6.8.2.2 Effects of SO₂

SO₂ enters vegetation in gaseous form through openings in the plant's leaf surface called stomata. Once inside the leaf, SO₂ contacts wet, cellular membranes, and sulfites and sulfates may be formed. The formation of these compounds can cause changes in the plant's metabolic system that will produce physiological dysfunctions (U.S. EPA, 1982b).

Short-term (1-hour) peak SO₂ concentrations are particularly important when assessing potential vegetation impacts (Houston, 1974). Laboratory experiments have demonstrated greater relative toxicity of short-term exposures at high SO₂ concentrations than long-term exposures with the same total treatment (Zahn, 1970; McLaughlin et al., 1979; Sij, Kanemasu, and Goltz, 1974; Wilhour et al., 1978; Miller et al., 1979; Sprugel et al., 1980; Houston, 1974; Berry, 1972; Temple, 1972).

The maximum SO₂ concentration increase from the proposed Plant (4.51 µg/m³ based on annual averaging) is far less than the lowest concentration of 240 µg/m³ (Miller et al., 1979; Sprugel et al., 1980) that has been shown to reduce yield in the most sensitive agricultural crop, soybean, and the 390 µg/m³ (Houston, 1974) forest species threshold.

6.8.2.3 Effects of PM/PM₁₀

Adverse impacts on vegetation from PM/PM₁₀ are most often associated with sustained accumulation of particles such as dust or fly ash on the leaf surface. Such particle accumulation on leaves can result in reduced gas exchange, increased leaf temperature, reduced photosynthesis, and eventual yellowing and tissue desiccation (Parish, 1910; Darley, 1966).

The maximum modeled PM/PM₁₀ impact from the proposed Plant is 6.15 µg/m³ (annual average). At less than 13 percent of the WAAQS, this increase in particulate concentration is not expected to cause plant injury.

6.9 REFERENCES FOR CHAPTER 6

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7.1 BACKGROUND

NOTE: The far field modeling analysis presented in this section is based on emissions and process parameters described in the original Permit Application dated June 19, 2007. This analysis is presented in its entirety to comprehensively describe the modeling conducted for the June 2007 permit application. The far field modeling analysis was supplemented on October 17, 2007 in response to comments from the WDEQ. These responses are included in Appendix J.

MBFP believes that this far field criteria pollutant modeling analysis should be considered to be sufficient with regard to criteria pollutants emitted by the proposed facility based on the revised process design. A comparison of revised emission rates and previously modeled emission rates is presented in Appendix I.

As discussed in Section 1.2 of this application, the project is a major stationary source under the PSD program and therefore has completed an analysis of potential long-range impacts in support of a requested air quality construction permit.

Air quality impact analysis for Class I and sensitive Class II areas within 300 km from the project was conducted using the EPA long-range dispersion model, CALPUFF. The CALPUFF analysis included 8 Class I areas and 1 Class II area. The nearest Class I area, which is Mount Zirkel Wilderness, is located approximately 93 km southwest from the facility. Class I and sensitive Class II areas within 300 km from the facility are listed in Table 7.1. There is one sensitive Class II area within 300 km from the facility, named Savage Run, which is located approximately 60 km south from the facility.

In addition, soils and vegetation analysis was conducted. Additional impact analysis was not conducted because modeling results did not show significant air quality impact on Class I and sensitive Class II areas. Therefore, visibility analysis for scenic and important views and impact analysis for water was not conducted and the additional analyses areas are not listed in the Table 7-1.

Table 7.1 – Class I Areas and Sensitive Class II Areas Within 300 km

	Areas
Class I Areas	Rocky Mountain National Park, Rawah Wilderness, Flat Tops Wilderness, Eagles nest Wilderness, Mount Zirkel Wilderness, Maroon Bell-Snowmass Wilderness, Bridger Wilderness, and Fitzpatrick Wilderness
Sensitive Class II Areas	Savage Run

CALPUFF modeling runs were completed for each Class I or Class II area using a worst-case emission inventory. Detailed descriptions of the emission inventories for the modeling analysis were shown in Section 7.2.2.

7.2 DESCRIPTION OF PROPOSED PROJECT

7.2.1 Site Location

The facility will be located approximately 7.5 miles north of Interstate 80, exit 260 (Elk Mountain) on County Road #3 in Section 29 of Township 21 north and Range 79 west in Carbon County, Wyoming. LULC shapefile plotted in ArcGIS shows that most of the area surrounded by the facility is shrub/brush. MBFP will be located in an area that is designated as attainment of all National Ambient Air Quality Standards (NAAQS). The project location for the site is shown in Figure 1.1.

7.2.2 Source Emissions

The facility will consist of the Plant and the Underground coal mine (Saddleback Hills). Construction of both the Plant and the Mine will take about three years. The combustion source at the site will be fuels with syngas during normal operation and pipeline quality natural gas during startup and in the event of a loss of fuel gas (syngas). The facility will require approximately 1000 hours to start all of the process. Once the facility is started, it will not shut down unless there are planned maintenance activity or in the event of a malfunction. The startup is discussed in more detail in Section 2.17 of this application.

Emissions sources will include three (3) combustions turbines, twelve (12) heaters, three (3) generators, one (1) firewater pump, one (1) Emergency Flare, one (1) CO₂ vent, and one (1) Sulfur Plant Incinerator. Detailed emission calculations for these sources are included in Appendix B.

7.2.3 Sources Included in CALPUFF Modeling

Required emissions in CALPUFF correspond with the needed analysis and include maximum short-term rates for increment and visibility impacts, as well as maximum annual emissions for species deposition and increment comparison. Because of the various operations involved and potential occurrence during a specific period, the CALPUFF modeled sources and emissions included potential overlapping operations.

The emission rate derivation is shown in Table 7.2 and the modeled emissions are shown in Table 7.3 (short-term) and Table 7.4 (annual). The overlapping scenarios include the Turbine/HRSG 3 aggregated NO_x emissions and the additive source emissions to account for normal and startup scenarios.

For example, in Table 7.2 the NO_x emission rates shown for source Turbine and HRSG Train 3 feature a higher rate than for the other two turbines. This is done to reflect startup scenarios that would include 18-hours of normal operations and 6-hours of startup operations. Aggregating the two and rating the hourly emissions for each type of operation returns the 24-hour emission rate shown. And the annual emission inventory includes both normal and startup sources, as operating with the annual hours provided.

The CALPUFF modeling also included speciation of emissions according to the National Park Service (NPS)'s Particulate Matter Speciation (PMS) method for natural gas combustion turbines. Applying the PMS methodology, 67% of total SO₂ was speciated into SO₂ and 33% of

total SO₂ was speciated into SO₄. Also, the total PM₁₀ emission was speciated into Elemental Carbon (EC) and Secondary Organic Aerosol (SOA). The SOA was speciated again into PM_{0.05}, PM_{0.01}, PM_{0.15}, PM_{0.20}, PM_{0.25}, and PM_{1.0} (indicated as PM0005, PM0010, PM0015, PM0020, PM0025, and PM0100 in the modeling, respectively). The SOA size distribution is shown in Table 7.5.

Table 7.2 – Maximum Emission Rate from All Sources

Pollutant	24-hour Averaged Emission Rate (lb/hr)									Annual Averaged Emission Rate Total Maximum		
	Startup	Normal	Startup	Normal	Startup	Normal	Total Maximum					
	24-hr	24-hr	3 and 24-hr	3 and 24-hr	24-hr	24-hr						
	NO _x	NO _x	SO ₂	SO ₂	PM ₁₀	PM ₁₀	NO _x	SO ₂	PM ₁₀	NO _x	SO ₂	PM ₁₀
Turbine and HRSG Train 1	134.56	18.15	0.04	0.04	10.00	10.00	18.15	0.04	10.00	17.51	0.04	10.00
Turbine and HRSG Train 2	134.56	18.15	0.04	0.04	10.00	10.00	18.15	0.04	10.00	17.51	0.04	10.00
Turbine and HRSG Train 3	134.56	18.15	0.04	0.04	10.00	10.00	47.25	0.04	10.00	17.51	0.04	10.00
Gasifier Preheater 1	0.74		0.01		0.11		0.74	0.01	0.11	0.04	5.04E-04	6.38E-03
Gasifier Preheater 2	0.74		0.01		0.11		0.74	0.01	0.11	0.04	5.04E-04	6.38E-03
Gasifier Preheater 3	0.74		0.01		0.11		0.74	0.01	0.11	0.04	5.04E-04	6.38E-03
Gasifier Preheater 4	0.74		0.01		0.11		0.74	0.01	0.11	0.04	5.04E-04	6.38E-03
Gasifier Preheater 5	0.74		0.01		0.11		0.74	0.01	0.11	0.04	5.04E-04	6.38E-03
H-3102 SRU Incinerator		0.13		9.51		2.04	0.13	9.51	2.04	0.13	9.51	2.04
H-5401 Frac Feed Heater	4.26	2.16	0.05	0.02	0.65	0.04	2.68	0.03	0.19	2.15	2.42E-02	0.07
H-5301 Cat Dewax Charge	0.19	0.10	0.00	0.00	0.03	0.24	0.12	0.00	0.24	0.10	1.08E-03	0.22
H-5201 Unicracker Feed	0.80	0.40	0.01	0.00	0.12	1.02	0.50	0.01	1.02	0.40	4.54E-03	0.91
H-5202 Unicracker Intermed.	2.17	1.10	0.03	0.01	0.33	2.75	1.36	0.02	2.75	1.10	1.23E-02	2.46
H-5101 Unionfiner Feed	0.25	0.13	0.00	0.00	0.04	0.32	0.16	0.00	0.32	0.13	1.42E-03	0.28
H-5102 Unionfiner Intermed	0.31	0.16	0.00	0.00	0.05	0.40	0.20	0.00	0.40	0.16	1.78E-03	0.35
Firewater Pump		6.02		0.01		0.08	6.02	0.01	0.08	0.18	3.27E-04	4.29E-05
Black-Start Generator 1										0.18	3.27E-04	4.29E-05
Black-Start Generator 2										0.18	3.27E-04	4.29E-05
Black-Start Generator 3										0.34	3.46E-04	4.34E-03
Total							98	10	38	58	10	36

Table 7.3 – 24 hour Averaged Emission Inventory for CALPUFF (3-hour and 24-hour SO₂, and 24-hour PM₁₀ and Visibility)

Sources (g/s)	SO ₂	SO ₄	NO _x	HNO ₃	NO ₃	INCPM	SOA						SOA	EC
							PM0005	PM0010	PM0015	PM0020	PM0025	PM0100		
Turbine and HRSG Train 1	3.54E-03	2.65E-03	2.29E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Turbine and HRSG Train 2	3.54E-03	2.65E-03	2.29E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Turbine and HRSG Train 3	3.59E-03	2.69E-03	5.95E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Gasifier Preheater 1	7.41E-04	5.56E-04	9.26E-02	0.00E+00	0.00E+00	1.41E-02	1.50E-03	2.50E-03	2.30E-03	1.50E-03	1.10E-03	1.10E-03	1.00E-02	3.52E-03
Gasifier Preheater 2	7.41E-04	5.56E-04	9.26E-02	0.00E+00	0.00E+00	1.41E-02	1.50E-03	2.50E-03	2.30E-03	1.50E-03	1.10E-03	1.10E-03	1.00E-02	3.52E-03
Gasifier Preheater 3	7.41E-04	5.56E-04	9.26E-02	0.00E+00	0.00E+00	1.41E-02	1.50E-03	2.50E-03	2.30E-03	1.50E-03	1.10E-03	1.10E-03	1.00E-02	3.52E-03
Gasifier Preheater 4	7.41E-04	5.56E-04	9.26E-02	0.00E+00	0.00E+00	1.41E-02	1.50E-03	2.50E-03	2.30E-03	1.50E-03	1.10E-03	1.10E-03	1.00E-02	3.52E-03
Gasifier Preheater 5	7.41E-04	5.56E-04	9.26E-02	0.00E+00	0.00E+00	1.41E-02	1.50E-03	2.50E-03	2.30E-03	1.50E-03	1.10E-03	1.10E-03	1.00E-02	3.52E-03
H-3102 SRU Incinerator	7.99E-01	5.99E-01	1.64E-02	0.00E+00	0.00E+00	2.57E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	6.42E-02
H-5401 Frac Feed Heater	2.59E-03	1.94E-03	3.38E-01	0.00E+00	0.00E+00	2.44E-02	2.45E-03	4.09E-03	3.76E-03	2.45E-03	1.80E-03	1.80E-03	1.63E-02	6.10E-03
H-5301 Cat Dewax Charge	1.16E-04	8.70E-05	1.52E-02	0.00E+00	0.00E+00	3.05E-02	3.42E-03	5.70E-03	5.25E-03	3.42E-03	2.51E-03	2.51E-03	2.28E-02	7.63E-03
H-5201 Unicracker Feed	4.86E-04	3.64E-04	6.35E-02	0.00E+00	0.00E+00	1.28E-01	1.43E-02	2.39E-02	2.20E-02	1.43E-02	1.05E-02	1.05E-02	9.56E-02	3.20E-02
H-5202 Unicracker Intermed.	1.32E-03	9.87E-04	1.72E-01	0.00E+00	0.00E+00	3.47E-01	3.88E-02	6.47E-02	5.96E-02	3.88E-02	2.85E-02	2.85E-02	2.59E-01	8.66E-02
Firewater Pump	1.52E-04	1.14E-04	1.99E-02	0.00E+00	0.00E+00	4.01E-02	4.50E-03	7.50E-03	6.90E-03	4.50E-03	3.30E-03	3.30E-03	3.00E-02	1.00E-02

SECTION SEVEN

Far Field Air Quality Impact Analysis

Table 7.4 – Annual Averaged Emission Inventory for CALPUFF (Annual NO_x, SO₂, and PM₁₀ and Deposition)

Sources (g/s)	SO ₂	SO ₄	NO _x	HNO ₃	NO ₃	INCPM	SOA						SOA	EC
							PM0005	PM0010	PM0015	PM0020	PM0025	PM0100		
Turbine and HRSG Train 1	3.42E-03	2.57E-03	2.21E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Turbine and HRSG Train 2	3.42E-03	2.57E-03	2.21E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Turbine and HRSG Train 3	3.42E-03	2.57E-03	2.21E+00	0.00E+00	0.00E+00	1.26E+00	1.41E-01	2.36E-01	2.17E-01	1.41E-01	1.04E-01	1.04E-01	9.42E-01	3.15E-01
Gasifier Preheater 1	4.23E-05	3.17E-05	5.29E-03	0.00E+00	0.00E+00	8.04E-04	8.57E-05	1.43E-04	1.31E-04	8.57E-05	6.28E-05	6.28E-05	5.71E-04	2.01E-04
Gasifier Preheater 2	4.23E-05	3.17E-05	5.29E-03	0.00E+00	0.00E+00	8.04E-04	8.57E-05	1.43E-04	1.31E-04	8.57E-05	6.28E-05	6.28E-05	5.71E-04	2.01E-04
Gasifier Preheater 3	4.23E-05	3.17E-05	5.29E-03	0.00E+00	0.00E+00	8.04E-04	8.57E-05	1.43E-04	1.31E-04	8.57E-05	6.28E-05	6.28E-05	5.71E-04	2.01E-04
Gasifier Preheater 4	4.23E-05	3.17E-05	5.29E-03	0.00E+00	0.00E+00	8.04E-04	8.57E-05	1.43E-04	1.31E-04	8.57E-05	6.28E-05	6.28E-05	5.71E-04	2.01E-04
Gasifier Preheater 5	4.23E-05	3.17E-05	5.29E-03	0.00E+00	0.00E+00	8.04E-04	8.57E-05	1.43E-04	1.31E-04	8.57E-05	6.28E-05	6.28E-05	5.71E-04	2.01E-04
H-3102 SRU Incinerator	7.99E-01	5.99E-01	1.64E-02	0.00E+00	0.00E+00	2.57E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	6.42E-02
H-5401 Frac Feed Heater	2.03E-03	1.52E-03	2.71E-01	0.00E+00	0.00E+00	9.35E-03	8.23E-04	1.37E-03	1.26E-03	8.23E-04	6.03E-04	6.03E-04	5.48E-03	2.34E-03
H-5301 Cat Dewax Charge	9.11E-05	6.83E-05	1.22E-02	0.00E+00	0.00E+00	2.73E-02	3.06E-03	5.09E-03	4.69E-03	3.06E-03	2.24E-03	2.24E-03	2.04E-02	6.81E-03
H-5201 Unicracker Feed	3.82E-04	2.86E-04	5.09E-02	0.00E+00	0.00E+00	1.14E-01	1.28E-02	2.13E-02	1.96E-02	1.28E-02	9.39E-03	9.39E-03	8.54E-02	2.86E-02
H-5202 Unicracker Intermed.	1.03E-03	7.76E-04	1.38E-01	0.00E+00	0.00E+00	3.09E-01	3.47E-02	5.78E-02	5.32E-02	3.47E-02	2.54E-02	2.54E-02	2.31E-01	7.74E-02
H-5101 Unionfiner Feed	1.20E-04	8.98E-05	1.60E-02	0.00E+00	0.00E+00	3.58E-02	4.02E-03	6.69E-03	6.16E-03	4.02E-03	2.95E-03	2.95E-03	2.68E-02	8.95E-03
H-5102 Unionfiner Intermed	1.49E-04	1.12E-04	1.99E-02	0.00E+00	0.00E+00	4.46E-02	5.01E-03	8.34E-03	7.68E-03	5.01E-03	3.67E-03	3.67E-03	3.34E-02	1.12E-02
Black-Start Generator 1	2.75E-05	2.06E-05	2.29E-02	0.00E+00	0.00E+00	5.40E-06	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.35E-06
Black-Start Generator 2	2.75E-05	2.06E-05	2.29E-02	0.00E+00	0.00E+00	5.40E-06	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.35E-06
Black-Start Generator 3	2.75E-05	2.06E-05	2.29E-02	0.00E+00	0.00E+00	5.40E-06	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.35E-06
Firewater Pump	2.91E-05	2.18E-05	4.33E-02	0.00E+00	0.00E+00	5.47E-04	5.83E-05	9.71E-05	8.93E-05	5.83E-05	4.27E-05	4.27E-05	3.88E-04	1.37E-04

Table 7.5 – Size Distribution of Secondary Organic Aerosols (SOA)

Species Name	Size Distribution (%)	Geometric Mass Mean Diameter (microns)	Geometric Std. Deviation (microns)
SO ₄	100	0.48	0.50
NO ₃	100	0.48	0.50
PM0005	15	0.05	0.00
PM0010	40	0.10	0.00
PM0015	63	0.15	0.00
PM0020	78	0.20	0.00
PM0025	89	0.25	0.00
PM0100	100	1.00	0.00

The 24-hour averaged emission rate was used for the 3-hour and 24-hour SO₂ and 24-hour PM₁₀ impact analyses, and visibility impairment impact analysis. The annual emission rate was used for the annual NO_x, annual SO₂, and annual PM₁₀ impact analyses as well as nitrogen and sulfur deposition analyses. The stack parameters of all sources are shown in Table 7.6.

7.2.4 Reference Reports

This air quality impact analysis modeling report was prepared based on written protocol comment guidance received from the WDEQ on May 5, 2007 as well as pre-application meeting with WDEQ on July 11, 2006, a conference call with representatives of the WDEQ on March 7, 2007, and protocol submitted to WDEQ on February 8, 2007. The following guidance documents were also consulted:

1. Wyoming Department of Environmental Quality/ Air Division Quality Requirements for Submitting Modeling Analyses (March 1, 2006)
2. Interagency Workgroup on Air Quality Modeling Phase 2 Summary Report and Recommendations for Modeling Long Range Transport Impacts (EPA-454/R-98-019) (IWAQM2) (December, 1998)
3. Federal Land Managers Air Quality Related Values Work Group Phase I report (FLAG) (USFS, NPS, USFWS, 2000)
4. U.S. Environmental Protection Agency (EPA) Guidelines on Air Quality Models (GAQM) (November 9, 2005)

Table 7.6 – Source Location and Parameter

Source Description	Source ID	UTM NAD27 Easting	UTM NAD27 Northing	LCC X	LCC Y	Base Elevation	Stack Height	Stack Temperature	Stack Velocity	Stack Diameter
		(m)	(m)	(km)	(km)	(m)	(m)	(K)	(m/s)	(m)
Turbine and HRSG Train 1	94.20548	391370.8502	4623838.482	94.2055	57.3291	2115.03	45.73	366.493	7.6476	5.79268
Turbine and HRSG Train 2	94.20554	391369.1877	4623777.21	94.2055	57.2678	2115.19	45.73	366.493	7.6476	5.79268
Turbine and HRSG Train 3	94.20561	391367.5348	4623716.29	94.2056	57.2068	2113.97	45.73	366.493	7.6476	5.79268
Gasifier Preheater 1	93.88945	391050.5564	4623693.356	93.8895	57.175	2117.34	25.91	422.06	7.44635	0.4065
Gasifier Preheater 2	93.88946	391050.2258	4623681.172	93.8895	57.1628	2116.41	25.91	422.06	7.44635	0.4065
Gasifier Preheater 3	93.88948	391049.8952	4623668.988	93.8895	57.1506	2115.6	25.91	422.06	7.44635	0.4065
Gasifier Preheater 4	93.88949	391049.5647	4623656.804	93.8895	57.1384	2114.91	25.91	422.06	7.44635	0.4065
Gasifier Preheater 5	93.88951	391049.2341	4623644.62	93.8895	57.1262	2114.5	25.91	422.06	7.44635	0.4065
H-3102 SRU Incinerator	93.96448	391137.24	4624096.22	93.9645	57.5798	2121.68	45.73	422.06	0.1285	4.57
H-5401 Frac Feed Heater	94.12435	391299.1329	4624173.23	94.1244	57.6616	2117.36	45.73	422.06	4.79348	1.21951
H-5301 Cat Dewax Charge	94.09279	391267.3293	4624164.97	94.0928	57.6525	2119.98	15.24	422.06	1.93335	0.4065
H-5201 Unicracker Feed	94.09533	391266.5292	4624046.993	94.0953	57.5345	2118.57	15.24	422.06	1.60033	0.91463
H-5202 Unicracker Intermed.	94.09777	391270.0084	4624083.457	94.0978	57.5711	2117.22	30.48	422.06	3.18521	1.06707
H-5101 Unionfiner Feed	94.12376	391294.9586	4624046.221	94.1238	57.5346	2116.81	15.24	422.06	2.02689	0.50813
H-5102 Unionfiner Intermed	94.11973	391292.8241	4624113.299	94.1197	57.6015	2115.13	15.24	422.06	2.54097	0.4065
Black-Start Generator 1	94.13621	391303.589	4623910.942	94.1362	57.3996	2117.18	6.097	767.604	1.96249	0.4065
Black-Start Generator 2	94.13669	391303.8135	4623901.81	94.1367	57.3905	2117.48	6.097	767.604	1.96249	0.4065
Black-Start Generator 3	94.13659	391303.4502	4623892.553	94.1366	57.3812	2117.58	6.097	767.604	1.96249	0.4065
Firewater Pump	94.12873	391286.31	4623564	94.1287	57.0523	2103.98	6.10	739.27	45	0.15

7.3 LONG-RANGE TRANSPORT MODELING METHOD

7.3.1 Long-Range Transport Modeling

A PSD analysis of increment and AQRV impacts on Class I and sensitive Class II areas will be performed if any Class I or sensitive Class II areas are located within 300 kilometers of the proposed project location. There are eight Class I areas within 300 km from the facility that will be accounted for this analysis. The nearest Class I area is the Mount Zirkel Wilderness, which is located approximately 93 km south from the project. The second nearest Class I area is the Rawah Wilderness, which is located approximately 102 km south from the project. Rocky Mountain NP and Flat Tops Wilderness Class I areas are located approximately 144 km and 192 km south from the facility, respectively. Eagles Nest Wilderness and Maroon Bell-Snowmass Wilderness Class I areas are located 214 km and 283 km south from the facility, respectively. Bridger Wilderness and Fitzpatrick Wilderness Class I areas are located 242 km and 294 km northwest from the facility, respectively. The sensitive Class II area is Savage Run which is located 60 km south from the facility. The locations of the Class I, sensitive Class II areas, and the facility are shown in Figure 7.1.

The analyses performed include the following:

- PSD Class I Increment modeling significance levels
- Visibility reduction thresholds,
- US National Park Service (USNPS) and US Fish and Wildlife Service (USFWS) deposition analysis thresholds (DAT), and
- Soil and Vegetation Analysis

Additional Air Quality Related Value (AQRV) impact analyses were not conducted because the modeling results did not demonstrate a significant impact on air quality in the Class I and sensitive Class II areas. Because there were no significant increment and visibility impacts on Class I and sensitive Class II areas, it was considered that none of visibility analysis for scenic and important views and impact analysis for water has significant impact.

7.3.2 Model Selection and Setup

To estimate air quality impacts at distances greater than 50 km, the CALPUFF model was used in conjunction with the CALMET diagnostic meteorological model. CALPUFF is a puff-type model that can incorporate three-dimensionally varying wind fields, wet and dry deposition, and atmospheric gas and particle phase chemistry.

The CALMET model is used to prepare the necessary gridded wind fields for use in the CALPUFF model. CALMET can accept as input; mesoscale meteorological data (MM5 data), surface station, upper air, precipitation, cloud cover, and over-water meteorological data (all in a variety of input formats). These data are merged and the effects of terrain and land cover types are estimated. This process results in the generation of gridded 3-D wind field that accounts for the effects of slope flows, terrain blocking effects, flow channelization, and spatially varying land use types.

The development of model inputs and options for both the CALMET and CALPUFF processors was based on guidance provided in following references:

- 1 Wyoming DEQ/Air Quality Division Requirements for Submitting Modeling Analyses (3/06)
- 2 Interagency Working Group on Air Quality Modeling (IWAQM) Phase 2 Summary Report and Recommendations for Modeling Long Range Transport Impacts (December 1998), and
- 3 Permit application PSD particulate matter speciation methodology developed by Don Shepherd, National Park Service (2006).

Key input and model options selected are discussed in the following sections.

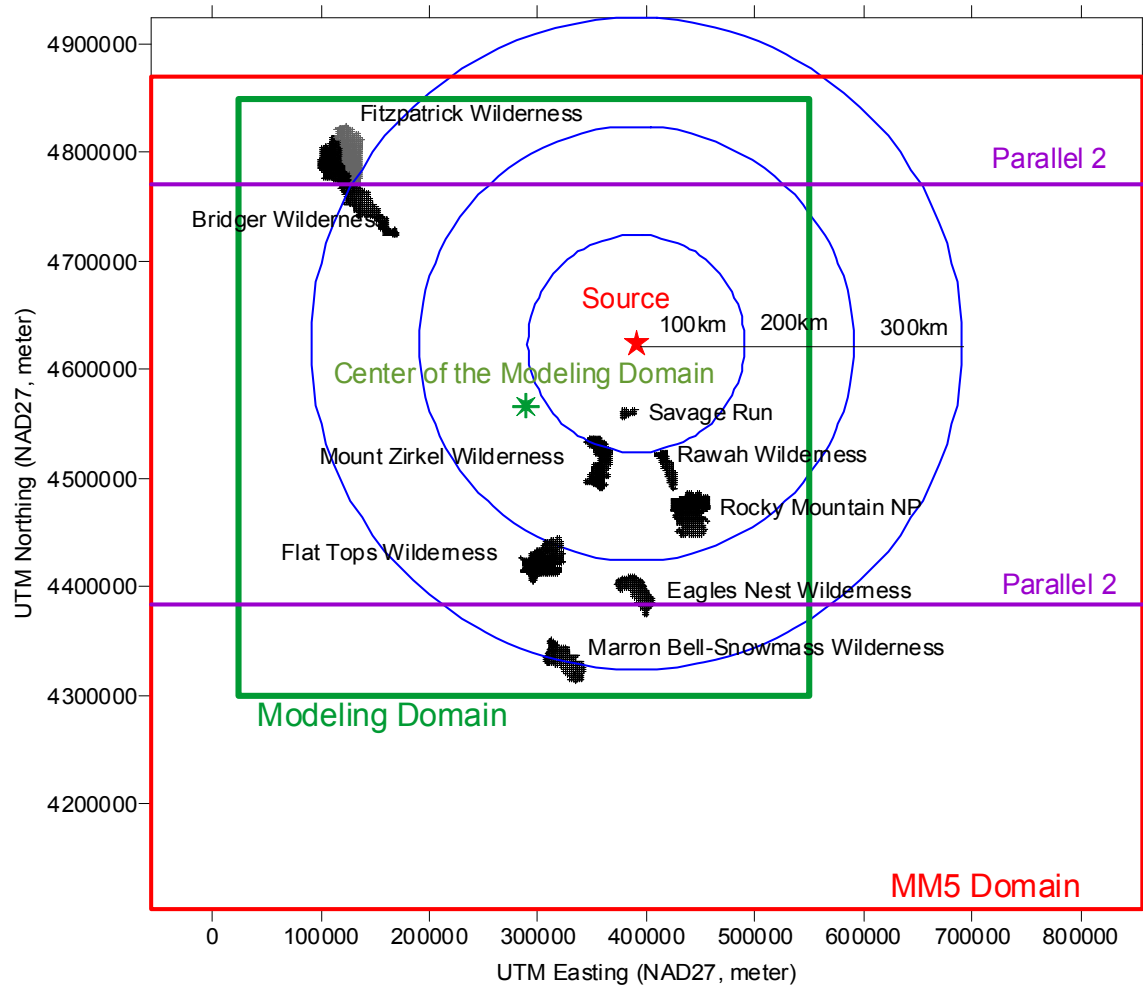
The EPA-approved version of the CALMET/CALPUFF/CALPOST system (CALPUFF of version 5.711a, CALMET of version 5.53a, and CALPOST of version 5.51) was used. Copies of all executable files used in the preparation of this modeling analysis are provided. As requested by the WDEQ, CALMET, CALPUFF, and CALPOST input and output files are provided electronically in Appendix E of the June 19, 2007 application.

7.3.3 Domain

The modeling domain was specified using the Lambert Conformal Conic (LCC) Project system in order to capture the earth curvature of the large modeling domain more accurately for this project. The false easting and northing at the projection origin were set to both zeros. The latitude and longitude of projection origin were set to 41.25 N and 107.44 W, respectively. Matching parallel of latitude 1 and 2 were defined as 39.57 N and 42.94 N, respectively. The modeling domain was defined using a grid-cell arrangement that is 131 cells in X (easting) direction and 137 cells in Y (northing) direction. The grid-cells are 4 kilometers wide. Therefore, the southwest corner of the grid cell (1,1) was set to -321.65 km and -272.07 km.

Approximately 130 km of buffer distance was set between the most east side of the Class I area and the east boundary of the modeling domain. Although 50 km of buffer distance meets the WDEQ's minimum criteria and there is no Class I area in the far east of the project location, 80 km of additional buffer distance was added to the 50 km of buffer distance to prevent the loss of mass outside the boundary under some meteorological scenario that might be associated with transport to nearby Class I areas. The modeling domain, origin of the modeling domain, and the parallels is shown in Figure 7.1 based on UTM coordinate.

Figure 7.1 – Relative Location of Modeling Domain, MM5 Domain, Class I and Sensitive Class II areas, and Source based on UTM Coordinates



7.3.4 LULC and TERREL Processing

The CALMET and CALPUFF models incorporate assumptions regarding land-use classification, leaf-area index, and surface roughness length to estimate deposition during transport. U.S. Geological Survey (USGS) 1:250,000 scale digital elevation models (DEMs) and Land Use Land Cover (LULC) classification files were obtained and used to develop the geophysical input files required by the CALMET model. U.S. Geological Survey (USGS) 1:250,000 scale digital elevation models (DEMs) data were obtained from the Lakes Environmental website, http://www.webgis.com/terr_us1deg.html. Using thirty nine (39) 1-degree DEM data files obtained, terrain pre-processor (TERREL) was processed to produce gridded fields of terrain elevation in the formats compatible with the CALMET.

LULC data (*.gz) were obtained from USGS 250K site, <http://edcftp.cr.usgs.gov/pub/data/LULC/250K/>. Land Use Data Preprocessors, CTGCOMP and CTGPROC were processed to compress twenty six (26) LULC data files obtained. The outputs

of TERREL and CTGPROC were combined in the geo-physical preprocessor (MAKEGEO) to prepare the CALMET geo-physical input file. These inputs include land use type, elevation, surface parameters (surface roughness, length, albedo, Bowen ratio, soil heat flux parameter, and vegetation leaf area index) and anthropogenic heat flux.

Input files for TERREL, CTGPROC, and MAKEGEO are supplied electronically in Appendix E of the June 19, 2007 application. The modeling domain is shown in Figure 7.2.

7.3.5 Hourly Surface and Precipitation Data

Three years of CALMET-ready hourly surface meteorological data and precipitation data for the project modeling domain were provided by WDEQ. The hourly surface data and precipitation data of the “SEWY” section among the data that WDEQ provided were used for the project CALMET modeling. Hourly surface data are from 30 different stations and precipitation data are from 108 different stations. The LCC coordinates of the surface meteorological stations and precipitation stations in the CALMET input files were modified based on the LCC projection.

7.3.6 Upper Air Sounding Data

Upper air sounding data were provided by WDEQ. Three years (2001, 2002, and 2003) of upper air data from Denver Stapleton International Airport (Station # 23062), Grand Junction Walker Field (Station # 23066), Riverton Municipal Airport (Station # 24061), and Rapid City (Station # 94043). The LCC coordinates of the upper air data stations in the CALMET input files were modified based on the LCC projection.

7.3.7 MM5 Data

Two years of MM5 data (2001 and 2002) were obtained from Colorado Department of Public Health and Environment (CDPHE) and one year of MM5 data (2003) was obtained from WDEQ. All three years MM5 data sets consist of a grid resolution of 36 kilometers. The 2001 and 2002 MM5 data consist by each month, but the 2003 MM5 data consist of one file as one year data. Three years of MM5 data were used for BART modeling for Western Regional Air Partnership (WRAP) by CDPHE and WDEQ.

7.3.8 CALMET

Pursuant to FLAG guidance, a three-year meteorological data set was developed using a combination of surface, upper-air, and mesoscale meteorological (MM) data. All surface and upper-air data were obtained from WDEQ. Surface, upper-air, and MM data points were combined and used in the CALMET model.

Monthly CALMET wind fields were generated using a combination of MM5 data sets augmented with the surface, precipitation, and upper air data. Per IWAQM guidance, the MM5 data are interpolated to the CALMET fine-scale grid to create the initial-guess wind fields (IPROG = 14 for MM5). The initial guess wind fields are then adjusted for kinematic terrain effects, slope flows, and terrain blocking effects using the fine-scale CALMET terrain and land use data. The resulting wind fields are referred to as the Step 1 wind field. The observational

NWS data are used to drive a diagnostic weighting between the Step 1 wind fields and the localized surface observations.

For all three years, ZIMAX (maximum overland mixing height) and the maximum ZFACE (top cell face height) was set as 3500 m as the WDEQ’s “SEWY” CALMET input was set up. Thus, 3500 m of XMAXZI (maximum mixing height) and 3500 m of ZFACE value in CALPUFF were used.

Based on the WDEQ’s “SEWY” CALMET input set up, 30 km of the maximum radius of influence over land in the surface layer (RMAX1), 50 km of the maximum radius of influence over land aloft (RMAX2), 5 km of the relative weighting of the first guess field and observations in the surface layer (R1), and 25 km of the relative weighting of the first guess field and observation in the layers ALOFT (R2) were used. 15 km of the TERRAD value was used per WDEQ’s “SEWY” CALMET input. CALMET input and model options are presented in Table 7.7.

Table 7.7 – CALMET Model Options

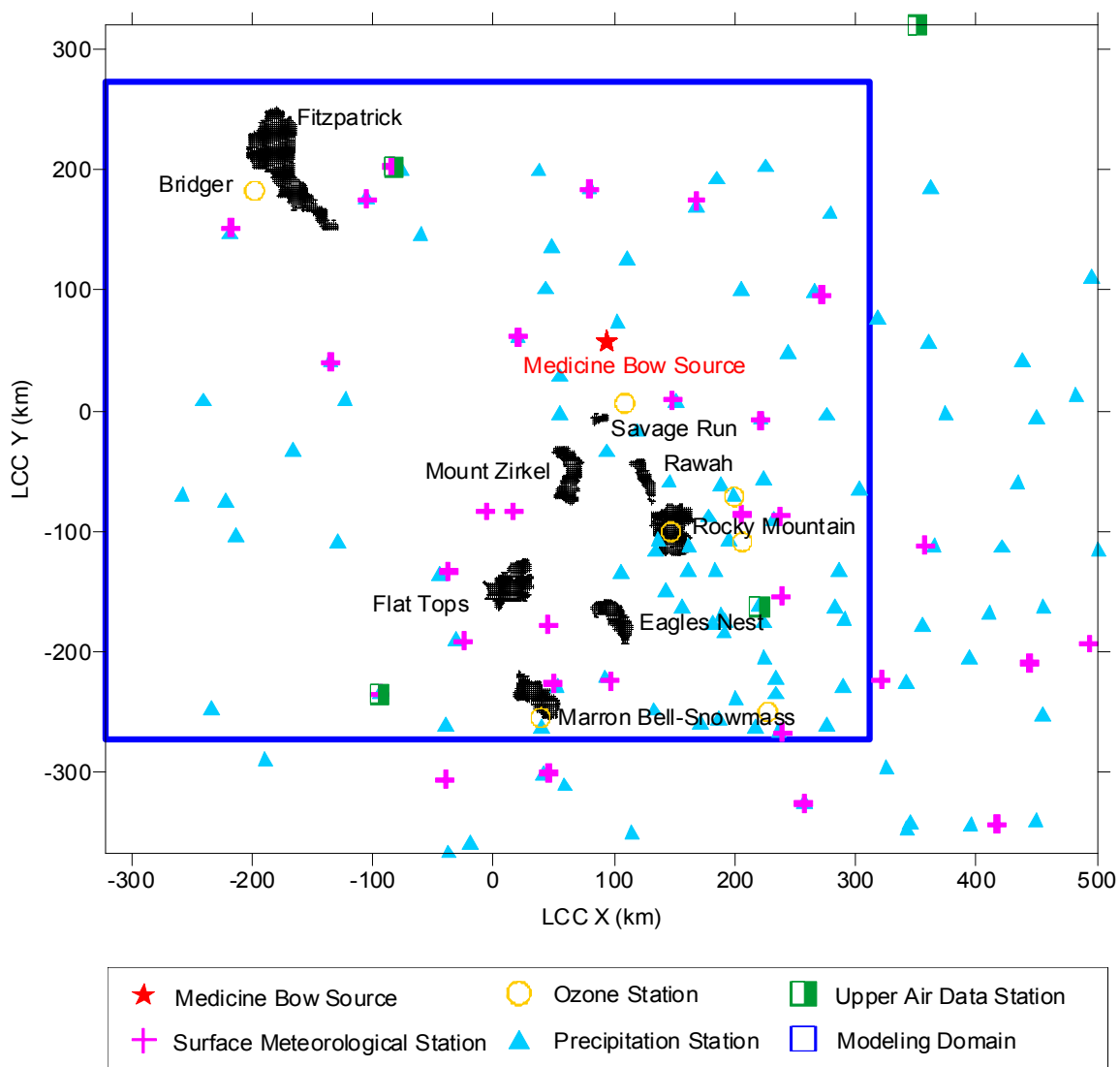
CALMET Variable	Specified Value	Comment
NUSTA	1	Number of Upper Air Stations
PMAP	LCC	Map Projection – Lambert Conformal Conic
FEAST	0	False Easting (km)
FNORTH	0	False Northing (km)
RLAT0	41.25 N	Latitude of Projection Origin
RLON0	107.44 W	Longitude of Projection Origin
XLAT1	39.57 N	Matching parallel of latitude (decimal degrees) for projection
XLAT2	42.94 N	Matching parallel of latitude (decimal degrees) for projection
DATUM	NAS-C	Datum for Output Coordinates
NX	131	Number of Grid Cells in the X-direction
NY	137	Number of Grid Cells in the Y-direction
DGRIDKM	4	Grid Cell Spacing (km)
XORIGKM	-321.65	Reference grid coordinate of southwest corner of grid cell (1,1) X coordinate
YORIGKM	-272.07	Reference grid coordinate of southwest corner of grid cell (1,1) Y coordinate
NZ	10	Number of Vertical Layers (0, 20, 40, 100, 200, 350, 500, 750, 1000, 2000, 3500 m)
ZIMAX	3500 m for years	It is consistent with XMAXZI = 3500 m in CALPUFF option
NOOBS	0	Use Surface, Overwater, and Upper Air Stations
NSSTA	30	Number of surface stations
NPSTA	108	Number of precipitation stations

Table 7.7 – CALMET Model Options

CALMET Variable	Specified Value	Comment
ICLOUD	0	Gridded Cloud not used
IWFCOD	1	Diagnostic Wind Module (1 = yes)
IFRADJ	1	Froude Number Adjustment (1 = yes)
IKINE	0	Kinematic Effects (0 = no)
IOBR	0	O'Brien Vertical Velocity Adjustment (0 = no)
ISLOPE	1	Slope Flow Effects (1 = yes)
IEXTRP	-4	Surface Wind Extrapolation – similarity theory, ignore layer 1
ICALM	0	Extrapolate calm surface winds (0 = no)
RMIN2	4	Minimum Distance from Surface Station to Upper Air for which Extrapolation is allowed
I PROG	14	MM5 Data Used as Initial Guess Field
RMAX1	30	Maximum Overland Radius of Influence at Surface (km)
RMAX2	50	Maximum Overland Radius of Influence Aloft (km)
RMAX3	50	Maximum Overwater Radius of Influence (km)
RMIN	0.1	Minimum Radius of Influence in Wind Field Interpolation (km)
R1	5	Relative weighting of the first guess field and observations in the SURFACE layer (R1 is the distance from an observational station at which the observation and first guess field are equally weighted)
R2	25	Relative weighting of the first guess field and observations in the layers ALOFT
TERRAD	15	Radius of Influence of Terrain Features

Locations of the hourly surface meteorological stations, upper air sounding monitoring stations, precipitation data monitoring stations, and ozone monitoring stations are shown in Figure 7.2.

Figure 7.2 – Modeling Domain with Receptors of Class I and Sensitive Class II Areas, Precipitation Data Monitoring Station, Ozone Monitoring Station, Surface Meteorological Data Monitoring Station, and Project Location



7.3.8.1 CALPUFF

Size parameters for dry deposition of nitrate, sulfate, and PM₁₀ particles were based on default CALPUFF model options. Chemical parameters for gaseous dry deposition and wet scavenging coefficients were based on default values presented in the CALPUFF User's Guide. Calculation of total nitrogen deposition includes the contribution of nitrogen resulting from the ammonium ion of the ammonium sulfate compound. For the CALPUFF runs that incorporate deposition and chemical transformation rates (i.e., deposition and visibility), the full chemistry option of CALPUFF was turned on (MCHEM = 1). The nighttime loss for SO₂, NO_x and nitric acid (HNO₃) was set at 0.2 percent per hour, 2 percent per hour and 2 percent per hour, respectively.

CALPUFF was also configured to allow predictions of SO₂, sulfate (SO₄), NO_x, HNO₃, nitrate (NO₃) and PM₁₀ using the MESOPUFF II chemical transformation module.

As described in Section 7.2, emissions were speciated in accordance with the National Park Service (NPS)'s Particulate Matter Speciation (PMS) guideline (<http://www2.nature.nps.gov/air/permits/ect/index.cfm>). In doing so, the sulfur emissions were speciated to relative sulfur constituents of SO₂ and SO₄ to better account for gas to particulate conversion and visibility effects.

CALPUFF input and model options are presented in Table 7.8. CALMET, CALPUFF, and CALPOST input files are provided electronically in Appendix E of the June 19, 2007 application.

Table 7.8 – CALPUFF Model Options

CALPUFF Variable	Specified Value	Comment
IBTZ	7	Base Time Zone
MGAUSS	1	Vertical Distribution Used In The Near Field
MCTADJ	3	Terrain Adjustment Method
MCTSG	0	Subgrid-Scale Complex Terrain Flag
MSLUG	0	Near-Field Puffs Modeled As Elongated 0
MTRANS	1	Transitional Plume Rise Modeled
MTIP	1	Stack Tip Downwash
MBDW	1	Building Downwash, 1= ISC method
MSHEAR	0	Vertical Wind Shear Modeled Above Stack Top
MSPLIT	0	Puff Splitting Allowed
MCHEM	1	Chemical Mechanism Flag
MWET	1	Wet Removal Modeled
MDRY	1	Dry Deposition Modeled
MDISP	3	Method Used To Compute Dispersion Coefficients
MROUGH	0	PG Sigma-Y,Z Adjusted For Roughness
MPARTL	1	Partial Plume Penetration Of Elevated Inversion (per IWAQM)
MTINV	0	Strength Of Temperature Inversion Provided In PROFILE.DAT Extended Records
MPDF	0	PDF Used For Dispersion Under Convective Conditions
MSGTIBL	0	Sub-Grid TIBL Module Used For Shore Line
MBCON	0	Boundary Conditions (Concentration) Modeled
MFOG	0	Configure For FOG Model Output
MREG	1	Test Options Specified To See If They Conform To Regulatory Values
PMAP	LCC	Map Projection
FEAST	0	False Easting (km)
FNORTH	0	False Northing (km)
RLAT0	41.25 N	Latitude of Projection Origin
RLON0	107.44 W	Longitude of Projection Origin

Table 7.8 – CALPUFF Model Options

CALPUFF Variable	Specified Value	Comment
XLAT1	39.57 N	Matching parallel of latitude (decimal degrees) for projection
XLAT2	42.94 N	Matching parallel of latitude (decimal degrees) for projection
NX	131	No. X Grid Cells
NY	137	No. Y Grid Cells
NZ	10	No. Vertical Layers
DGRIDK M	4	Grid Spacing (km)
ZFACE		0, 20, 40, 100, 200, 350, 500, 750, 1000, 2000, 3500
XORIGK M	-321.65	Reference grid coordinate of southwest corner of grid cell (1,1) X coordinate
YORIGK M	-272.07	Reference grid coordinate of southwest corner of grid cell (1,1) Y coordinate
RCUTR	30	Reference Cuticle Resistance
RGR	10	Reference Ground Resistance
REACTR	8	Reference Pollutant Reactivity
IVEG	1	Vegetation State In Unirrigated Areas
MOZ	1	Ozone Data Input Option (1= read hourly ozone concentration from the OZONE.DAT data file)
BCKO3	44	For O3 data missing
BCKNH3	2	Monthly ammonia concentrations
MHFTSZ	0	Switch For Using Heffter Equation For Sigma Z As Above
WSCALM	.5	Minimum Wind Speed (m/s) Allowed For Non-Calm Conditions
XMAXZI	3500 m	Maximum Mixing Height (m)
XMINZI	50 m	Minimum Mixing Height (m)

7.3.9 PSD Class I Increment Significance Analysis

CALMET/CALPUFF (Full CALPUFF) was used to model ambient air impacts of NO₂, PM₁₀, and SO₂ from the emission sources and the modeling results were compared to PSD Class I Increments modeling significance thresholds. The sources were modeled at full potential-to-emit (PTE) for this analysis. The full chemistry option of CALPUFF was turned on (MCHEM = 1, MESOPUFF II scheme), and a deposition option was turned on (MWET = 1 and MDRY = 1). 3-hour averaged SO₂ emission rates for all sources are same as 24-hour averaged SO₂ emission rates. Therefore, 24-hour averaged maximum SO₂ emission rate were modeled for 3-hour and 24-hour SO₂ increment analyses.

For 24-hour PM₁₀ increment analysis, the 24-hour averaged maximum PM₁₀ emission rate was modeled. The emission inventory for total PM was modeled as INCPM. The INCPM was treated as fine particulate matter in terms of geometric characteristics.

For the annual NO_x, SO₂, and PM₁₀ increment analyses, the annual emission rates estimated based on 8,760 hours of combination of normal operation and startup were used. For 24-hour and annual PM incremental analyses, the total PM emission (“INCPM” in the modeling) was modeled without speciation, and the INCPM was treated as fine particulate matter in terms of geometric characteristics.

7.3.10 Class I Area Visibility Reduction Analysis

Full CALPUFF was used to evaluate the potential for visibility reductions. All sources were modeled at full PTE for this analysis. Emissions of total SO₂ and PM₁₀ from the natural gas turbines were speciated based on National Park Service (NPS)’s Particulate Matter Speciation (PMS) guideline as described in Section 7.2.

The emissions of twelve chemical species, SO₂, SO₄, NO_x, HNO₃, NO₃, PM_{0.05}, PM_{0.01}, PM_{0.15}, PM_{0.20}, PM_{0.25}, PM_{1.0}, EC and PM₁₀, were modeled in CALPUFF to predict the visibility impact based on PMS for natural gas turbine. Because only SO₂ emissions estimates were provided, one-third of the estimated SO₂ emission was assumed to be SO₄ emissions, and the remaining two-thirds remained as SO₂ emissions. The total PM₁₀ emissions were speciated into Elemental Carbon (EC) and Secondary Organic Aerosol (SOA). The SOA is speciated again into PM_{0.05}, PM_{0.01}, PM_{0.15}, PM_{0.20}, PM_{0.25}, and PM_{1.0} (indicated as PM0005, PM0010, PM0015, PM0020, PM0025, and PM0100 in the modeling, respectively).

CALPOST was used to post process the modeled CALPUFF values. CALPOST was used to post-process the estimated 24-hour averaged ammonium nitrate, ammonium sulfate and PM concentrations into an extinction coefficient value for each day at each modeled receptor, using the three years of CALMET meteorological data. To do so, it required the use of extinction efficiency values.

All the PM species (PM_{0.05}, PM_{0.01}, PM_{0.15}, PM_{0.20}, PM_{0.25}, and PM_{1.0}) were grouped as PMF. The extinction efficiency of PMF was set as 4.0, which is equal to the extinction efficiency of SOA. Default extinction efficiencies of EC, soil, ammonium sulfate, and ammonium nitrate were used.

Background visibility and extinction coefficient values from the Federal Land Managers Air Quality Related Values Working Group (FLAG) Phase I Report (December 2000) were used for the visibility reduction analysis. Background values for hygroscopic concentration, without adjustment for relative humidity (RH), (0.6 µg/m³) and the non-hygroscopic concentration (4.5 µg/m³) are reported for western wilderness areas. Therefore, BKSO₄ = hygroscopic 0.6/3 = 0.2 and BKSOIL = non-hygroscopic = 4.5 were used. Modeled visibility reductions for each modeled year were compared to the level of acceptable change (LAC) of 5.0 percent.

7.3.11 Total Nitrogen and Sulfur Deposition Analyses

Full CALPUFF was used to evaluate the potential for nitrogen and sulfur deposition. All sources were modeled at full PTE for this analysis. The annual average emission rates were used for the annual averaged nitrogen and sulfur deposition analyses. The annual emission rates of all sources were estimated based on the combination of normal operation and startups. The annual emission rate was used for the annual NO_x, annual SO₂, and annual PM₁₀ impact analyses.

Since natural gas is the dominant fuel during the year, the total emissions of SO₂ and PM was speciated according to the NPS's PMS for natural gas combustion turbines. The emissions of twelve chemical species, SO₂, SO₄, NO_x, HNO₃, NO₃, PM_{0.05}, PM_{0.01}, PM_{0.15}, PM_{0.20}, PM_{0.25}, and PM_{1.0}, EC, and PM₁₀, were modeled in CALPUFF to predict the nitrogen and sulfur deposition.

The total deposition rates for each pollutant were obtained by summing the modeled wet and/or dry deposition rates as follows.

For S deposition, the wet and dry fluxes of sulfur dioxide and sulfate are calculated, normalized by the molecular weight of S, and expressed as total S. Total nitrogen deposition is the sum of N contributed by wet and dry fluxes of nitric acid (HNO₃), nitrate (NO₃⁻), ammonium sulfate ((NH₄)₂SO₄), and ammonium nitrate (NH₄NO₃) and the dry flux of nitrogen oxides (NO_x).

Per WDEQ's "SEWY" CALPUFF input set up, 2 parts per billion of background NH₃ was used. The total modeled nitrogen and sulfur deposition rates were compared to the USNPS/USFWS DATs for western states. The DAT for nitrogen and sulfur are each 0.005 kilogram per hectare per year (kg/ha-yr), which is 1.59E-11 g/m²/s.

7.4 MODELING RESULTS

7.4.1 CALPUFF Modeling Results

Three years of CALPUFF modeling results of Phase II are provided in Table 7.9 through Table 7.11. The modeled criteria pollutant increment concentrations were compared to the Class I area Significant Impact Levels (SIL). All pollutant for all Class I areas and sensitive Class II area are in compliance with the increment analysis threshold, SIL.

Modeled visibility reductions for each modeled year were compared to the level of acceptable extinction change (LAC) of 5.0 % at each modeled area for each year. Since the sensitive Class II area, which is Savage Run, is the sensitive area for all three primary criteria pollutants such as NO_x, SO₂, and PM₁₀, the visibility impact analysis was not applied to the sensitive Class II area.

None of the modeled results exceed the threshold values shown. The visibility impact is less than 5 percent and each criteria pollutant concentration is less than the corresponding threshold level. Deposition thresholds of total N and total S are both 0.005 kg/ha/yr, which is 1.59E-11 g/m²/s. Total N and S deposition impact do not exceeded the threshold.

None of the modeled results (criteria pollutant, deposition, visibility) exceeded the threshold. Therefore, no further analyses, including additional Air Quality Related Value (AQRV) impacts were conducted because the modeling results showed insignificant impact on air quality in the Class I and sensitive Class II areas.

7.4.2 Soil and Vegetation Analysis

Potential impact to soil and vegetation in Class I areas are evaluated on the basis of the model-predicted criteria pollutant concentrations, and the magnitude of predicted annual deposition of sulfur and nitrogen.

The predicted impacts are below significance levels and all threshold levels for soil and vegetation impact; therefore, the project can be expected to have negligible impacts.

Table 7.9 – 2001 CALPUFF Modeling Results

2001	Pollutant		Annual NO _x	3-hr SO ₂	24-hr SO ₂	Annual SO ₂	24-hr PM	Annual PM	Deposition N	Deposition S	No. of Days >5%	Max Extinction Change
	Unit		µg/m ³	µg/m ³	µg/m ³	µg/m ³	µg/m ³	Annual	g/m ² /s	g/m ² /s	Days	%
	Threshold		0.1	1	0.2	0.08	0.32	0.16	1.59E-11	1.59E-11	0	5
Class I Area	Bridger	BRID	1.57E-07	4.24E-04	9.19E-05	1.71E-06	5.61E-05	6.00E-07	3.62E-14	1.05E-14	0	0.28
	Eagles Nest	EANE	1.90E-05	2.06E-03	5.05E-04	7.02E-06	1.20E-04	1.77E-06	1.76E-13	6.12E-14	0	0.87
	Fitzpatrick	FITZ	2.38E-08	6.60E-04	1.23E-04	6.64E-07	4.53E-05	1.66E-07	5.95E-15	2.43E-15	0	0.17
	Flat Tops	FLTO	1.66E-05	1.31E-03	3.86E-04	7.29E-06	1.57E-04	1.94E-06	2.18E-13	5.34E-14	0	0.99
	Maroon Bell Snow	MABE	3.48E-06	9.24E-04	1.71E-04	2.09E-06	5.90E-05	5.04E-07	1.61E-13	3.74E-14	0	0.14
	Mount Zirkel	MOZI	1.98E-04	9.31E-03	3.50E-03	3.64E-05	2.31E-03	2.58E-05	4.71E-13	1.63E-13	0	2.96
	Rawah	RAWA	2.86E-04	1.32E-02	5.14E-03	5.19E-05	5.07E-03	5.51E-05	6.29E-13	2.14E-13	0	4.67
	Rocky Mountain	ROMO	1.36E-04	6.10E-03	2.99E-03	3.82E-05	2.33E-03	2.75E-05	4.23E-13	1.74E-13	0	1.96
Sensitive Class II Area	Savage Run	SAVA	3.68E-04	2.39E-02	6.56E-03	7.80E-05	6.12E-03	9.29E-05	1.24E-12	4.51E-13	0	4.73
Exceed?			No	No	No	No	No	No	No	No	No	No

Table 7-10 – 2002 CALPUFF Modeling Results

2002	Pollutant		Annual NO _x	3-hr SO ₂	24-hr SO ₂	Annual SO ₂	24-hr PM	Annual PM	Deposition N	Deposition S	No. of Days >5%	Max Extinction Change
	Unit		µg/m ³	µg/m ³	µg/m ³	µg/m ³	µg/m ³	Annual	g/m ² /s	g/m ² /s	Days	%
	Threshold		0.1	1	0.2	0.08	0.32	0.16	1.59E-11	1.59E-11	0	5
Class I Area	Bridger	BRID	9.83E-06	2.87E-03	3.76E-04	9.98E-06	4.50E-04	5.50E-06	2.17E-13	9.94E-14	0	0.19
	Eagles Nest	EANE	6.31E-06	1.02E-03	2.87E-04	6.27E-06	1.51E-04	3.57E-06	6.72E-14	2.58E-14	0	0.46
	Fitzpatrick	FITZ	2.38E-06	7.71E-04	2.40E-04	4.04E-06	1.80E-04	2.39E-06	7.86E-14	3.43E-14	0	0.15
	Flat Tops	FLTO	1.28E-05	1.37E-03	4.31E-04	9.82E-06	4.12E-04	6.59E-06	8.96E-14	3.40E-14	0	1.06
	Maroon Bell Snow	MABE	4.68E-06	1.66E-03	3.38E-04	3.19E-06	1.81E-04	2.19E-06	6.44E-14	1.96E-14	0	0.65
	Mount Zirkel	MOZI	7.31E-05	5.74E-03	1.92E-03	3.47E-05	2.18E-03	2.71E-05	2.44E-13	1.10E-13	0	1.41
	Rawah	RAWA	2.53E-04	1.11E-02	2.15E-03	7.06E-05	3.72E-03	5.29E-05	8.73E-13	3.33E-13	0	3.65
	Rocky Mountain	ROMO	1.22E-04	5.23E-03	1.28E-03	3.78E-05	1.24E-03	1.89E-05	6.99E-13	2.60E-13	0	1.78
Sensitive Class II Area	Savage Run	SAVA	2.92E-04	1.50E-02	3.96E-03	9.65E-05	6.33E-03	1.02E-04	8.37E-13	3.54E-13	0	2.87
Exceed?			No	No	No	No	No	No	No	No	No	No

Table 7-11 - 2003 CALPUFF Modeling Results

2003	Pollutant		Annual NO _x	3-hr SO ₂	24-hr SO ₂	Annual SO ₂	24-hr PM	Annual PM	Deposition N	Deposition S	No. of Days >5%	Max Extinction Change
	Unit		µg/m ³	µg/m ³	µg/m ³	µg/m ³	µg/m ³	Annual	g/m ² /s	g/m ² /s	Days	%
	Threshold		0.1	1	0.2	0.08	0.32	0.16	1.59E-11	1.59E-11	0	5
Class I Area	Bridger	BRID	5.21E-07	4.77E-04	1.25E-04	2.15E-06	4.05E-05	6.59E-07	8.20E-14	4.25E-14	0	0.08
	Eagles Nest	EANE	1.57E-05	1.96E-03	3.50E-04	7.34E-06	2.43E-04	3.97E-06	1.92E-13	6.62E-14	0	0.38
	Fitzpatrick	FITZ	2.95E-08	1.39E-04	6.09E-05	5.95E-07	2.75E-05	2.02E-07	3.37E-14	2.07E-14	0	0.03
	Flat Tops	FLTO	3.82E-05	4.68E-03	1.06E-03	1.46E-05	5.58E-04	8.47E-06	2.15E-13	9.21E-14	0	0.55
	Maroon Bell Snow	MABE	5.00E-06	1.21E-03	2.26E-04	3.71E-06	1.68E-04	2.15E-06	1.32E-13	5.22E-14	0	0.41
	Mount Zirkel	MOZI	3.71E-04	9.23E-03	2.80E-03	5.47E-05	5.09E-03	6.88E-05	1.26E-12	3.96E-13	0	1.45
	Rawah	RAWA	2.60E-04	1.30E-02	3.21E-03	7.55E-05	2.26E-03	5.39E-05	9.17E-13	3.54E-13	0	2.19
	Rocky Mountain	ROMO	1.33E-04	4.65E-03	1.65E-03	4.21E-05	9.70E-04	2.95E-05	7.16E-13	2.93E-13	0	0.96
Sensitive Class II Area	Savage Run	SAVA	6.72E-04	1.95E-02	5.56E-03	1.20E-04	6.49E-03	1.43E-04	2.04E-12	6.86E-13	0	4.17
Exceed?			No	No	No	No	No	No	No	No	No	No

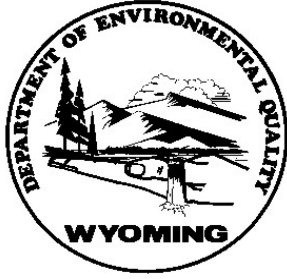
7.5 CONCLUSION

Conservatively modeling the proposed Medicine Bow emissions in CALPUFF resulted in modeled concentrations below the Class I Area threshold levels for deposition, significant impact, and visibility. Therefore, the proposed Medicine Bow sources will not have a significant impact on ambient air quality of Class I areas.

Since there were no significant increment, visibility impacts, or soil and vegetation on Class I areas, it was concluded that no further impacts would be likely and, therefore no additional Air Quality Related Value (AQRV) impact analyses were conducted

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Appendix A
Wyoming Permit Application Form



**DEPARTMENT OF ENVIRONMENTAL QUALITY
AIR QUALITY DIVISION**

PERMIT APPLICATION FORM

Date of Application December 31, 2007

1. Name of Firm or Institution: Medicine Bow Fuel & Power LLC

2. Mailing Address

<u>2 Riverway, Suite 1780</u>	<u>Houston</u>	<u>TX</u>
Number Street	City	State
<u>77056</u>	<u>713-425-6520</u>	
County Zip	Telephone	

3. Plant Location

<u>7.5 miles north of I-80, Exit 260</u>	<u>near Medicine Bow</u>	<u>WY</u>
Number Street	City	State
<u>Carbon</u>	<u>82329</u>	
County Zip	Telephone	
<u>Section 29, Township 21 North, Range 79 West</u>		

4. Name of owner or company official to contact regarding air pollution matters

<u>Tim Stamp</u>	<u>Site Contact</u>	<u>(307) 340-1978</u>
Name	Title	Telephone
<u>1620 Central Avenue</u>	<u>Cheyenne</u>	<u>WY 82001</u>
Number Street	City	State Zip

5. General nature of business

Hydrocarbon conversion through gasification & liquefaction technologies. Support activities include coal mining and electrical generation (no electricity export).

6. Permit application is made for: New Construction Modification
 Relocation Operation
7. Type of equipment to be constructed, modified, or relocated. (List each major piece of equipment separately.)

Refer to Table 3.2 of the application document for a list of major equipment with point source emissions. A list of all major equipment (with and without point source emissions) is provided in Appendix D of the application document.

8. If application is being made for operation of an existing source in a new location, list previous location and new location:

Previous Location: NA

New Location: NA

9. If application is being made for a crushing unit, is there: (mark all appropriate boxes)

Primary Crushing Control Equipment: Enclosed crushers, coal mixed with water, fogging

Secondary Crushing Control Equipment: N/A

Tertiary Crushing Control Equipment: N/A

Recrushing & Screening Control Equipment: N/A

Conveying Control Equipment: Enclosed conveyors, fogging system

Drying Control Equipment: N/A

Other Control Equipment: N/A

Proposed dates of operation (month/year) Facility startup December 2010; equipment to operate year-round.

10. Materials used in unit or process (include solid fuels):

Type of Material	Process Weight Average (lb/hr)	Process Weight Maximum (lb/hr)	Quantity/Year
Coal	666,600 (dry)	666,600 (dry)	3,200,000 ton/year

11. Air contaminants emitted:

Emission Point	Pollutant	[lb/hr]	[ton/yr]	Basis of Data
Refer to Section 3.0 and Appendix B of the application document.				

12. Air contaminant control equipment:

Emission Point	Type	Pollutant Removed	Efficiency
Active Coal Storage	Stacking Tubes	PM	See Note 1
Coal Handling	Enclosed conveyors; Fogger & Passive Engineering Design at transfer points	PM	See Note 1
Combustion Turbines (3, total)	Low NO _x burner, SCR, oxidation catalyst	NO _x , CO, VOC	See Note 1
Process Heaters (3, total) Auxiliary Boiler (1, total)	Low NO _x burner	NO _x	See Note 1
Storage Tanks	Internal Floating Roof (IFR)	VOC, HAPs	See Note 1
Startup/Shutdown/Malfunions	Flares (2, total)	VOC, HAPs, H ₂ S	98%

Notes for Item #12:

1. Refer to Section 3.0 (Emission Estimates), 4.0 (BACT Analysis), and Appendix B (Emission Calculations) of application document for control equipment efficiencies.

Combustion Turbines (Electricity Generation)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas **X (startup and as supplement during normal operations)** ;

D. If other, please specify **Fuel Gas Mixture** _____

Hourly fuel consumption (estimate for new equipment): _____

Size of combustion unit: **660-786 MMBtu/hr** _____

14. Operating Schedule: **24** hours/day; **7** days/week; **52** weeks/year

Peak production season (if any): **NA** _____

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

15. Fuel analysis:

	FUEL GAS MXTURE	NATURAL GAS
% Sulfur	0.1 ppmv	2.9 ppmv (2,000 gr/MMscf)
% Ash	0	0
BTU Value		1,020 Btu/scf assumed

Auxiliary Boiler

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas **X (startup and as supplement during normal operations)** ;

D. If other, please specify **Fuel Gas Mixture**

Hourly fuel consumption (estimate for new equipment): _____

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

Size of combustion unit: _____ **66 MM BTU heat input/hour**

14. Operating Schedule: **Full load for 760 hr/yr and 25% load for 8000 hr/yr**

Peak production season (if any): **NA**

15. Fuel analysis:

	FUEL GAS MXTURE	NATURAL GAS
% Sulfur	0.1 ppmv	2.9 ppmv (2,000 gr/MMscf)
% Ash	0	0
BTU Value		1,020 Btu/scf assumed

Catalyst Regeneration Heater (B-1)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas **X (startup and as supplement during normal operations)** ;

D. If other, please specify **during times of normal operation in standby, the fuel will be a Fuel Gas Mixture, mixed with Natural Gas**

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

Hourly fuel consumption (estimate for new equipment): _____

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

Size of combustion unit: _____ **21.53 MM BTU** heat input/hour

14. Operating Schedule: **@21.53 MMBtu/hr for 877 hr/yr and @ 3.58 MMBtu/hr for 7123 hr/yr**

Peak production season (if any): **NA**

15. Fuel analysis:

	FUEL GAS MXTURE	NATURAL GAS
% Sulfur	0.1 ppmv	2.9 ppmv (2,000 gr/MMscf)
% Ash	0	0
BTU Value		1,020 Btu/scf assumed

Reactivation Heater (B-2)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas **X (startup and as supplement during normal operations)** ;

D. If other, please specify **Fuel Gas Mixture**

Hourly fuel consumption (estimate for new equipment): _____

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

Size of combustion unit: _____ **12.45 MM BTU heat input/hour**

14. Operating Schedule: _____ **2,216** hours/year

Peak production season (if any): **NA**

15. Fuel analysis:

	FUEL GAS MXTURE	NATURAL GAS
% Sulfur	0.1 ppmv	2.9 ppmv (2,000 gr/MMscf)
% Ash	0	0
BTU Value		1,020 Btu/scf assumed

HGT Reactor Charge Heater (B-3)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas **X (startup and as supplement during normal operations)** ;

D. If other, please specify **Fuel Gas Mixture** _____

Hourly fuel consumption (estimate for new equipment): _____

Refer to Section 3 and Appendix B of the application document for details on fuel fired and amounts fired.

Size of combustion unit: _____ **2.22 MM BTU heat input/hour**

14. Operating Schedule: 24 hours/day; 7 days/week; 52 weeks/year

Peak production season (if any): NA _____

15. Fuel analysis:

	FUEL GAS MXTURE	NATURAL GAS
% Sulfur	0.1 ppmv	2.9 ppmv (2,000 gr/MMscf)
% Ash	0	0
BTU Value		1,020 Btu/scf assumed

Gasifier Preheaters (5 total, for cold startup only)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas X;

D. If other, please specify _____

Hourly fuel consumption (estimate for new equipment): _____

Refer to Section 3 and Appendix B of the application document for details on fuel consumption.

Size of combustion unit: _____ **21 MM** BTU heat input/hour

14. Operating Schedule: **As needed during normal operation**

Peak production season (if any): **500 hr/yr (each) during cold startup, as needed during normal operation**

15. Fuel analysis:

	COAL	FUEL OIL	NATURAL GAS
% Sulfur			2.9 ppmv (2,000 gr/MMscf)
% Ash			0
BTU Value			1,020 Btu/scf assumed

Black Start Generators (3, total)

13. Type of combustion unit: (check if applicable)

A. Coal _____

1. *Pulverized* :

General____; Dry Bottom____; Wet Bottom____; With Flyash Reinjection____;
Without Flyash Reinjection____; Other_____

2. *Spreader Stoker* :

With Flyash Reinjection____; Without Flyash Reinjection____; Cyclone____;
Hand-Fired____; Other_____

B. Fuel Oil

Horizontally Fired____; Tangentially Fired _____

C. Natural Gas X;

D. If other, please specify _____

Hourly fuel consumption (estimate for new equipment): _____

Size of combustion unit: _____ **19.5 MM** BTU heat input/hour
(assuming 2,889 bhp, 6,748 Btu/hp-hr)

14. Operating Schedule: As needed during normal operation

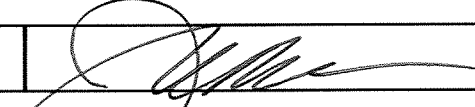
Peak production season (if any): Up to 250 hr/yr, each during cold startup

15. Fuel analysis:

	COAL	FUEL OIL	NATURAL GAS
% Sulfur			2.9 ppmv (2,000 gr/MMscf)
% Ash			0
BTU Value			1,020 Btu/scf assumed

19. Using a flow diagram:
- (1) Illustrate input of raw materials. **Refer to Figures 2.1 and 2.2.**
 - (2) Label production processes, process fuel combustion, process equipment, and air pollution control equipment. **Refer to Figures 2.1 and 2.2.**
 - (3) Illustrate locations of air contaminant release so that emission points under items 11, 12 and 17 can be identified. For refineries show normal pressure relief and venting systems. Attach extra pages as needed. **Refer to Figures 1.3 and 1.4.**
20. A site map should be included indicating the layout of facility at the site. All buildings, pieces of equipment, roads, pits, rivers and other such items should be shown on the layout. **Refer to Figures 1.1, 1.2, and 1.3.**
21. A location drawing should be included indicating location of the facility with respect to prominent highways, cities, towns, or other facilities (include UTM coordinates). **Refer to Figure 1.1.**

"I certify to the accuracy of the plans, specifications, and supplementary data submitted with this application. It is my Opinion that any new equipment installed in accordance with these submitted plans and operated in accordance with the manufacturer's recommendations will meet emission limitations specified in the Wyoming Air Quality Standards and Regulations."

Signature		Typed Name	Jude R. Rolfes		
Title	Senior Vice President		Company	Medicine Bow Fuel & Power LLC	
Mailing Address	Two Riverway, Suite 1780		Telephone No.	(713) 425-6526	
City	Houston		State	Texas	Zip 77056
P.E. Registration (if applicable)	016517				
State where registered	VA				

Appendix B
Emission Calculations

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Emission Summary Sheet**

Normal Operations (8760 hr/yr)

ID No.	Description	Usage	Operation (hr/yr)	Potential Emissions (tpy)					HAPs Emissions (tpy)														TOTALS				
				NO _x	CO	VOC	SO ₂	PM ₁₀	1,3-Butadiene	Acetaldehyde	Acrolein	2,2,4-Trimethylpentane	Benzene	Carbonyl Sulfide	Dichlorobenzene	Ethyl Benzene	Formaldehyde	Hexane	Mercury	Methanol	Naphthalene	PAH		Propylene Oxide	Toluene	Xylene	
CT-1	Turbine and HRSG Train 1	General Electric, 66 MW	8,760	75.86	46.19	6.59	10.79	43.80	1.37E-03	1.27E-01	2.03E-02	3.81E-02			1.02E-01	2.25E-01	4.33E-05	4.13E-03	6.98E-03	9.21E-02	4.13E-01	2.03E-01	2.03E-01	1.23E+00			
CT-2	Turbine and HRSG Train 2	General Electric, 66 MW	8,760	75.86	46.19	6.59	10.79	43.80	1.37E-03	1.27E-01	2.03E-02	3.81E-02			1.02E-01	2.25E-01	4.33E-05	4.13E-03	6.98E-03	9.21E-02	4.13E-01	2.03E-01	2.03E-01	1.23E+00			
CT-3	Turbine and HRSG Train 3	General Electric, 66 MW	8,760	75.86	46.19	6.59	10.79	43.80	1.37E-03	1.27E-01	2.03E-02	3.81E-02			1.02E-01	2.25E-01	4.33E-05	4.13E-03	6.98E-03	9.21E-02	4.13E-01	2.03E-01	2.03E-01	1.23E+00			
AB	Auxiliary Boiler	Heater, 66 MMBtu/hr ¹	8,760	14.17	23.81	1.56	0.17	2.15				5.95E-04	3.40E-04		2.13E-02	5.10E-01		1.73E-04				9.64E-04		5.33E-01			
B-1	Catalyst Regenerator Heater	Heater, 21.53 MMBtu/hr ¹	8,760	4.62	7.77	0.51	0.06	0.70				1.94E-04	1.11E-04		6.93E-03	1.66E-01		5.64E-05				3.14E-04		1.74E-01			
B-2	Reactivation Heater	Heater, 12 MMBtu/hr ¹	8,760	2.67	4.49	0.29	0.03	0.41				1.12E-04	6.42E-05		4.01E-03	9.62E-02		3.26E-05				1.82E-04		1.01E-01			
B-3	HGT Reactor Charge Heater	Heater, 2 MMBtu/hr ¹	8,760	0.48	0.80	0.05	0.01	0.07				2.00E-05	1.14E-05		7.15E-04	1.72E-02		5.82E-06				3.24E-05		1.79E-02			
Tanks	Storage Tanks	Product Storage	8,760			102.62						5.27E-01		3.79E-02	4.95E-01		2.39E+00				5.67E-01	1.60E-01	4.17E+00				
EL	Equipment Leaks	Fugitives	8,760			59.63						7.90E+00	2.34E-01				7.87E+00						1.60E+01				
CS	Coal Storage & Processing	Conveyance (point) & Fugitives	8,760					61.08																0.00E+00			
FW-Pump	Firewater Pump ²	Engine, 575 HP	500	1.51	0.09	0.34	1.52E-03	0.02	3.77E-05	7.39E-04	8.91E-05	8.99E-04			1.14E-03			8.17E-05		2.49E-03	3.94E-04	2.75E-04		6.14E-03			
FL-1	HP / Emergency Flare ³	Flare, 0.816 MMBtu/hr	8,760	0.49	0.98	2.97	2.10E-03																	0.00E+00			
FL-2	LP Flare ²	Flare, 0.204 MMBtu/hr	8,760	0.12	0.25	0.74	0.00																	0.00E+00			
Total Emissions					251.63	176.75	188.49	32.65	195.84	0.00	0.38	0.06	0.00	8.54	0.23	0.00	0.34	0.71	1.29	0.00	10.26	0.01	0.02	0.28	1.81	0.77	24.71

Notes:
¹ Emissions from auxiliary boiler and process heaters assume operation at full design capacity, firing natural gas; however, the equipment may not always fire at full load, and in many cases, will be firing a lower-BTU fuel gas mixture instead of
² SO₂ emissions from the Firewater Pump are based on burning ultra-low sulfur diesel (15 ppm).
³ Flare emissions include pilot emissions for 8760 hr/yr.

Malfunctions and Other Events

ID No.	Description	Usage	Operation (hours) ¹	Potential Emissions (tons)					HAPs Emissions (tpy)														TOTALS		
				NO _x	CO	VOC	SO ₂	PM ₁₀	1,3-Butadiene	Acetaldehyde	Acrolein	2,2,4-Trimethylpentane	Benzene	Carbonyl Sulfide	Dichlorobenzene	Ethyl Benzene	Formaldehyde	Hexane	Mercury	Methanol	Naphthalene	Propylene Oxide		Toluene	Xylene
CO2 VS	CO2 Vent Stack	CO2 Vent Stack	50		83.97	0.23								2.25E-01											2.25E-01
FL-1	HP / Emergency Flare	Flare, 0.816 MMBtu/hr	40	7.83	64.99	0.12	150.16																		0.00E+00
FL-2	LP Flare	Flare, 0.204 MMBtu/hr	8	1.15E-02	2.25E-04	6.79E-04	14.40																		
GP-1	Gasification Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05			9.69E-03

Notes:
¹ The hours shown are annual estimates, except for the Gasification Preheater which is based on 500 hours per preheating event for one gasifier.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Emission Summary Sheet**

Initial Year Including Cold Startup Emissions

This sheet includes total emissions from a cold startup (second set of emissions) and from the remainder of the initial year of operations. The total emissions shown at the bottom of this sheet provide the total emissions for the initial year (or any year with a cold startup).

Normal Operations (After Startup)

ID No.	Description	Usage	Normal Operating Hours After Startup (hr/yr)	Potential Emissions (tpy)					HAPs Emissions (tpy)														TOTALS			
				NO _x	CO	VOC	SO ₂	PM ₁₀	1,3-Butadiene	Acetaldehyde	Acrolein	2,2,4-Trimethylpentane	Benzene	Carbonyl Sulfide	Dichlorobenzene	Ethyl Benzene	Formaldehyde	Hexane	Mercury	Methanol	Naphthalene	PAH		Propylene Oxide	Toluene	Xylene
CT-1	Turbine and HRSG Train 1	General Electric, 66 MW	7,760	67.20	40.92	5.84	9.56	38.80	1.21E-03	1.12E-01	1.80E-02	3.37E-02			9.00E-02	2.00E-01	3.84E-05	3.66E-03	6.19E-03	8.16E-02	3.66E-01	1.80E-01	1.09E+00			
CT-2	Turbine and HRSG Train 2	General Electric, 66 MW	7,760	67.20	40.92	5.84	9.56	38.80	1.21E-03	1.12E-01	1.80E-02	3.37E-02			9.00E-02	2.00E-01	3.84E-05	3.66E-03	6.19E-03	8.16E-02	3.66E-01	1.80E-01	1.09E+00			
CT-3	Turbine and HRSG Train 3	General Electric, 66 MW	7,760	67.20	40.92	5.84	9.56	38.80	1.21E-03	1.12E-01	1.80E-02	3.37E-02			9.00E-02	2.00E-01	3.84E-05	3.66E-03	6.19E-03	8.16E-02	3.66E-01	1.80E-01	1.09E+00			
AB	Auxiliary Boiler	Heater, 66 MMBtu/hr ¹	8,760	14.17	23.81	1.56	0.17	2.15				5.95E-04	3.40E-04		2.13E-02	5.10E-01		1.73E-04				9.64E-04	5.33E-01			
B-1	Catalyst Regenerator Heater	Heater, 21.53 MMBtu/hr ^{1,2}	8,760	4.62	7.77	0.51	0.06	0.70				1.94E-04		1.11E-04	6.93E-03	1.66E-01		5.64E-05				3.14E-04	1.74E-01			
B-2	Reactivation Heater	Heater, 12 MMBtu/hr ¹	8,760	2.67	4.49	0.29	0.03	0.41				1.12E-04		6.42E-05	4.01E-03	9.62E-02		3.26E-05				1.82E-04	1.01E-01			
B-3	HGT Reactor Charge Heater	Heater, 2 MMBtu/hr ¹	8,760	0.48	0.80	0.05	0.01	0.07				2.00E-05	1.14E-05		7.15E-04	1.72E-02		5.82E-06				3.24E-05	1.79E-02			
Tanks	Storage Tanks	Product Storage	8,760			102.62						5.27E-01		3.79E-02		4.95E-01		2.39E+00				5.67E-01	1.60E-01			
EL	Equipment Leaks	Fugitives	8,760			59.63					7.90E+00	2.34E-01						7.87E+00					1.60E+01			
CS	Coal Storage & Processing	Conveyance (point) & Fugitives	8,760					61.08															0.00E+00			
FW-Pump	Firewater Pump ³	Engine, 575 HP	500	1.51	0.09	0.34	1.52E-03	0.02	3.77E-05	7.39E-04	8.91E-05	8.99E-04			1.14E-03					8.17E-05		2.49E-03	3.94E-04	2.75E-04	6.14E-03	
FL-1	HP / Emergency Flare ⁴	Flare Pilot, 0.816 MMBtu/hr	8,760	0.49	0.98	2.97	2.10E-03																	0.00E+00		
FL-2	LP Flare ⁴	Flare Pilot, 0.204 MMBtu/hr	8,760	0.12	0.25	0.74	0.00																	0.00E+00		
Total Emissions (Partial Year of Normal Operations)				225.65	160.94	186.23	28.95	180.84	3.67E-03	3.38E-01	5.41E-02	0.00E+00	8.53E+00	2.34E-01	5.27E-04	3.08E-01	6.33E-01	1.29E+00	1.15E-04	1.03E+01	1.13E-02	1.86E-02	2.47E-01	1.67E+00	7.00E-01	24.29

Notes:

¹ Emissions from auxiliary boiler and process heaters assume operation at full design capacity, firing natural gas; however, the equipment may not always fire at full load, and in many cases, will be firing a lower-BTU fuel gas mixture instead of natural gas.

During startup periods, the equipment will fire natural gas, and may or may not operate at full capacity. Emissions are based on operation at full load and 8,760 hrs/year as a conservative estimate.

² The catalyst regenerator heater (B-1) will not operate during startup conditions; it will operate only during times of normal facility operation. Therefore, in a startup year, the heater will operate less than 8,760 hrs. A full year of operation is assumed as a conservative emission estimate.

³ SO₂ emissions from the Firewater Pump are based on burning ultra-low sulfur diesel (15 ppm).

⁴ Flare emissions include pilot emissions for 8760 hr/yr.

Cold Startup

ID No.	Description	Usage	Startup Operating Hours (hr/yr)	Potential Emissions (tpy)					HAPs Emissions (tpy)														TOTALS			
				NO _x	CO	VOC	SO ₂	PM ₁₀	1,3-Butadiene	Acetaldehyde	Acrolein	2,2,4-Trimethylpentane	Benzene	Carbonyl Sulfide	Dichlorobenzene	Ethyl Benzene	Formaldehyde	Hexane	Mercury	Methanol	Naphthalene	PAH		Propylene Oxide	Toluene	Xylene
CT-1	Turbine and HRSG Train 1	General Electric, 66 MW	1,000	9.48	5.69	0.81	1.33	5.00	1.69E-04	1.57E-02	2.51E-03	4.71E-03			1.26E-02	2.79E-02	0.00E+00	5.10E-04	8.64E-04	1.14E-02	5.10E-02	2.51E-02	1.52E-01			
CT-2	Turbine and HRSG Train 2	General Electric, 66 MW	1,000	9.48	5.69	0.81	1.33	5.00	1.69E-04	1.57E-02	2.51E-03	4.71E-03			1.26E-02	2.79E-02	0.00E+00	5.10E-04	8.64E-04	1.14E-02	5.10E-02	2.51E-02	1.52E-01			
CT-3	Turbine and HRSG Train 3	General Electric, 66 MW	1,000	9.48	5.69	0.81	1.33	5.00	1.69E-04	1.57E-02	2.51E-03	4.71E-03			1.26E-02	2.79E-02	0.00E+00	5.10E-04	8.64E-04	1.14E-02	5.10E-02	2.51E-02	1.52E-01			
Gen-1	Black-Start Generator	Caterpillar, 2889 HP	360	1.15	2.79	1.03	2.06E-03	2.71E-04	9.37E-04	2.93E-02	1.80E-02	8.77E-04	7.44E-04			1.85E-01	3.90E-04					1.43E-03	6.46E-04	2.38E-01		
Gen-2	Black-Start Generator	Caterpillar, 2889 HP	360	1.15	2.79	1.03	2.06E-03	2.71E-04	9.37E-04	2.93E-02	1.80E-02	8.77E-04	7.44E-04			1.85E-01	3.90E-04					1.43E-03	6.46E-04	2.38E-01		
Gen-3	Black-Start Generator	Caterpillar, 2889 HP	360	1.15	2.79	1.03	2.06E-03	2.71E-04	9.37E-04	2.93E-02	1.80E-02	8.77E-04	7.44E-04			1.85E-01	3.90E-04					1.43E-03	6.46E-04	2.38E-01		
GP-1	Gasifier Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05	9.69E-03			
GP-2	Gasifier Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05	9.69E-03			
GP-3	Gasifier Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05	9.69E-03			
GP-4	Gasifier Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05	9.69E-03			
GP-5	Gasifier Preheater	Heater, 21.00 MMBtu/hr	500	0.26	0.43	0.03	3.09E-03	0.04				1.08E-05	6.18E-06		3.86E-04	9.26E-03						1.75E-05	9.69E-03			
CO2 VS	CO2 Vent Stack	CO2 Vent Stack	250			314.89	0.84					8.44E-01											8.44E-01			
FL-1	HP / Emergency Flare ¹	Venting to Flare, 0.816 MMBtu/hr	50	9.78	80.88	0.14	187.70																	0.00E+00		
FL-2	LP Flare ¹	Venting to Flare, 0.204 MMBtu/hr	20	0.03	0.19	0.00	36.01																	0.00E+00		
Total Emissions (Cold Startup Only, Partial Year)				42.99	423.55	6.64	227.74	15.20	3.32E-03	1.35E-01	6.16E-02	2.63E-03	1.64E-02	8.44E-01	3.09E-05	3.77E-02	6.41E-01	4.75E-02	0.00E+00	0.00E+00	1.53E-03	2.59E-03	3.42E-02	1.57E-01	7.73E-02	2.06

Notes:

¹ Flare operating hours include cold startup and malfunctions. Up to 50 hr/yr of venting to the HP Flare and up to 20 hr/yr of venting to the LP Flare are included. Pilot emissions are included above in the Normal Operations summary.

				Potential Emissions (tpy)					HAPs Emissions (tpy)																	
				NO _x	CO	VOC	SO ₂	PM ₁₀	1,3-Butadiene	Acetaldehyde	Acrolein	2,2,4-Trimethylpentane	Benzene	Carbonyl Sulfide	Dichlorobenzene	Ethyl Benzene	Formaldehyde	Hexane	Mercury	Methanol	Naphthalene	PAH	Propylene Oxide	Toluene	Xylene	TOTALS
TOTAL EMISSIONS FOR COLD STARTUP YEAR				268.64	584.48	192.87	256.69	196.04	0.01	0.47	0.12	0.00	8.54	1.08	0.00	0.35	1.27	1.33	0.00	10.26	0.01	0.02	0.28	1.82	0.78	26.35

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Initial Year (Cold Start and Remainder Normal Operations [Base Load])**

Source ID Number	Turbine and HRSG Train 1		
Equipment ID			
Turbine Usage	Power Generation		
Turbine Make	GE		
Turbine Model	7EA		
Serial Number	TBD		
Installation Date	TBD		
Engine Configuration	Turbine		
Emission Controls	SCR/Oxidation Catalyst		
Design Output	66 MW		
Site Operating Hours	7760 hr/yr		
Exhaust Temperature	300 °F		
	-12°F	45°F	85°F
Gas Heating Value	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation (Normal operations, Partial year)

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	67.20	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	40.92	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	5.84	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	9.56	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	38.80	Manf. Data ¹
Mercury	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	3.84E-05	Manf. Data ³
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.21E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.12E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	1.80E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.37E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	9.00E-02	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.00E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	3.66E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.19E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	8.16E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	3.66E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	1.80E-01	AP-42 ²

Exhaust Composition	Base Load, Temp. = -12°F			Base Load, Temp. = 45°F			Base Load, Temp. = 85°F	
	Mol. Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	
Argon	39.94	1.03	0.41	1.03	0.41	1.03	0.41	
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61	21.47	
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37	3.96	
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17	1.40	
Water	18.02	6.23	1.12	6.71	1.21	6.73	1.21	
		100.0	28.5	100.0	28.5	99.9	28.4	

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are very similar to the emissions produced during fuel gas combustion, so these emission factors should provide representative emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

The operating hours include 500 hours for malfunction and warm start-up.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - SSM Emissions, Natural Gas Firing (Cold Start-up)**

Source ID Number	Turbine and HRSG Train 1
Design Output	66 MW
Cold Operating Hours	6 hr/yr
Normal Operating Hours	994 hr/yr
Natural Gas Heating Value	21515 Btu/lb
Natural Gas Flow Rate	36,495 lb/hr
Natural Gas Heat Rate	785.2 MMBtu/hr
Gas Flow Rate	0.77 MMscf/hr

Potential Emissions from Natural Gas Operation (Cold Startup, Partial year)

Pollutant	Emission Factor (lb/MMBtu)	Emission Factor (ppmv, dry)	Estimated Emissions		Source of Emission Factor
			(lb/hr)	(tpy)	
NOx (cold)		25	77.56	0.23	Manf. Data ¹
NOx (normal)		6	18.61	9.25	Manf. Data ¹
CO (cold)		10	18.89	0.06	Manf. Data ¹
CO (normal)		6	11.33	5.63	Manf. Data ¹
VOC		1.4 (ppmv, wet)	1.62	0.81	Manf. Data ¹
SO2	0.0034		2.67	1.33	AP-42 ²
PM10 Total			10.00	5.00	Manf. Data ¹
Mercury		0.000E+00	0.00E+00	0.00E+00	AP-42 ²
1,3-Butadiene	4.30E-07		3.38E-04	1.69E-04	AP-42 ²
Acetaldehyde	4.00E-05		3.14E-02	1.57E-02	AP-42 ²
Acrolein	6.40E-06		5.03E-03	2.51E-03	AP-42 ²
Benzene	1.20E-05		9.42E-03	4.71E-03	AP-42 ²
Ethylbenzene	3.20E-05		2.51E-02	1.26E-02	AP-42 ²
Formaldehyde	7.10E-05		5.57E-02	2.79E-02	AP-42 ²
Naphthalene	1.30E-06		1.02E-03	5.10E-04	AP-42 ²
PAH	2.20E-06		1.73E-03	8.64E-04	AP-42 ²
Propylene Oxide	2.90E-05		2.28E-02	1.14E-02	AP-42 ²
Toluene	1.30E-04		1.02E-01	5.10E-02	AP-42 ²
Xylene	6.40E-05		5.03E-02	2.51E-02	AP-42 ²

Exhaust Composition Base Load, Temp. = 0°F

Component	Mol. Wt.	Volume %	Weighted Mol Wt.
Argon	39.94	0.9	0.36
Nitrogen	28.02	75.5	21.16
Oxygen	32.00	13.88	4.44
Carbon Dioxide	44.01	3.22	1.42
Water	18.02	6.5	1.17
		100.0	28.5

Calculation of dry mass flow rate:

Mass flow of exhaust =	2.06E+06	lb/hr	
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =			72132.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =			4688.6 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =			10012.0 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =			67444.3 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =			14.8%

¹ Criteria pollutant emission factors provided by the manufacturer. The NOx emission factor is corrected to 15% O₂. Cold operation emissions assume that the SCR / oxidation catalyst is not operating. Nitrogen injection is assumed; however, nitrogen may not be available until the Air separation Unit is operating.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note, no mercury emission factor is given for natural gas combustion and so is assumed as zero here.

Additional notes:

These emissions are calculated assuming an ambient temperature of -12°F, which produces the worst case emission estimate.

All natural gas heat rates, flow rates, and exhaust compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/18/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Normal Operations (Base Load)**

Source ID Number	Turbine and HRSG Train 1		
Equipment ID			
Turbine Usage	Power Generation		
Turbine Make	GE		
Turbine Model	7EA		
Serial Number	TBD		
Installation Date	TBD		
Engine Configuration	Turbine		
Emission Controls	SCR/Oxidation Catalyst		
Design Output	66 MW		
Site Operating Hours	8760 hr/yr		
Exhaust Temperature	300 °F		
Gas Heating Value	-12°F	45°F	85°F
	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	75.86	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	46.19	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	6.59	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	10.79	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	43.80	Manf. Data ¹
Mercury ³	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	4.33E-05	Manf. Data ³
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.37E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.27E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	2.03E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.81E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	1.02E-01	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.25E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	4.13E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.98E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	9.21E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	4.13E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	2.03E-01	AP-42 ²

Exhaust Composition	Component	Mol. Wt.	Base Load, Temp. = -12°F		Base Load, Temp. = 45°F		Base Load, Temp. = 85°F	
			Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.
Argon	39.94	1.03	0.41	1.03	0.41	1.03	0.41	
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61	21.47	
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37	3.96	
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17	1.40	
Water	18.02	6.23	1.12	6.71	1.21	6.73	1.21	
		100.0	28.5	100.0	28.5	99.9	28.4	

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are greater than or equal to the emissions produced during fuel gas combustion, so these emission factors should provide worst case emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Initial Year (Cold Start and Remainder Normal Operations [Base Load])

Source ID Number	Turbine and HRSG Train 2
Equipment ID	
Turbine Usage	Power Generation
Turbine Make	GE
Turbine Model	7EA
Serial Number	TBD
Installation Date	TBD
Engine Configuration	Turbine
Emission Controls	SCR/Oxidation Catalyst
Design Output	66 MW
Site Operating Hours	7760 hr/yr
Exhaust Temperature	300 °F

	-12°F	45°F	85°F
Gas Heating Value	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation (Normal operations, Partial year)

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	67.20	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	40.92	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	5.84	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	9.56	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	38.80	Manf. Data ¹
Mercury	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	3.84E-05	Manf. Data ³
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.21E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.12E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	1.80E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.37E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	9.00E-02	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.00E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	3.66E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.19E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	8.16E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	3.66E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	1.80E-01	AP-42 ²

Exhaust Composition	Base Load, Temp. = -12°F		Base Load, Temp. = 45°F		Base Load, Temp. = 85°F	
	Mol. Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %
Argon	39.94	1.03	0.41	1.03	0.41	1.03
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17
Water	18.02	6.23	1.12	6.71	1.21	6.73
		100.0	28.5	100.0	28.5	99.9
						28.4

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume 1, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are very similar to the emissions produced during fuel gas combustion, so these emission factors should provide representative emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

The operating hours include 500 hours for malfunction and warm start-up.

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - SSM Emissions, Natural Gas Firing (Cold Start-up)

Source ID Number	Turbine and HRSG Train 2
Design Output	66 MW
Cold Operating Hours	6 hr/yr
Normal Operating Hours	994 hr/yr
Natural Gas Heating Value	21515 Btu/lb
Natural Gas Flow Rate	36,495 lb/hr
Natural Gas Heat Rate	785.2 MMBtu/hr
Gas Flow Rate	0.77 MMscf/hr

Potential Emissions from Natural Gas Operation (Cold Startup, Partial year)

Pollutant	Emission Factor (lb/MMBtu)	Emission Factor (ppmv, dry)	Estimated Emissions		Source of Emission Factor
			(lb/hr)	(tpy)	
NOx (cold)		25	77.56	0.23	Manf. Data ¹
NOx (normal)		6	18.61	9.25	Manf. Data ¹
CO (cold)		10	18.89	0.06	Manf. Data ¹
CO (normal)		6	11.33	5.63	Manf. Data ¹
VOC		1.4 (ppmv, wet)	1.62	0.81	Manf. Data ¹
SO2	0.0034		2.67	1.33	Manf. Data ¹
PM10 Total			10.00	5.00	Manf. Data ¹
Mercury		0.000E+00	0.00E+00	0.00E+00	AP-42 ²
1,3-Butadiene	4.30E-07		3.38E-04	1.69E-04	AP-42 ²
Acetaldehyde	4.00E-05		3.14E-02	1.57E-02	AP-42 ²
Acrolein	6.40E-06		5.03E-03	2.51E-03	AP-42 ²
Benzene	1.20E-05		9.42E-03	4.71E-03	AP-42 ²
Ethylbenzene	3.20E-05		2.51E-02	1.26E-02	AP-42 ²
Formaldehyde	7.10E-05		5.57E-02	2.79E-02	AP-42 ²
Naphthalene	1.30E-06		1.02E-03	5.10E-04	AP-42 ²
PAH	2.20E-06		1.73E-03	8.64E-04	AP-42 ²
Propylene Oxide	2.90E-05		2.28E-02	1.14E-02	AP-42 ²
Toluene	1.30E-04		1.02E-01	5.10E-02	AP-42 ²
Xylene	6.40E-05		5.03E-02	2.51E-02	AP-42 ²

Exhaust Composition Base Load, Temp. = 0°F

Component	Mol. Wt.	Volume %	Weighted Mol Wt.
Argon	39.94	0.9	0.36
Nitrogen	28.02	75.5	21.16
Oxygen	32.00	13.88	4.44
Carbon Dioxide	44.01	3.22	1.42
Water	18.02	6.5	1.17
		100.0	28.5

Calculation of dry mass flow rate:

Mass flow of exhaust =	2.06E+06	lb/hr		
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =			72132.9	lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =			4688.6	lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =			10012.0	lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =			67444.3	lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =			14.8%	

¹ Criteria pollutant emission factors provided by the manufacturer. The NOx emission factor is corrected to 15% O₂. Cold operation emissions assume that the SCR / oxidation catalyst is not operating. Nitrogen injection is assumed.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note, no mercury emission factor is given for natural gas combustion and so is assumed as zero here.

Additional notes:

These emissions are calculated assuming an ambient temperature of -12°F, which produces the worst case emission estimate. All natural gas heat rates, flow rates, and exhaust compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/18/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Normal Operations (Base Load)**

Source ID Number	Turbine and HRSG Train 2	
Equipment ID		
Turbine Usage	Power Generation	
Turbine Make	GE	
Turbine Model	7EA	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Turbine	
Emission Controls	SCR/Oxidation Catalyst	
Design Output	66 MW	
Site Operating Hours	8760 hr/yr	
Exhaust Temperature	300 °F	

	-12°F	45°F	85°F
Gas Heating Value	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	75.86	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	46.19	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	6.59	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	10.79	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	43.80	Manf. Data ¹
Mercury ³	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	4.33E-05	Manf. Data ³
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.37E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.27E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	2.03E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.81E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	1.02E-01	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.25E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	4.13E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.98E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	9.21E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	4.13E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	2.03E-01	AP-42 ²

Exhaust Composition	Base Load, Temp. = -12°F		Base Load, Temp. = 45°F		Base Load, Temp. = 85°F	
	Mol. Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %
Argon	39.94	1.03	0.41	1.03	0.41	1.03
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17
Water	18.02	6.23	1.12	6.71	1.21	6.73
		100.0	28.5	100.0	28.5	99.9
						28.4

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are greater than or equal to the emissions produced during fuel gas combustion, so these emission factors should provide worst case emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Initial Year (Cold Start and Remainder Normal Operations [Base Load])

Source ID Number	Turbine and HRSG Train 3	
Equipment ID		
Turbine Usage	Power Generation	
Turbine Make	GE	
Turbine Model	7EA	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Turbine	
Emission Controls	SCR/Oxidation Catalyst	
Design Output	66 MW	
Site Operating Hours	7760 hr/yr	
Exhaust Temperature	300 °F	

	-12°F	45°F	85°F
Gas Heating Value	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation (Normal operations, Partial year)

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	67.20	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	40.92	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	5.84	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	9.56	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	38.80	Manf. Data ¹
Mercury	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	3.84E-05	Manf. Data ³
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.21E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.12E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	1.80E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.37E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	9.00E-02	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.00E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	3.66E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.19E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	8.16E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	3.66E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	1.80E-01	AP-42 ²

Exhaust Composition	Base Load, Temp. = -12°F		Base Load, Temp. = 45°F		Base Load, Temp. = 85°F	
	Mol. Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %
Argon	39.94	1.03	0.41	1.03	0.41	1.03
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17
Water	18.02	6.23	1.12	6.71	1.21	6.73
		100.0	28.5	100.0	28.5	99.9
						28.4

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are greater than or equal to the emissions produced during fuel gas combustion, so these emission factors should provide worst case emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

The operating hours include 500 hours for malfunction and warm start-up.

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - SSM Emissions, Natural Gas Firing (Cold Start-up)

Source ID Number	Turbine and HRSG Train 3
Design Output	66 MW
Cold Operating Hours	6 hr/yr
Normal Operating Hours	994 hr/yr
Natural Gas Heating Value	21515 Btu/lb
Natural Gas Flow Rate	36,495 lb/hr
Natural Gas Heat Rate	785.2 MMBtu/hr
Gas Flow Rate	0.77 MMscf/hr

Potential Emissions from Natural Gas Operation (Cold Startup, Partial year)

Pollutant	Emission Factor (lb/MMBtu)	Emission Factor (ppmv, dry)	Estimated Emissions		Source of Emission Factor
			(lb/hr)	(tpy)	
NOx (cold)		25	77.56	0.23	Manf. Data ¹
NOx (normal)		6	18.61	9.25	Manf. Data ¹
CO (cold)		10	18.89	0.06	Manf. Data ¹
CO (normal)		6	11.33	5.63	Manf. Data ¹
VOC		1.4 (ppmv, wet)	1.62	0.81	Manf. Data ¹
SO2	0.0034		2.67	1.33	Manf. Data ¹
PM10 Total			10.00	5.00	Manf. Data ¹
Mercury		0.000E+00	0.00E+00	0.00E+00	AP-42 ²
1,3-Butadiene	4.30E-07		3.38E-04	1.69E-04	AP-42 ²
Acetaldehyde	4.00E-05		3.14E-02	1.57E-02	AP-42 ²
Acrolein	6.40E-06		5.03E-03	2.51E-03	AP-42 ²
Benzene	1.20E-05		9.42E-03	4.71E-03	AP-42 ²
Ethylbenzene	3.20E-05		2.51E-02	1.26E-02	AP-42 ²
Formaldehyde	7.10E-05		5.57E-02	2.79E-02	AP-42 ²
Naphthalene	1.30E-06		1.02E-03	5.10E-04	AP-42 ²
PAH	2.20E-06		1.73E-03	8.64E-04	AP-42 ²
Propylene Oxide	2.90E-05		2.28E-02	1.14E-02	AP-42 ²
Toluene	1.30E-04		1.02E-01	5.10E-02	AP-42 ²
Xylene	6.40E-05		5.03E-02	2.51E-02	AP-42 ²

Exhaust Composition

Component	Mol. Wt.	Volume %	Base Load, Temp. = 0°F	
			Weighted Mol Wt.	
Argon	39.94	0.9	0.36	
Nitrogen	28.02	75.5	21.16	
Oxygen	32.00	13.88	4.44	
Carbon Dioxide	44.01	3.22	1.42	
Water	18.02	6.5	1.17	
		100.0	28.5	

Calculation of dry mass flow rate:

Mass flow of exhaust =	2.06E+06	lb/hr		
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =			72132.9	lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =			4688.6	lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =			10012.0	lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =			67444.3	lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =			14.8%	

¹ Criteria pollutant emission factors provided by the manufacturer. The NOx emission factor is corrected to 15% O₂. Cold operation emissions assume that the SCR / oxidation catalyst is not operating. Nitrogen injection is assumed.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note, no mercury emission factor is given for natural gas combustion and so is assumed as zero here.

Additional notes:

These emissions are calculated assuming an ambient temperature of -12°F, which produces the worst case emission estimate. All natural gas heat rates, flow rates, and exhaust compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/18/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Turbine Detail Sheet - Normal Operations (Base Load)**

Source ID Number	Turbine and HRSG Train 3	
Equipment ID		
Turbine Usage	Power Generation	
Turbine Make	GE	
Turbine Model	7EA	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Turbine	
Emission Controls	SCR/Oxidation Catalyst	
Design Output	66 MW	
Site Operating Hours	8760 hr/yr	
Exhaust Temperature	300 °F	

	-12°F	45°F	85°F
Gas Heating Value	16399.6 Btu/lb	16399.6 Btu/lb	16399.6 Btu/lb
Gas Flow Rate	47,910 lb/hr	44,450 lb/hr	40,240 lb/hr
Gas Heat Rate	785.7 MMBtu/hr	729.0 MMBtu/hr	659.9 MMBtu/hr

Potential Emissions from Fuel Gas Mixture Operation

Pollutant	Emission Factor (ppmv, dry)	Emission Factor (lb/MMBtu)	Estimated Hourly Emissions			Max Hourly Emissions (lb/hr)	Estimated Annual Emissions (tpy)	Source of Emission Factor
			-12°F (lb/hr)	45°F (lb/hr)	85°F (lb/hr)			
NOx	6	0.0234	18.40	17.44	16.12	18.40	75.86	Manf. Data ¹
CO	6	0.0143	11.20	10.62	9.81	11.20	46.19	Manf. Data ¹
VOC	1.4 (ppmv, wet)	0.0020	1.59	1.52	1.40	1.59	6.59	Manf. Data ¹
SO2		0.0034	2.67	2.48	2.24	2.67	10.79	AP-42 ²
PM10 Total		0.0127	10.00	10.00	10.00	10.00	43.80	Manf. Data ¹
Mercury ³	7.86E-07	1.34E-08	1.05E-05	9.96E-06	9.21E-06	1.05E-05	4.33E-05	Manf. Data ¹
1,3-Butadiene		4.30E-07	3.38E-04	3.13E-04	2.84E-04	3.38E-04	1.37E-03	AP-42 ²
Acetaldehyde		4.00E-05	3.14E-02	2.92E-02	2.64E-02	3.14E-02	1.27E-01	AP-42 ²
Acrolein		6.40E-06	5.03E-03	4.67E-03	4.22E-03	5.03E-03	2.03E-02	AP-42 ²
Benzene		1.20E-05	9.43E-03	8.75E-03	7.92E-03	9.43E-03	3.81E-02	AP-42 ²
Ethylbenzene		3.20E-05	2.51E-02	2.33E-02	2.11E-02	2.51E-02	1.02E-01	AP-42 ²
Formaldehyde		7.10E-05	5.58E-02	5.18E-02	4.69E-02	5.58E-02	2.25E-01	AP-42 ²
Naphthalene		1.30E-06	1.02E-03	9.48E-04	8.58E-04	1.02E-03	4.13E-03	AP-42 ²
PAH		2.20E-06	1.73E-03	1.60E-03	1.45E-03	1.73E-03	6.98E-03	AP-42 ²
Propylene Oxide		2.90E-05	2.28E-02	2.11E-02	1.91E-02	2.28E-02	9.21E-02	AP-42 ²
Toluene		1.30E-04	1.02E-01	9.48E-02	8.58E-02	1.02E-01	4.13E-01	AP-42 ²
Xylene		6.40E-05	5.03E-02	4.67E-02	4.22E-02	5.03E-02	2.03E-01	AP-42 ²

Exhaust Composition	Base Load, Temp. = -12°F		Base Load, Temp. = 45°F		Base Load, Temp. = 85°F	
	Mol. Wt.	Volume %	Weighted Mol Wt.	Volume %	Weighted Mol Wt.	Volume %
Argon	39.94	1.03	0.41	1.03	0.41	1.03
Nitrogen	28.02	77.34	21.67	76.82	21.52	76.61
Oxygen	32.00	12.08	3.87	12.22	3.91	12.37
Carbon Dioxide	44.01	3.32	1.46	3.23	1.42	3.17
Water	18.02	6.23	1.12	6.71	1.21	6.73
		100.0	28.5	100.0	28.5	99.9

Calculation of dry mass flow rate:	Base Load, Temp. = 0°F	Base Load, Temp. = 45°F	Base Load, Temp. = 80°F
Mass flow of exhaust =	2.03E+06 lb/hr	1.93E+06 lb/hr	1.78E+06 lb/hr
Molar flow of exhaust = Mass flow of exhaust / Mol Wt =	71079.6 lb-mol/hr	67738.0 lb-mol/hr	62614.9 lb-mol/hr
Molar flow of water = Vol.% H ₂ O * Exhaust molar flow =	4428.3 lb-mol/hr	4545.2 lb-mol/hr	4214.0 lb-mol/hr
Molar Flow of O ₂ = Vol.% O ₂ * Exhaust molar flow =	8586.4 lb-mol/hr	8277.6 lb-mol/hr	7745.5 lb-mol/hr
Molar flow of Exhaust, dry = Exhaust molar flow - H ₂ O molar flow =	66651.4 lb-mol/hr	63192.8 lb-mol/hr	58400.9 lb-mol/hr
Vol. % O ₂ , dry = O ₂ molar flow / Exhaust molar flow =	12.9%	13.1%	13.3%
total exhaust flow, acfm	499,773	476,277	440,256

¹ Criteria pollutant emission factors provided by the manufacturer, but in some cases have been adapted from natural gas combustion. The NOx emission factor is corrected to 15% O₂.

² EPA AP-42, Volume I, Fifth Edition - April 2000, Table 3.1-3, Emission Factors for Hazardous Air Pollutants from Natural Gas-Fired Stationary Gas Turbines. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are greater than or equal to the emissions produced during fuel gas combustion, so these emission factors should provide worst case emission estimates.

³ Mercury concentration in turbine exhaust gas is based on estimated mercury emission rates, as provided by the mercury guard bed manufacturer.

Additional notes:

All gas flow rates and compositions are based on information provided by GE. (Information provided by Paul Rood of SNC Lavalin via email on 12/17/07.)

Average VOC molecular weight assumed to be 46 lb-mol/lb.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Auxillary Boiler Detail Sheet**

Source ID Number	
Equipment Usage	Auxillary Boiler
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	Low Nox Burner
Design Heat Rate	66.00 MMBtu/hr
Operating Hours	8760 hrs/yr
Natural Gas Rates	
<i>Note: boiler will fire natural gas during cold start (760 hours); normally, it will operate at lower (25%) load and fire a lower-Btu fuel gas mixture (vent gas).</i>	
Fuel Heating Value	1,020 Btu/scf
NG Potential Fuel Usage	0.0647 MMscf/hr

Potential Emissions (firing natural gas at 100% load)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	3.24	14.17	AP-42 ¹
CO	84.00	0.08	5.44	23.81	AP-42 ¹
VOC	5.50	5.4E-03	0.36	1.56	AP-42 ²
SO2	0.60	5.9E-04	0.04	0.17	AP-42 ²
PM10	7.60	7.5E-03	0.49	2.15	AP-42 ²
Benzene	2.1E-03	2.1E-06	1.36E-04	5.95E-04	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	7.76E-05	3.40E-04	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	4.85E-03	2.13E-02	AP-42 ³
Hexane	1.8E+00	1.8E-03	1.16E-01	5.10E-01	AP-42 ³
Naphthalene	6.1E-04	6.0E-07	3.95E-05	1.73E-04	AP-42 ³
Toluene	3.4E-03	3.3E-06	2.20E-04	9.64E-04	AP-42 ³

1. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-1, Emission Factors for Nitrogen Oxides (NOx) and Carbon Monoxide (CO) from Natural Gas Combustion.
2. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-2, Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion.
3. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-3, Emission Factors for Speciated Organic Compounds from Natural Gas Combustion.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Heater Detail Sheet**

Source ID Number	Catalyst Regenerator
Equipment Usage	Process Heater
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	Low NOx Burner
Design Heat Rate	21.53 MMBtu/hr
<i>Note: will only fire 3.58 MMBtu/hr during standby operations, anticipated to be approximately 7,800 hr/yr</i>	
Operating Hours	8,760 hr/yr
Fuel Heating Value	1,020 Btu/scf
NG Potential Fuel Usage ¹	0.021 MMscf/hr

Potential Emissions (firing natural gas)¹

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.06	4.62	AP-42 ²
CO	84.00	0.08	1.77	7.77	AP-42 ²
VOC	5.50	5.4E-03	0.12	0.51	AP-42 ³
SO2	0.60	5.9E-04	0.01	0.06	AP-42 ³
PM10	7.60	7.5E-03	0.16	0.70	AP-42 ³
Benzene	2.1E-03	2.1E-06	4.43E-05	1.94E-04	AP-42 ⁴
Dichlorobenzene	1.2E-03	1.2E-06	2.53E-05	1.11E-04	AP-42 ⁴
Formaldehyde	7.5E-02	7.4E-05	1.58E-03	6.93E-03	AP-42 ⁴
Hexane	1.8E+00	1.8E-03	3.80E-02	1.66E-01	AP-42 ⁴
Naphthalene	6.1E-04	6.0E-07	1.29E-05	5.64E-05	AP-42 ⁴
Toluene	3.4E-03	3.3E-06	7.18E-05	3.14E-04	AP-42 ⁴

1. This heater will operate only on a fuel gas mixture, during normal operations. It will not operate during startup operations. PTE emission rates are calculated here based on natural gas firing, as a conservatively high estimate. The heating value of the fuel gas mixture will be lower than that for natural gas. Refer also to notes 3 and 4 below.
2. NOx emissions are estimated based on vendor specifications.
3. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-2, Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are than or equal to the emissions produced during fuel gas mixture combustion, so these emission should provide conservative emission estimates.
4. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-3, Emission Factors for Speciated Organic Compounds from Natural Gas Combustion. Note: These emission factors are for natural gas combustion, which is expected to produce emissions of these pollutants that are greater than or equal to the emissions produced during fuel gas mixture combustion, so these emission factors provide conservative emission estimates.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Heater Detail Sheet**

Source ID Number	Reactivation Heater (B-2)
Equipment Usage	Process Heater
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	Low NOx Burner
Design Heat Rate	12.45 MMBtu/hr
Expected Operating Hours	1,456 hr/yr normal 760 hr/yr cold start 2,216 annual hours
Operating Hours for PTE Emission Calculation	8,760 hr/yr
<u>Natural Gas Usage</u>	
<i>Note: heater will fire natural gas during cold start; however, during normal operations, it will fire a lower-Btu fuel gas mixture.</i>	
Fuel Heating Value	1,020 Btu/scf
NG Potential Fuel Usage	0.0122 MMscf/hr

Potential Emissions (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMbtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	0.61	2.67	AP-42 ¹
CO	84.00	0.08	1.03	4.49	AP-42 ¹
VOC	5.50	5.4E-03	0.07	0.29	AP-42 ²
SO2	0.60	5.9E-04	0.01	0.03	AP-42 ²
PM10	7.60	7.5E-03	0.09	0.41	AP-42 ²
Benzene	2.1E-03	2.1E-06	2.56E-05	1.12E-04	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	1.46E-05	6.42E-05	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	9.15E-04	4.01E-03	AP-42 ³
Hexane	1.8E+00	1.8E-03	2.20E-02	9.62E-02	AP-42 ³
Naphthalene	6.1E-04	6.0E-07	7.45E-06	3.26E-05	AP-42 ³
Toluene	3.4E-03	3.3E-06	4.15E-05	1.82E-04	AP-42 ³

1. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-1, Emission Factors for Nitrogen (NOx) and Carbon Monoxide (CO) from Natural Gas Combustion.
2. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-2, Emission Factors for Criteria and Greenhouse Gases from Natural Gas Combustion.
3. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-3, Emission Factors for Speciated Organic Compounds from Natural Gas Combustion.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Heater Detail Sheet**

Source ID Number	HGT reactor Charge Heater (B-3)
Equipment Usage	Process Heater
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	Low NOx Burner
Design Heat Rate	2.22 MMBtu/hr
Operating Hours	8,760 hr/yr
<u>Natural Gas Usage</u>	
<i>Note: heater will fire natural gas during cold start (760 hours); however, during normal operations, it will fire a lower-Btu fuel gas mixture.</i>	
Fuel Heating Value	1,020 Btu/scf
NG Potential Usage	0.0022 MMscf/hr

Potential Emissions (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	0.11	0.48	AP-42 ¹
CO	84.00	0.08	0.18	0.80	AP-42 ¹
VOC	5.50	5.4E-03	0.01	0.05	AP-42 ²
SO2	0.60	5.9E-04	0.00	0.01	AP-42 ²
PM10	7.60	7.5E-03	0.02	0.07	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.57E-06	2.00E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.61E-06	1.14E-05	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.63E-04	7.15E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.92E-03	1.72E-02	AP-42 ³
Naphthalene	6.1E-04	6.0E-07	1.33E-06	5.82E-06	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.40E-06	3.24E-05	AP-42 ³

1. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-1, Emission Factors for Nitrogen Oxides (NOx) and Carbon Monoxide (CO) from Natural Gas Combustion.
2. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-2, Emission Factors for Criteria and Greenhouse Gases from Natural Gas Combustion.
3. EPA AP-42, Volume I, Fifth Edition - July 1998, Table 1.4-3, Emission Factors for Speciated Organic Compounds from Natural Gas Combustion.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Stack Detail Sheet**

Source ID Number	CO2 Vent Stack	
Equipment Usage	Vent for Off-Spec CO2	
Equipment Make	TBD	
Equipment Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Emission Controls	None	
Potential Operation during initial startup	250 hr/yr	
Potential Operation during malfunctions	50 hr/yr	
Total Vent Stream Flowrate	21,731 lb-mol/hr	
	8,248,270 scf/hr	
Actual Vent Stream Flowrate		
assume T=40 deg F, P=50 psia	38,862 acfm	
Initial Startup		
Vent Gas Molar Flow Rate during startup	5,433 lb-mol/hr	
Vent Gas Molecular Weight	43.1 lb/lb-mol	
Vent Gas H2O Molar Flow Rate	0.20% lb-mol/hr	
Vent Gas Molar Flow Rate (dry)	5422 lb-mol/hr	
Vent Gas Flow Rate (dry)	2,057,945 scf/hr	
Malfunction		
Vent Gas Molar Flow Rate during malfunction	7,244 lb-mol/hr	
Vent Gas Molecular Weight	43.1 lb/lb-mol	
Vent Gas H2O Molar Flow Rate	0.20% lb-mol/hr	
Vent Gas Molar Flow Rate (dry)	7229 lb-mol/hr	
Vent Gas Flow Rate (dry)	2,743,926 scf/hr	

Stack Parameters, for Modeling		
Stack:	100	ft, height
	3	ft, diameter
Velocity:	91.68	ft/s
	27.94	m/s
Temperature	75	deg F
Vent Pressure	50	psia

Potential Emissions from SSM Operation:

Cold Startup Malfunction

Pollutant	Emission Factor ppmv	Estimated Hourly Emissions		Max Hourly Emissions (lb/hr)	Total Annual Emissions (tpy)	Total Annual Emissions (tpy)	Source of Emission Factor
		Initial Startup (lb/hr)	Malfunction (lb/hr)				
CO	16,560	2,519.09	3,358.79	3,358.79	314.89	83.97	Vendor ¹
VOC (COS)	20.7	6.75	9.00	9.00	0.84	0.23	Vendor ¹

¹ CO and VOC emissions are estimated based on vendor specifications.

Additional notes:

Vent gas molar flow rates are from information in email from James Knox, 1/25/08, based on updated UOP data.

VOC is in the form of carbonyl sulfide (COS), which is a HAP.

Annual emissions for this source have been estimated both for the first year of operation, which will include the initial startup emissions and malfunction emissions, and for subsequent years of operation, which will include only malfunction emissions.

The total potential flow rate from this source will only occur if all four gasifiers were operating at full load and both CO2 compressors were to fail. The flow rate at initial startup is estimated to be one-fourth of the total potential flow rate since at most only one gasifier will be operating at full load before the CO2 compression system is operational. The flow rate during a malfunction is estimated to be one-third of the total potential flow rate since at most only one of the three CO2 compressors could fail without a reduction in the production by the gasifiers.

Vent Gas MW Calculation (data from 1/25/08 email)						
Chemical	lb/hr	lb-mol/hr	mol frac	MW	MW*mol frac	ppmv
CO	10,076	360	0.016560	28	0.464	16,560
H2	615	307	0.014145	2	0.028	14,145
CO2	924,654	21,015	0.967051	44	42.550	967,051
H2O	638	35	0.001630	18	0.029	1,630
CH4	56	3	0.000160	16	0.003	160
Ar	322	8	0.000371	39.95	0.015	371
N2	37	1	0.000061	28	0.002	61
H2S	1	0	0.000001	34	0.000	1
COS	27	0	0.000021	60	0.001	21
total	936,425	21,731			43.092	

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
HP Flare Detail Sheet**

Source ID Number	Flare		
Equipment Usage	Emergency Flare/HP Flare		
Equipment Make	TBD		
Equipment Model	TBD		
Serial Number	TBD		
Installation Date	TBD		
Emission Controls	None		
Gas Flow Rate ¹	2,943,142 lb/hr	Syngas to flare (wet)	48" Diameter
Gas Heat Content ¹	2,000 Btu/lb		
Flare Firing Rate	5,886 MMBtu/hr	(low BTU gas)	
Hours of Operation	40 hrs/yr	Malfunctions	
	10 hrs/yr	Initial Year (Cold Starts)	
Pilot Fuel Flow Rate	800 scf/hr		
Pilot Fuel Heat Content	1,020 Btu/scf	Natural Gas (High BTU gas)	
Flare Pilot Firing Rate	0.816 MMBtu/hr		
Hours of Operation, Pilot	8,760 hrs/yr	Continuous pilot	

Estimated Flare Gas Composition During Coal Firing

Component	Flow Rate (lb/hr)	Mol Wt. lb/lb-mol
CO	750,294	28
H2	48,330	2
CO2	489,061	44
H2O	1,625,990	18
CH4	1,199	16
Ar	14,974	40
N2	6,305	28
H2S	3,922	34
COS	270	60
NH3	2,797	17
Total	2,943,142	

Potential Emissions ²

Pollutant	Emission Factors		Destruction Efficiency (%)	Estimated Emissions Pilot (Normal Operation) ⁸		Estimated Emissions Cold Start & Malfunctions		Estimated Emissions Malfunctions only	
	Low BTU gas (lb/MMBtu)	High BTU gas (lb/MMBtu)		(lb/hr)	(tpy)	(lb/hr)	(tpy)	(lb/hr)	(tpy)
NOx ³	0.0641	0.1380	98%	0.11	0.5	391.30	9.8	391.30	7.8
CO ⁴	0.5496	0.2755		0.22	1.0	3,235.10	80.9	3,249.31	65.0
VOC ^{5,6}				0.68	3.0	5.40	0.1	6.08	0.1
SO2 ⁷		0.0006		4.80E-04	2.1E-03	7,508.07	187.7	7,508.07	150.16

Notes:

- Flare gas composition, heat content, and flow rate are all from the Feasibility Study, dated 12/12/06.
- These emissions are based on the calculation methodology and emission factors presented in the TCEQ Guidance Document for Flares and Vapor Oxidizers (RG-109, October 2000). NOx, CO, and VOC emissions include constant pilot gas flow (natural gas).
- NOx emissions were calculated as a sum of the thermal and fuel generated NOx. Thermal NOx emissions were calculated using an emission factor from Table 4 (similar to CO) for an unassisted flare burning low Btu gas. Thermal NOx emissions from the continuous pilot were calculated using the Table 4 emission factor for high BTU gas. The fuel NOx emissions were calculated using the guidance in Table 4 that indicates NOx is 0.5 wt% of inlet NH3.
- The CO emission factor is from Table 4 in the TCEQ Guidance Document and is for an unassisted flare burning low Btu gas. CO emissions for the continuous pilot were calculated using the TCEQ Table 4 emission factor for high BTU gas.
- Fuel VOC emissions were calculated based on guidance in the TCEQ Guidance Document which indicates that 98% of VOCs entering the flare in the fuel will be combusted. The emissions are equal to 2 percent of the incoming flow of COS.
- VOCs from pilot gas combustion are calculated assuming natural gas density of 0.0424 lb/scf, and destruction efficiency of 98%
- SO2 emissions are a sum of the SO2 from the H2S combustion and from the COS combustion. Table 4 indicates that 98% of incoming H2S is converted to SO2, and since COS is a VOC, 98% of that compound will also be combusted and converted to SO2.
- Emissions from normal operations represent only the continuous pilot, since normal operation does not include high pressure vents to flare.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
LP Flare Detail Sheet**

Source ID Number	Flare		
Equipment Usage	Emergency Flare/LP Flare		
Equipment Make	TBD		
Equipment Model	TBD		
Serial Number	TBD		
Installation Date	TBD		
Emission Controls	None		
Gas Flow Rate ¹	3,989 lb/hr	Selexol Reflux Drum vent	24" diameter
Gas Heat Content ¹	8,831 Btu/lb		
Flare Firing Rate	35 MMBtu/hr	(low BTU gas)	
Hours of Operation	8 hrs/yr	Malfunctions	
	12 hrs/yr	Cold Starts	
Pilot Fuel Flow Rate	200 scf/hr		
Pilot Fuel Heat Content	1,020 Btu/scf	Natural Gas (High BTU gas)	
Flare Pilot Firing Rate	0.204 MMBtu/hr		
Hours of Operation, Pilot	8,760 hrs/yr	Continuous pilot	

Estimated Flare Gas Composition During Coal Firing

Component	Flow Rate (lb/hr)	Mol Wt. lb/lb-mol
CO	160	28
H2	399	2
CO2	1,157	44
H2O	199	18
CH4	0	16
Ar	0	40
N2	0	28
H2S	1,955	34
COS	0	60
NH3	120	17
Total	3,989	

Potential Emissions ²

Pollutant	Emission Factors		Destruction Efficiency (%)	Estimated Emissions Pilot (Normal Operation) ⁷		Estimated Emissions Cold Start		Estimated Emissions Cold Start & Malfunction ⁸		Estimated Emissions Malfunctions Only	
	Low BTU gas (lb/MMBtu)	High BTU gas (lb/MMBtu)		(lb/hr)	(tpy)	(lb/hr)	(tpy)	(lb/hr)	(tpy)	(lb/hr)	(tpy)
NOx ³	0.0641	0.1380		0.03	0.1	2.86	0.0	2.86	0.0	2.88	0.0
CO ⁴	0.5496	0.2755		0.06	0.2	19.36	0.1	19.36	0.2	0.06	0.0
VOC ⁵			98%	0.17	0.7	0.00	0.0	0.00	0.0	0.17	0.0
SO2 ⁶		0.0006		1.20E-04	5.3E-04	3,601.15	21.6	3,601.15	36.0	3,601.15	14.4

Notes:

- Flare gas composition and flow rate are from Flare RV Log, December 2007
- These emissions are based on the calculation methodology and emission factors presented in the TCEQ Guidance Document for Flares and Vapor Oxidizers (RG-109, October 2000). NOx, CO, and VOC emissions include constant pilot gas flow (natural gas).
- Fuel NOx emissions were calculated using TCEQ guidance (Table 4) that indicates NOx is 0.5 wt% of inlet NH3. Thermal NOx contribution from the process vent stream is assumed to be negligible; for the pilot gas, thermal NOx is calculated using the TCEQ Table 4 emission factor for high BTU gas.
- CO emissions for the continuous pilot were calculated using the TCEQ Table 4 emission factor for high BTU gas. TCEQ Table 4 emission factor for high BTU gas. CO emissions are from the pilot fuel only.
- VOCs from pilot gas combustion are calculated assuming natural gas density of 0.0424 lb/scf, and destruction efficiency of 98% .
- SO2 emissions are a sum of the SO2 from the H2S combustion and from the COS combustion. Table 4 indicates that 98% of incoming H2S is converted to SO2, and since COS is a VOC, 98% of that compound will also be combusted and converted to SO2.
- Emissions from normal operations represent only the continuous pilot, since normal operation does not include low pressure vents to flare.
- The initial year (i.e., cold start) emissions represent emissions from the low pressure vent gas to the flare. Emissions are estimated for the worst-case (high flow rate, high H2S content) vent stream directed to the LP Flare, and include both cold start and malfunction hours.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Preheater Detail Sheet**

Source ID Number	Gasifier Preheater 1	
Equipment Usage	Refractory Preheating	
Equipment Make	TBD	
Equipment Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Emission Controls	None	
Design Heat Rate	21.00	MMBtu/hr
<u>Cold Startup</u>		
Gas Heating Value	1020	Btu/scf
Gas Potential Operation	500	hr/yr
Gas Potential Fuel Usage	2.06E-02	MMscf/hr

Potential Emissions from Startup Operation (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.03	0.26	AP-42 ¹
CO	84.00	0.08	1.73	0.43	AP-42 ¹
VOC	5.50	5.4E-03	0.11	0.03	AP-42 ²
SO2	0.60	5.9E-04	0.01	3.09E-03	AP-42 ²
PM10	7.60	7.5E-03	0.16	0.04	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.32E-05	1.08E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.47E-05	6.18E-06	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.54E-03	3.86E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.71E-02	9.26E-03	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.00E-05	1.75E-05	AP-42 ³

¹ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-1. Emission Factors for Nitrogen Oxides and Carbon Monoxide from Natural Gas Combustion

² EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-2. Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion

³ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-3. Emission Factors For Speciated Organic Compounds From Natural Gas Combustion

Additional notes:

The average heating value for natural gas is used in these calculations (as provided in Section 1.4 of AP-42).

The PAH emission factor is a sum of all the constituent PAH emission factors in Table 1.4-3.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Preheater Detail Sheet**

Source ID Number	Gasifier Preheater 2
Equipment Usage	Refractory Preheating
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	None
Design Heat Rate	21.00 MMBtu/hr
<u>Cold Startup</u>	
Gas Heating Value	1020 Btu/scf
Gas Potential Operation	500 hr/yr
Gas Potential Fuel Usage	2.06E-02 MMscf/hr

Potential Emissions from Startup Operation (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.03	0.26	AP-42 ¹
CO	84.00	0.08	1.73	0.43	AP-42 ¹
VOC	5.50	5.4E-03	0.11	0.03	AP-42 ²
SO2	0.60	5.9E-04	0.01	3.09E-03	AP-42 ²
PM10	7.60	7.5E-03	0.16	0.04	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.32E-05	1.08E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.47E-05	6.18E-06	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.54E-03	3.86E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.71E-02	9.26E-03	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.00E-05	1.75E-05	AP-42 ³

¹ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-1. Emission Factors for Nitrogen Oxides and Carbon Monoxide from Natural Gas Combustion

² EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-2. Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion

³ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-3. Emission Factors For Speciated Organic Compounds From Natural Gas Combustion

Additional notes:

The average heating value for natural gas is used in these calculations (as provided in Section 1.4 of AP-42).

The PAH emission factor is a sum of all the constituent PAH emission factors in Table 1.4-3.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Preheater Detail Sheet**

Source ID Number	Gasifier Preheater 3	
Equipment Usage	Refractory Preheating	
Equipment Make	TBD	
Equipment Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Emission Controls	None	
Design Heat Rate	21.00	MMBtu/hr
<u>Cold Startup</u>		
Gas Heating Value	1020	Btu/scf
Gas Potential Operation	500	hr/yr
Gas Potential Fuel Usage	2.06E-02	MMscf/hr

Potential Emissions from Startup Operation (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.03	0.26	AP-42 ¹
CO	84.00	0.08	1.73	0.43	AP-42 ¹
VOC	5.50	5.4E-03	0.11	0.03	AP-42 ²
SO2	0.60	5.9E-04	0.01	3.09E-03	AP-42 ²
PM10	7.60	7.5E-03	0.16	0.04	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.32E-05	1.08E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.47E-05	6.18E-06	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.54E-03	3.86E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.71E-02	9.26E-03	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.00E-05	1.75E-05	AP-42 ³

¹ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-1. Emission Factors for Nitrogen Oxides and Carbon Monoxide from Natural Gas Combustion

² EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-2. Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion

³ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-3. Emission Factors For Speciated Organic Compounds From Natural Gas Combustion

Additional notes:

The average heating value for natural gas is used in these calculations (as provided in Section 1.4 of AP-42).

The PAH emission factor is a sum of all the constituent PAH emission factors in Table 1.4-3.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Preheater Detail Sheet**

Source ID Number	Gasifier Preheater 4	
Equipment Usage	Refractory Preheating	
Equipment Make	TBD	
Equipment Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Emission Controls	None	
Design Heat Rate	21.00	MMBtu/hr
<u>Cold Startup</u>		
Gas Heating Value	1020	Btu/scf
Gas Potential Operation	500	hr/yr
Gas Potential Fuel Usage	2.06E-02	MMscf/hr

Potential Emissions from Startup Operation (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.03	0.26	AP-42 ¹
CO	84.00	0.08	1.73	0.43	AP-42 ¹
VOC	5.50	5.4E-03	0.11	0.03	AP-42 ²
SO2	0.60	5.9E-04	0.01	3.09E-03	AP-42 ²
PM10	7.60	7.5E-03	0.16	0.04	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.32E-05	1.08E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.47E-05	6.18E-06	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.54E-03	3.86E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.71E-02	9.26E-03	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.00E-05	1.75E-05	AP-42 ³

¹ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-1. Emission Factors for Nitrogen Oxides and Carbon Monoxide from Natural Gas Combustion

² EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-2. Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion

³ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-3. Emission Factors For Speciated Organic Compounds From Natural Gas Combustion

Additional notes:

The average heating value for natural gas is used in these calculations (as provided in Section 1.4 of AP-42).

The PAH emission factor is a sum of all the constituent PAH emission factors in Table 1.4-3.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Preheater Detail Sheet**

Source ID Number	Gasifier Preheater 5
Equipment Usage	Refractory Preheating
Equipment Make	TBD
Equipment Model	TBD
Serial Number	TBD
Installation Date	TBD
Emission Controls	None
Design Heat Rate	21.00 MMBtu/hr
<u>Cold Startup</u>	
Gas Heating Value	1020 Btu/scf
Gas Potential Operation	500 hr/yr
Gas Potential Fuel Usage	2.06E-02 MMscf/hr

Potential Emissions from Startup Operation (firing natural gas)

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMscf)	(lb/MMBtu)	(lb/hr)	(tpy)	
NOx	50.00	0.05	1.03	0.26	AP-42 ¹
CO	84.00	0.08	1.73	0.43	AP-42 ¹
VOC	5.50	5.4E-03	0.11	0.03	AP-42 ²
SO2	0.60	5.9E-04	0.01	3.09E-03	AP-42 ²
PM10	7.60	7.5E-03	0.16	0.04	AP-42 ²
Benzene	2.1E-03	2.1E-06	4.32E-05	1.08E-05	AP-42 ³
Dichlorobenzene	1.2E-03	1.2E-06	2.47E-05	6.18E-06	AP-42 ³
Formaldehyde	7.5E-02	7.4E-05	1.54E-03	3.86E-04	AP-42 ³
Hexane	1.8E+00	1.8E-03	3.71E-02	9.26E-03	AP-42 ³
Toluene	3.4E-03	3.3E-06	7.00E-05	1.75E-05	AP-42 ³

¹ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-1. Emission Factors for Nitrogen Oxides and Carbon Monoxide from Natural Gas Combustion

² EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-2. Emission Factors for Criteria Pollutants and Greenhouse Gases from Natural Gas Combustion

³ EPA AP-42, Volume I, Fifth Edition - September 1998, Table 1.4-3. Emission Factors For Speciated Organic Compounds From Natural Gas Combustion

Additional notes:

The average heating value for natural gas is used in these calculations (as provided in Section 1.4 of AP-42).

The PAH emission factor is a sum of all the constituent PAH emission factors in Table 1.4-3.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Generator Detail Sheet**

Source ID Number	Black-Start Generator 1	
Engine Usage	Startup Generators	
Engine Make	Caterpillar	
Engine Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Natural Gas	
Emission Controls	None	
Design Rating	1650	ekW
Site Rated Horsepower	2889	BHP
Fuel Heating Value	1020	Btu/scf
Heat Rate	19.49	MMBtu/hr
Engine Heat Rate	6748	Btu/hp-hr
Potential Operation	360	hr/yr
Potential Fuel Usage	6.88	MMscf/yr

At 100% load (worst case emissions)

Potential Emissions

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMBtu)	(g/hp-hr)	(lb/hr)	(tpy)	
NOx		1	6.37	1.15	Manf. Data ¹
CO		2.43	15.48	2.79	Manf. Data ¹
VOC		0.9	5.73	1.03	Manf. Data ¹
SO2	0.000588		0.0115	0.002	AP-42 ²
PM10 Total	0.000077		0.0015	0.00027	AP-42 ²
1,3-Butadiene	2.67E-04		5.21E-03	9.37E-04	AP-42 ²
2,2,4-Trimethylpentane	2.50E-04		4.87E-03	8.77E-04	AP-42 ²
Acetaldehyde	8.36E-03		1.63E-01	2.93E-02	AP-42 ²
Acrolein	5.14E-03		1.00E-01	1.80E-02	AP-42 ²
Benzene	4.40E-04		8.58E-03	1.54E-03	AP-42 ²
Biphenyl	2.12E-04		4.13E-03	7.44E-04	AP-42 ²
Ethylbenzene	3.97E-05		7.74E-04	1.39E-04	AP-42 ²
Formaldehyde	5.28E-02		1.03E+00	1.85E-01	AP-42 ²
Methanol	2.50E-03		4.87E-02	8.77E-03	AP-42 ²
n-Hexane	1.11E-04		2.16E-03	3.90E-04	AP-42 ²
Toluene	4.08E-04		7.95E-03	1.43E-03	AP-42 ²
Xylene	1.84E-04		3.59E-03	6.46E-04	AP-42 ²

¹ Manufacturers Specification.

² EPA AP-42, Volume I, Fifth Edition - October 1996, Table 3.2-2, Uncontrolled Emission Factors for 4-Stroke Lean-Burn Engines.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Generator Detail Sheet**

Source ID Number	Black-Start Generator 2	
Engine Usage	Startup Generators	
Engine Make	Caterpillar	
Engine Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Natural Gas	
Emission Controls	None	
Design Rating	1650	ekW
Site Rated Horsepower	2889	BHP
Fuel Heating Value	1020	Btu/scf
Heat Rate	19.49	MMBtu/hr
Engine Heat Rate	6748	Btu/hp-hr
Potential Operation	360	hr/yr
Potential Fuel Usage	6.88	MMscf/yr

At 100% load (worst case emissions)

Potential Emissions

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMBtu)	(g/hp-hr)	(lb/hr)	(tpy)	
NOx		1	6.37	1.15	Manf. Data ¹
CO		2.43	15.48	2.79	Manf. Data ¹
VOC		0.9	5.73	1.03	Manf. Data ¹
SO2	0.000588		0.0115	0.002	AP-42 ²
PM10 Total	0.000077		0.0015	0.00027	AP-42 ²
1,3-Butadiene	2.67E-04		5.21E-03	9.37E-04	AP-42 ²
2,2,4-Trimethylpentane	2.50E-04		4.87E-03	8.77E-04	AP-42 ²
Acetaldehyde	8.36E-03		1.63E-01	2.93E-02	AP-42 ²
Acrolein	5.14E-03		1.00E-01	1.80E-02	AP-42 ²
Benzene	4.40E-04		8.58E-03	1.54E-03	AP-42 ²
Biphenyl	2.12E-04		4.13E-03	7.44E-04	AP-42 ²
Ethylbenzene	3.97E-05		7.74E-04	1.39E-04	AP-42 ²
Formaldehyde	5.28E-02		1.03E+00	1.85E-01	AP-42 ²
Methanol	2.50E-03		4.87E-02	8.77E-03	AP-42 ²
n-Hexane	1.11E-04		2.16E-03	3.90E-04	AP-42 ²
Toluene	4.08E-04		7.95E-03	1.43E-03	AP-42 ²
Xylene	1.84E-04		3.59E-03	6.46E-04	AP-42 ²

¹ Manufacturers Specification.

² EPA AP-42, Volume I, Fifth Edition - October 1996, Table 3.2-2, Uncontrolled Emission Factors for 4-Stroke Lean-Burn Engines.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Generator Detail Sheet**

Source ID Number	Black-Start Generator 3	
Engine Usage	Startup Generators	
Engine Make	Caterpillar	
Engine Model	TBD	
Serial Number	TBD	
Installation Date	TBD	
Engine Configuration	Natural Gas	
Emission Controls	None	
Design Rating	1650	ekW
Site Rated Horsepower	2889	BHP
Fuel Heating Value	1020	Btu/scf
Heat Rate	19.49	MMBtu/hr
Engine Heat Rate	6748	Btu/hp-hr
Potential Operation	360	hr/yr
Potential Fuel Usage	6.88	MMscf/yr

At 100% load (worst case emissions)

Potential Emissions

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMBtu)	(g/hp-hr)	(lb/hr)	(tpy)	
NOx		1	6.37	1.14645	Manf. Data ¹
CO		2.43	15.48	2.78588	Manf. Data ¹
VOC		0.9	5.73	1.03181	Manf. Data ¹
SO2	0.000588		0.0115	0.002	AP-42 ²
PM10 Total	0.000077		0.0015	0.00027	AP-42 ²
1,3-Butadiene	2.67E-04		5.21E-03	9.37E-04	AP-42 ²
2,2,4-Trimethylpentane	2.50E-04		4.87E-03	8.77E-04	AP-42 ²
Acetaldehyde	8.36E-03		1.63E-01	2.93E-02	AP-42 ²
Acrolein	5.14E-03		1.00E-01	1.80E-02	AP-42 ²
Benzene	4.40E-04		8.58E-03	1.54E-03	AP-42 ²
Biphenyl	2.12E-04		4.13E-03	7.44E-04	AP-42 ²
Ethylbenzene	3.97E-05		7.74E-04	1.39E-04	AP-42 ²
Formaldehyde	5.28E-02		1.03E+00	1.85E-01	AP-42 ²
Methanol	2.50E-03		4.87E-02	8.77E-03	AP-42 ²
n-Hexane	1.11E-04		2.16E-03	3.90E-04	AP-42 ²
Toluene	4.08E-04		7.95E-03	1.43E-03	AP-42 ²
Xylene	1.84E-04		3.59E-03	6.46E-04	AP-42 ²

¹ Manufacturers Specification.

² EPA AP-42, Volume I, Fifth Edition - October 1996, Table 3.2-2, Uncontrolled Emission Factors for 4-Stroke Lean-Burn Engines.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Engine Detail Sheet**

Source ID Number	Firewater Pump
Engine Usage	Firewater Pump Engine
Engine Make	TBD
Engine Model	TBD
Serial Number	TBD
Installation Date	TBD
Engine Configuration	Fuel Oil
Emission Controls	None
Design Rating	575 BHP
Fuel Heating Value	18300 Btu/lb
Fuel Density	7.34 lb/gal
Heat Rate	3.85 MMBtu/hr
Potential Operation	500 hr/yr
Potential Fuel Usage	28.70 gal/hr

Potential Emissions from Fuel Oil Operation

Pollutant	Emission Factor		Estimated Emissions		Source of Emission Factor
	(lb/MMBtu)	(g/hp-hr)	(lb/hr)	(tpy)	
NOx		4.75	6.02	1.51	Vendor ¹
CO		0.29	0.37	0.09	Vendor ¹
VOC	0.35		1.35	0.34	AP-42 ²
SO2			6.06E-03	1.52E-03	Eng. Est. ³
PM10 Total		0.06	7.61E-02	0.02	Vendor ¹
1,3-Butadiene	3.91E-05		1.51E-04	3.77E-05	AP-42 ⁴
Acetaldehyde	7.67E-04		2.96E-03	7.39E-04	AP-42 ⁴
Acrolein	9.25E-05		3.57E-04	8.91E-05	AP-42 ⁴
Benzene	9.33E-04		3.60E-03	8.99E-04	AP-42 ⁴
Formaldehyde	1.18E-03		4.55E-03	1.14E-03	AP-42 ⁴
Naphthalene	8.48E-05		3.27E-04	8.17E-05	AP-42 ⁴
Propylene	2.58E-03		9.94E-03	2.49E-03	AP-42 ⁴
Toluene	4.09E-04		1.58E-03	3.94E-04	AP-42 ⁴
Xylene	2.85E-04		1.10E-03	2.75E-04	AP-42 ⁴
Total HAPs			2.46E-02	6.14E-03	

¹ NOx, PM, and CO emissions are estimated based on vendor specifications.

² EPA AP-42, Volume I, Fifth Edition - October 1996, Table 3.3-1, Emission Factors for Uncontrolled Gasoline and Diesel Industrial Engines.

³ SO2 emissions are estimated based on 15 ppm S and assuming that 100% of S is converted to SO2.

⁴ EPA AP-42, Volume I, Fifth Edition - October 1996, Table 3.3-2, Speciated Organic Compound Emission Factors for Uncontrolled Diesel Engines.

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
 Tank Detail Sheet

Potential VOC Emissions

Source ID	Source Name	Tank Capacity (gal)	Annual Throughput (gal/yr)	Product	Total VOC Emissions (lb/yr)	VOC Emission Rates		HAP Emission Rates						TOTAL	
						(lb/hr)	(tpy)	Hexane (lb/yr)	Benzene (lb/yr)	Toluene (lb/yr)	Ethylbenzene (lb/yr)	Xylene (-m) (lb/yr)	Methanol (lb/yr)	(lb/yr)	(tpy)
TBD	Slops Tank	7,000	42,000	Misc.	606.6	0.07	0.3	19.65	4.69	4.24	0.33	1.39	0	30.30	0.0
TBD	Methanol Tank #1	6,341,984	25,367,936	Methanol	2,285	0.26	1.1	0	0	0	0	0	2,285	2284.56	1.1
TBD	Methanol Tank #2	6,341,984	25,367,936	Methanol	2,285	0.26	1.1	0	0	0	0	0	2,285	2284.56	1.1
TBD	Gasoline Product #1	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #2	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #3	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #4	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #5	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #6	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #7	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Gasoline Product #8	6,341,984	36,254,859	Product Gasoline	23,511	2.68	11.8	110.01	118.82	128.05	8.54	35.98	0	401.40	0.2
TBD	Heavy Gasoline Tank	4,763,841	36,761,340	Heavy Gasoline	9,637	1.10	4.8	80.89	87.32	94.76	6.48	27.56	0	297.01	0.1
TBD	Methanol Off-Spec Tank	5,000	30,000	Methanol	206	0.02	0.1	0	0	0	0	0	205.86	205.86	0.1
TBD	Gasoline Off-Spec Tank	5,000	30,000	Product Gasoline	2,143	0.24	1.1	10.01	10.8	11.43	0.72	3.04	0.00	36	0.0
TOTAL						23.43	102.6	0.495315	0.526685	0.567415	0.037925	0.159915	2.38749		4.175
						HAP-Specific TPY									

Notes:
 All emissions were calculated using the EPA TANKS Program, version 4.09.d.
 Annual hours of operation were assumed to be 8760.

Insignificant Emission Sources - Tanks

TBD	MTG Water Tank
TBD	Liquid Sulfur Storage Tank #1
TBD	Liquid Sulfur Storage Tank #2
03T-002	Grey Water Tank
03T-003	Slurry Additive Tank
01T-104	Mill Discharge Tank
01T-105	Slurry Tank
02T-001	Injector Coolant Tank
03T-001	Settler
03T-004	Filter Feed Tank
03T-005	Filtrate Tank
TBD	Glycol Storage Tank

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Mine - Mine Emissions Summary (PM₁₀)**

Project Year	Notes	South Portal PM ₁₀ (tpy)			East Portal PM ₁₀ (tpy)			Totals PM ₁₀ (tpy)			
		point	road haul	other fugitives	point	road haul	other fugitives	point	road haul	other fugitives	Grand Total
Year 1	No plant operations during this year	0	26.80	3.04	0	0	0	0.00	26.80	3.04	29.84
Year 2	No plant operations during this year	0	109.31	5.17	0	0	0	0.00	109.31	5.17	114.48
Year 3	Partial plant operations during this year	0	71.63	4.20	0.10	0	10.51	0.10	71.63	14.70	86.43
Year 4	Plant operations have begun.	0	0	0	0.86	0	60.23	0.86	0.00	60.23	61.08

Additional Comments:

South Portal emission sources are fugitives and road haul (transport) to Seminole II processing area.

At South Portal, fugitives are from coal stackout, wind erosion from stockpile, and truck loading via front-end loader.

South Portal emissions are due to mine development activity; after plant operations commence, emissions from South Portal are expected to cease.

East Portal emissions are fugitive and point sources. No transport from East Portal to offsite processing is planned.

Fugitive emissions are from stackout, wind erosion from stockpiles, and dozer reclaim to conveyor belt (from emergency stockpile).

Point source emissions are from conveyor drop points, controlled with water fogger.

East Portal Conveyors C1 through C10 will be completely enclosed.

East Portal Conveyors C6 through C10 will have three-quarter (3/4) cover, rather than being completely enclosed.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Mine - Emissions from On-Site Fuel Combustion**

Development Year	Carbon Basin Mine (2004 Application)			Percentage to SBH Mine ¹	SBH Mine South Portal			SBH Mine East Portal ²
	NOx [tpy]	CO [tpy]	SO2 [tpy]		NOx [tpy]	CO [tpy]	SO2 [tpy]	NOx, CO, SO2 [tpy]
Year 1	238.7	200.4	4.1	5%	12	10	0.2	0
Year 2	238.7	200.4	4.1	5%	12	10	0.2	0
Year 3	238.7	200.4	4.1	5%	12	10	0.2	0
Year 4	238.7	200.4	4.1	5%	0	0	0	Negl

Notes

1. Percentage assumed attributable to on-site fuel combustion at SBH Mine. Previous fuel consumption values were based on coal transfer operations at SBH Mine to support mine plan to transport and sell coal at offsite location. Due to mine plan changes, the amount of fuel consumption is expected to be less than originally planned.
2. During development years, no on-site fuel combustion is expected at East Portal. Once normal Plant operations begin in Year 4, some on-site fuel combustion may occur as a result of moving coal from the emergency (dead) stockpile to the conveying system. However, this is expected to be an infrequent activity; thus, annual emissions will be negligible.

Saddleback Hills Mine – Emission Calculations

Year One

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Detail Sheet - South Portal Emissions
Fugitive Emissions**

Production Rates are based on the 2007 Mine Plan Revisions
Total production, Years 1-3 2,538,000 tons

Development Emission Summary (South Portal)

Year	Coal Handling Emissions	Transportation Fugitive Emissions
	PM10 (tpy)	PM10 (tpy)
1	3.04	26.8
2	5.17	109.3
3	4.20	71.6

Year 1

Year 1 - Page 1of 2

Production rate = 218,000 tpy; All coal to be sold at Hanna, WY

Emissions from handling coal at South Portal - stackout and truck loading

Coal Stacker	Coal Dumping to Stockpile	Temporary, portable stacker (stacking tube)	Fugitive
	Emission Factor	0.017 Lb/Ton	WDEQ Emission Factor
	% Suspended	0.75	WDEQ Emission Factor
	Control Factor	50.00%	Estimated
	Material Dumped	218,000 Tons/Yr	Total Coal Through Storage
	TSP Emissions	0.69 Tons/Yr	$E=(EF \times \% \text{ sus} \times MD/2000) \times (1-CF)$
	PM-10 Emissions	0.21 Tons/Yr	30% of TSP
Coal Stockpile	Wind Erosion on Stockpiles	Water	Fugitive
	Emission Factor	1.2 Lb/Acre/Hr	WDEQ Emission Factor
	Pile Size	1.0 Acres	Estimate (assume 10,000 ton pile)
	Fraction Suspended	0.75	WDEQ Emission Factor
	Hours	8,760 Hours	Total Annual
	Ave. Wind Speed	5.03 meters/Sec	Adjusted for in-pit
	Wet Days	60	Seminole Mine 5-Year Average
	Control Factor	50%	assumed
	TSP Emissions	8.28 Tons/Yr	$E=(EF \times AWS \times \% \text{ sus} \times PS \times ((365-WD)/365) \times (1-CF))/2000$
	PM-10 Emissions	2.48 Tons/Yr	
Dozer Reclaim	Cat D11 Dozer to Trucks	No emission controls	Fugitive
	Emission Factor	8.0 Lb/Hr	WDEQ 2002 Guidance
	Total Throughput	218,000 Tons/Yr	Total Coal Through Storage
	Dozed Throughput	218,000 Tons/Yr	
	Dozer Productivity	750 Tons/Hr	Estimate
	Operating Hrs	291 Hrs	Productivity/Throughput
	TSP Emissions	1.16 Tons/Yr	$E=(EF \times Op \text{ Hrs})/2000$
	PM-10 Emissions	0.35 Tons/Yr	30% of TSP
Total South Portal PM ₁₀ Emissions		Conversions:	
		453.6 g/lb	8760 hr/yr
	PM ₁₀ = 3.04 tpy	2000 lb/ton	3600 sec/hr
	0.087 g/sec		

Production rate = 218,000 tpy; All coal to be sold at Hanna, WY

Emissions from the transport of coal with highway trucks on plant roads (South Portal to Seminole II)

These roadways are reconstructed gravel roads for the purpose of connecting loadout with public roads using Wyoming hauling emission factor, with a tire factor adjustment for highway hauler

Input Data

Plant road silt content (s) = 5.1 %
 Tire factor = 3.5

Mean speed 40 mph
 Grader hrs - Carbon Basin Road 975 hrs
 Grader hrs - Seminole II Road 2,000 hrs

Reference

AP-42 table 13.2.2-1 (gravel-upgraded roadway)
 Assuming each pair is equivalent to a single large truck tire, a truck and pup combination have 14 equivalent tires, for a tire factor of 14/4 = 3.5
 Mine estimate

Conversions:

453.6 g/lb 8760 hr/yr
 2000 lb/ton 3600 s/hr

CARBON BASIN COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 0.218 MMtpy
 Vehicle Miles Traveled 18,686 VMT
 RT Haul Distance 6.0 miles
PM-10 Emissions (tpy) 12.97

SEMINOLE II COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 0.218 MMtpy
 Vehicle Miles Traveled 12,457 VMT
 RT Haul Distance 4.0 miles
PM-10 Emissions (tpy) 8.648

CARBON BASIN TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 975
PM-10 Emissions (tpy) 1.70

SEMINOLE II TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 2,000
PM-10 Emissions (tpy) 3.48

CARBON BASIN TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 14.7 48.9
 g/s 0.4 1.4

SEMINOLE II TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 12.1 40.4
 g/s 0.3 1.2

Saddleback Hills Mine – Emission Calculations

Year Two

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Detail Sheet - South Portal Emissions
Fugitive Emissions**

Production Rates are based on the 2007 Mine Plan Revisions
Total production, Years 1-3 2,538,000 tons

Development Emission Summary (South Portal)

Year	Coal Handling Emissions	Transportation Fugitive Emissions
	PM10 (tpy)	PM10 (tpy)
1	3.04	26.8
2	5.17	109.3
3	4.20	71.6

Year 2

Year 2 - Page 1of 2

Production rate = 1,050,000 tpy; All coal to be sold at Hanna, WY

Emissions from handling coal at South Portal - stackout and truck loading

Coal Stacker	Coal Dumping to Stockpile	Temporary, portable stacker (stacking tube)	Fugitive
	Emission Factor	0.017 Lb/Ton	WDEQ Emission Factor
	% Suspended	0.75	WDEQ Emission Factor
	Control Factor	50.00%	Estimated
	Material Dumped	1,050,000 Tons/Yr	Total Coal Through Storage
	TSP Emissions	3.35 Tons/Yr	$E=(EF \times \% \text{ sus} \times MD/2000) \times (1-CF)$
	PM-10 Emissions	1.00 Tons/Yr	30% of TSP
Coal Stockpile	Wind Erosion on Stockpiles	Water	Fugitive
	Emission Factor	1.2 Lb/Acre/Hr	WDEQ Emission Factor
	Pile Size	1.0 Acres	Estimate (assume 10,000 ton pile)
	Fraction Suspended	0.75	WDEQ Emission Factor
	Hours	8,760 Hours	Total Annual
	Ave. Wind Speed	5.03 meters/Sec	Adjusted for in-pit
	Wet Days	60	Seminole Mine 5-Year Average
	Control Factor	50%	assumed
	TSP Emissions	8.28 Tons/Yr	$E=(EF \times AWS \times \% \text{ sus} \times PS \times ((365-WD)/365) \times (1-CF))/2000$
	PM-10 Emissions	2.48 Tons/Yr	
Dozer Reclaim	Cat D11 Dozer to Trucks	No emission controls	Fugitive
	Emission Factor	8.0 Lb/Hr	WDEQ 2002 Guidance
	Total Throughput	1,050,000 Tons/Yr	Total Coal Through Storage
	Dozed Throughput	1,050,000 Tons/Yr	
	Dozer Productivity	750 Tons/Hr	Estimate
	Operating Hrs	1,400 Hrs	Productivity/Throughput
	TSP Emissions	5.60 Tons/Yr	$E=(EF \times \text{Op Hrs})/2000$
	PM-10 Emissions	1.68 Tons/Yr	30% of TSP
Total South Portal PM ₁₀ Emissions		Conversions:	
		453.6 g/lb	8760 hr/yr
PM ₁₀ =	5.17 tpy	2000 lb/ton	3600 sec/hr
	0.149 g/sec		

Production rate = 1,050,000 tpy; All coal to be sold at Hanna, WY

Emissions from the transport of coal with highway trucks on plant roads (South Portal to Seminole II)

These roadways are reconstructed gravel roads for the purpose of connecting loadout with public roads using Wyoming hauling emission factor, with a tire factor adjustment for highway hauler

Input Data

Plant road silt content (s) = 5.1 %
 Tire factor = 3.5

Mean speed 40 mph
 Grader hrs - Carbon Basin Road 975 hrs
 Grader hrs - Seminole II Road 2,000 hrs

Reference

AP-42 table 13.2.2-1 (gravel-upgraded roadway)
 Assuming each pair is equivalent to a single large truck tire, a truck and pup combination have 14 equivalent tires, for a tire factor of 14/4 = 3.5
 Mine estimate

Conversions:

453.6 g/lb 8760 hr/yr
 2000 lb/ton 3600 s/hr

CARBON BASIN COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 1.050 MMtpy
 Vehicle Miles Traveled 90,000 VMT
 RT Haul Distance 6.0 miles
PM-10 Emissions (tpy) 62.48

SEMINOLE II COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 1.050 MMtpy
 Vehicle Miles Traveled 60,000 VMT
 RT Haul Distance 4.0 miles
PM-10 Emissions (tpy) 41.652

CARBON BASIN TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 975
PM-10 Emissions (tpy) 1.70

SEMINOLE II TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 2,000
PM-10 Emissions (tpy) 3.48

CARBON BASIN TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 64.2 213.9
 g/s 1.8 6.2

SEMINOLE II TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 45.1 150.5
 g/s 1.3 4.3

Saddleback Hills Mine – Emission Calculations

Year Three

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Detail Sheet - South Portal Emissions
Fugitive Emissions**

Production Rates are based on the 2007 Mine Plan Revisions

Total production, Years 1-3 2,538,000 tons

Development Emission Summary (South Portal)

Year	Coal Handling Emissions	Transportation Fugitive Emissions
	PM10 (tpy)	PM10 (tpy)
1	3.04	26.8
2	5.17	109.3
3	4.20	71.6

Year 3

Year 3 - Page 1 of 2

Production rate = 1,270,000 tpy; 600,000 tons to be sent via underground tunnel to East Portal
670,000 tpy of remaining coal to be sold at Hanna, WY from South Portal

Emissions from handling coal at South Portal - stackout and truck loading

Coal Stacker	Coal Dumping to Stockpile	Temporary, portable stacker (stacking tube)	Fugitive
	Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
	% Suspended	0.75	<i>WDEQ Emission Factor</i>
	Control Factor	50.00%	<i>Estimated</i>
	Material Dumped	670,000 Tons/Yr	<i>Total Coal Through Storage</i>
	TSP Emissions	2.14 Tons/Yr	<i>E=(EFx% sus x MD/2000)x(1-CF)</i>
	PM-10 Emissions	0.64 Tons/Yr	<i>30% of TSP</i>
Coal Stockpile	Wind Erosion on Stockpiles	Water	Fugitive
	Emission Factor	1.2 Lb/Acre/Hr	<i>WDEQ Emission Factor</i>
	Pile Size	1.0 Acres	<i>Estimate (assume 10,000 ton pile)</i>
	Fraction Suspended	0.75	<i>WDEQ Emission Factor</i>
	Hours	8,760 Hours	<i>Total Annual</i>
	Ave. Wind Speed	5.03 meters/Sec	<i>Adjusted for in-pit</i>
	Wet Days	60	<i>Seminole Mine 5-Year Average</i>
	Control Factor	50%	<i>assumed</i>
	TSP Emissions	8.28 Tons/Yr	<i>E=(EF x AWS x %sus x PS x</i>
	PM-10 Emissions	2.48 Tons/Yr	<i>((365-WD)/365) x (1-CF))/2000</i>
Dozer Reclaim	Cat D11 Dozer to Trucks	No emission controls	Fugitive
	Emission Factor	8.0 Lb/Hr	<i>WDEQ 2002 Guidance</i>
	Total Throughput	670,000 Tons/Yr	<i>Total Coal Through Storage</i>
	Dozed Throughput	670,000 Tons/Yr	
	Dozer Productivity	750 Tons/Hr	<i>Estimate</i>
	Operating Hrs	893 Hrs	<i>Productivity/Throughput</i>
	TSP Emissions	3.57 Tons/Yr	<i>E=(EF x Op Hrs)/2000</i>
	PM-10 Emissions	1.07 Tons/Yr	<i>30% of TSP</i>
Total South Portal PM ₁₀ Emissions		Conversions:	
		453.6 g/lb	8760 hr/yr
	PM ₁₀ = 4.20 tpy	2000 lb/ton	3600 sec/hr
	0.121 g/sec		

Production rate = 1,270,000 tpy; 600,000 tons to be sent via underground tunnel to East Portal

Emissions from the transport of coal with highway trucks on plant roads (South Portal to Seminoe II)

These roadways are reconstructed gravel roads for the purpose of connecting loadout with public roads using Wyoming hauling emission factor, with a tire factor adjustment for highway hauler

Input Data

Plant road silt content (s) = 5.1%
 Tire factor = 3.5

Mean speed 40 mph
 Grader hrs - Carbon Basin Road 975 hrs
 Grader hrs - Seminoe II Road 2,000 hrs

Reference

AP-42 table 13.2.2-1 (gravel-upgraded roadway)
 Assuming each pair is equivalent to a single large truck tire, a truck and pup combination have 14 equivalent tires, for a tire factor of 14/4 = 3.5
 Mine estimate

Conversions:

453.6 g/lb 8760 hr/yr
 2000 lb/ton 3600 s/hr

CARBON BASIN COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 0.670 MMtpy
 Vehicle Miles Traveled 57,429 VMT
 RT Haul Distance 6.0 miles
PM-10 Emissions (tpy) 39.87

SEMINOE II COAL TRANSPORT ROAD

Emission Factor 11.57 lb/VMT
 Number of Wet Days 100
 Truck Capacity 70 tons
 Truck Speed 40 mph
 Surface Silt Content 5.1 %
 Tire Correction Factor 3.5
 Percent Suspended (%) 62
 Control Factor (%) 60
 Control Method Water/Chemicals

Coal Hauled 0.670 MMtpy
 Vehicle Miles Traveled 38,286 VMT
 RT Haul Distance 4.0 miles
PM-10 Emissions (tpy) 26.578

CARBON BASIN TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 975
PM-10 Emissions (tpy) 1.70

SEMINOE II TRANSPORT ROAD REPAIR

Emission Factor 32 lb/hr
 Number of Wet Days 100
 Control Factor (%) 50
 Control Method Water
 Grader Hours/Year 2,000
PM-10 Emissions (tpy) 3.48

CARBON BASIN TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 41.6 138.6
 g/s 1.2 4.0

SEMINOE II TRANSPORT ROAD EMISSIONS

TOTALS PM10 TSP
 tpy 30.1 100.2
 g/s 0.9 2.9

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Mine - East Portal, Point Source Emissions
Point Source Emissions**

Production Rates are based on the 2007 Mine Plan Revisions

Activity at East Portal: bring coal out from underground mine; material handling, screening, send to gasifier (following Plant startup in Year 4). During Development Years 1-3, coal will be stockpiled; at start of Plant operations, coal will be directed to gasifier.

Year 3

Production rate = 300,000 tpy; to "run of mine" stockpiles (no coal to plant gasifiers)
300,000 tpy; to emergency (bankers) stockpile (no coal to plant gasifiers)

Material Handling Emissions (coal):	
$PM_{10} = k (.0032) ((U/5)^{1.3} / (M/2)^{1.4})$	AP-42, Section 13.2.4 Aggregate handling factors
k = 0.35	AP-42, Section 13.2.4 Aggregate handling factors
U = 14 mph	Seminole II Measurement
M = 6.9 %	AP-42 Table 13.2.4-1, Western Sfc Coal Mining, median
$PM_{10} = 8E-04 \text{ lb/ton}$	
Conveying by belt from conveyor C1 to C2 (material transfer) Controlled by fogger	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.023 \text{ tpy}$	
Conveying by belt from conveyor C2 to C3 (material transfer) Controlled by fogger	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.02 \text{ tpy}$	
Conveying by belt from conveyor C3 to C4 (material transfer) Controlled by fogger	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Conveying by belt from reclaim conveyor C5 to C6 (material transfer) Controlled by fogger	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Conveying by belt from conveyor C6 to Screener (material transfer) Controlled by fogger	
<i>Note: Conveyor C6 will be 3/4 covered, rather than completely enclosed.</i>	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Conveying by belt from Screener to conveyor C7 (material transfer) Controlled by fogger	
<i>Note: Conveyor C7 will be 3/4 covered, rather than completely enclosed.</i>	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Conveying by belt from conveyor C7 to C8 (material transfer) Controlled by fogger	
<i>Note: Conveyors C7, C8, & C9 will be 3/4 covered, rather than completely enclosed.</i>	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Conveying by belt from hopper to conveyor C10 material transfer) Controlled by fogger	
<i>Note: Conveyor C10 will be 3/4 covered, rather than completely enclosed.</i>	
Fogger control value = 90%	AP-42, Section 13.2.4.4
$PM_{10} = 0.01 \text{ tpy}$	
Total East Portal Point Source PM_{10} Emissions	
$PM_{10} = 0.10 \text{ tpy}$	Conversions:
0.003 g/sec	453.6 g/lb 8760 hr/yr
	2000 lb/ton 3600 sec/hr

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
 Saddleback Hills Mine, East Portal Fugitive Emissions
 Fugitive Emission Calculations
 (BACT Option: In-Pit Stacking Tubes)

Emissions starting in Year 3: 300,000 tons to "run of mine" stockpiles
 300,000 tons to emergency (bankers) stockpile

Emission Source	Description	Control	Additional Information
Dozer Reclaim	Cat D11 Dozer	None	
	Emission Factor	8.0 Lb/Hr	WDEQ 2002 Guidance
	Total Throughput	0 Tons/Yr	No dozer reclaim in Year 3
	Dozed Throughput	0 Tons/Yr	
	Dozer Productivity	750 Tons/Hr	Estimate for 300,000 Ton Pile
	Operating Hrs	0 Hrs	Productivity/Throughput
	TSP Emissions	0.00 Tons/Yr	$E=(EF \times Op \text{ Hrs})/2000$
	PM-10 Emissions	0.00 Tons/Yr	30% of TSP
Coal Stacker	Coal Dumping to Stockpile	Stacking Tubes	
	Emission Factor	0.017 Lb/Ton	WDEQ Emission Factor
	% Suspended	0.75	WDEQ Emission Factor
	Control Factor	50.00%	Estimated
	Material Dumped	600,000 Tons/Yr	Total Coal Through Storage
	TSP Emissions	1.91 Tons/Yr	$E=(EF \times \% \text{ sus} \times MD/2000) \times (1-CF)$
	PM-10 Emissions	0.57 Tons/Yr	30% of TSP
Coal Reclaim	Vibratory & Pile Activator Feeder	Passive Control	
	Emission Factor	0.017 Lb/Ton	WDEQ Emission Factor
	% Suspended	0.75	WDEQ Emission Factor
	Control Factor	100.00%	Estimated
	Material Reclaimed	300,000 Tons/Yr	Coal directed on to Emergency Pile
	TSP Emissions	0.00 Tons/Yr	$E=(EF \times \% \text{ sus} \times MR/2000) \times (1-CF)$
	PM-10 Emissions	0.00 Tons/Yr	30% of TSP
Coal Stockpile	Wind Erosion on Stockpiles	Water	
	Emission Factor	1.2 Lb/Acre/Hr	WDEQ Emission Factor
	Pile Size	2.0 Acres	Estimated
	Fraction Suspended	0.75	WDEQ Emission Factor
	Hours	8,760 Hours	Total Annual
	Ave. Wind Speed	5.03 meters/Sec	Adjusted for in-pit
	Wet Days	60	Seminole Mine 5-Year Average
	Control Factor	0.00%	
	TSP Emissions	33.10 Tons/Yr	$E=(EF \times AWS \times \% \text{ sus} \times PS \times ((365-WD)/365) \times (1-CF))/2000$
	PM-10 Emissions	9.93 Tons/Yr	
TOTAL PM-10 EMISSIONS	10.5 Tons/Yr		

Saddleback Hills Mine – Emission Calculations

Year Four

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Mine - East Portal, Point Source Emissions
Point Source Emissions**

Production Rates are based on the 2007 Mine Plan Revisions

Activity at East Portal: bring coal out from underground mine; material handling, screening, send to gasifier (following Plant startup in Year 4). During Development Years 1-3, coal will be stockpiled; at start of Plant operations, coal will be directed to gasifier.

Year 4

Production rate = 3,250,000 tpy

Material Handling Emissions (coal):		
$PM_{10} = k (U/5)^{1.3} / (M/2)^{1.4}$		AP-42, Section 13.2.4 Aggregate handling factors
k = 0.35		AP-42, Section 13.2.4 Aggregate handling factors
U = 14 mph		Seminole II Measurement
M = 6.9 %		AP-42 Table 13.2.4-1, Western Sfc Coal Mining, median
$PM_{10} = 8E-04 \text{ lb/ton}$		
Conveying by belt from conveyor C1 to C2 (material transfer) Controlled by fogger		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.123 \text{ tpy}$		
Conveying by belt from conveyor C2 to C3 (material transfer) Controlled by fogger		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from conveyor C3 to C4 (material transfer) Controlled by fogger		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from reclaim conveyor C5 to C6 (material transfer) Controlled by fogger		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from conveyor C6 to Screener (material transfer) Controlled by fogger		
<i>Note: Conveyor C6 will be 3/4 covered, rather than completely enclosed.</i>		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from Screener to conveyor C7 (material transfer) Controlled by fogger		
<i>Note: Conveyor C7 will be 3/4 covered, rather than completely enclosed.</i>		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from conveyor C7 to C9 (material transfer) Controlled by fogger		
<i>Note: Conveyors C7, C8, & C9 will be 3/4 covered, rather than completely enclosed.</i>		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.12 \text{ tpy}$		
Conveying by belt from hopper to conveyor C10 material transfer) Controlled by fogger		
<i>Note: Conveyor C10 will be 3/4 covered, rather than completely enclosed.</i>		
<i>Assume only 1.2 Mmtpy directed from emergency stockpile to reclaim hopper</i>		
Fogger control value = 90%		AP-42, Section 13.2.4.4
$PM_{10} = 0.05 \text{ tpy}$		
Total East Portal Point Source PM_{10} Emissions		
$PM_{10} = 0.86 \text{ tpy}$	Conversions:	
0.025 g/sec	453.6 g/lb	8760 hr/yr
	2000 lb/ton	3600 sec/hr

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Saddleback Hills Mine, East Portal Fugitive Emissions
Fugitive Emission Calculations
(BACT Option: In-Pit Stacking Tubes)

Emissions starting in Year 4

All coal through processing equipment to IGL Plant gasifiers

Emission Source	Description	Control	Additional Information
Dozer Reclaim	Cat D11 Dozer	None	
	Emission Factor	8.0 Lb/Hr	<i>WDEQ 2002 Guidance</i>
	Total Throughput	3,250,000 Tons/Yr	<i>Total Coal Through Storage</i>
	Dozed Throughput	1,500,000 Tons/Yr	<i>Portion to Dead Storage</i>
	Dozer Productivity	750 Tons/Hr	<i>Estimate for 300,000 Ton Pile</i>
	Operating Hrs	2,000 Hrs	<i>Productivity/Throughput</i>
	TSP Emissions	8.00 Tons/Yr	$E=(EF \times Op \text{ Hrs})/2000$
	PM-10 Emissions	2.40 Tons/Yr	30% of TSP
Coal Stacker	Coal Dumping to Stockpile	Stacking Tubes	
	Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
	% Suspended	0.75	<i>WDEQ Emission Factor</i>
	Control Factor	50.00%	<i>Estimated</i>
	Material Dumped	3,250,000 Tons/Yr	<i>Total Coal Through Storage</i>
	TSP Emissions	10.36 Tons/Yr	$E=(EF \times \% \text{ sus} \times MD/2000) \times (1-CF)$
	PM-10 Emissions	3.11 Tons/Yr	30% of TSP
Coal Reclaim	Vibratory & Pile Activator Feeder	Passive Control	
	Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
	% Suspended	0.75	<i>WDEQ Emission Factor</i>
	Control Factor	100.00%	<i>Estimated</i>
	Material Reclaimed	3,250,000 Tons/Yr	<i>Total Coal Through Storage</i>
	TSP Emissions	0.00 Tons/Yr	$E=(EF \times \% \text{ sus} \times MR/2000) \times (1-CF)$
	PM-10 Emissions	0.00 Tons/Yr	30% of TSP
Coal Stockpile	Wind Erosion on Stockpiles	Water	
	Emission Factor	1.2 Lb/Acre/Hr	<i>WDEQ Emission Factor</i>
	Pile Size	11.0 Acres	<i>Calculated from Pile Size</i>
	Fraction Suspended	0.75	<i>WDEQ Emission Factor</i>
	Hours	8,760 Hours	<i>Total Annual</i>
	Ave. Wind Speed	5.03 meters/Sec	<i>Adjusted for in-pit</i>
	Wet Days	60	<i>Seminole Mine 5-Year Average</i>
	Control Factor	0.00%	
	TSP Emissions	182.40 Tons/Yr	$E=(EF \times AWS \times \% \text{ sus} \times PS \times ((365-WD)/365) \times (1-CF))/2000$
	PM-10 Emissions	54.72 Tons/Yr	
TOTAL PM-10 EMISSIONS	60.2 Tons/Yr		

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
 Equipment Leaks Emission Summary

Process Stream	Service Type	Controlled Emissions		Uncontrolled Emissions	
		SOCMI Factors		SOCMI Factors	
		VOC Emissions (ton/yr)	HAP Emissions (ton/yr)	VOC Emissions (ton/yr)	HAP Emissions (ton/yr)
Acid Gas	Gas	0.09	0.09	0.12	0.12
Flare KO Drum Drainage	Gas	4.50	1.45	6.70	2.16
Gasifier Vent	Gas	0.14	0.14	0.22	0.22
Gasoline (Gas)	Gas	9.30	3.00	12.38	3.99
Gasoline (Light Liquid)	Light Liquid	10.42	3.36	36.22	11.67
Gasoline (Heavy Liquid)	Heavy Liquid	0.17	0.05	0.26	0.09
LPG	Light Liquid	0.77	0.00	2.21	0.00
Methanol Gas	Gas	0.99	0.99	1.28	1.28
Methanol Pure Liquid	Light Liquid	0.47	0.47	1.44	1.44
Methanol Product (MeOH 1)	Light Liquid	5.94	5.93	14.90	14.86
Methanol Product (MeOH 2)	Light Liquid	0.06	0.06	0.54	0.54
Methanol Product (MeOH 3)	Light Liquid	0.06	0.06	0.54	0.54
Methanol Product (MeOH 5)	Gas	0.35	0.35	0.50	0.50
Mixed Fuel Gas	Gas	0.40	0.01	1.77	0.06
MTG Fuel Gas	Gas	3.88	0.04	5.44	0.06
Propylene	Gas	22.11	0.00	24.36	0.00
Total		59.63	16.00	108.86	37.52

Individual HAPs	Controlled Emissions		Uncontrolled Emissions	
	SOCMI Factors		SOCMI Factors	
	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
Carbonyl Sulfide (COS)	0.05	0.23	0.08	0.35
Methanol (MeOH)	1.80	7.87	4.39	19.22
C6 - C10 Aromatics (Assumed to be Benzene)	1.80	7.90	4.10	17.96
Total	3.65	16.00	8.57	37.52

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Equipment Leaks HAP Emission Summary**

Controlled Emissions (SOCMI Factors)

Process Stream	COS		MeOH		Benzene*	
	(lb/hr)	(ton/yr)	(lb/hr)	(ton/yr)	(lb/hr)	(ton/yr)
Acid Gas	1.96E-02	8.58E-02	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Flare KO Drum Drainage	1.16E-03	5.10E-03	0.00E+00	0.00E+00	3.30E-01	1.45E+00
Gasifier Vent	3.26E-02	1.43E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Gasoline (Gas)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	6.84E-01	3.00E+00
Gasoline (Light Liquid)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	7.67E-01	3.36E+00
Gasoline (Heavy Liquid)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.22E-02	5.33E-02
LPG	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Methanol Gas	0.00E+00	0.00E+00	2.26E-01	9.90E-01	0.00E+00	0.00E+00
Methanol Pure Liquid	0.00E+00	0.00E+00	1.08E-01	4.74E-01	0.00E+00	0.00E+00
Methanol Product (MeOH 1)	0.00E+00	0.00E+00	1.35E+00	5.93E+00	0.00E+00	0.00E+00
Methanol Product (MeOH 2)	0.00E+00	0.00E+00	1.29E-02	5.64E-02	0.00E+00	0.00E+00
Methanol Product (MeOH 3)	0.00E+00	0.00E+00	1.28E-02	5.62E-02	0.00E+00	0.00E+00
Methanol Product (MeOH 5)	0.00E+00	0.00E+00	8.02E-02	3.51E-01	0.00E+00	0.00E+00
Mixed Fuel Gas	0.00E+00	0.00E+00	3.23E-03	1.42E-02	0.00E+00	0.00E+00
MTG Fuel Gas	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.01E-02	4.41E-02
Propylene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total	5.33E-02	2.34E-01	1.80E+00	7.87E+00	1.80E+00	7.90E+00

* Benzene is assumed from emissions of C6-C10 aromatics.

Uncontrolled HAP Summary

Uncontrolled Emissions (SOCMI Factors)

Process Stream	COS		MeOH		Benzene*	
	(lb/hr)	(ton/yr)	(lb/hr)	(ton/yr)	(lb/hr)	(ton/yr)
Acid Gas	2.79E-02	1.22E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Flare KO Drum Drainage	1.73E-03	7.59E-03	0.00E+00	0.00E+00	4.92E-01	2.15E+00
Gasifier Vent	4.92E-02	2.15E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Gasoline (Gas)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	9.10E-01	3.99E+00
Gasoline (Light Liquid)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	2.66E+00	1.17E+01
Gasoline (Heavy Liquid)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.94E-02	8.51E-02
LPG	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Methanol Gas	0.00E+00	0.00E+00	2.92E-01	1.28E+00	0.00E+00	0.00E+00
Methanol Pure Liquid	0.00E+00	0.00E+00	3.28E-01	1.44E+00	0.00E+00	0.00E+00
Methanol Product (MeOH 1)	0.00E+00	0.00E+00	3.39E+00	1.49E+01	0.00E+00	0.00E+00
Methanol Product (MeOH 2)	0.00E+00	0.00E+00	1.23E-01	5.40E-01	0.00E+00	0.00E+00
Methanol Product (MeOH 3)	0.00E+00	0.00E+00	1.23E-01	5.38E-01	0.00E+00	0.00E+00
Methanol Product (MeOH 5)	0.00E+00	0.00E+00	1.15E-01	5.02E-01	0.00E+00	0.00E+00
Mixed Fuel Gas	0.00E+00	0.00E+00	1.44E-02	6.32E-02	0.00E+00	0.00E+00
MTG Fuel Gas	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.41E-02	6.18E-02
Propylene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total	7.88E-02	3.45E-01	4.39E+00	1.92E+01	4.10E+00	1.80E+01

* Benzene is assumed from emissions of C6-C10 aromatics.

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Equipment Leaks
Pump LDAR Control Effectiveness Calculation**

Leak detection and repair (LDAR) control effectiveness factors for valves and connectors are based on "HON reg neg" factors from *Protocol for Equipment Leak Emission Estimates* (Table G-1) [EPA-453/R-95-017]. These factors assume leak definitions of 500 ppmv for valves and connectors, which equate to the leak definitions expected to be used at MBFP. However, the HON reg neg leak definition for pumps in light liquid service is 1,000 ppmv, which is more stringent than the 2,000 ppmv leak definition planned for the LDAR program to be implemented at MBFP. Consequently, the LDAR control effectiveness factor for a 2,000 ppmv pump leak definition is calculated below. All table numbers refer to the *Protocol for Equipment Leak Emission Estimates (Protocol)*.

Values Used in Pump LDAR Control Effectiveness Calculation

Source Type	SOCMI	
Equipment Type	Pumps in Light Liquid Service	
LDAR Program	Monthly monitoring with a leak definition of 2,000 ppmv for pumps	
Initial Leak Fraction (LKFRAC)	14.9%	
Occurrence Rate (OC)	7.0%	
Recurrence Rate (R) *	0.0%	* Similar to the calculations shown in the <i>Protocol</i> for the HON reg neg
Unsuccessful Repair Rate *	0.0%	LDAR control effectiveness calculations, the recurrence rate and unsuccessful
Successful Repair Rate (FR)	100.0%	repair rates are assumed to be zero. (Table G-3, footnote a)

Initial Leak Frequency

Table 5-4 $ALR = (0.13 \times LKFRAC) + 6.7 \times 10^{-4}$
 Where: ALR = Average SOCMI leak rate (kg/hr/source)
 $ALR_{initial} = 0.02$ kg/hr/source (from Figure 5-3)

Solving for LKFRAC $LKFRAC = (0.02 - 6.7 \times 10^{-4})/0.13$
 $LKFRAC = 0.149$

Occurrence Rate

Table G-3 $OC = 0.47 \times LKFRAC$
 $OC = 0.47 \times 0.149$
 $OC = 0.070$

Steady-State Leak Frequency

Table 5-9 $Y_i = Z_i - (FR \times Z_i) + (FR \times Z_i \times R)$
 $Z_{i+1} = OC \times (1 - Y_i) + Y_i$
 Where: Y_i = Leak frequency immediately after LDAR monitoring
 Z_i = Leak frequency immediately before LDAR monitoring

Monitoring Cycle	Y_i Leak Frequency After Monitoring (%)	Z_i Leak Frequency Before Monitoring (%)
1	0.0%	14.9%
2	0.0%	7.0%
3	0.0%	7.0%
4	0.0%	7.0%
5	0.0%	7.0%
6	0.0%	7.0%
Steady-State Leak Frequency	3.5%	

Average of leak frequency before and after 6th monitoring cycle

Average Steady-State Leak Rate $ALR_{ss} = (0.13 \times LKFRAC) + 6.7 \times 10^{-4}$
 (based on 3.5% steady-state leak frequency) $ALR_{ss} = (0.13 \times 0.035) + 6.7 \times 10^{-4}$
 $ALR_{ss} = 0.00522$ kg/hr/source

Pump LDAR Control Effectiveness $CE = (ALR_{initial} - ALR_{ss})/ALR_{initial} \times 100\%$
 $CE = (0.02 - 0.00522)/0.02 \times 100\%$
 $CE = 73.9\%$

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Acid Gas Process Stream

Stream Name: Acid Gas
 Service Type: Gas
 Hours of Operation: 8760
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	55.94%	1.27E-02	47.86%
H2O	7732-18-5	N	N	18.02	3.37%	1.87E-03	7.05%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	40.16%	1.18E-02	44.37%
COS	463-58-1	Y	Y	60.07	0.28%	4.68E-05	0.18%
NH3	7664-41-7	N	N	17.03	0.25%	1.45E-04	0.55%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	2.66E-02	100.00%

Assumed Octane
 Assumed Octene
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	0.28%
Weight % VOC	0.28%
Weight % HAP	0.28%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	204	0.0003	0.0003	8760	2.64E-03	3.30E-02
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		27	0.0079	0.0079	8760	7.62E-02	7.62E-02
Connectors	0.00183	93.00%	130	0.0000	0.0000	8760	4.52E-04	6.45E-03
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		16	0.0007	0.0007	8760	6.51E-03	6.51E-03
Totals				0.01	0.01		0.09	0.12

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).

² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table G-1). Assumes monthly monitoring with leak definition of 500 ppmv for valves and connectors.

³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.28%	0.28%	8760	1.96E-02	8.58E-02	2.79E-02	1.22E-01
Cl2	0.00%	0.28%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	0.28%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	0.28%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	0.00%	0.28%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.02	0.09	0.03	0.12

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Flare KO Drum Drainage Process Stream

Stream Name: Flare KO Drum Drainage
Service Type: Gas
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	22.46%	8.02E-03	29.34%
H2	1333-74-0	N	N	2.02	1.16%	5.77E-03	21.11%
CO2	124-38-9	N	N	44.01	18.13%	4.12E-03	15.08%
H2O	7732-18-5	N	N	18.02	7.50%	4.16E-03	15.23%
CH4	74-82-8	N	N	16.04	0.03%	2.05E-05	0.07%
Ar	7440-37-1	N	N	39.95	0.37%	9.29E-05	0.34%
N2	7727-37-9	N	N	28.01	0.12%	4.25E-05	0.16%
H2S	7783-06-4	N	N	34.08	0.16%	4.72E-05	0.17%
COS	463-58-1	Y	Y	60.07	0.06%	9.44E-06	0.03%
NH3	7664-41-7	N	N	17.03	0.01%	3.15E-06	0.01%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	23.93%	2.09E-03	7.66%
C4 - C12 Olefins	N/A	Y	N	112.21	4.20%	3.74E-04	1.37%
C6 - C10 Naphthenes	N/A	Y	N	112.21	5.77%	5.14E-04	1.88%
C6 - C10 Aromatics	N/A	Y	Y	78.11	16.11%	2.06E-03	7.54%
TOTALS					100.00%	2.73E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	50.09%
Weight % VOC	50.06%
Weight % HAP	16.16%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	68	0.0163	0.0163	8760	1.57E-01	1.96E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		8	0.4167	0.4165	8760	4.02E+00	4.02E+00
Connectors	0.00183	93.00%	48	0.0031	0.0031	8760	2.97E-02	4.24E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		4	0.0301	0.0300	8760	2.90E-01	2.90E-01
Totals				0.47	0.47		4.50	6.70

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.06%	50.06%	8760	1.16E-03	5.10E-03	1.73E-03	7.59E-03
Cl2	0.00%	50.06%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	50.06%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	50.06%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	16.11%	50.06%	8760	3.30E-01	1.45E+00	4.92E-01	2.15E+00
Total				0.33	1.45	0.49	2.16

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Gasifier Vent Process Stream**

Stream Name: Gasifier Vent
 Service Type: Gas
 Hours of Operation: 8760
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	44.91%	1.60E-02	35.98%
H2	1333-74-0	N	N	2.02	2.33%	1.15E-02	25.89%
CO2	124-38-9	N	N	44.01	36.27%	8.24E-03	18.49%
H2O	7732-18-5	N	N	18.02	15.00%	8.33E-03	18.68%
CH4	74-82-8	N	N	16.04	0.07%	4.09E-05	0.09%
Ar	7440-37-1	N	N	39.95	0.74%	1.86E-04	0.42%
N2	7727-37-9	N	N	28.01	0.24%	8.50E-05	0.19%
H2S	7783-06-4	N	N	34.08	0.32%	9.45E-05	0.21%
COS	463-58-1	Y	Y	60.07	0.11%	1.89E-05	0.04%
NH3	7664-41-7	N	N	17.03	0.01%	6.30E-06	0.01%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	4.46E-02	100.00%

Assumed Octane
 Assumed Octane
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	0.18%
Weight % VOC	0.11%
Weight % HAP	0.11%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	957	0.0008	0.0005	8760	5.01E-03	6.26E-02
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		112	0.0209	0.0132	8760	1.28E-01	1.28E-01
Connectors	0.00183	93.00%	804	0.0002	0.0001	8760	1.13E-03	1.61E-02
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		55	0.0015	0.0009	8760	9.04E-03	9.04E-03
Totals				0.02	0.01		0.14	0.22

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.11%	0.11%	8760	3.26E-02	1.43E-01	4.92E-02	2.15E-01
Cl2	0.00%	0.11%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	0.11%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	0.11%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	0.00%	0.11%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.03	0.14	0.05	0.22

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Gasoline (Gas) Process Stream

Stream Name: Gasoline (Gas)
Service Type: Gas
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	47.85%	4.19E-03	41.52%
C4 - C12 Olefins	N/A	Y	N	112.21	8.39%	7.48E-04	7.41%
C6 - C10 Naphthenes	N/A	Y	N	112.21	11.54%	1.03E-03	10.19%
C6 - C10 Aromatics	N/A	Y	Y	78.11	32.21%	4.12E-03	40.87%
TOTALS					100.00%	1.01E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	100.00%
Weight % VOC	100.00%
Weight % HAP	32.21%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	50	0.0239	0.0239	8760	2.31E-01	2.88E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		9	0.9360	0.9360	8760	9.04E+00	9.04E+00
Connectors	0.00183	93.00%	26	0.0033	0.0033	8760	3.22E-02	4.59E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Totals				0.96	0.96		9.30	12.38

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	100.00%	8760	0.00E+00	8760	0.00E+00	0.00E+00
Cl2	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	32.21%	100.00%	8760	6.84E-01	3.00E+00	9.10E-01	3.99E+00
Total				0.68	3.00	0.91	3.99

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Gasoline (Light Liquid) Process Stream**

Stream Name: Gasoline (Light Liquid)
Service Type: Light Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	47.85%	4.19E-03	41.52%
C4 - C12 Olefins	N/A	Y	N	112.21	8.39%	7.48E-04	7.41%
C6 - C10 Naphthenes	N/A	Y	N	112.21	11.54%	1.03E-03	10.19%
C6 - C10 Aromatics	N/A	Y	Y	78.11	32.21%	4.12E-03	40.87%
TOTALS					100.00%	1.01E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	100.00%
Weight % VOC	100.00%
Weight % HAP	32.21%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	487	0.2355	0.2355	8760	2.27E+00	1.89E+01
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	24	0.1247	0.1247	8760	1.20E+00	4.61E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	348	0.0446	0.0446	8760	4.30E-01	6.15E+00
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		45	0.6750	0.6750	8760	6.52E+00	6.52E+00
Totals				1.08	1.08		10.42	36.22

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	32.21%	100.00%	8760	7.67E-01	3.36E+00	2.66E+00	1.17E+01
Total				0.77	3.36	2.66	11.67

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Gasoline (Heavy Liquid) Process Stream

Stream Name: Gasoline (Heavy Liquid)
Service Type: Heavy Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	47.85%	4.19E-03	41.52%
C4 - C12 Olefins	N/A	Y	N	112.21	8.39%	7.48E-04	7.41%
C6 - C10 Naphthenes	N/A	Y	N	112.21	11.54%	1.03E-03	10.19%
C6 - C10 Aromatics	N/A	Y	Y	78.11	32.21%	4.12E-03	40.87%
TOTALS					100.00%	1.01E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	100.00%
Weight % VOC	100.00%
Weight % HAP	32.21%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		6	0.0014	0.0014	8760	1.33E-02	1.33E-02
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	6	0.0008	0.0008	8760	7.42E-03	1.06E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		1	0.0150	0.0150	8760	1.45E-01	1.45E-01
Totals				0.02	0.02		0.17	0.26

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	100.00%	8760	0.00E+00	8760	0.00E+00	0.00E+00
Cl2	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	32.21%	100.00%	8760	1.22E-02	5.33E-02	1.94E-02	8.51E-02
Total				0.01	0.05	0.02	0.09

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
LPG Process Stream

Stream Name: LPG
Service Type: Light Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	8.34%	2.98E-03	13.04%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	3.60%	5.00E-04	2.19%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	21.86%	7.79E-03	34.13%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	37.82%	6.51E-03	28.49%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	28.38%	5.06E-03	22.15%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	2.28E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	91.66%
Weight % VOC	91.66%
Weight % HAP	0.00%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	28	0.0124	0.0124	8760	1.20E-01	9.98E-01
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	2	0.0095	0.0095	8760	9.19E-02	3.52E-01
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	20	0.0023	0.0023	8760	2.27E-02	3.24E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		4	0.0550	0.0550	8760	5.31E-01	5.31E-01
Totals				0.08	0.08		0.77	2.21

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	91.66%	8760	0.00E+00	8760	0.00E+00	0.00E+00
Cl2	0.00%	91.66%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	91.66%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	91.66%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	0.00%	91.66%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.00	0.00	0.00	0.00

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Gas Process Stream**

Stream Name: **Methanol Gas**
 Service Type: **Gas**
 Hours of Operation: **8760**
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.02%	6.44E-06	0.02%
H2	1333-74-0	N	N	2.02	0.00%	3.19E-06	0.01%
CO2	124-38-9	N	N	44.01	0.30%	6.92E-05	0.22%
H2O	7732-18-5	N	N	18.02	3.16%	1.75E-03	5.49%
CH4	74-82-8	N	N	16.04	0.03%	1.59E-05	0.05%
Ar	7440-37-1	N	N	39.95	0.06%	1.61E-05	0.05%
N2	7727-37-9	N	N	28.01	0.03%	1.14E-05	0.04%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	96.19%	3.00E-02	94.01%
Ethanol	64-17-5	Y	N	46.07	0.05%	1.04E-05	0.03%
Dimethyl Ether	115-10-6	Y	N	46.07	0.03%	7.31E-06	0.02%
Methyl Acetate	79-20-9	Y	N	74.08	0.08%	1.10E-05	0.03%
Propanol	71-23-8	Y	N	60.10	0.02%	4.00E-06	0.01%
Butanol	71-36-3	Y	N	74.12	0.02%	2.60E-06	0.01%
Acetone	67-64-1	Y	N	58.08	0.00%	3.31E-07	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	1.33E-07	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	3.19E-02	100.00%

Assumed Octane
 Assumed Octane
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	96.42%
Weight % VOC	96.40%
Weight % HAP	96.19%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	5	0.0023	0.0023	8760	2.22E-02	2.78E-01
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		1	0.1003	0.1003	8760	9.68E-01	9.68E-01
Connectors	0.00183	93.00%	2	0.0002	0.0002	8760	2.38E-03	3.41E-02
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Totals				0.10	0.10		0.99	1.28

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	96.19%	96.40%	8760	2.26E-01	9.90E-01	2.92E-01	1.28E+00
C6 - C10 Aromatics	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.23	0.99	0.29	1.28

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Pure Liquid Process Stream

Stream Name: **Methanol Pure Liquid**
 Service Type: **Light Liquid**
 Hours of Operation: **8760**
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	100.00%	3.12E-02	100.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	3.12E-02	100.00%

Assumed Octane
 Assumed Octane
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	100.00%
Weight % VOC	100.00%
Weight % HAP	100.00%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	16	0.0077	0.0077	8760	7.47E-02	6.22E-01
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	2	0.0104	0.0104	8760	1.00E-01	3.84E-01
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	8	0.0010	0.0010	8760	9.89E-03	1.41E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		2	0.0300	0.0300	8760	2.90E-01	2.90E-01
Totals				0.05	0.05		0.47	1.44

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	100.00%	100.00%	8760	1.08E-01	4.74E-01	3.28E-01	1.44E+00
C6 - C10 Aromatics	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.11	0.47	0.33	1.44

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Product (MeOH 1) Process Stream**

Stream Name: Methanol Product (MeOH 1)
Service Type: Light Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.02%	6.44E-06	0.02%
H2	1333-74-0	N	N	2.02	0.00%	3.19E-06	0.01%
CO2	124-38-9	N	N	44.01	0.30%	6.92E-05	0.22%
H2O	7732-18-5	N	N	18.02	3.16%	1.75E-03	5.49%
CH4	74-82-8	N	N	16.04	0.03%	1.59E-05	0.05%
Ar	7440-37-1	N	N	39.95	0.06%	1.61E-05	0.05%
N2	7727-37-9	N	N	28.01	0.03%	1.14E-05	0.04%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	96.19%	3.00E-02	94.01%
Ethanol	64-17-5	Y	N	46.07	0.05%	1.04E-05	0.03%
Dimethyl Ether	115-10-6	Y	N	46.07	0.03%	7.31E-06	0.02%
Methyl Acetate	79-20-9	Y	N	74.08	0.08%	1.10E-05	0.03%
Propanol	71-23-8	Y	N	60.10	0.02%	4.00E-06	0.01%
Butanol	71-36-3	Y	N	74.12	0.02%	2.60E-06	0.01%
Acetone	67-64-1	Y	N	58.08	0.00%	3.31E-07	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	1.33E-07	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	3.19E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	96.42%
Weight % VOC	96.40%
Weight % HAP	96.19%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	134	0.0625	0.0625	8760	6.03E-01	5.03E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	22	0.1102	0.1101	8760	1.06E+00	4.07E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	96	0.0119	0.0119	8760	1.14E-01	1.63E+00
Open-ended Lines	0.00170		16	0.0262	0.0262	8760	2.53E-01	2.53E-01
Sampling Connections	0.01500		28	0.4050	0.4049	8760	3.91E+00	3.91E+00
Totals				0.62	0.62		5.94	14.90

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	96.19%	96.40%	8760	1.35E+00	5.93E+00	3.39E+00	1.49E+01
C6 - C10 Aromatics	0.00%	96.40%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				1.35	5.93	3.39	14.86

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Product (MeOH 2) Process Stream**

Stream Name: Methanol Product (MeOH 2)
Service Type: Light Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.08%	2.89E-05	0.09%
H2	1333-74-0	N	N	2.02	0.02%	1.09E-04	0.34%
CO2	124-38-9	N	N	44.01	0.42%	9.63E-05	0.30%
H2O	7732-18-5	N	N	18.02	3.32%	1.84E-03	5.74%
CH4	74-82-8	N	N	16.04	0.08%	4.81E-05	0.15%
Ar	7440-37-1	N	N	39.95	0.44%	1.09E-04	0.34%
N2	7727-37-9	N	N	28.01	0.18%	6.42E-05	0.20%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	95.46%	2.98E-02	92.84%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	3.21E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	95.54%
Weight % VOC	95.46%
Weight % HAP	95.46%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Light Liquids	0.00403	88.00%	10	0.0046	0.0046	8760	4.46E-02	3.71E-01
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Connectors	0.00183	93.00%	10	0.0012	0.0012	8760	1.18E-02	1.69E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Totals				0.01	0.01		0.06	0.54

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	95.46%	8760	0.00E+00	8760	0.00E+00	0.00E+00
Cl2	0.00%	95.46%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	95.46%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	95.46%	95.46%	8760	1.29E-02	5.64E-02	1.23E-01	5.40E-01
C6 - C10 Aromatics	0.00%	95.46%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.01	0.06	0.12	0.54

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Product (MeOH 3) Process Stream**

Stream Name: Methanol Product (MeOH 3)
Service Type: Light Liquid
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.07%	2.57E-05	0.08%
H2	1333-74-0	N	N	2.02	0.02%	1.16E-04	0.36%
CO2	124-38-9	N	N	44.01	0.42%	9.65E-05	0.30%
H2O	7732-18-5	N	N	18.02	3.62%	2.01E-03	6.25%
CH4	74-82-8	N	N	16.04	0.08%	5.15E-05	0.16%
Ar	7440-37-1	N	N	39.95	0.46%	1.16E-04	0.36%
N2	7727-37-9	N	N	28.01	0.19%	6.76E-05	0.21%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	95.12%	2.97E-02	92.28%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	3.22E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	95.21%
Weight % VOC	95.12%
Weight % HAP	95.12%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions			Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	0	0.0000	0.0000	8760	0.00E+00
Valves-Light Liquids	0.00403	88.00%	10	0.0046	0.0046	8760	4.44E-02
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00
Relief Valves-Gas/Vapor	0.10400		0	0.0000	0.0000	8760	0.00E+00
Connectors	0.00183	93.00%	10	0.0012	0.0012	8760	1.18E-02
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00
Sampling Connections	0.01500		0	0.0000	0.0000	8760	0.00E+00
Totals				0.01	0.01		0.06

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	95.12%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	95.12%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	95.12%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	95.12%	95.12%	8760	1.28E-02	5.62E-02	1.23E-01	5.38E-01
C6 - C10 Aromatics	0.00%	95.12%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.01	0.06	0.12	0.54

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Methanol Product (MeOH 5) Process Stream**

Stream Name: Methanol Product (MeOH 5)
Service Type: Gas
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	15.02%	5.36E-03	7.09%
H2	1333-74-0	N	N	2.02	9.73%	4.83E-02	63.83%
CO2	124-38-9	N	N	44.01	3.93%	8.92E-04	1.18%
H2O	7732-18-5	N	N	18.02	0.05%	3.03E-05	0.04%
CH4	74-82-8	N	N	16.04	2.78%	1.73E-03	2.29%
Ar	7440-37-1	N	N	39.95	47.22%	1.18E-02	15.63%
N2	7727-37-9	N	N	28.01	19.58%	6.99E-03	9.24%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-58-1	Y	Y	32.04	1.70%	5.29E-04	0.70%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	0.00%	0.00E+00	0.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	7.56E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	4.47%
Weight % VOC	1.70%
Weight % HAP	1.70%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	125	0.0027	0.0010	8760	9.78E-03	1.22E-01
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		16	0.0745	0.0282	8760	2.72E-01	2.72E-01
Connectors	0.00183	93.00%	136	0.0008	0.0003	8760	2.85E-03	4.08E-02
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		27	0.0181	0.0069	8760	6.63E-02	6.63E-02
Totals				0.10	0.04		0.35	0.50

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	1.70%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	1.70%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	1.70%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	1.70%	1.70%	8760	8.02E-02	3.51E-01	1.15E-01	5.02E-01
C6 - C10 Aromatics	0.00%	1.70%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.08	0.35	0.11	0.50

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Mixed Fuel Gas Process Stream**

Stream Name: **Mixed Fuel Gas**
 Service Type: **Gas**
 Hours of Operation: **8760**
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	1.88%	6.70E-04	1.36%
H2	1333-74-0	N	N	2.02	2.06%	1.02E-02	20.76%
CO2	124-38-9	N	N	44.01	3.38%	7.68E-04	1.56%
H2O	7732-18-5	N	N	18.02	0.01%	7.40E-06	0.02%
CH4	74-82-8	N	N	16.04	39.92%	2.49E-02	50.67%
Ar	7440-37-1	N	N	39.95	15.43%	3.86E-03	7.87%
N2	7727-37-9	N	N	28.01	7.59%	2.71E-03	5.52%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.99%	3.09E-04	0.63%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	2.02%	6.73E-04	1.37%
Ethylene	74-85-1	Y	N	28.05	0.20%	6.96E-05	0.14%
Propane	74-98-6	Y	N	44.10	7.00%	1.59E-03	3.23%
Propylene	115-07-1	Y	N	42.08	0.36%	8.56E-05	0.17%
Isobutane	75-28-5	Y	N	58.12	16.30%	2.80E-03	5.71%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	2.32%	4.14E-04	0.84%
Isopentane	78-78-4	Y	N	72.15	0.47%	6.53E-05	0.13%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.08%	6.80E-06	0.01%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	4.91E-02	100.00%

Assumed Octane
 Assumed Octane
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	69.65%
Weight % VOC	27.71%
Weight % HAP	0.99%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	90	0.0299	0.0119	8760	1.15E-01	1.44E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Relief Valves-Gas/Vapor	0.10400		1	0.0724	0.0288	8760	2.78E-01	2.78E-01
Connectors	0.00183	93.00%	11	0.0010	0.0004	8760	3.77E-03	5.39E-02
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Totals				0.10	0.04		0.40	1.77

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	27.71%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	27.71%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	27.71%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.99%	27.71%	8760	3.23E-03	1.42E-02	1.44E-02	6.32E-02
C6 - C10 Aromatics	0.00%	27.71%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.00	0.01	0.01	0.06

**Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
MTG Fuel Gas Process Stream**

Stream Name: **MTG Fuel Gas**
 Service Type: **Gas**
 Hours of Operation: **8760**
 This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	34.27%	1.22E-02	34.25%
H2	1333-74-0	N	N	2.02	0.01%	6.11E-05	0.17%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.39%	2.17E-04	0.61%
CH4	74-82-8	N	N	16.04	22.67%	1.41E-02	39.56%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	8.92%	2.97E-03	8.31%
Ethylene	74-85-1	Y	N	28.05	5.69%	2.03E-03	5.68%
Propane	74-98-6	Y	N	44.10	6.95%	1.58E-03	4.41%
Propylene	115-07-1	Y	N	42.08	0.30%	7.24E-05	0.20%
Isobutane	75-28-5	Y	N	58.12	2.52%	4.34E-04	1.21%
N-Butane	106-97-8	Y	N	58.12	0.43%	7.48E-05	0.21%
Butylene	25167-67-3	Y	N	56.11	0.78%	1.39E-04	0.39%
Isopentane	78-78-4	Y	N	72.15	5.20%	7.21E-04	2.02%
C4 - C12 Paraffins	N/A	Y	N	114.23	7.48%	6.54E-04	1.83%
C4 - C12 Olefins	N/A	Y	N	112.21	2.69%	2.39E-04	0.67%
C6 - C10 Naphthenes	N/A	Y	N	112.21	1.31%	1.17E-04	0.33%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.38%	4.91E-05	0.14%
TOTALS					100.00%	3.57E-02	100.00%

Assumed Octane
 Assumed Octane
 Assumed Cyclooctane
 Assumed Benzene

Weight % TOC	65.33%
Weight % VOC	33.74%
Weight % HAP	0.38%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	60	0.0187	0.0097	8760	9.33E-02	1.17E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		4	0.5958	0.3077	8760	2.97E+00	2.97E+00
Relief Valves-Gas/Vapor	0.10400		2	0.1359	0.0702	8760	6.77E-01	6.77E-01
Connectors	0.00183	93.00%	88	0.0074	0.0038	8760	3.67E-02	5.24E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		2	0.0196	0.0101	8760	9.77E-02	9.77E-02
Totals				0.78	0.40		3.88	5.44

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	33.74%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	33.74%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	33.74%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	33.74%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	0.38%	33.74%	8760	1.01E-02	4.41E-02	1.41E-02	6.18E-02
Total				0.01	0.04	0.01	0.06

Medicine Bow Fuel & Power Industrial Gasification & Liquefaction Plant
Propylene Process Stream

Stream Name: Propylene
Service Type: Gas
Hours of Operation: 8760
This piping is included in the LDAR program.

Chemical Name	CAS Number	VOC	HAP	Molecular Weight (lb/lb-mol)	Weight %	Mole Fraction	Mole Percent
CO	630-08-0	N	N	28.01	0.00%	0.00E+00	0.00%
H2	1333-74-0	N	N	2.02	0.00%	0.00E+00	0.00%
CO2	124-38-9	N	N	44.01	0.00%	0.00E+00	0.00%
H2O	7732-18-5	N	N	18.02	0.00%	0.00E+00	0.00%
CH4	74-82-8	N	N	16.04	0.00%	0.00E+00	0.00%
Ar	7440-37-1	N	N	39.95	0.00%	0.00E+00	0.00%
N2	7727-37-9	N	N	28.01	0.00%	0.00E+00	0.00%
H2S	7783-06-4	N	N	34.08	0.00%	0.00E+00	0.00%
COS	463-58-1	Y	Y	60.07	0.00%	0.00E+00	0.00%
NH3	7664-41-7	N	N	17.03	0.00%	0.00E+00	0.00%
O2	7782-44-7	N	N	32.00	0.00%	0.00E+00	0.00%
SO2	7446-09-5	N	N	64.06	0.00%	0.00E+00	0.00%
Cl2	7782-50-5	N	Y	70.91	0.00%	0.00E+00	0.00%
HCl	7647-01-0	N	Y	36.46	0.00%	0.00E+00	0.00%
MeOH	67-56-1	Y	Y	32.04	0.00%	0.00E+00	0.00%
Ethanol	64-17-5	Y	N	46.07	0.00%	0.00E+00	0.00%
Dimethyl Ether	115-10-6	Y	N	46.07	0.00%	0.00E+00	0.00%
Methyl Acetate	79-20-9	Y	N	74.08	0.00%	0.00E+00	0.00%
Propanol	71-23-8	Y	N	60.10	0.00%	0.00E+00	0.00%
Butanol	71-36-3	Y	N	74.12	0.00%	0.00E+00	0.00%
Acetone	67-64-1	Y	N	58.08	0.00%	0.00E+00	0.00%
MEK	78-93-3	Y	N	72.11	0.00%	0.00E+00	0.00%
Ethane	74-84-0	N	N	30.07	0.00%	0.00E+00	0.00%
Ethylene	74-85-1	Y	N	28.05	0.00%	0.00E+00	0.00%
Propane	74-98-6	Y	N	44.10	0.00%	0.00E+00	0.00%
Propylene	115-07-1	Y	N	42.08	100.00%	2.38E-02	100.00%
Isobutane	75-28-5	Y	N	58.12	0.00%	0.00E+00	0.00%
N-Butane	106-97-8	Y	N	58.12	0.00%	0.00E+00	0.00%
Butylene	25167-67-3	Y	N	56.11	0.00%	0.00E+00	0.00%
Isopentane	78-78-4	Y	N	72.15	0.00%	0.00E+00	0.00%
C4 - C12 Paraffins	N/A	Y	N	114.23	0.00%	0.00E+00	0.00%
C4 - C12 Olefins	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Naphthenes	N/A	Y	N	112.21	0.00%	0.00E+00	0.00%
C6 - C10 Aromatics	N/A	Y	Y	78.11	0.00%	0.00E+00	0.00%
TOTALS					100.00%	2.38E-02	100.00%

Assumed Octane
Assumed Octane
Assumed Cyclooctane
Assumed Benzene

Weight % TOC	100.00%
Weight % VOC	100.00%
Weight % HAP	0.00%

Fugitive Emissions - SOCM1 Factors				Controlled Emissions				Uncontrolled Emissions
Equipment Type	SOCM1 Emission Factor ¹ (kg/hr-source)	% Control With LDAR ^{2,3}	Source Count	TOC Emission Rate (kg/hr)	VOC Emission Rate (kg/hr)	Hours of Operation	VOC Emissions (tpy)	VOC Emissions (tpy)
Valves-Gas	0.00597	92.00%	40	0.0191	0.0191	8760	1.84E-01	2.31E+00
Valves-Light Liquids	0.00403	88.00%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Valves-Heavy Liquids	0.00023		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Light Liquids	0.01990	73.90%	0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Pump Seals-Heavy Liquids	0.00862		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Compressor Seals-Gas	0.22800		8	1.8240	1.8240	8760	1.76E+01	1.76E+01
Relief Valves-Gas/Vapor	0.10400		4	0.4160	0.4160	8760	4.02E+00	4.02E+00
Connectors	0.00183	93.00%	8	0.0010	0.0010	8760	9.89E-03	1.41E-01
Open-ended Lines	0.00170		0	0.0000	0.0000	8760	0.00E+00	0.00E+00
Sampling Connections	0.01500		2	0.0300	0.0300	8760	2.90E-01	2.90E-01
Totals				2.29	2.29		22.11	24.36

¹ EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 2-1).
² EPA-453/R-95-017 Protocol for Equipment Leak Emission Estimates (Table 5-2). Assumes monthly monitoring with leak definition of 10,000 ppmv.
³ Assumes monthly monitoring with leak definition of 2,000 ppmv for pumps in light liquid service. See Pump LDAR Control Effectiveness Calculation page.

HAP Emissions - SOCM1 Factors				Controlled Emissions		Uncontrolled Emissions	
HAP	Individual HAP Weight %	VOC Weight %	Hours of Operation	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)	HAP Emissions (lb/hr)	HAP Emissions (ton/yr)
COS	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cl2	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
HCl	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
MeOH	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
C6 - C10 Aromatics	0.00%	100.00%	8760	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Total				0.00	0.00	0.00	0.00

Appendix C
Manufacturer Specifications

DKRW**ESTIMATED PERFORMANCE PG7121**

Load Condition		BASE	BASE	50%	BASE	50%	BASE	50%
Exhaust Pressure Loss	in H2O	15.0	15.0	15.0	15.0	15.0	15.0	15.0
Ambient Temperature	deg F	85.	85.	85.	45.	45.	0.	0.
Ambient Relative Humidity	%	16.0	16.0	16.0	80.0	80.0	80.0	80.0
Fuel Type		Methane	Methane	Methane	Methane	Methane	Methane	Methane
Fuel LHV	BTU/lb	21,515	21,515	21,515	21,515	21,515	21,515	21,515
Fuel Temperature	deg F	80	80	80	80	80	80	80

EMISSIONS

NOx	ppmvd @ 15% O2	154.	25.	25.	25.	25.	25.	25.
NOx AS NO2	lb/hr	389.	70.	41.	78.	45.	80.	46.
CO	ppmvd	10.	25.	*****	25.	*****	25.	*****
CO	lb/hr	16.	39.	*****	42.	*****	47.	*****
UHC	ppmvw	7.	7.	-999.	7.	-999.	7.	-999.
<u>UHC</u>	lb/hr	7.	7.	804.	8.	859.	8.	919.
Particulates	lb/hr	5	5	5	5	5	5	5

(PM10 Front-half Filterable Only)

*** See Combustion For Emissions

EXHAUST ANALYSIS % VOL.

Argon	0.90	0.87	0.89	0.85	0.89	0.87	0.91
Nitrogen	75.07	72.36	74.29	72.29	74.26	73.42	75.11
Oxygen	14.04	12.89	15.19	12.76	15.14	13.47	15.53
Carbon Dioxide	3.08	3.29	2.40	3.35	2.42	3.14	2.34
Water	6.92	10.60	7.23	10.75	7.29	9.10	6.12

SITE CONDITIONS

Elevation	ft	7355.0
Site Pressure	psia	11.2
Inlet Loss	in H2O	3.50
Exhaust Loss	in H2O	15.00 @ ISO Conditions
Application		Air-Cooled Generator
<u>Power Factor (lag)</u>		0.8
Combustion System		Quiet Combustor

Emission information based on GE recommended measurement methods. NOx emissions are corrected to 15% O2 without heat rate correction and are not corrected to ISO reference condition per 40CFR 60.335(a)(1)(i). NOx levels shown will be controlled by algorithms within the SPEEDTRONIC control system.

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MEDICINE BOW - NITROGEN INJECTION WITH AIR EXTRACTION
ESTIMATED PERFORMANCE PG7121 - IGCC. PRELIMINARY. FOR STUDY PURPOSES ONLY

Load Condition		BASE	BASE	BASE
Inlet Loss	in H2O	3.5	3.5	3.5
Exhaust Pressure Loss	in H2O	14.0	14.0	14.0
Ambient Temperature	deg F	45.	-12.	85.
Ambient Relative Humidity	%	60.0	80.0	18.0

EMISSIONS

NOx	ppmvd @ 15% O2	25.	25.	25.
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LHV	BTU/lb	16399.6	16399.6	16399.6
Flow Rate	lb/h	44,450. ✓	47,910.	40,240.
Pressure	psia	335.	335.	335.
Temperature	°F	300.	300.	300.

EXHAUST ANALYSIS % VOL.

Argon		1.03	1.03	1.03
Nitrogen		76.82	77.34	76.71
Oxygen		12.22	12.08	12.37
Carbon Dioxide		3.23	3.32	3.17
Water		6.71	6.23	6.73

SITE CONDITIONS

Elevation	ft	7354.9		
Site Pressure	psia	11.2		
Exhaust Loss	in H2O	14.00 @ ISO Conditions		
Application		Air-Cooled Generator		
Power Factor (lag)		0.8		
Combustion System		IGCC Combustor		

Appendix D
Major Equipment List and SCCs

IGL Plant Source Classification Codes

Emission Unit	SCC Code
Auxiliary Boiler	10200602
Black-Start Generators (3)	20100201
Catalyst Regenerator	30600106
CO2 Vent Stack	N/A
Coal Storage	30501009
Firewater Pump	20200102
Flares	30490024
Fugitives	30600811
Gasifier Preheaters (5)	30600105
Gasoline Storage Tanks	2501000120
HGT Reactor Charge Heater	30600106
Methanol Storage Tanks	2510000260
Reactivation Heater	30600106
Turbine and HRSG Trains (3)	20100301

IGL Plant Equipment List

Equipment Type	Equipment Name	Equipment Tag	No. of Identical Items
Centrifugal pump	PROCESS CONDENSATE PUMP	P-13001A	1
Centrifugal pump	PROCESS CONDENSATE PUMP SPARE	P-13001B	1
Centrifugal pump	SWS BOTTOMS PUMP	P-13002A	1
Centrifugal pump	SWS BOTTOMS PUMP SPARE	P-13002B	1
Centrifugal pump	AMMONIA STRIPPER BOTTOMS PUMP	P-13004A	1
Centrifugal pump	AMMONIA STRIPPER BOTTOMS PUMP SPARE	P-13004B	1
Centrifugal pump	SOUR KO DRUM PUMP	P-13005A	1
Centrifugal pump	SOUR KO DRUM PUMP SPARE	P-13005B	1
Centrifugal pump	SOUR SHIFT PC PUMP	P-13007A	1
Centrifugal pump	SOUR SHIFT PC PUMP SPARE	P-13007B	1
Aircooler	COS HYDROLYSIS RXTR EFFLUENT	AC-13001	1
Aircooler	SWS PUMPAROUND CLR	AC-13002	1
Aircooler	SOUR GAS COOLER	AC-13003	1
Aircooler	BLOWDOWN WATER COOLER	AC-13004	1
Aircooler	SOUR SHIFT REACTOR EFFLUENT CONDENSER	AC-13006	1
Shell and tube	LP STEAM GENERATOR	E-13002	1
Shell and tube	COS HYDROLYSIS PREHEATER	E-13003	1
Shell and tube	LP BFW PREHEATER	E-13004	1
Shell and tube	HG GUARD BED PREHEATER	E-13005	1
Shell and tube	SWS REBOILER	E-13006	1
Shell and tube	AMMONIA STRIPPER REBOILER	E-13007	1
Shell and tube	SOUR SHIFT LP STEAM GEN	E-13008	1
Shell and tube	SOUR SHIFT FEED/EFFLUENT	E-13009	1
Shell and tube	SOUR SHIFT MP STEAM GEN	E-13010	1
Shell and tube	AMMONIA STRIP FEED PRE	E-13011	1
Shell and tube	MP STEAM GENERATOR	E-13011	1
Shell and tube	1ST MP BFW PREHEATER	E-13012	1
Shell and tube	1ST MP BFW PREHEATER	E-13013	1
Shell and tube	VLP STEAM GEN	E-13014	1
Shell and tube	SWS FEED PREHEATER	E-13015	1
Shell and tube	SHIFTED HG GB PREHTR	E-13016	1
Tower	SOUR WATER STRIPPER	T-13001	1
Tower	AMMONIA STRIPPER	T-13002	1
Reactor	COS HYDROLYSIS REACTOR	R-13001	1
Reactor	CO SHIFT REACTOR	R-13002	3
KO Drum	HOT SYNGAS KO DRUM	V-13001	1
KO Drum	COLD SYNGAS KO DRUM	V-13002	1
KO Drum	SOUR GAS KO DRUM	V-13003	1
Tank	CAUSTIC INJECTION DRUM	V-13004	1
KO Drum	SOUR SHIFT KO DRUM	V-13005	1
KO Drum	COLD SHIFTED KO DRUM	V-13006	1
Aircooler	No 1 Vac Flash Overhead Con	03E-303	1
Aircooler	No 2 Vac Flash Overhead Con	03E-303	1
Aircooler	HP Flash Trim Air Cooler	03E-302	1
Aircooler	No 1 Vac Flash Overhead Con	03E-203	1
Aircooler	No 2 Vac Flash Overhead Con	03E-203	1
Aircooler	HP Flash Trim Air Cooler	03E-202	1
Aircooler	No 1 Vac Flash Overhead Con	03E-103	1

IGL Plant Equipment List

Aircooler	No 2 Vac Flash Overhead Con	03E-103	1
Aircooler	HP Flash Trim Air Cooler	03E-102	1
Aircooler	Quench Water Startup Clr	03E-005	1
Aircooler	HP Flash Trim Air Cooler	03E-402	1
Aircooler	No 1 Vac Flash Overhead Con	03E-403	1
Aircooler	No 2 Vac Flash Overhead Con	03E-403	1
Aircooler	HP Flash Trim Air Cooler	03E-502	1
Aircooler	No 1 Vac Flash Overhead Con	03E-503	1
Aircooler	No 2 Vac Flash Overhead Con	03E-503	1
Centrifugal pump	Injector Coolant Pump	02-P001A/B/C	3
Centrifugal pump	Lockhopper Circ. Pump	02-P102A/B	10
Centrifugal pump	Slag Sump Pump	02P-103A/B	10
Centrifugal pump	Preheat Water Pump	02-P-104A	10
Centrifugal pump	Vac. Flash Cond. Pump	03P-104A/B	10
Centrifugal pump	Slurry Transfer Pump	01P-103	1
Centrifugal pump	Slurry Transfer Pump	01P-203	1
Centrifugal pump	Slurry Transfer Pump	01P-303	1
Centrifugal pump	Scrubber Feed Pump	03P-002	6
Centrifugal pump	Settler Bottoms Pump	03-P005	4
Centrifugal pump	Grey Water Discharge Pump	03P-006	4
Centrifugal pump	Filter Feed Pump	03P-008A/B/C	3
Centrifugal pump	Grinding Water Pumps	03P-009A/B/C	3
Centrifugal pump	Quench Water Pump	03P-101A/B	10
Centrifugal pump	Vac Flash Bottoms Pump	03P-103A/B	10
Centrifugal pump	Grinding Sump Pump	01P-005A/B	2
Centrifugal pump	Fines Area Sump Pump	03P-007A/B	2
Ejector	Startup aspirator	02X-105	5
Filter	Quench water Strainer	02F-102	10
GE Quench Gasifier	Quench Gasifier	02R-101	1
GE Quench Gasifier	Quench Gasifier	02R-101	1
GE Quench Gasifier	Quench Gasifier	02R-101	1
GE Quench Gasifier	Quench Gasifier	02R-101	1
GE Quench Gasifier	Quench Gasifier	02R-101	1
KO Drum	Gasifier Seal Pot	02V-102	5
KO Drum	Aspirator Separator	02V-103	5
KO Drum	Injector Coolant Gas Sepr	02V-105	5
KO Drum	HP Flash Drum	03V-103	5
KO Drum	LP Flash Drum	03V-105	5
KO Drum	Vacuum Flash Drum No 1	03V-106	5
KO Drum	Vacuum Flash Drum No2	03V-108	5
KO Drum	HP Flash OH Drum	03V-104	5
KO Drum	No 1 Vac Flash OH Drum	03V-107	5
KO Drum	No 1 Vac Flash OH Drum	03V-109	5
KO Drum	Lockhopper	02V-106	5
KO Drum	Lockhopper Flush Drum	02V-107	5
Other	Slag Crusher	02X-103	5
Shell and tube	HP Flash OH Condenser	03E-101	5
Shell and tube	HP Flash OH Condenser	03E-201	5
Shell and tube	HP Flash OH Condenser	03E-301	5
Shell and tube	HP Flash OH Condenser	03E-401	5
Shell and tube	HP Flash OH Condenser	03E-501	5

IGL Plant Equipment List

Tank	Grey Water Tank	03T-002	1
Tank	Slurry Additive Tank	03T-003	1
Tank	Mill Discharge tank	01T-104	3
Tank	Slurry Tank	01T-105	3
Tank	Injector Coolant Tank	02T-001	1
Tank	Settler	03T-001	2
Tank	Filter Feed Tank	03T-004	1
Tank	Filtrate Tank	03T-005	1
Tower	Syngas Scrubber	03V-101	5
	Slurry Additive Tank Agit	01A-001	1
	Grinding Sump Agitator	01A-004	1
	Mill Discharge Tank Agitr	01A-102	3
	Slurry Tank Agitator	01A-103	5
	Grind Mill Disch HVAC Fan	01C-101	3
	Trommel Screen	01F-101	5
	Fluxant feed Conveyor	01L-101	3
	Grinding Sump	01T-106	1
	Fluxant Weigh Feeder	01W-101	3
	Slag Sump Agitator	02A-102	5
	Oxygen Filter	02F-101	10
	Slurry Vibrating Screen	02F-102	3
	Coarse Slag Screen	02F-103	5
	Slag Drag Conveyor	02L-101	5
	Slag Sump	02T-102	5
	Oxygen Silencer	02X-101	5
	Feed Injector	02X-102	10
	Preheat Burner	02X-104	5
	Settler Rake	03A-001	2
	Fines Sump Agitator	03A-002	1
	Filter Feed tank Agitator	03A-003	1
	filtrate Tank Agitator	03A-004	1
	Fines Filter Press	03F-001	3
	Fines Sump	03T-003	1
	Nozzle Scrubber	03X-101	5
	Gasifier Refractory	02R-101-int	5
Filter	Crude Methanol Filter	H-321 A/B	2
Filter	Crude Methanol Filter	H-322 A/B	2
Compressor	Syngas Compressor	J-111	1
Compressor	Loop Circulator	J-121	1
Aircooler	Syngas Comp Spilback	E-211	1
aircooler	Loop condenser No.1	E-221	1
aircooler	Loop condenser No.2	E-222	1
Shell and tube	Syngas purifict preheater	E-111	1
Shell and tube	loop interchanger no.1	E-121	1
Shell and tube	loop interchanger no.2	E-123	1
Reactor	Syngas purification vessl	D-111	2
Reactor	Methano Synthesis Reactor	D-121	1
Reactor	Methano Synthesis Reactor	D-122	1
KO Drum	Syngas KO Drum	D-311	1
Reactor	PSA Unit - 5 drums total	L-121	5
KO Drum	Methanol Catchpot No.1	D-321	1

IGL Plant Equipment List

KO Drum	Methanol Catchpot No.2	D-322	1
KO Drum	Letdown Vessel	D-323	1
Centrifugal pump	MeOH Charge	P-01 A/B	2
Centrifugal pump	Deethanizer Feed	P-02 A/B	2
Centrifugal pump	MTG Process Water	P-03 A/B	2
Centrifugal pump	Deethanizer Ovhd Cooler	P-04 A/B	2
Centrifugal pump	Stabilizer OVHD	P-05 A/B	2
Centrifugal pump	Lean oil Supply	P-06 A/B	2
Centrifugal pump	Splitter OVHD	P-07 A/B	2
Centrifugal pump	Splitter BTTMS	P-08 A/B	2
Centrifugal pump	Absorber BTTMS	P-09 A/B	2
Centrifugal pump	MeOH Recovery OVHD	P-10 A/B	2
Centrifugal pump	MeOH BTTMS	P-11 A/B	2
Centrifugal pump	HGT Charge	P-351 A/B	2
Centrifugal pump	Stripper OVHD	P-352 A/B	2
Tower	Deethanizer	C-1	1
Furnace	Regeneration Heater	B-1	1
Furnace	Reactivation Heater	B-2	1
Furnace	HGT Reactor Charge	B-351	1
Compressor	MTG Recycle gas	K-1	1
Compressor	Regeneration Air	K-2	1
Compressor	Regeneration Gas	K-3	1
Compressor	HGT Recycle	K-351 A/B	1
Aircooler	MTG Reactor Effluent Coolers	EA-1	1
Aircooler	Regeneration Cooler	EA-2	1
Aircooler	Deethanizer Ovhd Condenser	EA-3	1
Aircooler	Stabilizer OVHD Condenser	EA-4	1
Aircooler	LPG Cooler	EA-5	1
Aircooler	Lean Oil Cooler	EA-6	1
Aircooler	Splitter OVHD Condenser	EA-7	1
Aircooler	Light Gasoline Cooler	EA-8	1
Aircooler	Heavy Gasoline Cooler	EA-9	1
Aircooler	MeOH Recovery Condenser	EA-10	1
Aircooler	LT Separator Feed Cooler	EA-351	1
Aircooler	Stripper OVHD Condenser	EA-352	1
Shell and Tube	MeOH Preheater	E-1	1
Shell and Tube	MeOH Vaporizer	E-2	1
Shell and Tube	MeOH Supper Heater	E-3	1
Shell and Tube	Recycle Gas/Effluent HX	E-4	1
Shell and Tube	HP Steam Generator	E-5	1
Shell and Tube	Regeneration Gas Interchanger	E-6	1
Shell and Tube	Deethanizer Reboiler	E-7	1
Shell and Tube	Deethanizer Feed / Bttms	E-8	1
Shell and Tube	Stabilizer Reboiler	E-9	1
Shell and Tube	Splitter Reboiler	E-10	1
Shell and Tube	HGT Feed/ Stripper BTTMS	E-351	1
Shell and Tube	HGT Feed / reactor Effluent	E-352	1
Shell and Tube	HGT Recycle Gas / HT Separator	E-353	1
Shell and Tube	Cold Stripper Feed / LT Sep Feed	E-354	1
Shell and Tube	Stripper Reboiler	E-355	1
Shell and Tube	Treated Heavy Gasoline Cooler	E-356	1

IGL Plant Equipment List

KO Drum	MeOH Flash Drum	D-1	1
KO Drum	HP Steam Drum	D-2	1
KO Drum	Product Separator	D-3	1
KO Drum	Deethanizer OH Drum	D-6	1
KO Drum	Stabilizer OH Drum	D-7	1
KO Drum	Splitter OH Drum	D-8	1
KO Drum	Absorber Feed KO Drum	D-9	1
KO Drum	MeOH OVHD Drum	D-11	1
KO Drum	HGT Feed Surge Drum	D-351	1
KO Drum	Low Temp Separator	D-353	1
KO Drum	Stripper OH Drum	D-355	1
Tower	Absoorber	C-4	1
Tower	MeOH Recovery Column	C-5	1
Tower	Product Stripper	C-351	1
Tower	Gasoline Splitter	C-3	
KO Drum	MTG Process Water Flash Drum	D-4	1
KO Drum	Regeneration Gas Separator	D-5	1
KO Drum	Absorber OVHD KO Drum	D-10	1
KO Drum	Height Temp. Separator	D-352	1
KO Drum	HGT Recycle Gas KO Drum	D-354	1
Centrifugal pump	MTG Water Pump		2
Centrifugal pump	Methanol Transfer Pump		2
Centrifugal pump	Gasoline Send-Out Pump		3
Centrifugal pump	Sulfur Send-Out Pump		2
Centrifugal pump	Slops Tank Transfer Pump		1
Centrifugal pump	Acid Gas Wash Drum Pump	P-31001 A/B	2
Centrifugal pump	Contact Cond. Circ. Pump	P-31005 AB	2
Centrifugal pump	Desuperheater Circ. Pump	P-31006	1
Centrifugal pump	Sulfur Degassing Pump	P-31003 A/B	2
Centrifugal pump	Sulfur Transfer Pump	P-31004 A/B	2
Ejector	Sulfur Pit Vent Ejector	EJ-31001	1
Ejector	Degassing Vent Ejector	EJ-31002	1
Fan	Start-Up Blower	BL-31002	1
Furnace	Claus Reaction Furnace	H-31001	1
Aircooler	Waste Steam Condenser	AC-31006	1
Aircooler	Spent Caustic Cooler	AC-31009	1
Aircooler	Contact Cond. H2O Cooler	AC-31014	1
Shell and Tube	No. 1 Condenser	E-31002	1
Shell and Tube	No. 2 Condenser	E-31003	1
Shell and Tube	No. 3 Condenser	E-31004	1
Shell and Tube	No. 4 Condenser	E-31005	1
Shell and Tube	No. 1 Reheater	E-31007	1
Shell and Tube	No. 2 Reheater	E-31008	1
Shell and Tube	No. 3 Reheater	E-31009	1
Shell and Tube	Hydrogen. Effl. Cooler*	E-31012	1
Shell and Tube	Hydrogenator Preheater	E-31013	1
Shell and Tube	Waste Heat Boiler	E-31001	1
Tower	Sulfur Degasser	T-31xxx	1
Tower	Desuperhtr/Contact Cond.	T-31003	1
KO Drum	Acid Gas KO Drum	V-31001	1
KO Drum	Steam Drum	V-31003	1

IGL Plant Equipment List

KO Drum	*Claus Converter	R-31001/2/3	1
KO Drum	S/U Blower KO Drum	V-31xxx	1
Centrifugal pump	HP Lean Solvent Pump	P-21001	4
Centrifugal pump	Reflux Pump	P-21002	4
Centrifugal pump	Loaded Solvent Pump	P-21003	4
Centrifugal pump	Semi-Lean Solvent Pump	P-21004	4
Centrifugal pump	LP Lean Solvent Pump	P-21005	4
Centrifugal pump	Semi-Lean Pump Unshifted	P-21xxx	4
Centrifugal pump	H2S Pump for Unshifted	P-21xxx	4
Centrifugal pump	Hydraulic Turbine 1	P-21xxx	4
Centrifugal pump	Hydraulic Turbine 2	P-21xxx	4
Compressor	Stripping Gas Compressor	K-21001	2
Compressor	H2S Flash Gas Comp. 2	K-21002	2
Compressor	CO2 Recycle Compressor	K-21003	2
Compressor	H2S Flash Gas Comp. 1	K-21xxx	2
Compressor	TG Comp. Stage 1	K-21010	2
Compressor	TG Comp. Stage 2	K-21011	2
Aircooler	H2S Recycle Gas Cooler 2	E-21007	2
Aircooler	H2S Flash Gas Cooler 2	E-21003	2
Aircooler	H2S Recycle Gas Cooler	E-21006	2
Aircooler	Reflux Condenser	E-21005	2
Aircooler	CO2 Recycle Gas Cooler	E-21011	2
Aircooler	H2S Flash Gas Cooler 1	E-21010	2
Aircooler	Shifted Feed Gas Cooler	E-21xxx	2
Aircooler	TG Compressor Cooler 1	E-21xxx	2
Shell and Tube	Feed / Product Exchanger	E-21001	2
Shell and Tube	Lean / Rich Exchanger	E-21002	2
Shell and Tube	Lean Solvent Chiller	E-21008	2
Shell and Tube	Loaded Solvent Chiller	E-21009	2
KO Drum	H2S Rich MP Flash Drum	V-21001	2
KO Drum	Flash Gas KO Drum	V-21002	2
KO Drum	Reflux Drum	V-21003	2
KO Drum	CO2 Recycle Flash Drum	V-21004	2
KO Drum	CO2 MP Flash Drum	V-21005	2
KO Drum	CO2 LP Flash Drum	V-21006	2
KO Drum	H2S Rich LP Flash Drum	V-21xxx	2
Shell and Tube	Stripper Reboiler	E-21004	1
Tower	H2S Absorber Shifted Gas	C-21001	2
Tower	H2S Concentrator	C-21002	2
Tower	H2S Stripper	C-21003	2
Tower	CO2 Absorber Shifted Gas	C-21004	2
Tower	CO2 Absorber Unshifted Gas	C-21005	2
Tower	H2S Absorber Unshifted Gas	C-21006	2
Other	Refrigeration Package A/B	Z-21001AB	2
Tank	Methanol Tanks		2
Tank	Gasoline Product Tanks		8
Tank	MTG Water Tank		1
Tank	Liquid Sulfur Storage Tk.		2
Tank	Slops Tank		1
Tank	Off-spec methanol tank		1
Tank	Off-spec gasoline tank		1

IGL Plant Equipment List

Tank	Heavy Gasoline Tank	1
KO Drum	METHANOL LET DOWN DRUM	1
KO Drum	Flare KO Drums	4
Tank	LPG Tanks	2
Flare	Flare Stack	2
Centrifugal pump	MTG Water Pump	2
Centrifugal pump	Methanol Transfer Pump	2
Centrifugal pump	Gasoline Send-Out Pump	3
Centrifugal pump	Sulfur Send-Out Pump	2
Centrifugal pump	Slops Tank Transfer Pump	1
Centrifugal pump	Flare KO Drum Pump	4
	Air Separation Unit	2
	Power Plant	1
	Auxiliary Boiler	1
	Fire Protection	1
	Set Up Transformers	1
	Switchyard	1
	Water Treatment System	1

Along with the equipment listed above, there will be several conveyors that will be used to transfer coal from the mine to the coal storage, and from storage to the plant. There will also be conveyors to move slag from the gasifiers to the slag storage area.

Appendix E

BACT Review of Recent NO_x Limits for Combined Cycle Combustion Turbines Fueled With Other Gaseous Fuels

BACT Review of Recent NO_x Limits for Combined Cycle Combustion Turbines Fueled With Other Gaseous Fuels

Facility	Fuel	Capacity	NO _x Emission Limit	Pollution Control Method	Basis	Permit Date (Permit Number)
Bayport Energy Center LP, Bayport Energy Center	Mixture of low-sulfur fuel gas and NG	225 MMBtu/hr	3.5 ppmvd (3-hour), 1.9 ppmvd (annual)	Dry low- NO _x combustors and low- NO _x duct burners	BACT-PSD	10/20/2003 (P1031)
Union Carbide Corp., Texas City Operations	Primary fuel gas	14.2 MW	25 ppmvd @ 15% O ₂ (each)	Low- NO _x combustor	Other case-by-case	1/23/2003 (PSD-TX-841)
Tampa Electric Company TECO-Polk Power	Syngas from petcoke and coal	190 MW	15 ppmvd @ 15% O ₂ (each)	Combustion improvement, nitrogen diluent injection	Other case-by-case	12/23/2002 (PSD-FL-194)
Exxon Mobil, Exxon Mobil Shute Creek	Proprietary mix of process gas, sales gas, and hydrogen	35.8 MW	8 ppmvd @ 15% O ₂ (30-day rolling average)	Proprietary low-BTU fuel and low- NO _x burners	BACT-PSD	6/19/2002 (MD-771)
Global Energy, Inc., Lima Energy Company	Syngas	170 MW	15 ppmvd @ 15% O ₂	Dilution prior to combustion and dilution injection into combustion zone	BACT-PSD	3/26/2002 (03-13445)
Kentucky Pioneer Energy, LLC, Kentucky Pioneer Energy, LLC - Trapp	Synthesis gas	197 MW	15 ppmvd @ 15% O ₂	Steam injection	BACT-PSD	6/7/2001 (V-00-049)
Borden Chemicals and Plastics Operating, LP (COGEN III Unit)	NG / acetylene	473 MMBtu/hr	62 ppmvd @ 15% O ₂	Steam injection	RACT	5/29/2001 (PSD-LA-539)
Borden Chemicals and Plastics Operating, LP (COGEN II Unit)	NG / acetylene	471 MMBtu/hr	51 ppm @ 15% O ₂	Steam injection	BACT-PSD (prior determination)	5/29/2001 (PSD-LA-535 [M-2])
Valero Refining Co. - Texas City	Refinery fuel gas	Not available	27 ppm @ 15% O ₂	Not available	Other case-by-case	2/23/2000 (PSD-TX-822M2)
Sweeny Cogeneration	Residue gas	121.3 MW (each)	15 ppm @ 15% O ₂	Dry low- NO _x burners	Other case-by-	9/30/1998

Facility	Fuel	Capacity	NO _x Emission Limit	Pollution Control Method	Basis	Permit Date (Permit Number)
Limited Partnership			(natural gas only), 25 ppm @ 15% O ₂ (natural gas and residue gas)		case	(PSD-TX-857)
Star Enterprise	Syngas or LSDF	90 MW (each)	16 ppm @ 15% O ₂	Nitrogen injection (firing syngas), steam injection (firing LSDF)	LAER	3/30/1998 (APC-97/0503-CONST)

Note:

Information was obtained from the RACT/BACT/LAER Clearinghouse based on process type 15.250 (Large Combustion Turbines [>25 MW], Combined-Cycle & Cogeneration [>25 MW], Other Gaseous Fuel & Gaseous Fuel Mixtures). The search period included the ten-year period from 9/28/1997 to 9/28/2007.

Acronyms:

LAER = Lowest achievable emission rate

LSDF = Low-sulfur diesel fuel

MMBtu/hr = Million British thermal units per hour

MW = Megawatt

NG = Natural gas

NO_x = Nitrogen oxides

O₂ = Oxygen

PSD = Prevention of significant deterioration

RACT = Reasonable available control technology

Appendix F
Coal Storage BACT Cost Analysis

**Best Available Control Technology (BACT) Analysis
Coal Storage and Reclaim Facility
Saddle Back Hills Mine
Carbon County, Wyoming**

Arch Coal Company

April 5, 2007

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Summary

Three options were considered to provide 300,000 tons of live coal storage, as required by the longwall operation and the companion coal-to-liquids plant.

1. Stacking tubes located in the pit excavated for the underground portal (reference drawing no. 6945-L010)
2. Stacking tubes located on the surface next to the pit (reference drawing no. 6945-L020)
3. Covered slot storage (reference drawing no. 6945-L030)

The first two options differ in the placement of the stacking tubes. In Option 1 the storage facility is on the pit floor, with the excavated spoils placed in a large berm on the west and north sides of the pit. This configuration is intended to reduce storage pile erosion and resulting PM₁₀ emissions, by sheltering the pile from prevailing winds. Support for this approach is provided at the end of this document.

Option 1 would reduce PM₁₀ emissions by roughly 25% relative to Option 2. With a calculated, incremental PM₁₀ emissions control cost of \$6,902 per ton, Option 1 is proposed as BACT. Option 3 would eliminate PM₁₀ emissions from the storage facility, but the additional capital cost would result in an incremental PM₁₀ emissions control cost of \$54,119 per ton relative to Option 1. This option is therefore considered infeasible.

Analysis

Table 1 presents a top-down comparison between first Options 3 and 1, then between Options 1 and 2. Facility designs and capital costs for all three options were developed by Roberts & Schaefer. Operating costs were provided by Arch Coal Company. A mine life of 20 years was used in the analysis, along with a discount rate of 8% per year. Capital and operating costs were converted to levelized annual costs to enable direct comparison between options. PM₁₀ emissions were projected for each option based on emission factors approved by the Wyoming Department of Environmental Quality. Incremental emissions control costs between any two options were obtained from dividing differential levelized costs by differential emissions.

Tables 2 and 3 show the calculation of PM₁₀ emissions for Options 1 and 2, respectively (Option 3 would generate no emissions). The maximum production throughput is assumed to be 3.2 million tons per year. Sources of emissions for both options include the stacking tubes, dozer activity to groom the storage pile and assist the reclaim operations, and wind erosion from the storage pile. The reclaim system is designed with passive controls (100% control) to eliminate emissions from that source.

TABLE 1

Saddleback Hills Mine Storage System

BACT Analysis: In-Pit Tube Stacker vs. Covered Slot Storage

	Option 3: Covered Slot Storage	Option 1: Tube Stacker in Pit	Option 3 vs. Option 1 Comparison
Capital Cost	\$157,200,000	\$84,700,000	
Mine Life (Years)	20	20	
Discount Rate (annual cost of capital)	8.0%	8.0%	
Net Present Value of Annual O&M Cost	\$0	\$7,363,611	
Levelized Annual Cost	\$7,860,000	\$4,603,181	
Annual PM-10 Emissions (tpy)	0.0	60.2	
Differential Emissions Control (tpy)			60.2
Differential Technology Cost per Year			\$3,256,819
Incremental Control Cost (per ton PM-10)			\$54,119

BACT Analysis: In-Pit Tube Stacker vs. Surface Tube Stacker

	Option 1: Tube Stacker in Pit	Option 2: Tube Stacker Surf.	Option 1 vs. Option 2 Comparison
Capital Cost	\$84,700,000	\$82,200,000	
Mine Life (Years)	20	20	
Discount Rate (annual cost of capital)	8.0%	8.0%	
Net Present Value of Annual O&M Cost	\$7,363,611	\$7,363,611	
Levelized Annual Cost	\$4,603,181	\$4,478,181	
Annual PM-10 Emissions (tpy)	60.2	78.3	
Differential Emissions Control (tpy)			18.1
Differential Technology Cost per Year			\$125,000
Incremental Control Cost (per ton PM-10)			\$6,902

Common assumptions used for Options 1 and 2 are:

1. All emission sources except wind erosion are identical for both options
2. Dozer operations on the storage pile average 2,000 hours per year
3. Stacking tubes are credited with 50% emissions control in comparison to a free drop
4. Maximum storage pile extent is 11 acres
5. The number of wet days (defined as having 0.01" of precipitation or more) per year is 60, taken from five years of meteorological data at the nearby Seminole mine.

TABLE 2**BACT Option 1 (In-Pit Stacking Tubes) PM-10 Emissions**

Emission Source	Type	Description	Control	Additional Information
Dozer Reclaim	Fugitive	Cat D11 Dozer	None	
		Emission Factor	8.0 Lb/Hr	<i>WDEQ 2002 Guidance</i>
		Total Throughput	3,200,000 Tons/Yr	<i>Total Coal Through Storage</i>
		Dozed Throughput	1,500,000 Tons/Yr	<i>Portion to Dead Storage</i>
		Dozer Productivity	750 Tons/Hr	<i>Estimate for 300,000 Ton Pile</i>
		Operating Hrs	2,000 Hrs	<i>Productivity/Throughput</i>
		TSP Emissions	8.00 Tons/Yr	<i>E=(EF x Op Hrs)/2000</i>
		PM-10 Emissions	2.40 Tons/Yr	<i>30% of TSP</i>
Coal Stacker	Fugitive	Coal Dumping to Stockpile	Stacking Tubes	
		Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
		% Suspended	0.75	<i>WDEQ Emission Factor</i>
		Control Factor	50.00%	<i>Estimated</i>
		Material Dumped	3,200,000 Tons/Yr	<i>Total Coal Through Storage</i>
		TSP Emissions	10.20 Tons/Yr	<i>E=(EFx% sus x MD/2000)x(1-CF)</i>
		PM-10 Emissions	3.06 Tons/Yr	<i>30% of TSP</i>
Coal Reclaim	Fugitive	Vibratory & Pile Activator Feeder	Passive Control	
		Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
		% Suspended	0.75	<i>WDEQ Emission Factor</i>
		Control Factor	100.00%	<i>Estimated</i>
		Material Reclaimed	3,200,000 Tons/Yr	<i>Total Coal Through Storage</i>
		TSP Emissions	0.00 Tons/Yr	<i>E=(EFx% sus x MR/2000)x(1-CF)</i>
		PM-10 Emissions	0.00 Tons/Yr	<i>30% of TSP</i>
Coal Stockpile	Fugitive	Wind Erosion on Stockpiles	Water	
		Emission Factor	1.2 Lb/Acre/Hr	<i>WDEQ Emission Factor</i>
		Pile Size	11.0 Acres	<i>Calculated from Pile Size</i>
		Fraction Suspended	0.75	<i>WDEQ Emission Factor</i>
		Hours	8,760 Hours	<i>Total Annual</i>
		Ave. Wind Speed	5.03 meters/Sec	<i>Adjusted for in-pit</i>
		Wet Days	60	<i>Seminole Mine 5-Year Average</i>
		Control Factor	0.00%	
		TSP Emissions	182.40 Tons/Yr	<i>E=(EF x AWS x %sus x PS x</i>
		PM-10 Emissions	54.72 Tons/Yr	<i>((365-WD)/365) x (1-CF))/2000</i>
TOTAL PM-10 EMISSIONS			60.2 Tons/Yr	

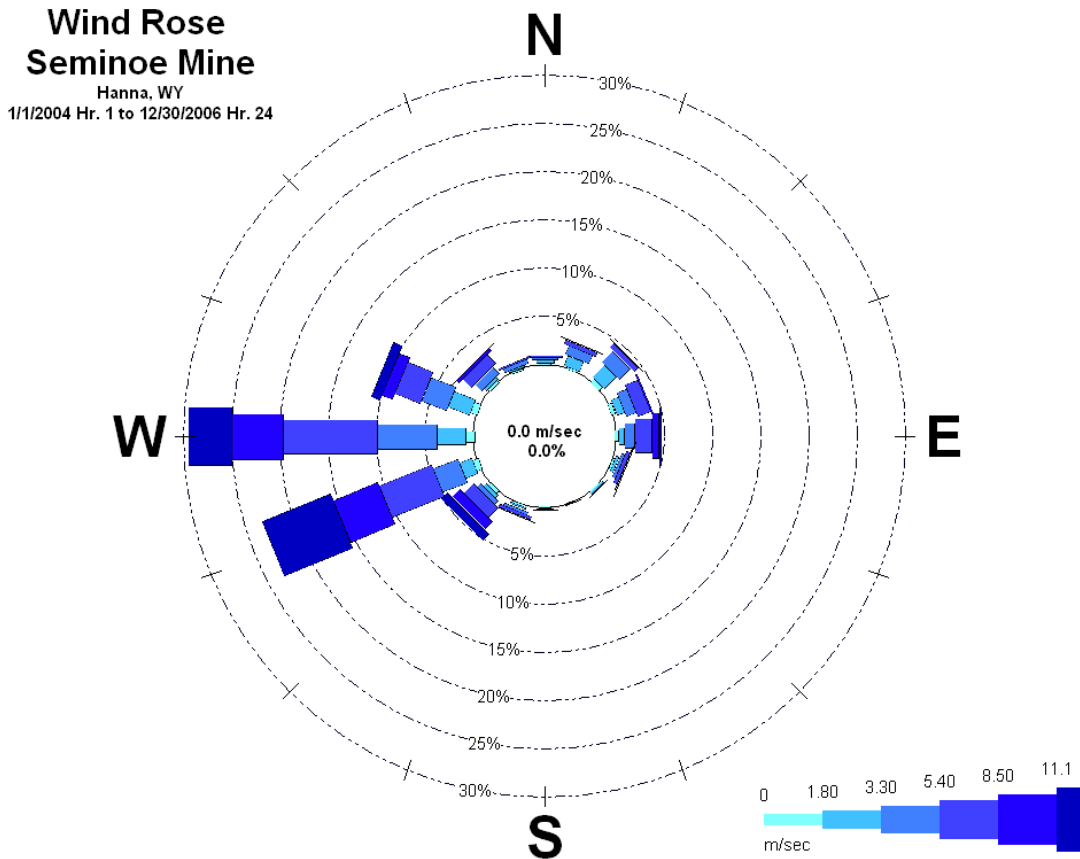
The difference in emissions between Options 1 and 2 is due entirely to the sheltering effect of locating the storage facility in the pit and shielding it with a spoil berm on the windward side. Average wind speed at ground level is assumed to be 6.7 meters per second, based on monitoring history at the nearby Seminole Mine. The assumption of a 25% reduction in average wind speed under Option 1 results in a PM₁₀ emissions reduction of 18.1 tons per year.

TABLE 3**BACT Option 2 (On-Surface Tube Stacker) PM-10 Emissions**

Emission Source	Type	Description	Control	Additional Information
Dozer Reclaim	Fugitive	Cat D11 Dozer	None	
		Emission Factor	8.0 Lb/Hr	<i>WDEQ 2002 Guidance</i>
		Total Throughput	3,200,000 Tons/Yr	<i>Total Coal Through Storage</i>
		Dozed Throughput	1,500,000 Tons/Yr	<i>Portion to Dead Storage</i>
		Dozer Productivity	750 Tons/Hr	<i>Estimate for 300,000 Ton Pile</i>
		Operating Hrs	2,000 Hrs	<i>Productivity/Throughput</i>
		TSP Emissions	8.00 Tons/Yr	$E=(EF \times Op \text{ Hrs})/2000$
		PM-10 Emissions	2.40 Tons/Yr	30% of TSP
Coal Stacker	Fugitive	Coal Dumping to Stockpile	Stacking Tubes	
		Emission Factor	0.017 Lb/Ton	<i>WDEQ Emission Factor</i>
		% Suspended	0.75	<i>WDEQ Emission Factor</i>
		Control Factor	50.00%	<i>Estimated</i>
		Material Dumped	3,200,000 Tons/Yr	<i>Total Coal Through Storage</i>
		TSP Emissions	10.20 Tons/Yr	$E=(EF \times \% \text{ sus} \times MD/2000) \times (1-CF)$
		PM-10 Emissions	3.06 Tons/Yr	30% of TSP
		Coal Reclaim	Fugitive	Vibratory & Pile Activator Feeder
Emission Factor	0.017 Lb/Ton			<i>WDEQ Emission Factor</i>
% Suspended	0.75			<i>WDEQ Emission Factor</i>
Control Factor	100.00%			<i>Estimated</i>
Material Reclaimed	3,200,000 Tons/Yr			<i>Total Coal Through Storage</i>
TSP Emissions	0.00 Tons/Yr			$E=(EF \times \% \text{ sus} \times MR/2000) \times (1-CF)$
PM-10 Emissions	0.00 Tons/Yr			30% of TSP
Coal Stockpile	Fugitive			Wind Erosion on Stockpiles
		Emission Factor	1.2 Lb/Acre/Hr	<i>WDEQ Emission Factor</i>
		Pile Size	11.0 Acres	<i>Calculated from Pile Size</i>
		Fraction Suspended	0.75	<i>WDEQ Emission Factor</i>
		Hours	8,760 Hours	<i>Total Annual</i>
		Ave. Wind Speed	6.70 meters/Sec	<i>Avg wind speed at surface</i>
		Wet Days	60	<i>Seminole Mine 5-Year Average</i>
		Control Factor	0.00%	
		TSP Emissions	242.77 Tons/Yr	$E=(EF \times AWS \times \% \text{ sus} \times PS \times ((365-WD)/365) \times (1-CF))/2000$
		PM-10 Emissions	72.83 Tons/Yr	
		TOTAL PM-10 EMISSIONS		78.3 Tons/Yr

The assumed reduction in wind speed is based on anticipated wind shielding from the pit walls and surrounding spoil pile, as shown on drawing no. 6945-L010. The spoil berm would extend in an "L" shape from the southwestern corner of the pit to the northeastern end of the pit. The top of the berm would be at 7,081 ft. elevation, with the pushed storage pile top at 7,060 ft. elevation. The prevailing winds in this area are from the west and the west-southwest, as typified by the most recent 3-year summary from the Seminole Mine (see Figure 1 below). The combination of berm, highwall and natural terrain would afford some shielding against wind originating anywhere between southwest and east-northeast. As implied by Figure 1, this constitutes the majority of the winds in this area.

FIGURE 1



In 2004, Inter-Mountain Laboratories conducted a study of the wind sheltering effect in an existing pit at the Bridger Coal Mine. This study was driven by a proposal to locate a 240,000-ton storage pile and stacking tube facility near the portal of an underground mine. The proposal was subsequently approved. The Bridger pit is oriented in an east-west direction, while prevailing winds are from the west-southwest. It is approximately 200 ft. from the pit floor to the top of the highwall on the north side of the pit. A spoil pile and access ramp border the south side of the pit.

In order to assess the degree of wind shelter provided by the Bridger pit, a wind monitor was placed in the pit near the probable storage site. For reference, a second wind monitor was placed at the top of the highwall several hundred feet northeast of the proposed storage site. After monitoring ten-minute average wind speeds at both these sites from 12/31/2003 to 2/06/2004, the data were collected and analyzed. During this period, wind speeds averaged as follows:

Highwall 10-minute averages ----- 10.0 mph
In-Pit 10-minute averages ----- 5.5 mph

Regular met station hourly averages ----- 9.5 mph

At the regular met station, three-year wind speeds (1/1/2002 through 12/31/2004) averaged 10.3 mph. Given this longer time period, the in-pit average wind speed was compared to the met station average (rather than the highwall average) over the 5-week interval. In making this comparison, a statistical analysis revealed less variability in wind speed ratios than wind speed differences. For these reasons, the ratio of in-pit average wind speed to met station average wind speed over the 5-week monitoring period was applied to the three-year average wind speed:

$$\frac{5.5}{9.5} * 10.3 = 5.96 \frac{\text{miles}}{\text{hour}} * \frac{88}{60} * 0.3048 = 2.66 \frac{\text{meters}}{\text{second}} \text{ (58\% of the 3-year surface average)}$$

Since the accepted PM₁₀ emissions factor for wind erosion is directly proportional to average wind speed, in-pit storage in the Bridger case would control roughly 40% of stockpile erosion emissions.

Additional research results were consulted to confirm the effect of wind shields. The University of Nebraska and U.S. Soil Conservation Service examined the influence of windbreaks on average wind speeds (University of Nebraska Extension EC 91-1763-B). Tests showed a 30% reduction in wind speed at a downwind distance of 10 times the height of a solid barrier.

An erosion study conducted by the U.S. Environmental Protection Agency led to an assumed 50% reduction in wind speed (and a 75% reduction in emissions due to nonlinear effects). The study utilized a 3-sided enclosure with 50% porosity (Sierra Research, 2003, *Final BACM Technological And Economic Feasibility Analysis*, report prepared for the San Joaquin Valley Unified Air Pollution Control District, March 21).

In relation to these other studies, a more conservative 25% reduction in wind speed was claimed for the Saddleback in-pit storage option. The Bridger pit is roughly twice as deep as the combination of pit and spoil berm at Saddleback (although the pit orientation relative to prevailing wind is quite similar). The University of Nebraska study oriented the wind barrier perpendicular to the wind direction, which would apply only to a portion of the winds at Saddleback. The EPA study used a 3-sided enclosure, whereas the Saddleback berm is configured more like a 2-sided enclosure.

SADDLEBACK HILLS MINE SURFACE FACILITY

Preliminary Cost Estimates

The following preliminary cost estimates, with an accuracy of $\pm 20\%$, are based on three active storage options that were considered:

- Option 1 reflects a 300,000 ton active storage pile with stacking tubes and live reclaim located in a sheltered area located between the high wall and an earthen berm.
- Option 2 reflects a 300,000 ton active storage pile with stacking tubes and live reclaim located in an open area that is un-sheltered from wind erosion.
- Option 3 reflects 300,000 ton totally enclosed slot storage with 100% live storage.

	<u>Option #1</u>	<u>Option #2</u>	<u>Option #3</u>
Ancillary Buildings			
Road and Ditches & Civil	\$30,746,100	\$30,742,800	\$30,654,000
Material Handling	\$8,554,700	\$5,096,400	\$5,030,400
Enclosed Slot Storage	\$45,399,200	\$46,360,800	\$43,701,600
Total	\$0	\$0	\$77,814,000
+20%	\$84,700,000	\$82,200,000	\$157,200,000
-20%	\$101,640,000	\$98,640,000	\$188,640,000
	\$67,760,000	\$65,760,000	\$125,760,000



ENGINEERS & CONTRACTORS

ROBERT & SCHAEFER
Company

DEQ 000078-000273

Appendix G
Mercury Removal Costs

COSTS FOR MERCURY REMOVAL SYSTEM

Client	DKRW
Service	Mercury Guard Beds
Equipment ID	R-2801 A/B
Capacity, MMscfd (each vessel)	304.00
Flow Rate, Nm ³ /hr (each vessel)	334,927
Hg Inlet Concentration, µg/Nm ³	91.22
Hg Outlet Concentration, µg/Nm ³	0.02
Hg Mass Removed, µg/Nm ³	91.20
Hg Removal Efficiency, %	99.98
Hg Mass Removed, lb/hr (each vessel)	0.067
Hg Mass Removed, ton/yr (each vessel)	0.295
Hg Mass Removed, ton/yr (both vessels)	0.590

Total Capital Cost		Cost in Estimated
Itemized Expenditures		
CAPITAL COSTS:		
Carbon Adsorbent Cost		\$ 135,000
Equipment Installed Cost		\$ 1,000,000
Total Installed Cost (TIC)		\$ 1,135,000
OPERATING COSTS:		
Catalyst Replacement (every 10 years)		\$ 13,500
Annual Operating Costs		\$ 13,500
TOTAL ANNUAL COSTS:		
Capital Recovery Factor (9.1%, 20 yr life)		
Annualized Total Capital Investment	0.1103	x TIC \$ 125,223
Total Annual Costs,\$/yr		\$ 138,723
HG REMOVAL:		
Hg Removed, ton/yr	0.590	
Cost of Hg Removed, \$/ton		\$ 235,164

All costs are based on a mercury guard bed design provided by SME Associates.

Hg-A**VAPOR PHASE MERCURY FLITRATION**

Prepared for SNC

PROJECT: DKRW Energy CGTL

ITEM: Hg Capture

SME Associates

13231 Champion Forest Dr, Suite 201

Houston, Tx. 77069

Phone (281)440-7350

Fax (281)440-7353

This design was prepared 13-Jun-06

Call Daren Scott if questions arise

DESIGN CONDITIONS

FLOWRATE:	304	MMSCFD/VESSEL
FLOWRATE:	694,414	#/FT3
FLOWRATE:	334,927.4	NM3/HR
MOLECULAR WEIGHT:	20.805	
OPERATING PRESSURE:	945	PSIA
DESIGN TEMPERATURE:	120	oF
H2O SATURATION TEMP:	120	oF
H2O RELATIVE HUMIDITY:	100	%
COMPRESSIBILITY USED:	0.987	Z
DENSITY:	3.20	#/FT3
VISCOSITY:	0.017	cp
INLET Hg CONTENT:	97	PPB(WT)
" " " :	10.06	PPB(VOL)
" " " :	91,223	NANOGRAMS /NM3 (ng/Nm3)
" " " :	91.22	MICROGRAMS/NM3 (ug/Nm3)
" " " :	0.0912	MILLIGRAMS/NM3 (mg/Nm3)
" " " :	2.412	GRAMS/MMSCF
" " " :	1.617	#/DAY HG
OUTLET Hg CONTENT:	<0.02	MICROGRAMS/NM3 (ug/Nm3)

DESIGN VARIABLES

MAXIMUM SUPERFICIAL VELOCITY:	50	FPM
MINIMUM CONTACT TIME:	15	SEC.
EXTRUDATE SIZE:	4	mm
LOADING USED:	20	

SELECTION

VESSEL ID; USED:	9.5	FT.
SUPERFICIAL VELOCITY:	51.0	FPM
ADSORBENT BED HEIGHT:	12.70	FT.
L/D:	1.34	
CONTACT TIME:	14.95	SEC.
NUMBER OF DRUMS:	180	DRUMS
NUMBER OF PALLETS:	45	PALLETS
AMOUNT OF ADSORBENT:	30600	LBS.
VESSEL HEIGHT USED:	16	FT.
EST LIFE OF ADSORBENT:	10.4	YRS.
HgA BED PD:	8.9	PSI.
FLOW DIRECTION:	DOWN	FLOW
PIPE SIZE; USED:	14	IN.
CERAMIC SUPPORT BALLS:	6	IN.(RECOMMENDED)
CERAMIC HOLD-DOWN BALLS:	6	IN.(RECOMMENDED)

THIS DESIGN PLAN HAS BEEN PREPARED IN ACCORDANCE WITH GUIDELINES PROVIDED BY SME ASSOCIATES REFLECTING ITS PAST EXPERIENCE AND LABORATORY TESTING OF THIS PRODUCT. PLEASE DO NOT DEVIATE FROM THIS DESIGN PLAN WITHOUT CONSULTING US FIRST. NO SPECIFIC WARRANTY, EXCEPT FITNESS FOR PURPOSE, IS OFFERED. THIS DESIGN IS NOT A LICENSE TO USE PATENTS OWNED BY OTHERS.

DEQ 000078 000276



Robert Moss <rross@dkrwaf.com>

11/12/2007 10:30 AM

To "Susan_Bassett@URSCorp.com"
<Susan_Bassett@URSCorp.com>

cc

bcc

Subject FW: Mercury Removal from Syngas

Susan,

Attached is the vendor sheet (different than the one you sent this morning). Also, note below that there are two carbon beds and no third bed.

Bob Moss
Development Engineer
DKRW Advanced Fuels
713-425-6533 (O)
713-670-4544 (M)
rross@dkrwaf.com
www.dkrwaf.com
www.dkrwenergy.com

This e-mail is the property of DKRW Energy LLC and/or its relevant affiliate and may contain confidential and privileged material for the sole use of the intended recipient (s). Any review, use, distribution or disclosure by others is strictly prohibited. If you are not the intended recipient (or authorized to receive for the recipient), please contact the sender or reply to DKRW Energy LLC at info@DKRWenergy.com and delete all copies of the message. This e-mail (and any attachments hereto) are not intended to be an offer (or an acceptance) and do not create or evidence a binding and enforceable contract between DKRW Energy LLC (or any of its affiliates) and the intended recipient or any other party, and may not be relied on by anyone as the basis of a contract by estoppel or otherwise. Thank you.

From: Bonnell, Leo [<mailto:Leo.Bonnell@snc-lavalin-gds.com>]
Sent: Monday, November 12, 2007 10:38 AM
To: Robert Moss
Cc: Ray Birch
Subject: FW: Mercury Removal from Syngas

Robert,

Attached is the vendor data sheet for the Mercury Guard Beds that was used for the Feasibility Study.

I had forgotten, but with the long 10 year bed life claimed by the vendor, for the F.S. we decided not to put a spare guard bed in. So we would have 2 X 50% capacity beds with the total carbon adsorbent cost of \$135,000.

SNC estimated the purchased costs of the two guard beds to be \$400,000 for both. The "all-in" installed cost estimates were not broken down by item, but based on the data we developed they should be about 2.5 X the purchased costs, or \$ 1 million TIC for the two beds (excluding adsorbent).

Hope this will be helpful.

Regards,
Leo Bonnell
Process Director
SNC-Lavalin Houston

DEQ 000078-000277

Tel. 713-295-4815
leo.bonnell@snclavalin-gds.com

From: Daren Scott [mailto:dscott@sme-llc.com]
Sent: Tuesday, June 13, 2006 3:33 PM
To: Bonnell, Leo
Cc: Birch, Ray; Daren Scott
Subject: RE: Mercury Removal from Syngas
Leo,

Attached is a quick design which would require approx 60,000 lbs of HgA at \$2.25/lb. The lead time would be 16-20 weeks.

I divided the flow into 2 to bring the vessel size to a reasonable value and even at this you have 2- 10' dia vessels. The other option would be to use a single 14' dia vessel.

Most of the required data is on the data sheet but FYI this would give you a 10 year life on the carbon, the maximum temperature is 180F and we have no problems with any of the gas components.

Sincerely;

Daren Scott

SME Associates, LLC
Ph: 281-440-7350
Fx: 281-440-7353
Cell: 832-257-6281
dscott@sme-llc.com

From: Bonnell, Leo [mailto:Leo.Bonnell@snclavalin-gds.com]
Sent: Tuesday, June 13, 2006 1:09 PM
To: Daren Scott
Cc: Birch, Ray
Subject: Mercury Removal from Syngas

To: Daren Scott, SME Associates Inc.

Daren,

As I mentioned today, SNC is doing a feasibility study, and later FEED package, for a coal-to-liquids project in Wyoming for DKRW Energy (www.dkrwenergy.com). The syngas contains mercury from the coal that must be removed prior to desulfurizing and syngas conversion.

Can you give us a budget quote for a mercury removal adsorbent bed for this applicati

Flow and composition of the feed syngas:

Temp = 120 deg F
Pressure = 945 psia
Total Flow (lbmoles/hr) = 66,600
Composition (mole %, dry)
CO = 38.0

DEQ 000078-000278

H2 = 40.0
CO2 = 20.0
CH4 = 0.1
N2 = 1.75
H2S = 0.15
C2+ = nil

Water = saturated
NH3 = 100 ppm
Mercury = 10 ppb by volume

Note that the Hg level is based on the highest of several local coal samples.
The long-term average is likely to be less.

Thanks for your help.

Regards,
Leo Bonnell
Process Engineering Consultant
SNC-Lavalin GDS, Inc.
9009 West Loop South, Houston, TX 77096
Office: 713-295-4815 Fax: 713-667-9241



Mercury Guard Bed Design.pdf

Appendix H
Incremental NO_x Removal Cost for SCR

INCREMENTAL NO_x REMOVAL COST FOR SCR

NO_x Removal Cost to 6 ppm (76% Removal)

Catalyst Cost ¹	650,000 USD
Catalyst Life ¹	3 years
Power Usage	160 KW
Cost of Power ¹	0.07 \$/kW-hr
Hours per year	8760 hr/yr
<u>Interest Rate</u>	7.00 %
Ammonia Usage ¹	46.20 gal/hr
Cost of Ammonia ¹	0.50 USD/gal
Uncontrolled NO _x	316.08 ton/yr
NO _x Emissions	75.86 ton/yr
Tons NO _x Removed	240.22 ton/yr

Costs¹

Ammonia	202,295 USD/yr
Vaporizer Power	98,015 USD/yr
Catalyst	240,890 USD/yr
Total	541,200 USD/yr

(Based on normal operations, prior to SCR, fuel gas mixture)

NO_x Removal Cost to 4 ppm (84% Removal)

Catalyst Cost ¹	750,000 USD
Catalyst Life ¹	3 years
Power Usage	173 KW
Cost of Power ¹	0.07 \$/kW-hr
Hours per year	8760 hr/yr
<u>Interest Rate</u>	7.00 %
Ammonia Usage ¹	50.00 gal/hr
Cost of Ammonia ¹	0.50 USD/gal
Uncontrolled NO _x	316.08 ton/yr
NO _x Emissions	50.57 ton/yr
Tons NO _x Removed	265.51 ton/yr

Costs¹

Ammonia	219,152 USD/yr
Vaporizer Power	106,183 USD/yr
Catalyst	277,950 USD/yr
Total	603,285 USD/yr

(Based on normal operations, prior to SCR, fuel gas mixture)

Incremental Cost to Reduce NO_x From 6 ppm to 4 ppm

Annual Cost for Achieving 4 ppm	603,285 USD/yr
Annual Cost for Achieving 6 ppm	541,200 USD/yr
Incremental Cost	62,085 USD/yr

NO _x Removed When Achieving 4 ppm	265.51 tons/yr
NO _x Removed When Achieving 6 ppm	240.22 tons/yr
Incremental NO_x Removal	25.29 tons/yr

Incremental Cost 2,455 USD/ton

¹ Information provided by Paul Rood, Process Engineer at SNC Lavalin, on November 16, 2007.

Appendix I
Analysis of Criteria Pollutant Far Field Modeling Sufficiency

Appendix I

Analysis of Criteria Pollutant Far Field Modeling Sufficiency

1.1 INTRODUCTION

Medicine Bow Fuel & Power LLC (MBFP) believes that the far field criteria pollutant modeling performed for the June 19, 2007 permit application remains sufficient for the revised permit application. The following pollutant-specific discussions compare modeled emission rates to emissions rates included in this revised application.

Emissions from the industrial gasification and liquefaction plant (the Plant) have been revised due to a number of process and equipment changes. Emission unit changes are summarized in Table I-1. The combustion turbines are the largest emitters of nitrogen oxides (NO_x), carbon monoxide (CO), and sulfur dioxide (SO₂) during normal operations. The turbines are also the largest point source emitters of particulate matter with a diameter of less than 10 microns (PM₁₀). Combustion turbine stack parameters are not expected to change significantly. Consequently, prior far field modeling of turbine emissions should be adequate.

With regard to other emission sources, many units do not change. However, the Sulfur Recovery Unit (SRU) incinerator has been removed from the process. Furthermore, many process heaters have been deleted while a few new process heaters have been added.

Table I-1 – Emission Unit Changes

Description	Identification	Size
<i>Equipment with no Capacity Changes</i>		
Combustion Turbine 1	CT-1	66 MW
Combustion Turbine 2	CT-2	66 MW
Combustion Turbine 3	CT-3	66 MW
Black Start Generator 1 ¹	Gen-1	2889 hp
Black Start Generator 2 ¹	Gen-2	2889 hp
Black Start Generator 3 ¹	Gen-3	2889 hp
Firewater Pump Engine ¹	FW-Pump	575 hp
CO ₂ Vent Stack ¹	CO ₂ VS	N/A
High Pressure Flare	FL-1	0.2 MMBtu/hr (for pilot)
<i>Added Equipment</i>		
Auxiliary Boiler ²	AB	66.0 MMBtu/hr
Catalyst Regenerator ^{1, 3}	B-1	21.5 MMBtu/hr
Reactivation Heater ¹	B-2	12.5 MMBtu/hr
HGT Reactor Charge Heater ¹	B-3	2.2 MMBtu/hr
Low Pressure Flare	FL-2	0.2 MMBtu/hr (for pilot)

Table I-1 – Emission Unit Changes

Description	Identification	Size
<i>Removed Equipment</i>		
Fractionation Feed Heater	H-5401	87 MMBtu/hr
Catalytic Dewaxing Charge Unit	H-5301	3.9 MMBtu/hr
Unicracker Feed Heater	H-5201	16.3 MMBtu/hr
Unicracker Intermediate Heater	H-5202	44.2 MMBtu/hr
Unionfiner Feed Heater	H-5101	5.1 MMBtu/hr
Unionfiner Intermediate Heater	H-5102	6.4 MMBtu/hr
Sulfur Recovery Unit Incinerator	H-3102	11.2 MMBtu/hr
<i>Modified Equipment</i>		
Gasifier Preheater 1 ^{1,4}	GP-1	21 MMBtu/hr
Gasifier Preheater 2 ^{1,4}	GP-2	21 MMBtu/hr
Gasifier Preheater 3 ^{1,4}	GP-3	21 MMBtu/hr
Gasifier Preheater 4 ^{1,4}	GP-4	21 MMBtu/hr
Gasifier Preheater 5 ^{1,4}	GP-5	21 MMBtu/hr
<p>1. This equipment operates less than 8,760 hr/yr. However, in some cases, potential emissions are calculated based on 8,760 hr/yr in order to simplify compliance.</p> <p>2. The auxiliary boiler usually operates on standby at 25% load to prevent freeze ups if there is a Plant shutdown. The equivalent continuous heat input rate would be approximately 21 MMBtu/hr.</p> <p>3. The catalyst regenerator operates only during catalyst regeneration; the average equivalent continuous rate will be approximately 9 MMBtu/hr.</p> <p>4. Gasifier preheater heat input capacity was increased from 15 MMBtu/hr to 21 MMBtu/hr for each preheater.</p>		

Table I-2 summarizes proposed maximum emission rates within this revised application and compares them to modeled emission rates. Emission rates are given in terms of grams per second (g/sec) for easy comparison to modeled rates. Emission rates do not include the following malfunctions: emergency venting to the High Pressure or Low Pressure Flares and CO₂ venting during the first plant startup and as a result of malfunctions thereafter.

Table I-2 – Revised Emissions Compared to CALPUFF Modeled Emissions

Pollutant	Revised Plant-Wide Emission Rate (tpy)	Revised Plant-Wide Annual Emission Rate (g/sec)	Modeled Plant-Wide Annual Emission Rate (g/sec)	Emission Rate Increase From Modeled Rates (g/sec)
NO _x	251.63 ¹	7.24	7.28	-0.04
SO ₂	32.65 ¹	0.94	0.81	0.13
PM/PM ₁₀	194.93 ²	5.61	8.96	-3.35

- Does not include emergency venting to the High Pressure Flare or startup, shutdown, or malfunction (SSM) venting to the Low Pressure Flare.

1.2 FAR FIELD MODELING

Far field modeling was performed in 2006 using CALPUFF to predict air quality impacts relating to visibility and nitrogen and sulfur deposition. The modeled pollutants that contribute to these air quality impacts are NO_x, SO₂, and PM₁₀. Plant-wide gram per second emissions of NO_x, and PM₁₀ decreased. However, SO₂ emissions increased slightly.

1.2.1 NO_x Modeling

As shown in Table I-2, maximum Plant-wide NO_x emission rates are approximately 0.04 g/sec less than the emission rates used for CALPUFF modeling. The largest NO_x emitters at the Plant continue to be the three combustion turbines. These turbines account for more than 90 percent of total annual emissions during normal operations.

Since there is a decrease in emissions and equipment changes will occur in largely the same areas as the modeled emission sources, MBFP believes that no additional NO_x modeling is necessary.

1.2.2 SO₂ Modeling

Removal of the Sulfur Recovery Unit (SRU) incinerator deleted the largest single source of normal operation SO₂ emissions from the original process. However, this reduction in SO₂ emissions has been largely offset by increases in SO₂ emissions from the three combustion turbines. The combustion turbine emission increases derive in part from firing more natural gas, which has a greater sulfur concentration than the syngas that was originally expected to be fired in the turbines. In addition, the SO₂ emission factor for natural gas firing that was used in the emission calculations submitted with the original June 19, 2007 permit application was too low.

As shown in Table I-2, modeled Plant SO₂ emissions are slightly less than revised emission estimates, with modeled emissions of 0.81 g/sec, compared to revised emissions of 0.94 g/sec. Based on previous CALPUFF modeling, SO₂ emissions result in low deposition and are a minor component of visibility impacts.

1.2.3 PM/PM₁₀ Modeling

While coal storage PM₁₀ emissions have not changed (because coal usage has not changed), PM₁₀ emissions from combustion sources have decreased substantially. Removal of the SRU incinerator accounts for a large share of the PM₁₀ emission decrease. The modeled emission rate for Plant point sources was 8.96 g/sec compared to revised estimated emissions of 5.61 g/sec (including Plant point and area source emissions from coal storage).

Appendix J
Responses to WDEQ July 17, 2007 Far Field Modeling Comments

Responses to WDEQ July 17, 2007 Far Field Modeling Comments

1. CALMET Files on DVD

Comment. An examination of the terrain and landuse output files shows that both include blocks of missing data (see figure below showing terrain for the modeling domain). The applicant should obtain complete data for the domain, revise the MAKEGEO portion of the CALMET processing and submit the revised input/output files to the Division. [graphic has been deleted]

Response. The files are included within the MAKEGEO file folder.

2. Section 7: Far-Field Air Quality Impact Analysis

Comment. The letter from the Division dated March 5, 2007 provided comments on the CALPUFF protocol, including item B.6 which requested an analysis of the final CALMET wind field: *"At a minimum, the analysis should include an examination of the wind flows for selected times and vertical layers. The flows produced by CALMET should be compared to observed flows as seen in archived weather maps and/or compared to expected flows (e.g., downslope winds during stable conditions at night). Other parameters such as precipitation can also be compared to observed conditions."* No analysis was provided with the application.

Response. After running CALMET, the resulting data fields were analyzed using the PRTMET utility to illustrate the assimilated wind and temperature fields within the domain for quality assurance purposes. PRTMET enables the user to extract meteorological data fields such as wind speed and direction, temperature, and mixing height on an hourly "snapshot" or average basis.

Part of the quality assurance process determined whether wind patterns were influenced by terrain; this is a good indication of whether meteorological data is properly located relative to the terrain. Figure 1 shows area contours, with pink shaded areas representing high terrain. PRTMET quality assurance graphics are included in Figures 2 through 9 for an approximate 10 km grid to demonstrate that the selection of CALMET control options resulted in a reasonable simulation of the meteorology within the domain. Particularly good instances of terrain influenced flow can be seen in Figure 2 (March 19, 2003 – hour 3) at the following locations:

East -220, North -200
East -220, North -20
East 150, North 150
East 75, North 0

Another good example of terrain influenced flow can be seen in Figure 6 (June 19, 2003 – hour 3) at the following locations:

- East -275, North 75
- East 50, North -125
- East 75, North 0
- East -275, North -25

The time for one of the hourly wind field vector snapshots was chosen based on the worst visibility impairment day from CALPUFF modeling. The largest extinction change occurred at the Savage Run sensitive Class II area on March 19, 2003. Meteorological conditions on March 19, 2003 were unusual due to a major winter storm. Appendix N includes “Mesoscale Model Simulations in Quasi-Forecast Mode of the Great Western Storm of 16-20 March 2003.” This document summarizes meteorological conditions during that time. The document is also available on the CD-ROM as “Meso_Model_Great_Storm_2004.pdf.”

Since March 19th conditions represent winds flowing toward Class I areas in Colorado, the other snapshot was chosen based on the worst visibility impairment day for Class I areas in Wyoming such as the Bridger Wilderness area and the Fitzpatrick Wilderness area. The largest extinction change in both Class I areas in Wyoming occurred on June 19, 2003.

These snapshot days also represent one day for summer (June 19, 2003) and one day for winter (March 19, 2003). Two hours on each day were plotted: 0300 Mountain Standard Time (MST) and 1500 MST. Furthermore, for each time period, a surface wind field, corresponding to Level 1, and an upper air wind field, corresponding to Level 8, was plotted. Plots developed in this study are shown in Table 1. These wind fields appeared to accurately capture terrain, slope, and seasonal effects expected within the modeling domain, and demonstrated generally smooth translations and continuous Mesoscale flow. These characteristics validated the spatial behavior of the meteorological data set throughout the modeling domain.

Table 1 - List of Wind Vector Plots

Date	March 19, 2003	June 19, 2003
Hour	3,15	3,15
Vertical layer	1,8	1,8

Windroses from the CALMET model output and the surface observation station data sets indicated general agreement in wind directions, frequencies, and speeds. Windroses for March 2003 from several surface observation stations such as Aspen, Laramie General Brees Field (Laramie), Craig-Moffat stations were plotted and are shown in Figures 11 through 13. The locations of the selected stations are shown in the Figure 10. The list of windroses developed in this study is included in Table 2. Windrose plots from surface observation stations and the CALMET-predicted output are shown in Figures 11 through 13 and indicate good agreement between surface observations and CALMET predicted output.

Table 2 - List of Windroses (March 1 –March 31, 2003)

Station Name	Data Period (Total Count) March 1 – March 31, 2003		Location of the Station	
	Observation	CALMET-Predicted	Observation (Latitude, Longitude)	CALMET-Predicted (Grid Cell)
Aspen	672 hours	743 hours	39.217N, 106.867W	93, 12
Laramie	715 hours	743 hours	41.313N, 105.674W	118, 71
Craig-Moffat	684 hours	743 hours	40.5N, 107.533W	79, 48

Figure 1 - 3-D Terrain Elevation Contours

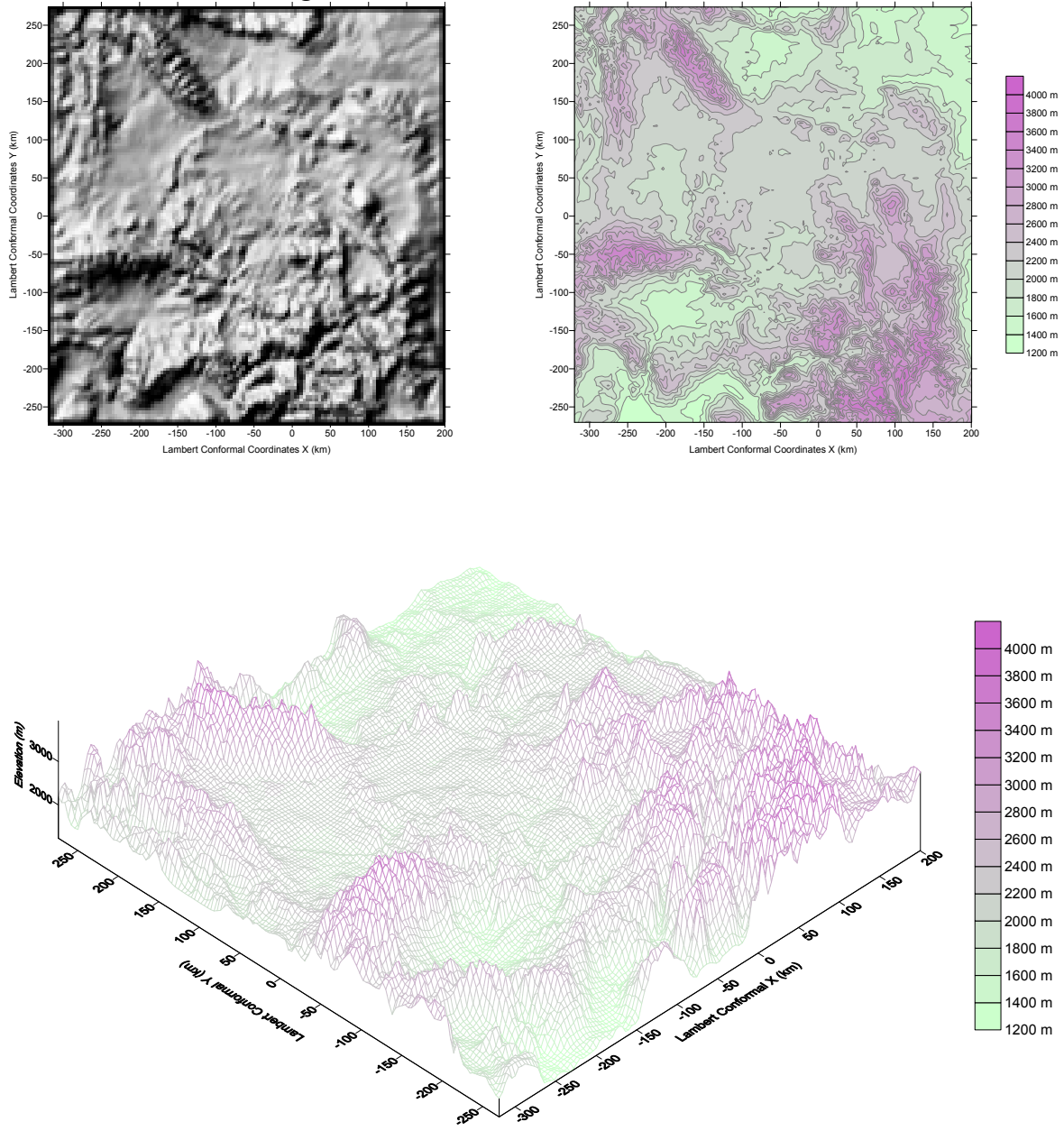


Figure 2 - Surface Air Windfield March 19, 2003, Hour 3, Layer 1

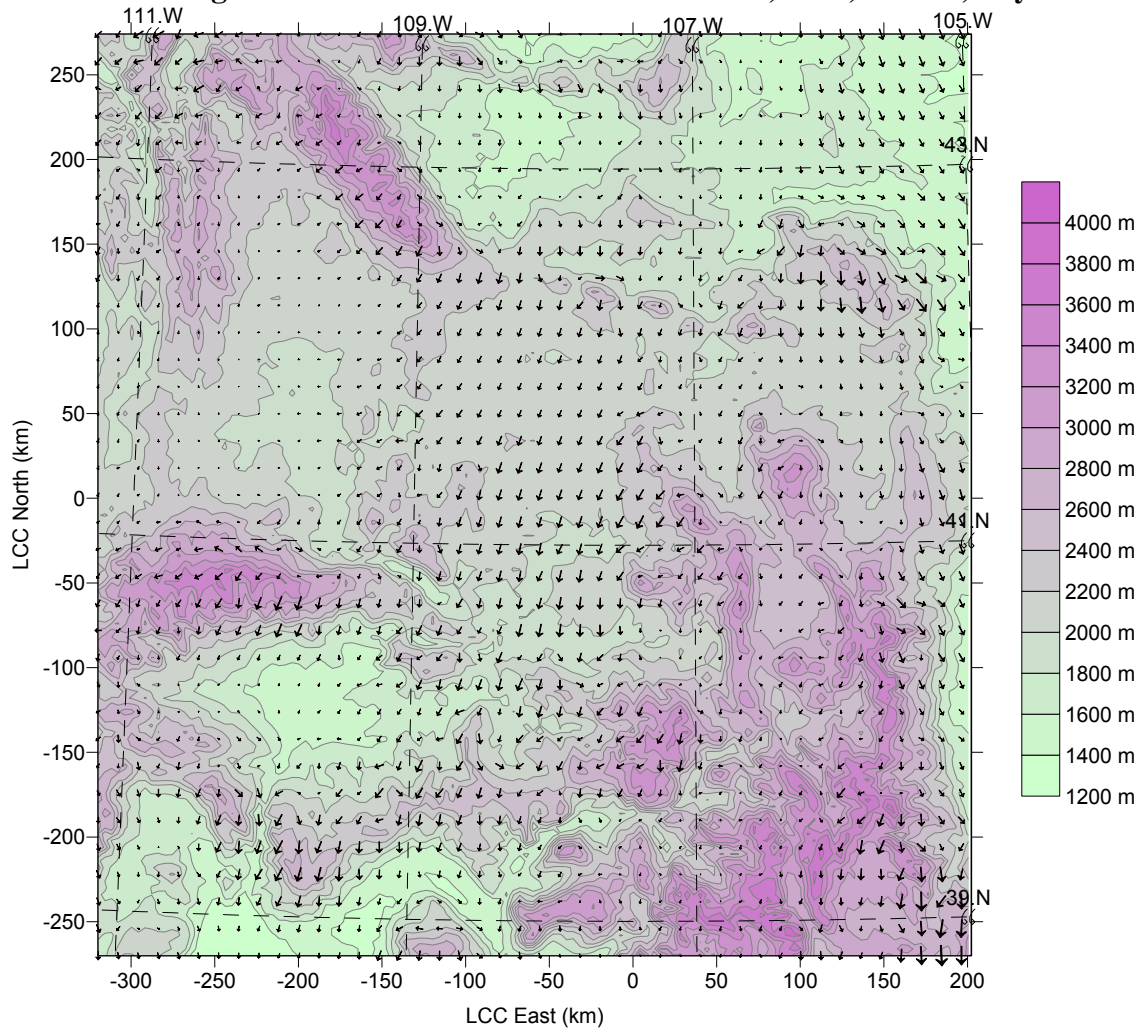


Figure 3 - Surface Air Windfield March 19, 2003, Hour 15, Layer 1

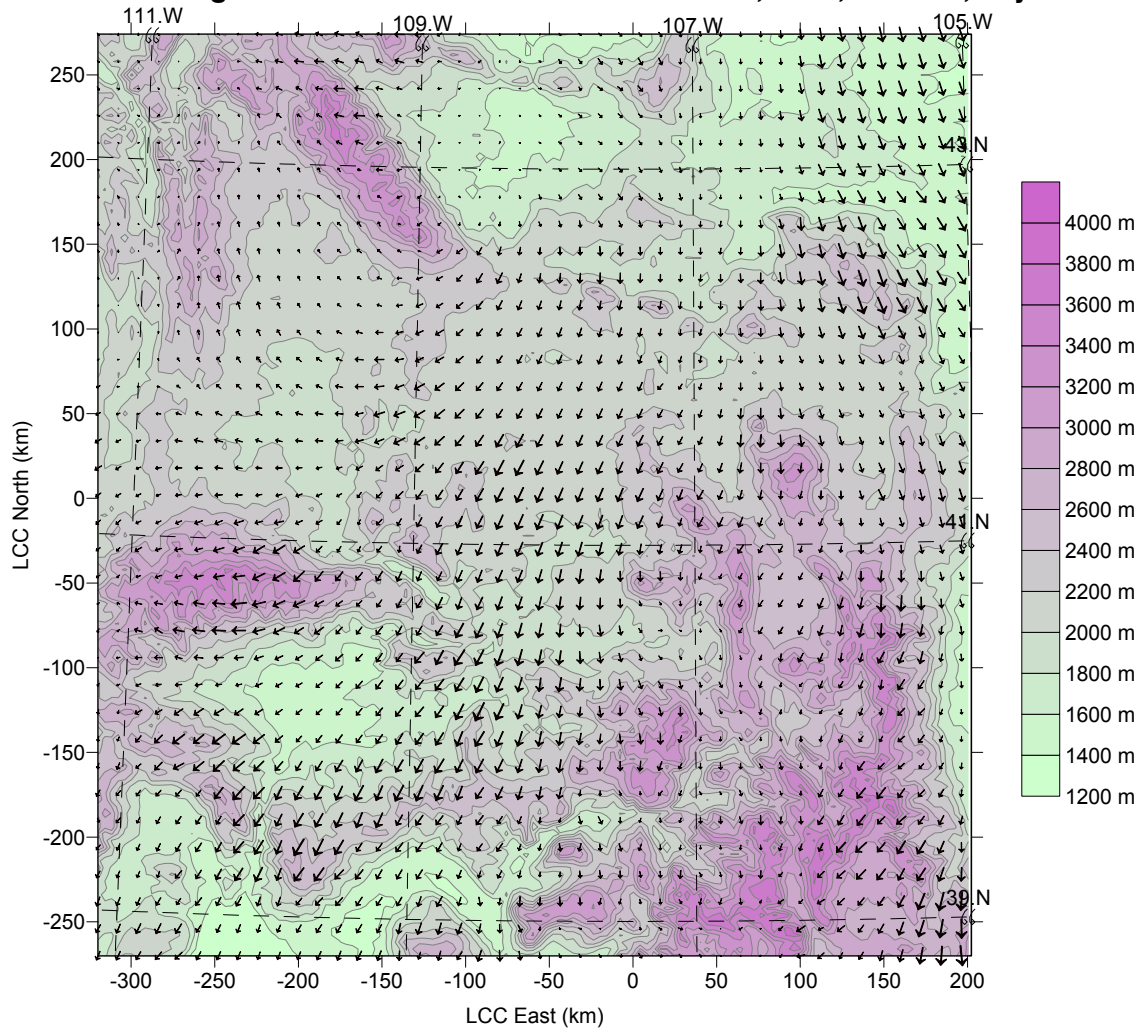


Figure 4 - Upper Air Windfield March 19, 2003, Hour 3, Layer 8

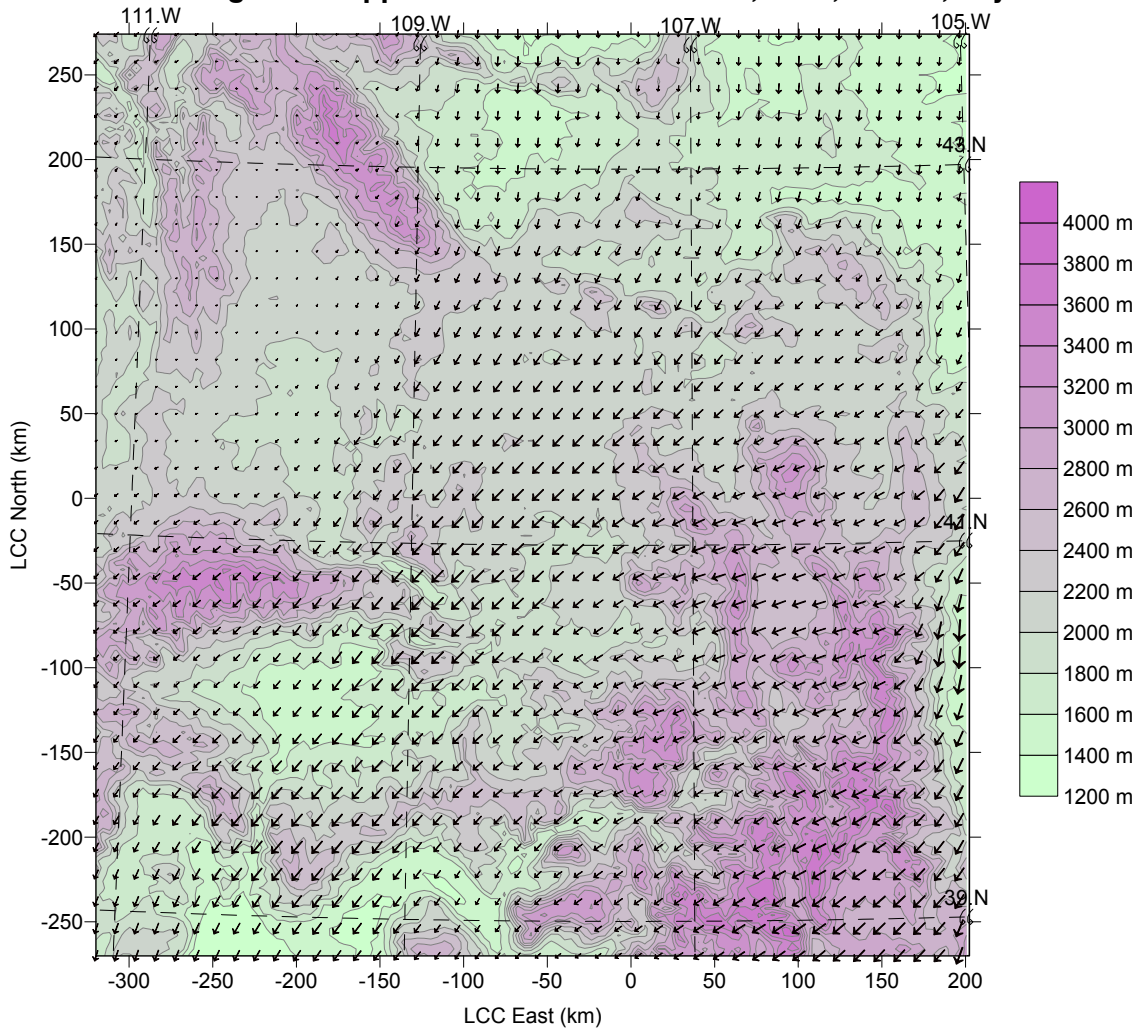


Figure 5 - Upper Air Windfield March 19, 2003, Hour 15, Layer 8

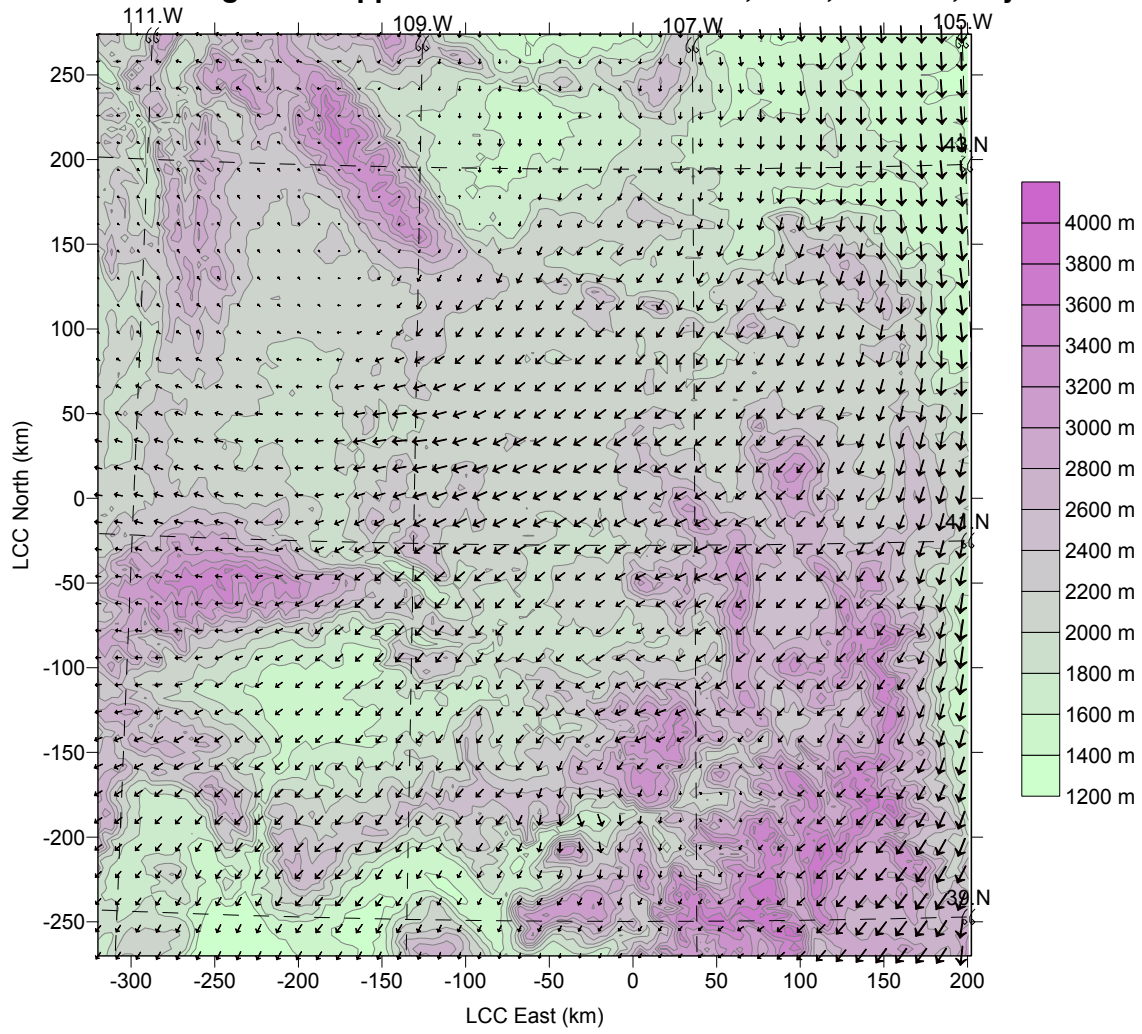


Figure 6 - Surface Air Windfield June 19, 2003, Hour 3, Layer 1

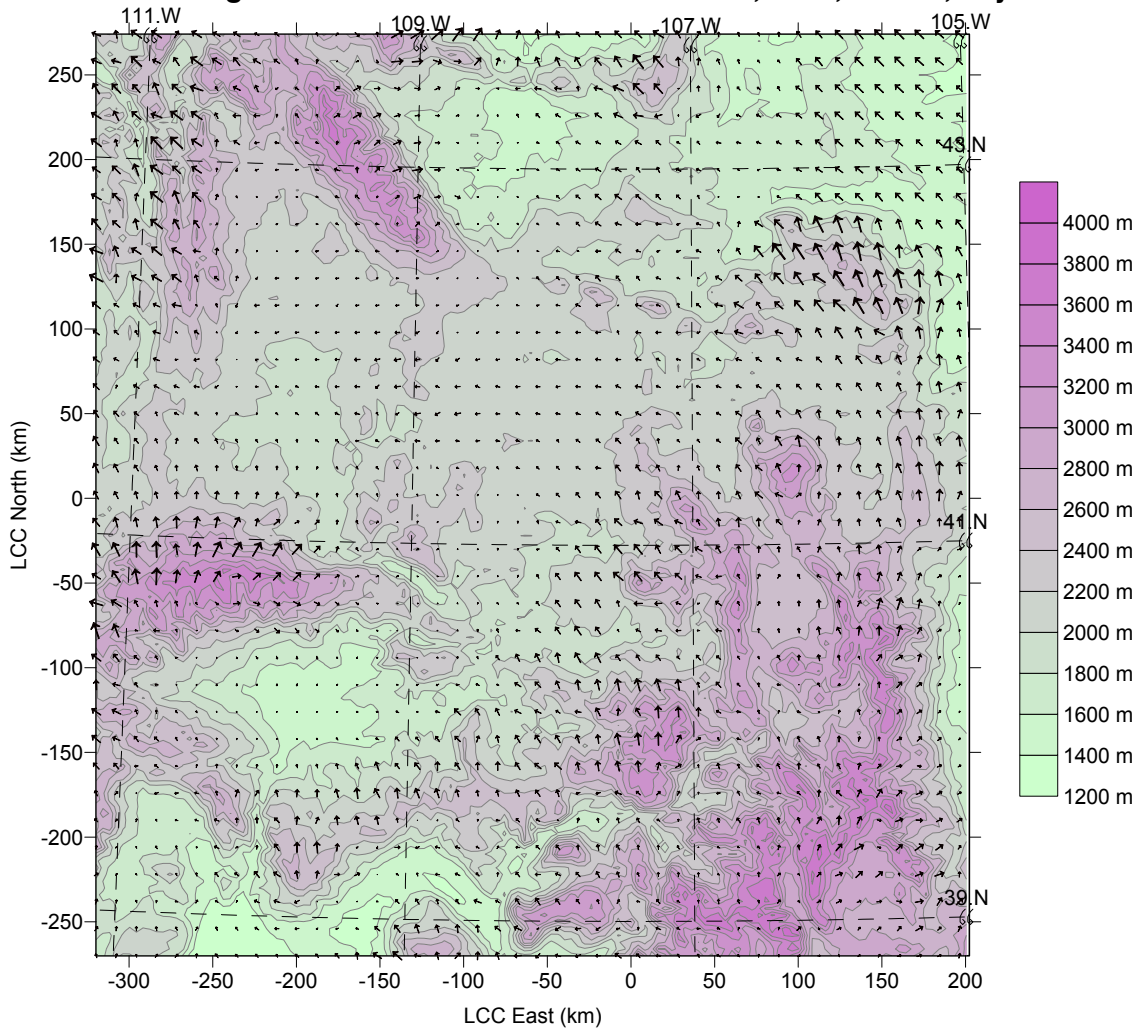


Figure 7 - Surface Air Windfield June 19, 2003, Hour 15, Layer 1

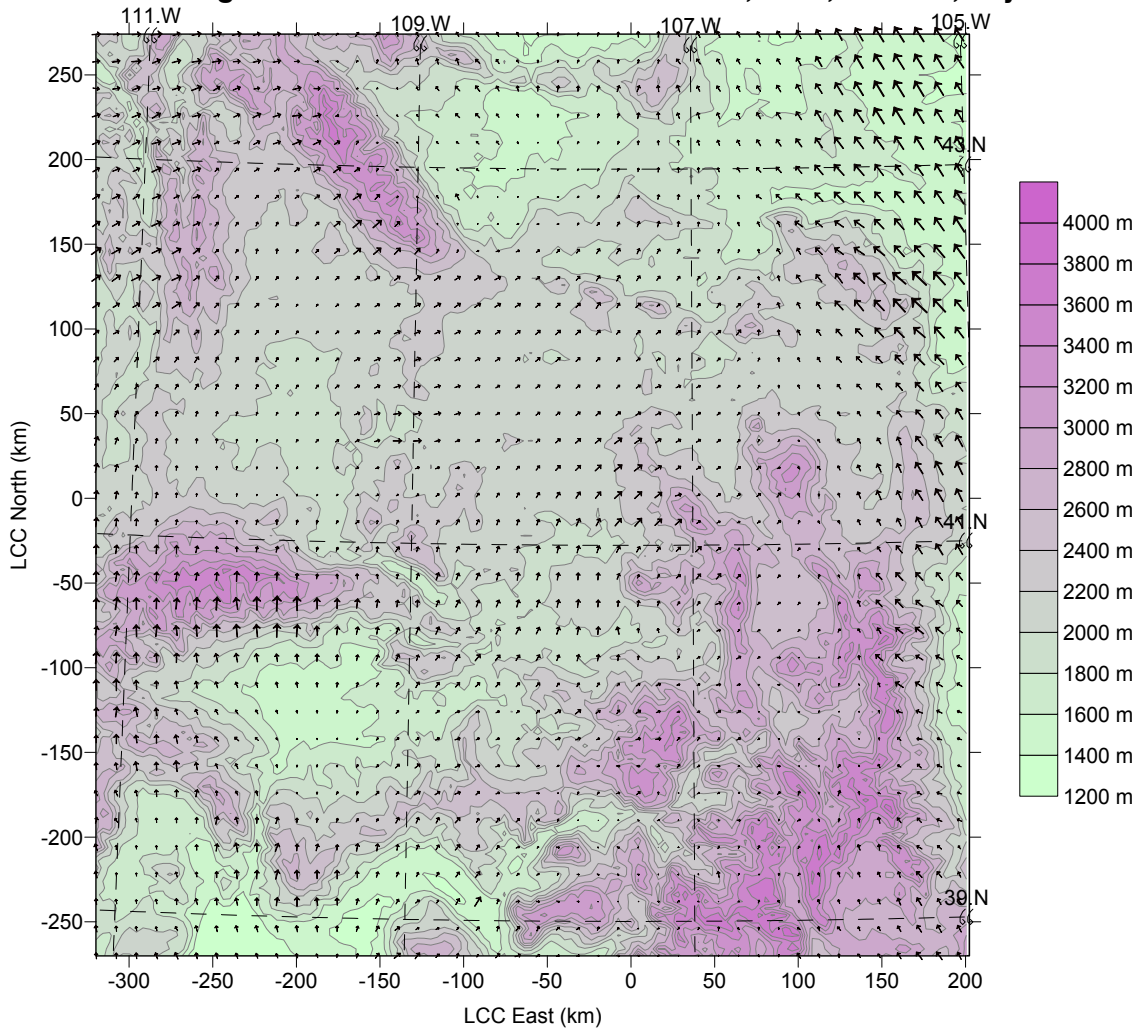


Figure 8 - Upper Air Windfield June 19, 2003, Hour 3, Layer 8

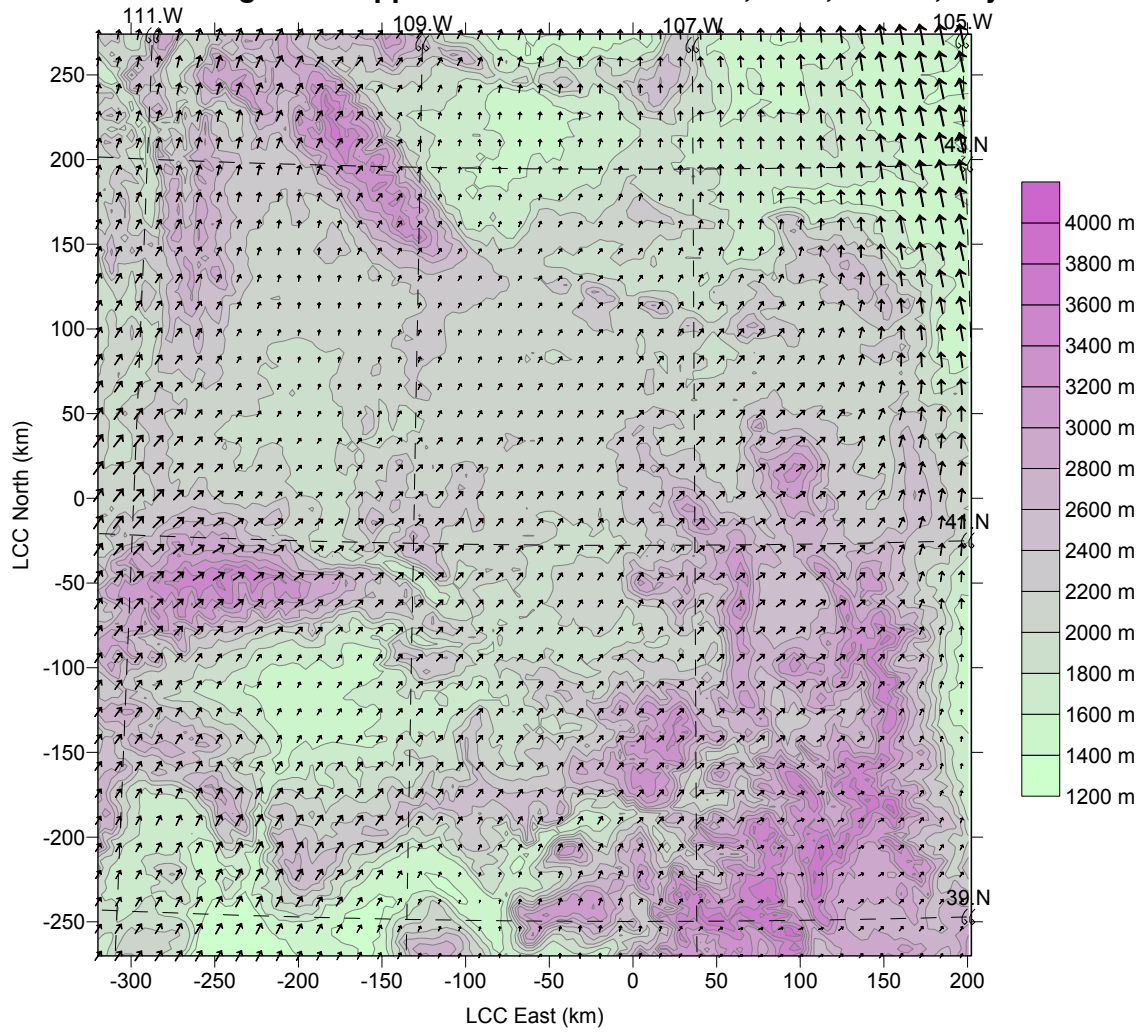


Figure 9 - Upper Air Windfield June 19, 2003, Hour 15, Layer 8

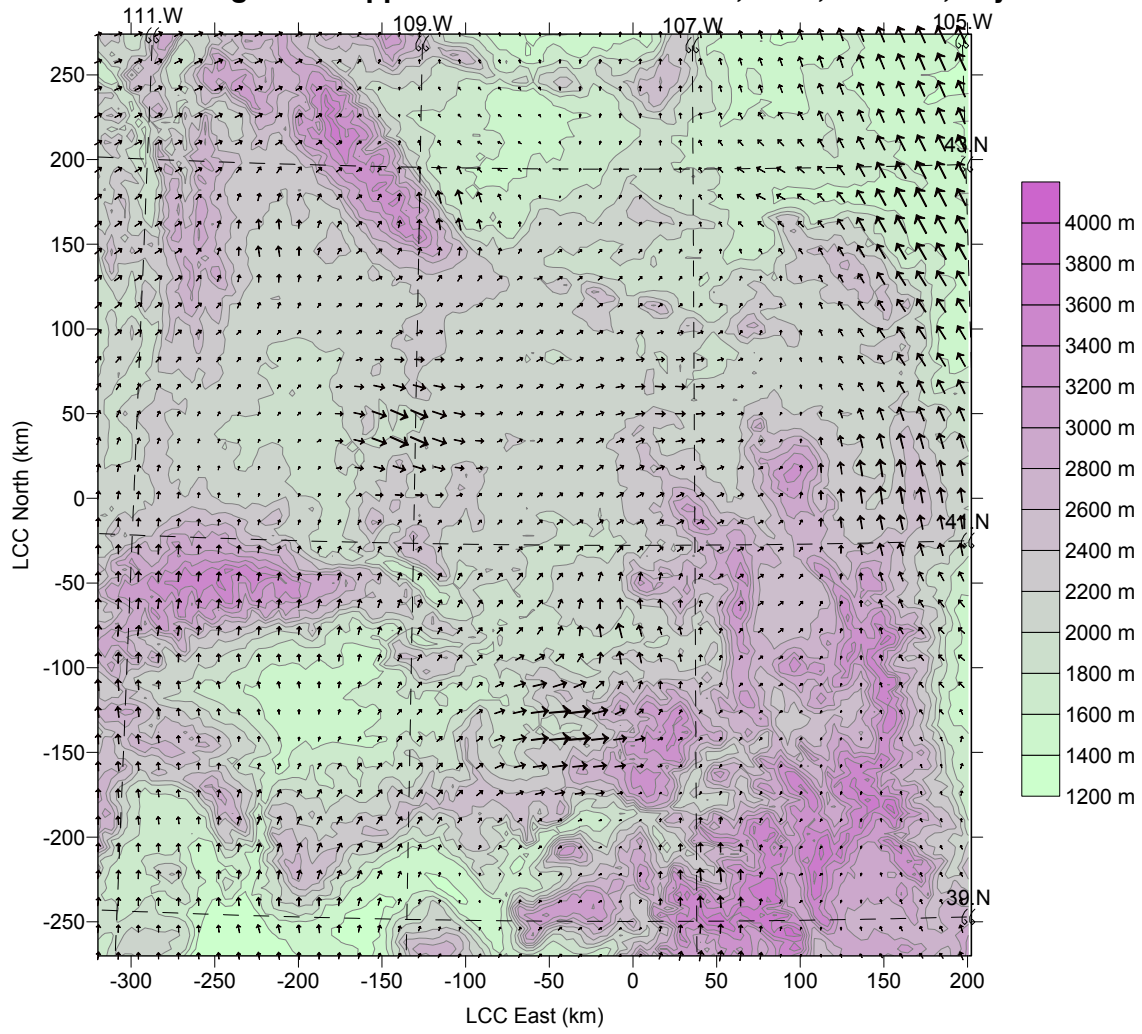


Figure 10- Location of Selected Surface Observation Stations

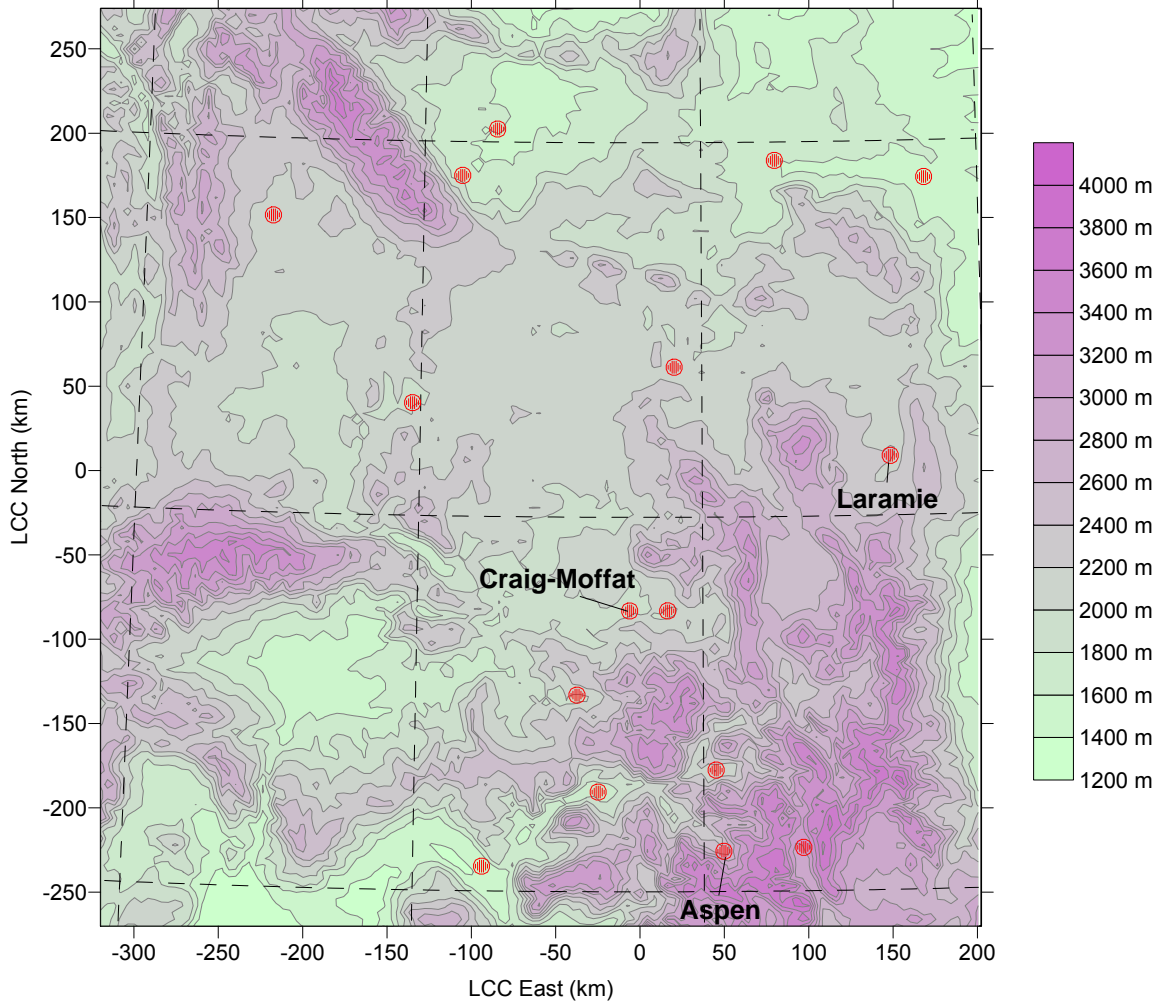
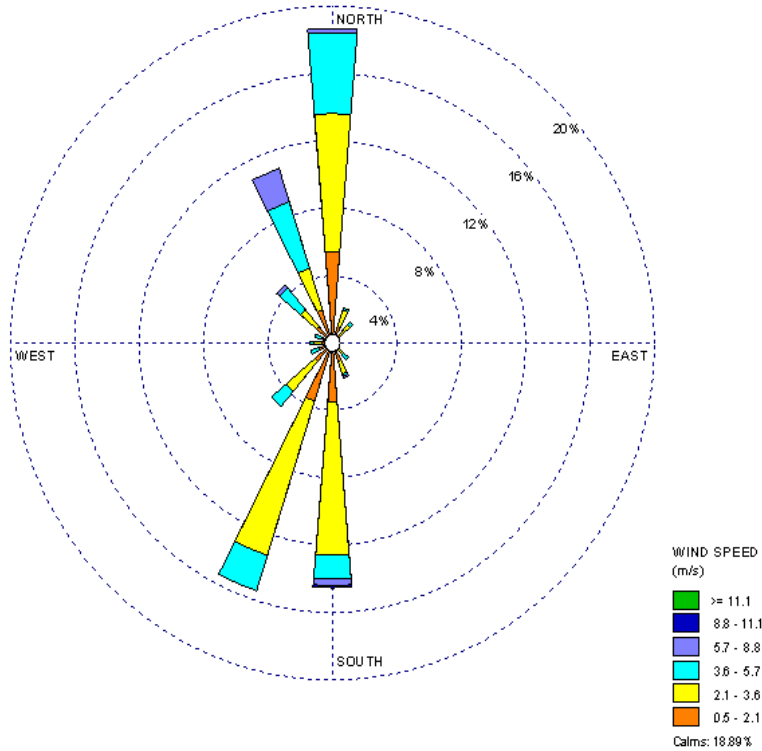


Figure 11 - Aspen field Windroses (March, 2003)

Aspen Field Observation Station Windrose -672 hours



Aspen Field CALMET-predicted Windrose (grid cell:93, 12)-743 hours

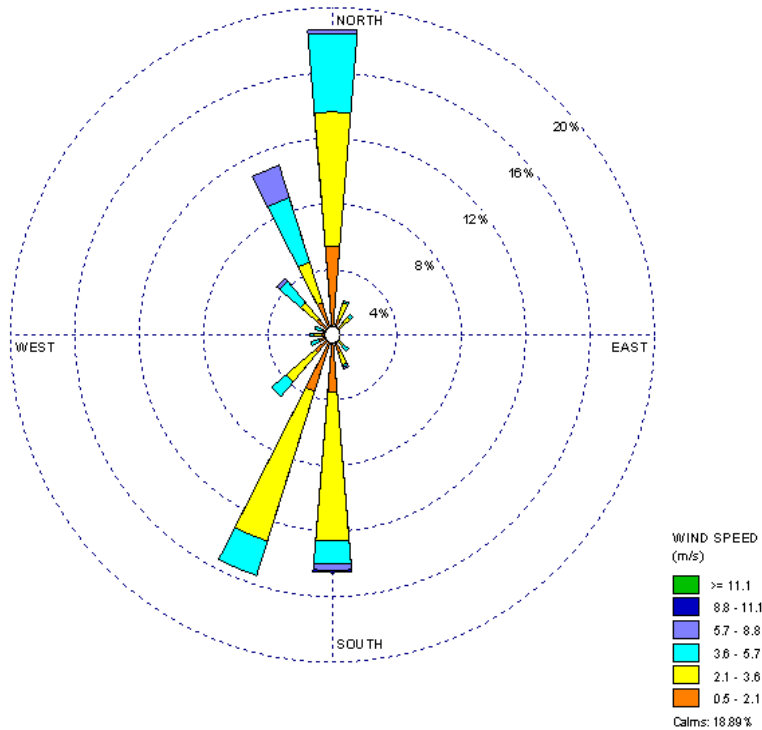
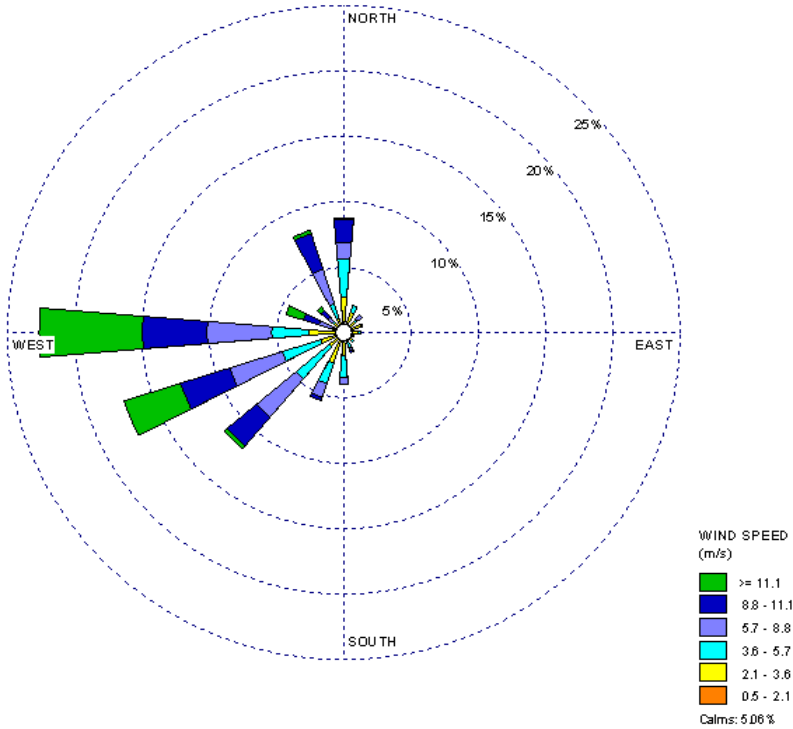


Figure 12 - Laramie field Windroses (March, 2003)

Laramie Field Observation Station Windrose-715 hours



Laramie Field CALMET-predicted Windrose (grid cell:118, 71)-743 hours

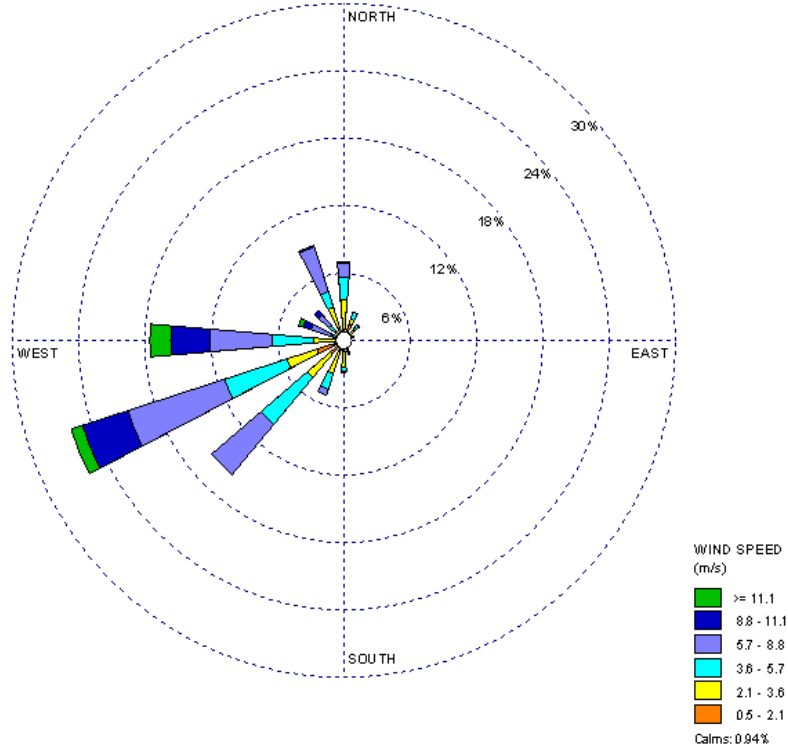
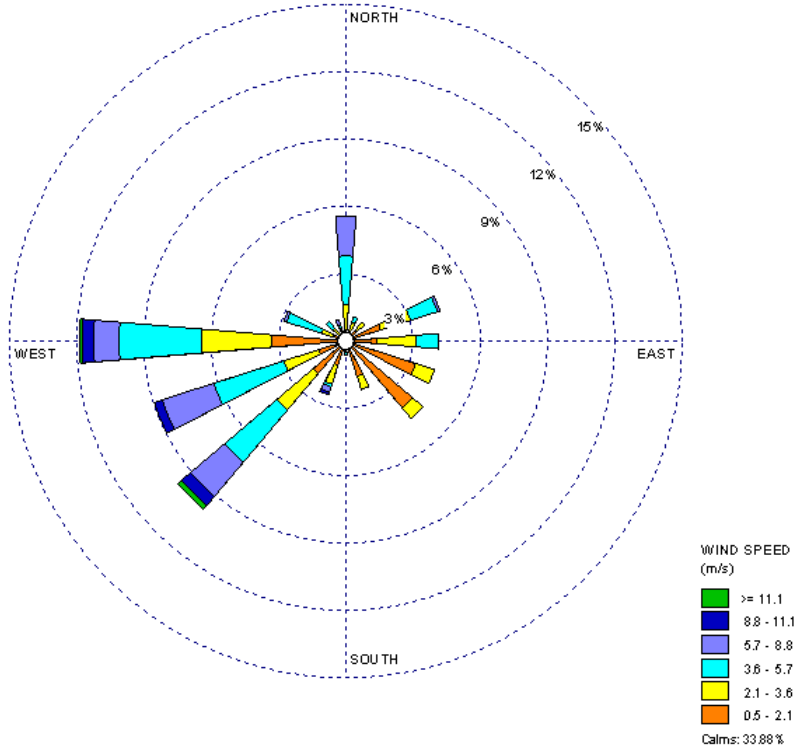
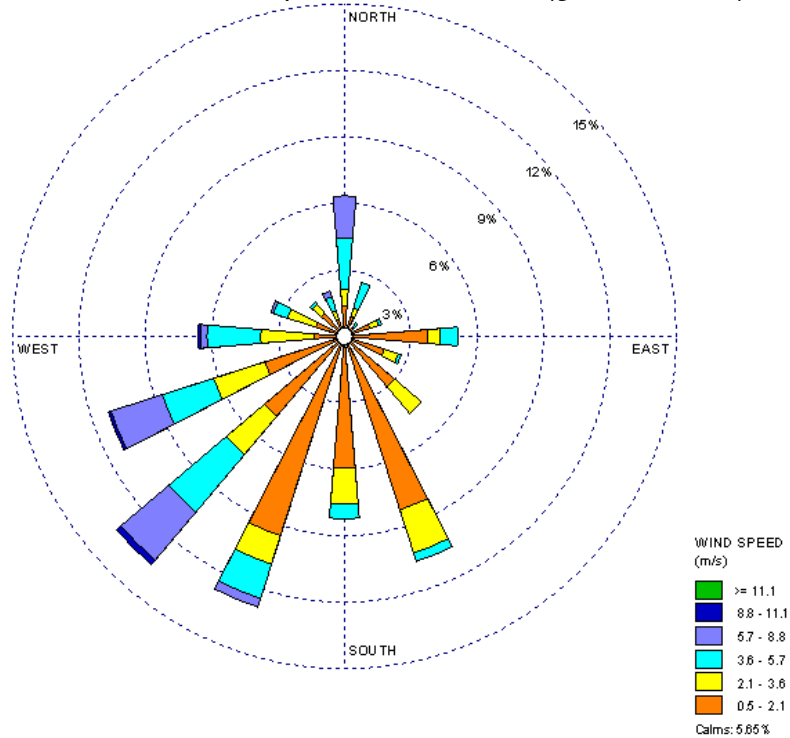


Figure 13 - Laramie field Windroses (March, 2003)

Craig-Moffat Field Observation Station Windrose-684 hours



Craig-Moffat Field CALMET-predicted Windrose (grid cell:79,48)-743 hours



Appendix K
NRCS Irrigated and Nonirrigated Yields by Map Unit
for Carbon County, Wyoming

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
4: Canburn	4w	4w	---	---	---	---	---	---
8: Gerrard	6w	6w	---	1.00	---	1.00	---	---
9: Grieves variant	4w	3w	---	---	---	---	---	---
9H: Grieves variant, alkali	4w	4w	---	---	---	---	---	---
13: Rhoamett	6e	4s	---	3.50	---	3.50	---	---
15A: Poposhia	4e	3e	---	3.00	---	3.00	---	4.0
15B: Poposhia	4e	3e	---	3.00	---	3.00	---	4.0
18A: Alcova	4e	3e	---	---	---	---	---	---
18B: Alcova	4e	3e	---	---	---	---	---	---
18C: Alcova	4e	4e	---	---	---	2.50	---	5.0
18H: Alcova, saline	6e	4e	---	---	---	3.50	---	7.0
20: Debone	7s	4s	---	---	---	---	---	---
22: Edlin	4e	3e	---	3.50	---	3.50	---	5.0
29: Canburn variant	4w	4w	---	---	---	---	---	---
31A: Tisworth	6s	4s	---	---	---	---	---	---

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
31B: Tisworth	6s	4s	---	---	---	---	---	---
34: Tresano	6e	6e	---	---	---	---	---	---
38A: Rock River	4e	3e	---	---	---	3.00	---	4.5
38B: Rock River	4e	3e	---	---	---	3.00	---	4.5
38C: Rock River	4e	4e	---	---	---	2.00	---	3.5
38H: Rock River, saline	6s	4s	---	---	---	3.00	---	4.5
40: Fiveoh	4e	6e	---	---	---	---	---	---
40H: Fiveoh, saline	6e	4s	---	---	---	---	---	---
43B: Grieves	4e	3e	---	3.00	---	2.00	---	5.0
45: Yetull variant	4e	6e	---	---	---	---	---	---
49: Firth variant	4w	3w	2.00	4.00	1.00	3.00	---	---
51W: Patent variant	4w	4w	---	---	---	---	---	---
52: Laney	6s	4s	---	2.00	---	---	---	---
52S: Laney variant Slickspots	6s 8	4s 8	---	---	---	---	---	---
53A: Pinelli	4e	3e	---	---	---	---	---	---

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
59: Absher variant	6s	4s	---	---	---	---	---	---
69: Kiltabar	4s	4s	---	---	---	---	---	---
78A: Ryan Park	4e	3e	---	---	---	3.00	---	4.0
78B: Ryan Park	4e	3e	---	---	---	3.00	---	4.0
79D: Blackhall	7e	7e	---	---	---	---	---	---
86: Ansel	6e	6e	---	---	---	---	---	---
101: Echemoor Clayburn	6e 6e	4e 4e	---	---	---	---	---	---
102: Echemoor Inchau	6e 7e	4e 6e	---	---	---	---	---	---
105: Starman Barrett	7e 7e	7e 7e	---	---	---	---	---	---
107: Starman Vabem	7e 7e	7e 7e	---	---	---	---	---	---
108: Lymanson Youga	6e 6e	6e 4e	---	---	---	---	---	---
109: Lymanson Roxal	6e 7e	4e 6e	---	---	---	---	---	---
111: Vabem Inchau	7e 6e	7e 6e	---	---	---	---	---	---

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
118A:			---	---	---	3.50	---	7.0
Alcova	4e	3e						
Rock River	4e	3e						
118B:			---	---	---	3.00	---	6.0
Alcova	4e	3e						
Rock River	4e	3e						
128:			---	---	---	---	---	---
McFadden	6e	4e						
Brownsto	6e	4e						
Blackhall	7e	7e						
135B:			---	3.50	---	3.50	---	7.0
Cushool	6e	4e						
138A:			---	---	---	---	---	---
Rawlins	4e	3e						
Bosler	4e	3e						
138B:			---	---	---	---	---	---
Rawlins	4e	3e						
Bosler	4e	3e						
140:			---	---	---	---	---	---
Tisworth	6s	4s						
Poposhia	4e	3e						
141:			---	4.00	---	4.00	---	7.0
McFadden	6e	4e						
Brownsto	6s	4s						
144:			---	3.00	---	3.00	---	5.0
McFadden	6e	4e						
Blackhall	7e	6e						
147:			---	---	---	---	---	---
Rogert	7s	6e						
Quander	7s	6e						
Rock outcrop	8	8						
200:			---	---	---	---	---	---
Patent variant	4w	4w						
Hagga	4w	4w						

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
208:			---	---	---	---	---	---
Pinelli	6e	4e						
Forelle	6e	4e						
209:			---	---	---	---	---	---
Chaperton	6e	4e						
Boettcher	6e	4e						
210:			---	---	---	---	---	---
Absher variant	6s	4s						
217:			---	---	---	---	---	---
Dahlquist	6s	4s						
Cragosen	7s	6s						
218A:			---	---	---	3.50	---	7.0
Alcova	6e	3e						
Rawlins	6e	3e						
218B:			---	---	---	3.00	---	6.0
Alcova	6e	3e						
Rawlins	6e	3e						
221:			---	---	---	---	---	---
Blazon	7e	6s						
Chaperton	6e	4s						
224A:			---	4.00	---	4.00	---	7.0
McFadden	6e	3e						
Brownsto	6s	4s						
224B:			---	3.50	---	3.50	---	6.0
McFadden	6e	3e						
Brownsto	6s	4s						
224w:			---	4.00	---	4.00	---	7.0
McFadden, wet	6w	3w						
Brownsto, wet	6w	4w						
225:			---	3.00	---	3.00	---	7.0
Cushool	6e	4s						
Rock River	4e	3e						

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
229:			---	3.50	---	3.50	---	7.0
Cushool	6e	4s						
Cushool variant	6e	4s						
235:			---	---	---	---	---	---
Blazon	7e	6e						
Blazon, THIN SOLUM	7e	6e						
236:			---	---	---	---	---	---
Cushool	6e	6e						
Worfman	7e	6e						
Blackhall	7e	6e						
237:			---	---	---	---	---	---
Seaverson	7e	6e						
Blazon	7e	6e						
244:			---	---	---	3.00	---	4.5
Rock River	4e	3e						
251:			---	3.00	---	---	---	7.0
Grieves	6e	4e						
Blackhall	7e	6e						
252:			---	---	---	---	---	---
Blazon	7e	6e						
Blazon, thin solum	7e	7e						
Rentsac	7s	7e						
253:			---	---	---	---	---	---
Blazon	7e	6e						
Cushool	6e	4e						
254:			---	---	---	---	---	---
Abston	6s	4s						
Seaverson	7s	6s						
255:			---	---	---	---	---	---
Ponded soils	8	8						
256:			---	3.50	---	3.50	---	6.0
McFadden	6e	4e						
Brownsto	6s	6e						
Rawlins	6e	4e						

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
257:			---	---	---	---	---	---
Havre variant	4e	3s						
Glendive variant	4e	3s						
258:			---	---	---	3.00	---	4.5
Rock River	4e	3e						
Cushool	4e	4e						
260:			---	---	---	2.00	---	3.0
Ryan Park	6e	6e						
Rock River	4e	4e						
261:			---	5.00	---	3.00	---	---
Luhon	6e	4e						
Rock River	4e	3e						
262:			---	---	---	---	---	---
Rentsac	7s	7s						
Thermopolis	7e	6e						
263:			---	3.50	---	3.50	---	5.0
Edlin	4e	4e						
Carmody	6e	4e						
264:			---	---	---	---	---	---
Rentsac	7s	7e						
Rock outcrop	8	---						
272:			---	---	---	---	---	---
Rawlins	6e	4e						
275:			---	3.00	---	3.00	---	4.0
Poposhia	4e	3e						
Chaperton	6e	4e						
278:			---	---	---	3.00	---	4.0
Ryan Park	4e	3e						
Elk Mountain	6e	4e						
279:			---	---	---	---	---	---
Blackhall	7e	7e						
Grieves	4e	4e						

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
280:			---	---	---	---	---	---
Hazton variant	7e	7e						
Baggott variant	7e	7e						
282:			---	---	---	---	---	---
Tisworth	6s	6s						
284:			---	---	---	---	---	---
Blackhall	7e	7e						
Carmody	6e	4e						
Rock outcrop	8	8						
286:			---	---	---	---	---	---
Tisworth	6s	6s						
296:			---	---	---	---	---	---
Pinelli	6e	4e						
Boettcher	6e	4e						
332:			---	---	---	---	---	---
Chaperton, dry	6e	4e						
Hatermus	7e	6e						
Haterton	7e	6e						
333:			---	4.00	---	3.50	---	7.0
Sagecreek, alkali	6s	6s						
Sagecreek	4e	4e						
334:			---	3.50	---	3.00	---	6.0
Sagecreek, alkali	6s	6s						
336:			---	---	---	---	---	---
Haterton, thin solum	7e	7e						
Hatermus	7e	7e						
Haterton	7e	7e						
380:			---	---	---	---	---	---
Hazton variant	7e	6e						
Burgess	6e	4e						
400:			2.00	4.00	1.00	3.00	---	---
Firth variant	4w	3w						
Canburn variant	4w	3w						

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
483: Sandbranch	6s	6s	---	---	---	---	---	---
495: Chaperton, dry Sagecreek	6e 4e	4e 4e	---	---	---	---	---	---
502: Hagga, saline, alkali	4w	4w	---	---	---	3.00	---	---
703: Havre	4e	4e	---	---	---	---	---	---
761: Glendive variant	4e	4e	---	---	---	---	---	---
911: Forelle Diamondville	4e 4e	4e 4e	---	2.50	---	2.50	---	5.0
912: Evanston	4e	3e	---	3.00	---	3.00	---	6.0
928: Grieves variant Gerrard	4w 6w	4w 4w	---	---	---	---	---	---
931: Forelle	4e	3e	---	3.00	---	3.00	---	6.0
1202: Delplain variant Morling	7e 7e	7e 6e	---	---	---	---	---	---
1209: Zillman Peyton variant	6s 6e	6e 6e	---	---	---	---	---	---
1217: Zillman variant Highpoint	6s 7e	6e 6e	---	---	---	---	---	---

Irrigated and Nonirrigated Yields by Map Unit

Carbon County Area, Wyoming

Map symbol and soil name	Land capability		Alfalfa hay		Grass hay		Pasture	
	N	I	N	I	N	I	N	I
			Tons	Tons	Tons	Tons	AUM	AUM
1251:			---	---	---	---	---	---
McFadden	6e	4e						
Blackhall	7e	7e						
Edlin	4e	4e						
1252:			---	---	---	---	---	---
Rentsac	7e	7e						
Blazon	7e	7e						
Rubble land	8	8						
1255:			---	---	---	---	---	---
Blackhall	7e	7e						
Rentsac	7e	7e						
1256:			---	---	---	---	---	---
Rawlins	4e	4e						
Rock River	4e	4e						
1260:			---	3.00	---	3.00	---	5.0
McFadden	6e	4e						
Edlin	6e	4e						
1912:			---	---	---	---	---	---
Peyton variant	4e	4e						
Evanston variant	4e	4e						
2080:			---	---	---	---	---	---
Pinelli variant	4e	3e						
Forelle	6e	3e						
2199:			---	---	---	---	---	---
Anchutz	4e	3e						
9120:			---	---	---	---	---	---
Evanston variant	4e	3e						
Evanston	4e	3e						
W:			---	---	---	---	---	---
Water	---	---						

Appendix L
NRCS Acreage and Proportionate Extent of the Soils
for Carbon County, Wyoming

Acreage and Proportionate Extent of the Soils

Carbon County Area, Wyoming

Map symbol	Map unit name	Acres	Percent
4	Canburn loam, 0 to 2 percent slopes	397	*
8	Gerrard loam, 0 to 2 percent slopes	992	*
9	Grieves variant fine sandy loam, 0 to 3 percent slopes	3,759	*
9H	Grieves variant fine sandy loam, alkali, 0 to 3 percent slopes	1,283	*
13	Rhoamett silty clay, 0 to 2 percent slopes	388	*
15A	Poposhia loam, 0 to 2 percent slopes	20	*
15B	Poposhia loam, 2 to 6 percent slopes	320	*
18A	Alcova sandy loam, 0 to 2 percent slopes	1,000	*
18B	Alcova sandy loam, 2 to 6 percent slopes	1,000	*
18C	Alcova sandy loam, 6 to 12 percent slopes	1,000	*
18H	Alcova sandy loam, saline, 0 to 3 percent slopes	1,000	*
20	Debone silty clay loam, 0 to 3 percent slopes	1,300	*
22	Edlin sandy loam, 0 to 10 percent slopes	1,356	*
29	Canburn variant fine sandy loam, 0 to 2 percent slopes	1,046	*
31A	Tisworth sandy loam, 0 to 2 percent slopes	836	*
31B	Tisworth sandy loam, 2 to 6 percent slopes	332	*
34	Tresano sandy loam, 0 to 20 percent slopes	3,320	*
38A	Rock River sandy loam, 0 to 2 percent slopes	5,115	0.1
38B	Rock River sandy loam, 2 to 6 percent slopes	4,206	*
38C	Rock River sandy loam, 6 to 12 percent slopes	418	*
38H	Rock River sandy loam, saline, 0 to 3 percent slopes	861	*
40	Fiveoh very fine sandy loam, 2 to 10 percent slopes	3,232	*
40H	Fiveoh loam, saline, 0 to 3 percent slopes	412	*
43B	Grieves fine sandy loam, 2 to 6 percent slopes	833	*
45	Yetull variant loamy sand, 2 to 20 percent slopes	966	*
49	Firth variant fine sandy loam, 0 to 2 percent slopes	2,986	*
51W	Patent variant very fine sandy loam, 0 to 3 percent slopes	3,137	*
52	Laney loam, 0 to 6 percent slopes	1,260	*
52S	Laney variant-Slickspots complex, 3 to 10 percent slopes	700	*
53A	Pinelli loam, 0 to 2 percent slopes	262	*
59	Absher variant silty clay loam, 0 to 2 percent slopes	2,518	*
69	Kiltabar loam, 0 to 3 percent slopes	320	*
78A	Ryan Park sandy loam, 0 to 2 percent slopes	928	*
78B	Ryan Park sandy loam, 2 to 6 percent slopes	1,922	*
79D	Blackhall sandy loam, 6 to 30 percent slopes	160	*
86	Ansel loam, 10 to 30 percent slopes	260	*
101	Echemoor-Clayburn association, 0 to 10 percent slopes	260	*
102	Echemoor-Inchau association, 3 to 10 percent slopes	405	*
105	Starman-Barrett complex, 6 to 40 percent slopes	1,940	*
107	Starman-Vabem complex, 10 to 40 percent slopes	20	*
108	Lymanson-Youga association, 3 to 20 percent slopes	430	*
109	Lymanson-Roxal association, 3 to 20 percent slopes	300	*
111	Vabem-Inchau association, 6 to 30 percent slopes	1,580	*

Acreage and Proportionate Extent of the Soils

Carbon County Area, Wyoming

Map symbol	Map unit name	Acres	Percent
118A	Alcova-Rock River sandy loams, 0 to 2 percent slopes	5,657	0.1
118B	Alcova-Rock River sandy loams, 2 to 6 percent slopes	191	*
128	McFadden-Brownsto-Blackhall complex, 6 to 20 percent slopes	1,760	*
135B	Cushool sandy loam, 0 to 6 percent slopes	2,028	*
138A	Rawlins-Bosler complex, 0 to 2 percent slopes	1,548	*
138B	Rawlins-Bosler complex, 2 to 6 percent slopes	2,140	*
140	Tisworth-Poposhia complex, 0 to 8 percent slopes	3,581	*
141	McFadden-Brownsto complex, 0 to 8 percent slopes	5,400	0.1
144	McFadden-Blackhall sandy loams, 2 to 15 percent slopes	4,013	*
147	Rogert-Quander-Rock outcrop complex, 20 to 50 percent slopes	1,000	*
200	Patent variant-Hagga complex, 0 to 3 percent slopes	4,899	0.1
208	Pinelli-Forelle association, 3 to 15 percent slopes	2,235	*
209	Chaperton-Boettcher association, 3 to 10 percent slopes	3,480	*
210	Absher variant very fine sandy loam, 0 to 6 percent slopes	13,321	0.3
217	Dahlquist-Cragosen association, 6 to 40 percent slopes, eroded	9,080	0.2
218A	Alcova-Rawlins complex, 0 to 2 percent slopes	1,660	*
218B	Alcova-Rawlins complex, 2 to 6 percent slopes	1,000	*
221	Blazon-Chaperton association, 6 to 12 percent slopes	2,223	*
224A	McFadden-Brownsto complex, 0 to 2 percent slopes	2,040	*
224B	McFadden-Brownsto complex, 2 to 6 percent slopes	2,345	*
224w	McFadden-Brownsto complex, wet, 0 to 3 percent slopes	760	*
225	Cushool-Rock River association, 3 to 10 percent slopes	3,000	*
229	Cushool-Cushool variant association, 3 to 9 percent slopes	1,440	*
235	Blazon-Blazon thin solum loams, 6 to 40 percent slopes	2,304	*
236	Cushool-Worfman-Blackhall sandy loams, 6 to 30 percent slopes	9,556	0.2
237	Seaverson-Blazon complex, 3 to 30 percent slopes, eroded	7,308	0.2
244	Rock River sandy loam, 0 to 6 percent slopes	1,517	*
251	Grieves-Blackhall association, 3 to 30 percent slopes	10,647	0.2
252	Blazon, thin solum-Blazon-Rentsac complex, 10 to 50 percent slopes, eroded	32,645	0.7
253	Blazon-Cushool association, 2 to 20 percent slopes	12,280	0.3
254	Abston-Seaverson complex, 0 to 6 percent slopes	18,974	0.4
255	Playa lakes	660	*
256	McFadden-Brownsto-Rawlins complex 6 to 20 percent slopes	9,836	0.2
257	Havre variant-Glendive variant complex, 0 to 3 percent slopes	5,226	0.1
258	Rock River-Cushool sandy loams, 0 to 12 percent slopes	11,927	0.3
260	Ryan Park-Rock River association, 2 to 20 percent slopes	12,181	0.3
261	Luhon-Rock River association, 0 to 10 percent slopes	7,013	0.2
262	Thermopolis-Rentsac complex, 10 to 30 percent slopes	5,199	0.1
263	Edlin-Carmody sandy loams, 3 to 15 percent slopes	2,448	*
264	Rentsac-rock outcrop complex, 5 to 50 percent slopes	980	*
272	Rawlins gravelly loamy sand, 0 to 10 percent slopes	18,434	0.4
275	Poposhia-Chaperton loams, 0 to 5 percent slopes	4,161	*
278	Ryan Park-Elk Mountain loamy fine sands, 2 to 7 percent slopes	6,000	0.1

Acreage and Proportionate Extent of the Soils

Carbon County Area, Wyoming

Map symbol	Map unit name	Acres	Percent
279	Blackhall-Grieves fine sandy loams, 10 to 40 percent slopes	2,240	*
280	Hazton variant-Baggott variant gravelly sandy loams, 5 to 50 percent slopes	2,500	*
282	Tisworth loam, 0 to 5 percent slopes	4,795	0.1
284	Blackhall-Carmody-Rock outcrop complex, 3 to 50 percent slopes	1,968	*
286	Tisworth fine sandy loam, 1 to 5 percent slopes	4,398	*
296	Pinelli-Boettcher clay loams, 2 to 20 percent slopes	650	*
332	Chaperton,dry-Haterton-Hatermus loams, 2 to 15 percent slopes	9,900	0.2
333	Sagecreek alkali-Sagecreek loams, 0 to 10 percent slopes	12,720	0.3
334	Sagecreek loam, alkali, 1 to 8 percent slopes	3,680	*
336	Haterton,thin solum-Hatermus-Haterton loams, 8 to 30 percent slopes	3,185	*
380	Hazton variant-Burgess association, 5 to 30 percent slopes	1,000	*
400	Firth variant-Canburn variant complex, 0 to 3 percent slopes	2,675	*
483	Sandbranch fine sandy loam, 0 to 6 percent slopes	3,175	*
495	Chaperton,dry-Sagecreek loams, 2 to 10 percent slopes	650	*
502	Hagga loam, saline, alkali, 0 to 2 percent slopes	643	*
703	Havre loam, 0 to 3 percent slopes	2,170	*
761	Glendive variant fine sandy loam, 0 to 3 percent slopes	603	*
911	Forelle-Diamondville loams, 3 to 15 percent slopes	80	*
912	Evanston loam, 0 to 6 percent slopes	2,496	*
928	Grieves variant-Gerrard complex, 0 to 3 percent slopes	4,589	0.1
931	Forelle loam, 0 to 6 percent slopes	4,800	0.1
1202	Delplain variant-Morling complex, 6 to 30 percent slopes	2,974	*
1209	Zillman-Peyton variant associaiton, 10 to 50 percent slopes	7,000	0.2
1217	Zillman variant-Highpoint association, 10 to 60 percent slopes	5,042	0.1
1251	McFadden-Blackhall-Edlin sandy loams, 5 to 50 percent slopes	20,463	0.5
1252	Rentsac-Blazon-Rubble land association, 10 to 50 percent slopes	20,816	0.5
1255	Blackhall-Rentsac complex, 10 to 40 percent slopes	6,335	0.1
1256	Rawlins-Rock River association, 0 to 15 percent slopes	13,968	0.3
1260	McFadden-Edlin association, 2 to 20 percent slopes	17,571	0.4
1912	Peyton variant-Evanston variant fine sandy loams, 0 to 6 percent slopes	9,350	0.2
2080	Pinelli variant-Forelle association, 0 to 10 percent slopes	2,858	*
2199	Anchutz sandy loam, 0 to 6 percent slopes	15,980	0.4
9120	Evanston variant-Evanston complex, 0 to 6 percent slopes	5,011	0.1
W	Water	36,203	0.8
Total		541,365	12.2

* Less than 0.1 percent.

Appendix M
NRCS Rangeland Productivity and Plant Composition

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
4:						
Canburn	SUBIRRIGATED (10-14SE)	4,300	3,700	3,000	Basin wildrye	20
					Bluejoint	15
					Northern reedgrass	15
					Prairie cordgrass	15
					Nebraska sedge	10
					Canada wildrye	5
					Other perennial forbs	5
					Slender wheatgrass	5
					Tufted hairgrass	5
					Western wheatgrass	1
8:						
Gerrard	WETLAND (10-14SE)	6,000	5,000	3,500	Nebraska sedge	30
					Northern reedgrass	10
					Willow	10
					American bistort	5
					American mannagrass	5
					Arrowgrass	5
					Baltic rush	5
					Blueeyed grass	5
					Clustered field sedge	5
					Common reed	5
					Horsetail	5
					Tufted hairgrass	5
					Water hemlock	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
9:						
Grieves variant	LOWLAND (10-14SE)	3,000	2,300	1,600	Western wheatgrass	20
					Basin wildrye	10
					Narrowleaf cottonwood	10
					Needleandthread	10
					Silver sagebrush	10
					Big sagebrush	5
					Canby bluegrass	5
					Indian ricegrass	5
					Prairie Junegrass	5
					Yellow rabbitbrush	5
9H:						
Grieves variant, alkali	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	Alkali sacaton	15
					Basin wildrye	15
					Greasewood	15
					Indian ricegrass	5
					Inland saltgrass	5
					Western wheatgrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
13:						
Rhoamett	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	Streambank wheatgrass	40
					Green needlegrass	10
					Birdfoot sagebrush	5
					Bluebunch wheatgrass	5
					Muttongrass	5
					Other perennial forbs	5
					Other perennial grasses	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Truckee rabbitbrush	5
					Winterfat	5
15A:						
Poposhia	LOAMY (10-14SE)	1,400	1,100	600	---	---
15B:						
Poposhia	LOAMY (10-14SE)	1,400	1,100	600	---	---
18A:						
Alcova	SANDY (10-14SE)	1,500	1,200	700	---	---
18B:						
Alcova	SANDY (10-14SE)	1,500	1,200	700	---	---
18C:						
Alcova	SANDY (10-14SE)	1,500	1,200	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
18H:						
Alcova, saline	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
20:						
Debone	SALINE UPLAND (7-9GR)	650	500	300	Greasewood	20
					Basin wildrye	10
					Fourwing saltbush	10
					Indian ricegrass	10
					Western wheatgrass	10
					Alkali sacaton	5
					Bottlebrush squirreltail	5
					Bud sagebrush	5
22:						
Edlin	SANDY (10-14SE)	1,500	1,200	700	---	---
29:						
Canburn variant	SUBIRRIGATED (10-14SE)	4,000	3,000	2,500	Basin wildrye	40
					Tufted hairgrass	20
					Western wheatgrass	10
					Nebraska sedge	5
					Northern reedgrass	5
					Slender wheatgrass	5
					Willow	5
					Canada wildrye	3

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
31A:						
Tisworth	SALINE UPLAND (10-14SE)	650	500	300	Fourwing saltbush	50
					Streambank wheatgrass	15
					Indian ricegrass	10
					Winterfat	10
					Bottlebrush squirreltail	5
					Greasewood	5
					Other perennial forbs	5
31B:						
Tisworth	SALINE UPLAND (10-14SE)	650	500	300	Fourwing saltbush	50
					Streambank wheatgrass	15
					Indian ricegrass	10
					Winterfat	10
					Bottlebrush squirreltail	5
					Greasewood	5
					Other perennial forbs	5
34:						
Tresano	LOAMY (7-9GR)	1,400	1,100	600	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
38A:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5
38B:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5
38C:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
38H:						
Rock River, saline	SANDY (10-14SE)	1,500	1,200	700	---	---
40:						
Fiveoh	SANDY (10-14SE)	1,500	1,200	700	---	---
40H:						
Fiveoh, saline	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
43B:						
Grieves	SANDY (10-14SE)	1,500	1,200	700	---	---
45:						
Yetull variant	SANDS (10-14SE)	1,700	1,400	900	---	---
49:						
Firth variant	LOWLAND (10-14SE)	3,000	2,300	1,600	---	---
51W:						
Patent variant	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
52:						
Laney	SALINE UPLAND (7-9GR)	600	450	300	Fourwing saltbush	50
					Other shrubs	15
					Bottlebrush squirreltail	10
					Indian ricegrass	10
					Other perennial grasses	10
					Bud sagebrush	5
					Sandberg bluegrass	5
					Streambank wheatgrass	5
					Winterfat	5
52S:						
Laney variant	SALINE LOWLAND, DRAINED (7-9GR)	2,000	1,200	800	---	---
Slickspots	---	---	---	---	---	---
53A:						
Pinelli	LOAMY (10-14SE)	1,400	1,100	600	Streambank wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
59:						
Absher variant	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
69:						
Kiltabar	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
78A:						
Ryan Park	SANDY (10-14SE)	1,500	1,200	700	---	---
78B:						
Ryan Park	SANDY (10-14SE)	1,500	1,200	700	---	---
79D:						
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
86:						
Ansel	---	---	---	---	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
101:						
Echemoor	LOAMY (15-19SE)	2,000	1,500	800	Bluebunch wheatgrass	20
					Griffith wheatgrass	10
					Idaho fescue	10
					Needleandthread	10
					Basin wildrye	5
					Big sagebrush	5
					Parry's danthonia	5
					Prairie Junegrass	5
					Spike fescue	5
					Threetip sagebrush	5
Clayburn	LOAMY (15-19SE)	2,400	2,000	1,400	Idaho fescue	20
					Streambank wheatgrass	15
					Antelope bitterbrush	10
					Big sagebrush	10
					Canby bluegrass	10
					Other perennial grasses	10
					Spike fescue	10
					Mountain brome	5
					Other perennial forbs	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
102:						
Echemoor	LOAMY (15-19SE)	2,000	1,500	800	Bluebunch wheatgrass	20
					Griffith wheatgrass	10
					Idaho fescue	10
					Needleandthread	10
					Basin wildrye	5
					Big sagebrush	5
					Parry's danthonia	5
					Prairie Junegrass	5
					Spike fescue	5
					Threetip sagebrush	5
Inchau	LOAMY (15-19SE)	2,000	1,500	800	---	---
105:						
Starman	VERY SHALLOW (15-19SE)	600	500	300	Bluebunch wheatgrass	20
					Mountain mahogany	15
					Antelope bitterbrush	10
					Idaho fescue	10
					Needleandthread	10
					Black sagebrush	5
					Juniper	5
					Prairie Junegrass	5
					Sandberg bluegrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
105: Barrett	SHALLOW LOAMY (15-19SE)	1,200	900	700	Needleandthread	20
					Bluebunch wheatgrass	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Threadleaf sedge	10
					Sandberg bluegrass	5
					Winterfat	5
107: Starman	VERY SHALLOW (15-19SE)	600	500	300	Bluebunch wheatgrass	20
					Mountain mahogany	15
					Antelope bitterbrush	10
					Idaho fescue	10
					Needleandthread	10
					Black sagebrush	5
					Juniper	5
					Prairie Junegrass	5
					Sandberg bluegrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
107: Vabem	SHALLOW LOAMY (15-19SE)	1,400	1,100	800	Bluebunch wheatgrass	20
					Griffith wheatgrass	10
					Threetip sagebrush	10
					Antelope bitterbrush	5
					Black sagebrush	5
					Curleaf mountain mahogany	5
					Idaho fescue	5
					Mountain muhly	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Spike fescue	5
108: Lymanson	LOAMY (15-19SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Black sagebrush	10
					Muttongrass	10
					Needleandthread	10
					Prairie sagewort	5
Youga	LOAMY (15-19SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
109:						
Lymanson	LOAMY (15-19SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Black sagebrush	10
					Muttongrass	10
					Needleandthread	10
					Prairie sagewort	5
Roxal	SHALLOW LOAMY (15-19SE)	1,400	1,100	800	---	---
111:						
Vabem	SHALLOW LOAMY (15-19SE)	1,400	1,100	800	---	---
Inchau	LOAMY (15-19SE)	1,200	900	700	---	---
118A:						
Alcova	LOAMY (10-14SE)	1,400	1,100	600	---	---
Rock River	LOAMY (10-14SE)	1,400	1,100	600	---	---
118B:						
Alcova	LOAMY (10-14SE)	1,400	1,100	600	---	---
Rock River	LOAMY (10-14SE)	1,400	1,100	600	---	---
128:						
McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Brownsto	GRAVELLY (10-14SE)	650	450	300	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
128:						
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
135B:						
Cushool	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Indian ricegrass	20
					Streambank wheatgrass	20
					Silver sagebrush	10
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Plains reedgrass	5
					Sandberg bluegrass	5
138A:						
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
138B:						
Bosler	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	30
					Streambank wheatgrass	20
					Indian ricegrass	15
					Silver sagebrush	10
					Threadleaf sedge	10
					Bluebunch wheatgrass	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
138B:						
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
138B: Bosler	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	30
					Streambank wheatgrass	20
					Indian ricegrass	15
					Silver sagebrush	10
					Threadleaf sedge	10
					Bluebunch wheatgrass	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
140: Tisworth	SALINE UPLAND (10-14SE)	650	500	300	Fourwing saltbush	50
					Streambank wheatgrass	15
					Indian ricegrass	10
					Winterfat	10
					Bottlebrush squirreltail	5
					Greasewood	5
					Other perennial forbs	5
Poposhia	LOAMY (10-14SE)	1,400	1,100	600	---	---
141: McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Brownsto	GRAVELLY (10-14SE)	650	450	300	---	---
144: McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
144: Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
147: Rogert	VERY SHALLOW (15-19SE)	1,400	1,000	700	Western wheatgrass	20
					Antelope bitterbrush	5
					Bluebunch wheatgrass	5
					Needleandthread	5
					Prairie Junegrass	5
Quander	SHALLOW LOAMY (15-19SE)	1,500	1,200	1,000	Bluebunch wheatgrass	25
					Antelope bitterbrush	15
					Big sagebrush	15
					Idaho fescue	15
					Muttongrass	10
					Common snowberry	5
					Saskatoon serviceberry	5
Rock outcrop	---	---	---	---	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
200:						
Patent variant	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
Hagga	SALINE SUBIRRIGATED (10-14SE)	2,500	2,000	1,500	Western wheatgrass	40
					Sedge	25
					Basin wildrye	5
					Rush	5
					Slender wheatgrass	5
208:						
Pinelli	LOAMY (10-14SE)	1,400	1,100	600	---	---
Forelle	LOAMY (10-14SE)	1,400	1,100	600	---	---
209:						
Chaperton	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Prairie Junegrass	5
					Sandberg bluegrass	5
Boettcher	LOAMY (10-14SE)	1,400	1,100	600	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
210:						
Absher variant	SALINE UPLAND (10-14SE)	650	500	300	Gardner's saltbush	40
					Indian ricegrass	10
					Western wheatgrass	10
					Birdfoot sagebrush	5
					Bottlebrush squirreltail	5
					Sandberg bluegrass	5
					Desert biscuitroot	1
					Spiny phlox	1
217:						
Dahlquist	GRAVELLY (10-14SE)	650	450	300	---	---
Cragosen	GRAVELLY (10-14SE)	650	450	300	---	---
218A:						
Alcova	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	20
					Indian ricegrass	10
					Threadleaf sedge	10
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Prairie Junegrass	5
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
218B:						
Alcova	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	20
					Indian ricegrass	10
					Threadleaf sedge	10
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Prairie Junegrass	5
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
221:						
Blazon	SHALLOW LOAMY (10-14 SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Muttongrass	10
					Black sagebrush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
					Yellow rabbitbrush	5
Chaperton	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Prairie Junegrass	5
					Sandberg bluegrass	5
224A:						
McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
224A: Brownsto	GRAVELLY (10-14SE)	650	450	300	---	---
224B: McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Brownsto	GRAVELLY (10-14SE)	650	450	300	---	---
224w: McFadden, wet	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Brownsto, wet	GRAVELLY (10-14SE)	650	450	300	---	---
225: Cushool	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Indian ricegrass	20
					Streambank wheatgrass	20
					Silver sagebrush	10
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Plains reedgrass	5
					Sandberg bluegrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
225:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5
229:						
Cushool	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Indian ricegrass	20
					Streambank wheatgrass	20
					Silver sagebrush	10
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Plains reedgrass	5
					Sandberg bluegrass	5
Cushool variant	SANDY (10-14SE)	1,500	1,200	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
235:						
Blazon	SHALLOW LOAMY (10-14 SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Muttongrass	10
					Black sagebrush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
					Yellow rabbitbrush	5
Blazon, THIN SOLUM	SHALE (10-14SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Muttongrass	10
					Black sagebrush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
					Yellow rabbitbrush	5
236:						
Cushool	SANDY (10-14SE)	1,500	1,200	700	---	---
Worfman	SHALLOW SANDY (10-14SE)	1,200	900	700	Bluebunch wheatgrass	20
					Indian ricegrass	10
					Needleandthread	10
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Skunkbush sumac	5
					Truckee rabbitbrush	5
					Western wheatgrass	5
					Winterfat	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
236:						
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
237:						
Seaverson	SALINE UPLAND (10-14SE)	650	500	300	Gardner's saltbush	40
					Indian ricegrass	15
					Western wheatgrass	15
					Bottlebrush squirreltail	10
					Birdfoot sagebrush	5
					Desert biscuitroot	5
					Sandberg bluegrass	5
Blazon	SHALLOW LOAMY (10-14 SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Muttongrass	10
					Black sagebrush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
					Yellow rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac	Pct	
244:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5
251:						
Grieves	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Other perennial forbs	5
					Other perennial grasses	5
					Truckee rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
251: Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
252: Blazon	SHALLOW LOAMY (10-14 SE)	1,400	1,100	800	---	---
Blazon, thin solum	SHALE (10-14SE)	400	300	200	---	---
Rentsac	VERY SHALLOW (10-14SE)	600	450	250	---	---
253: Blazon	SHALLOW LOAMY (10-14 SE)	1,400	1,100	800	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Muttongrass	10
					Black sagebrush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
					Yellow rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
253:						
Cushool	LOAMY (10-14SE)	1,400	1,100	600	Streambank wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Other perennial forbs	5
					Other perennial grasses	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
254:						
Abston	IMPERVIOUS CLAY (10-14SE)	600	400	250	---	---
Seaverson	IMPERVIOUS CLAY (10-14SE)	600	400	250	Birdfoot sagebrush	25
					Western wheatgrass	25
					Bottlebrush squirreltail	5
					Desert biscuitroot	5
					Gardner's saltbush	5
					Indian ricegrass	5
					Sandberg bluegrass	5
255:						
Ponded soils	---	---	---	---	---	---
256:						
McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
256:						
Brownsto	GRAVELLY (10-14SE)	650	450	300	Bluebunch wheatgrass	25
					Needleandthread	10
					Big sagebrush	5
					Black sagebrush	5
					Blue grama	5
					Bottlebrush squirreltail	5
					Indian ricegrass	5
					Sandberg bluegrass	5
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
257:						
Havre variant	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
Glendive variant	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	---	---
258:						
Rock River	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Blue grama	5
					Canby bluegrass	5
					Indian ricegrass	5
					Truckee rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
258: Cushool	LOAMY (10-14SE)	1,400	1,100	600	Streambank wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Other perennial forbs	5
					Other perennial grasses	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
260: Ryan Park	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	30
					Streambank wheatgrass	15
					Indian ricegrass	10
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Spineless horsebrush	5
					Threadleaf sedge	5
					Winterfat	5
					Yellow rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
260:						
Rock River	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Truckee rabbitbrush	5
261:						
Luhon	SHALLOW LOAMY (10-14 SE)	1,500	1,200	700	Other perennial grasses	25
					Streambank wheatgrass	20
					Other perennial forbs	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Needleandthread	10
					Other shrubs	10
Rock River	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Blue grama	5
					Canby bluegrass	5
					Indian ricegrass	5
					Truckee rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
262:						
Rentsac	VERY SHALLOW (10-14SE)	600	450	250	---	---
Thermopolis	SHALLOW LOAMY (10-14 SE)	1,200	900	700	Bluebunch wheatgrass	20
					Western wheatgrass	20
					Needleandthread	10
					Big sagebrush	5
					Indian ricegrass	5
					Prairie Junegrass	5
263:						
Edlin	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	20
					Indian ricegrass	15
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Bottlebrush squirreltail	5
					Silver sagebrush	5
					Threadleaf sedge	5
Carmody	SANDY (10-14SE)	1,500	1,200	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
264:						
Rentsac	SHALLOW LOAMY (10-14 SE)	1,000	800	500	Bluebunch wheatgrass	50
					Needleandthread	15
					Western wheatgrass	10
					Blue grama	5
					Other perennial forbs	5
					Other perennial grasses	5
					Other shrubs	5
Rock outcrop	---	---	---	---	---	---
272:						
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
275:						
Poposhia	LOAMY (10-14SE)	1,400	1,100	600	---	---
Chaperton	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Prairie Junegrass	5
					Sandberg bluegrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
278:						
Ryan Park	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	30
					Streambank wheatgrass	15
					Indian ricegrass	10
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Spineless horsebrush	5
					Threadleaf sedge	5
					Winterfat	5
					Yellow rabbitbrush	5
Elk Mountain	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	30
					Streambank wheatgrass	15
					Indian ricegrass	10
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Spineless horsebrush	5
					Threadleaf sedge	5
					Truckee rabbitbrush	5
					Winterfat	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
279:						
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
Grieves	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Indian ricegrass	10
					Bottlebrush squirreltail	5
					Other perennial forbs	5
					Other perennial grasses	5
					Truckee rabbitbrush	5
280:						
Hazton variant	ROCKY HILLS (15-19SE)	1,150	900	550	---	---
Baggott variant	SHALLOW LOAMY (15-19SE)	1,400	1,100	800	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
282:						
Tisworth	SALINE LOWLAND (10-14SE)	2,500	1,800	1,200	Alkali sacaton	30
					Basin wildrye	15
					Greasewood	15
					Western wheatgrass	10
					Fourwing saltbush	5
					Inland saltgrass	5
					Winterfat	5
284:						
Blackhall	SHALLOW BREAKS (10-14SE)	1,200	1,000	800	Utah juniper	45
					Bluebunch wheatgrass	15
					Needleandthread	15
					Big sagebrush	5
					Indian ricegrass	5
					Prairie Junegrass	5
					Western wheatgrass	5
Carmody	SHALLOW BREAKS (10-14SE)	1,300	1,100	800	Utah juniper	45
					Bluebunch wheatgrass	15
					Needleandthread	15
					Big sagebrush	5
					Indian ricegrass	5
					Prairie Junegrass	5
					Western wheatgrass	5
Rock outcrop	---	---	---	---	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
286:						
Tisworth	SALINE UPLAND (10-14SE)	650	500	300	Fourwing saltbush	50
					Streambank wheatgrass	15
					Indian ricegrass	10
					Winterfat	10
					Bottlebrush squirreltail	5
					Greasewood	5
					Other perennial forbs	5
296:						
Pinelli	CLAYEY (10-14SE)	1,400	1,100	600	Streambank wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
Boettcher	CLAYEY (10-14SE)	1,400	1,200	800	Western wheatgrass	35
					Green needlegrass	25
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Sandberg bluegrass	5
					Unknowns	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac	Pct	
332:						
Chaperton, dry	LOAMY (7-9GR)	700	500	300	Western wheatgrass	25
					Big sagebrush	10
					Needleandthread	10
					Prairie Junegrass	10
					Sandberg bluegrass	10
Hatermus	SALINE UPLAND (7-9GR)	450	350	200	Bluebunch wheatgrass	30
					Streambank wheatgrass	15
					Indian ricegrass	10
					Sandberg bluegrass	10
					Big sagebrush	5
					Black sagebrush	5
					Needleandthread	5
					Needleleaf sedge	5
Haterton	SHALLOW LOAMY (7-9GR)	450	350	200	Bluebunch wheatgrass	30
					Streambank wheatgrass	15
					Fourwing saltbush	10
					Indian ricegrass	10
					Big sagebrush	5
					Black sagebrush	5
					Needleandthread	5
					Needleleaf sedge	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
333:						
Sagecreek, alkali	SALINE UPLAND (7-9GR)	600	450	300	Fourwing saltbush	35
					Bottlebrush squirreltail	20
					Indian ricegrass	15
					Western wheatgrass	15
					Needleandthread	5
					Sandberg bluegrass	5
Sagecreek	LOAMY (7-9GR)	700	500	300	Streambank wheatgrass	30
					Needleandthread	20
					Big sagebrush	10
					Indian ricegrass	10
					Bluebunch wheatgrass	5
					Fourwing saltbush	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Truckee rabbitbrush	5
					Winterfat	5
334:						
Sagecreek, alkali	SALINE UPLAND (7-9GR)	600	450	300	Fourwing saltbush	35
					Bottlebrush squirreltail	20
					Indian ricegrass	15
					Western wheatgrass	15
					Needleandthread	5
					Sandberg bluegrass	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
336:						
Haterton, thin solum	SHALLOW LOAMY (7-9GR)	450	350	200	Bluebunch wheatgrass	30
					Streambank wheatgrass	15
					Fourwing saltbush	10
					Indian ricegrass	10
					Big sagebrush	5
					Black sagebrush	5
					Needleandthread	5
					Needleleaf sedge	5
Hatermus	SALINE UPLAND (7-9GR)	600	450	300	---	---
Haterton	SHALLOW LOAMY (7-9GR)	450	350	200	Bluebunch wheatgrass	30
					Streambank wheatgrass	15
					Fourwing saltbush	10
					Indian ricegrass	10
					Big sagebrush	5
					Black sagebrush	5
					Needleandthread	5
					Needleleaf sedge	5
380:						
Hazton variant	ROCKY HILLS (15-19SE)	1,150	900	550	---	---
Burgess	LOAMY (15-19SE)	2,000	1,500	800	---	---
400:						
Firth variant	LOWLAND (10-14SE)	3,000	2,300	1,600	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
400: Canburn variant	WETLAND (10-14SE)	6,000	5,000	3,500	Nebraska sedge	35
					Northern reedgrass	10
					Tufted hairgrass	10
					Arrowgrass	5
					Willow	5
					Iris	1
483: Sandbranch	SALINE UPLAND (7-9GR)	600	450	300	Fourwing saltbush	50
					Bottlebrush squirreltail	10
					Indian ricegrass	10
					Bud sagebrush	5
					Greasewood	5
					Streambank wheatgrass	5
495: Chaperton, dry	LOAMY (7-9GR)	700	500	300	Western wheatgrass	25
					Big sagebrush	10
					Needleandthread	10
					Prairie Junegrass	10
					Sandberg bluegrass	10

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
495: Sagecreek	LOAMY (7-9GR)	700	500	300	Streambank wheatgrass	30
					Needleandthread	20
					Big sagebrush	10
					Indian ricegrass	10
					Bluebunch wheatgrass	5
					Fourwing saltbush	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
					Truckee rabbitbrush	5
					Winterfat	5
502: Hagga, saline, alkali	SALINE SUBIRRIGATED (10-14SE)	2,500	2,000	1,500	Western wheatgrass	40
					Sedge	25
					Basin wildrye	5
					Rush	5
					Slender wheatgrass	5
703: Havre	LOWLAND (10-14SE)	3,000	2,300	1,600	---	---
761: Glendive variant	LOWLAND (10-14SE)	3,000	2,300	1,600	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac	Pct	
911:						
Forelle	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
Diamondville	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	30
					Needleandthread	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Green needlegrass	5
					Muttongrass	5
					Yellow rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
912:						
Evanston	LOAMY (10-14SE)	1,400	1,100	600	Streambank wheatgrass	20
					Big sagebrush	15
					Needleandthread	15
					Bluebunch wheatgrass	10
					Canby bluegrass	10
					Letterman's needlegrass	10
					Indian ricegrass	5
					Other perennial forbs	5
					Other perennial grasses	5
					Prairie Junegrass	5
928:						
Grieves variant	LOWLAND (10-14SE)	3,000	2,300	1,600	Western wheatgrass	20
					Basin wildrye	10
					Narrowleaf cottonwood	10
					Needleandthread	10
					Silver sagebrush	10
					Big sagebrush	5
					Canby bluegrass	5
					Indian ricegrass	5
					Prairie Junegrass	5
					Yellow rabbitbrush	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
928: Gerrard	WETLAND (10-14SE)	3,700	3,100	2,500	Tufted hairgrass Slender wheatgrass Nebraska sedge Bluejoint Western wheatgrass Baltic rush	30 20 15 10 10 5
931: Forelle	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass Big sagebrush Needleandthread Bluebunch wheatgrass Green needlegrass Needleleaf sedge Plains reedgrass Prairie Junegrass Sandberg bluegrass	40 10 10 5 5 5 5 5 5
1202: Delplain variant	SHALE (10-14SE)	400	300	200	---	---
Morling	SHALE (10-14SE)	400	300	200	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
1209:						
Zillman	COARSE UPLAND (15-19SE)	1,200	800	500	Bluebunch wheatgrass	30
					Needleandthread	15
					Black sagebrush	10
					Bottlebrush squirreltail	10
					Streambank wheatgrass	10
					Antelope bitterbrush	5
					Big sagebrush	5
					Other perennial forbs	5
					Other perennial grasses	5
					Sandberg bluegrass	5
Peyton variant	LOAMY (15-19SE)	2,200	1,600	1,100	Bluebunch wheatgrass	25
					Western wheatgrass	25
					Green needlegrass	15
					Needleandthread	10
					Little bluestem	5
					Other perennial grasses	5
					Other shrubs	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
1217:						
Zillman variant	COARSE UPLAND (15-19SE)	1,600	1,200	800	Bluebunch wheatgrass	15
					Big sagebrush	10
					Western wheatgrass	10
					Antelope bitterbrush	5
					Canby bluegrass	5
					Indian ricegrass	5
					Needleandthread	5
					Penstemon	1
					Serviceberry	1
					Spiny phlox	1
Highpoint	GRAVELLY (10-14SE)	650	450	300	---	---
1251:						
McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Edlin	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	20
					Indian ricegrass	15
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Bottlebrush squirreltail	5
					Silver sagebrush	5
					Threadleaf sedge	5

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
1252:						
Rentsac	VERY SHALLOW (10-14SE)	600	450	250	Bluebunch wheatgrass	50
					Needleandthread	15
					Western wheatgrass	10
					Blue grama	5
					Other perennial forbs	5
					Other perennial grasses	5
					Other shrubs	5
Blazon	SHALE (10-14SE)	400	300	200	---	---
Rubble land	---	---	---	---	---	---
1255:						
Blackhall	SHALLOW SANDY (10-14SE)	1,200	900	700	Needleandthread	25
					Bluebunch wheatgrass	20
					Sedge	15
					Black sagebrush	10
					Indian ricegrass	10
					Muttongrass	10
					Western wheatgrass	10
					Big sagebrush	5
					Prairie Junegrass	5
Rentsac	VERY SHALLOW (10-14SE)	600	450	250	---	---
1256:						
Rawlins	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac		Pct
1256:						
Rock River	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	35
					Needleandthread	15
					Big sagebrush	10
					Bluebunch wheatgrass	10
					Blue grama	5
					Canby bluegrass	5
					Indian ricegrass	5
					Truckee rabbitbrush	5
1260:						
McFadden	SHALLOW SANDY (10-14SE)	1,200	900	700	---	---
Edlin	SANDY (10-14SE)	1,500	1,200	700	Needleandthread	25
					Streambank wheatgrass	20
					Indian ricegrass	15
					Big sagebrush	5
					Bluebunch wheatgrass	5
					Bottlebrush squirreltail	5
					Silver sagebrush	5
					Threadleaf sedge	5
1912:						
Peyton variant	LOAMY (15-19SE)	2,000	1,500	800	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac	Pct	
1912:						
Evanston variant	LOAMY (15-19SE)	2,000	1,500	800	Big sagebrush	10
					Bluebunch wheatgrass	10
					Idaho fescue	10
					Needleandthread	10
					Streambank wheatgrass	10
					Canby bluegrass	5
2080:						
Pinelli variant	LOAMY (10-14SE)	1,400	1,100	600	Idaho fescue	15
					Streambank wheatgrass	15
					Green needlegrass	10
					Big sagebrush	5
					Needleandthread	5
					Prairie Junegrass	5
Forelle	LOAMY (10-14SE)	1,400	1,100	600	Western wheatgrass	40
					Big sagebrush	10
					Needleandthread	10
					Bluebunch wheatgrass	5
					Green needlegrass	5
					Needleleaf sedge	5
					Plains reedgrass	5
					Prairie Junegrass	5
					Sandberg bluegrass	5
2199:						
Anchutz	SALINE LOAMY (10-14SE)	900	700	500	---	---

Rangeland Productivity and Plant Composition

Carbon County Area, Wyoming

Map symbol and soil name	Ecological site	Total dry-weight production			Characteristic vegetation	Rangeland composition
		Favorable year	Normal year	Unfavorable year		
		Lb/Ac	Lb/Ac	Lb/Ac	Pct	
9120:						
Evanston variant	LOAMY (15-19SE)	2,000	1,500	800	Big sagebrush	10
					Bluebunch wheatgrass	10
					Idaho fescue	10
					Needleandthread	10
					Streambank wheatgrass	10
					Canby bluegrass	5
Evanston	LOAMY (15-19SE)	1,800	1,500	900	Big sagebrush	10
					Needleandthread	10
					Western wheatgrass	10
					Mountain snowberry	5
					Muttongrass	5
					Prairie Junegrass	5
					Saskatoon serviceberry	5
W:						
Water	---	---	---	---	---	---

Appendix N
Mesoscale Model Simulations in Quasi-Forecast Mode
of the Great Western Storm of 16-20 March 2003

5.3A MESOSCALE MODEL SIMULATIONS IN QUASI-FORECAST MODE OF THE GREAT WESTERN STORM OF 16-20 MARCH 2003

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1. INTRODUCTION

A massive snowstorm crippled large portions of the central Rockies and adjacent plains during the period 16-20 March 2003. Snowfall accumulation in the foothills and mountains exceeded four feet in relatively large regions, while on the plains amounts above two feet were common (Fig. 1; also see Poulos et al. 2003). The large impacts of this historic storm are well documented. This paper examines experimental meso- γ scale model simulations of the event, utilizing larger-scale model-generated boundary conditions, from a forecasting standpoint.

Public forecasts of this event were generally accurate up to several days before the storm hit. NCEP model guidance provided initial alarms (in the form of ensemble forecasts) up to one week prior to the storm (Szoke et al. 2004). As the potential event approached, Eta model forecasts were trending towards a large precipitation event, and by about two days before the onset of snowfall along Colorado's Front Range very large precipitation totals (five or more inches) were output by this model for portions of the region during the period of 17-20 March. Accuracy of these forecasts was perhaps unprecedented in the area, for such a large event, primarily because the orographic forcing was so strong. The Eta forecasts clearly provided a crucial asset towards forecast operations prior to the storm. The model, however, did show some shortcomings regarding the precipitation type distribution, and of course was limited by its relatively large grid spacing, a required feature given the domain size of that model.

The crippling nature of the subsequent storm period, in terms of disrupting transportation and other day-to-day activities, has shown that even if a very large snowfall potential is emphasized in, say, a 2-4 day forecast, society is still vulnerable to this type of storm. Insurance claims and a paralyzed international airport attest to this fact. Importantly, the current challenge is to increase the resolution and details of the forecast to minimize this vulnerability, as much as currently possible.

Close examination of snowfall totals revealed extremely sharp gradients in snowfall, on the order of several feet within a horizontal distance of 15 miles or less. Many of these sharp gradient regions coincided with strong gradients in elevation; however some did not. For example, an area on the plains/foothills interface just north of Denver accumulated only 3-6 inches of wet snowfall, while 15-25 miles to the south, 24-36 inches fell, and areas another 20 miles to the south recorded nearly four feet. Meanwhile, 20-30 miles *north* of the aforementioned area of snowfall minimum, 24-36 inches fell. *All of these locations are at the same approximate elevation.* The current configuration of NWS forecast zones along the urban corridor is not designed to handle these types of gradients, nor is the current configuration of the Eta model. As NWS forecasts evolve towards gridded forecast fields, this issue will be addressed to some degree.

The purpose of this study is to closely examine the causes of extreme snowfall and wind variations in this storm from a mesoscale modeling standpoint in order to better predict them in the future. The MM5 was run in quasi-forecast mode (with Eta forecasts initialized at 00 UTC 17 Mar.) utilizing non-hydrostatic and multiple-grid configurations, with the smallest grid exhibiting 1-2 km horizontal grid spacing. The primary reason for utilizing such a small grid spacing is the presence of steep and variable topography throughout the foothills and higher terrain of the Front Range. The "workstation" Eta was run (non-hydrostatically) utilizing Eta analyses *and* 3-hr. forecasts at the boundaries. The smallest grid contained 2 km grid spacing.

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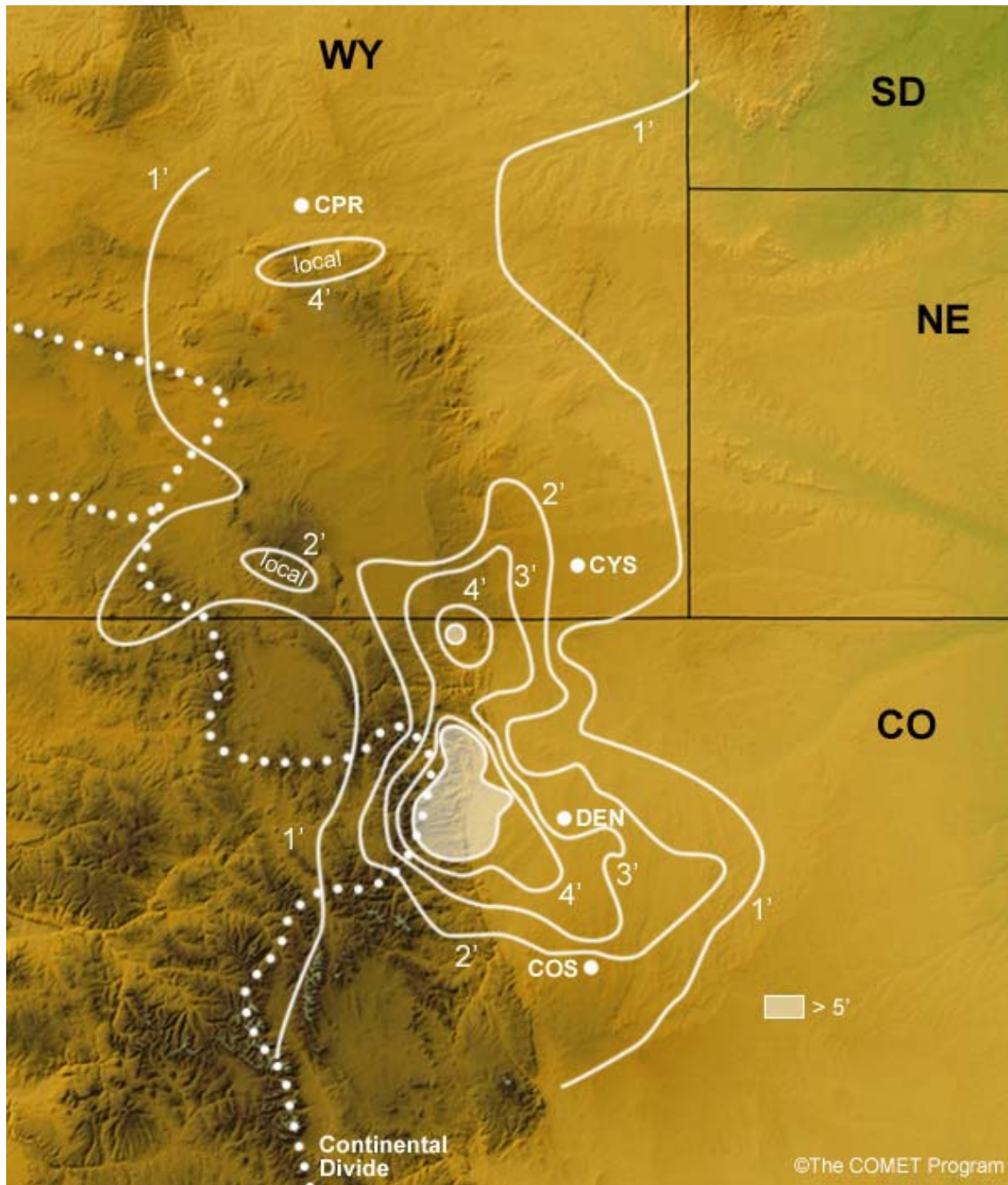


Fig. 1 Snowfall totals (in feet) for a portion of the Front Range region for 17-20 March 2003. Significant snows fell in other regions of the Rocky Mountains to the west of this area (see Meyers et al., 2004).

Preliminary indications are that both mesoscale models produce generally accurate precipitation distributions, and both produce cooler (but still above freezing) low-level conditions along the urban corridor for much of the storm evolution when compared to the operational Eta forecasts. The MM5 forecasts appear to capture better detail in the precipitation distributions, as expected, and exhibit low-level

temperatures closer to freezing in critical areas near the rain/snow line. Comparisons with operational profiler winds show some problems with the strength of the mid-level upslope, a critical component of the storm, and one perhaps related to the relatively warm low-level conditions along the urban corridor. This component is likely a primary factor in determining precipitation rates, in the sense of the warm conveyor

belt running up and over the barrier jet, and thus a critical determinant of surface precipitation type. It appears that an accurate initial analysis and subsequent prediction of the depth of the barrier jet is a crucial requirement to an accurate precipitation forecast. Another important feature of the mid-level easterly flow is its strong variation through the 3-4 day period as synoptic waves passed through the region, and these variations will be compared to the barrier jet depth and distributions of precipitation rates in the near future.

Initially it also appears that relatively subtle terrain features along the plains/foothills interface interacted with the barrier jet to contribute significantly to low-level vertical motion fields, and likely play a role in the cause of the snow minima discussed above.

2. STORM DYNAMICS OVERVIEW

During the period 15-17 March, significant troughing built into the central and southern Rockies and the Great Basin as intense mid- and upper-level jet energy impacted the California coast from the west-northwest. The amplification of the pattern increased rapidly as ridging built over the upper Midwest and mid-Atlantic regions. By 00 UTC 19 March, a strong, deep cutoff low pressure system was established over the southern Rockies and central/southern plains (Fig. 2). For a period of about 48 hours, a classic warm conveyor belt out in front of the cutoff set up and transported large amounts of moisture directly from the Gulf of Mexico northwestward into

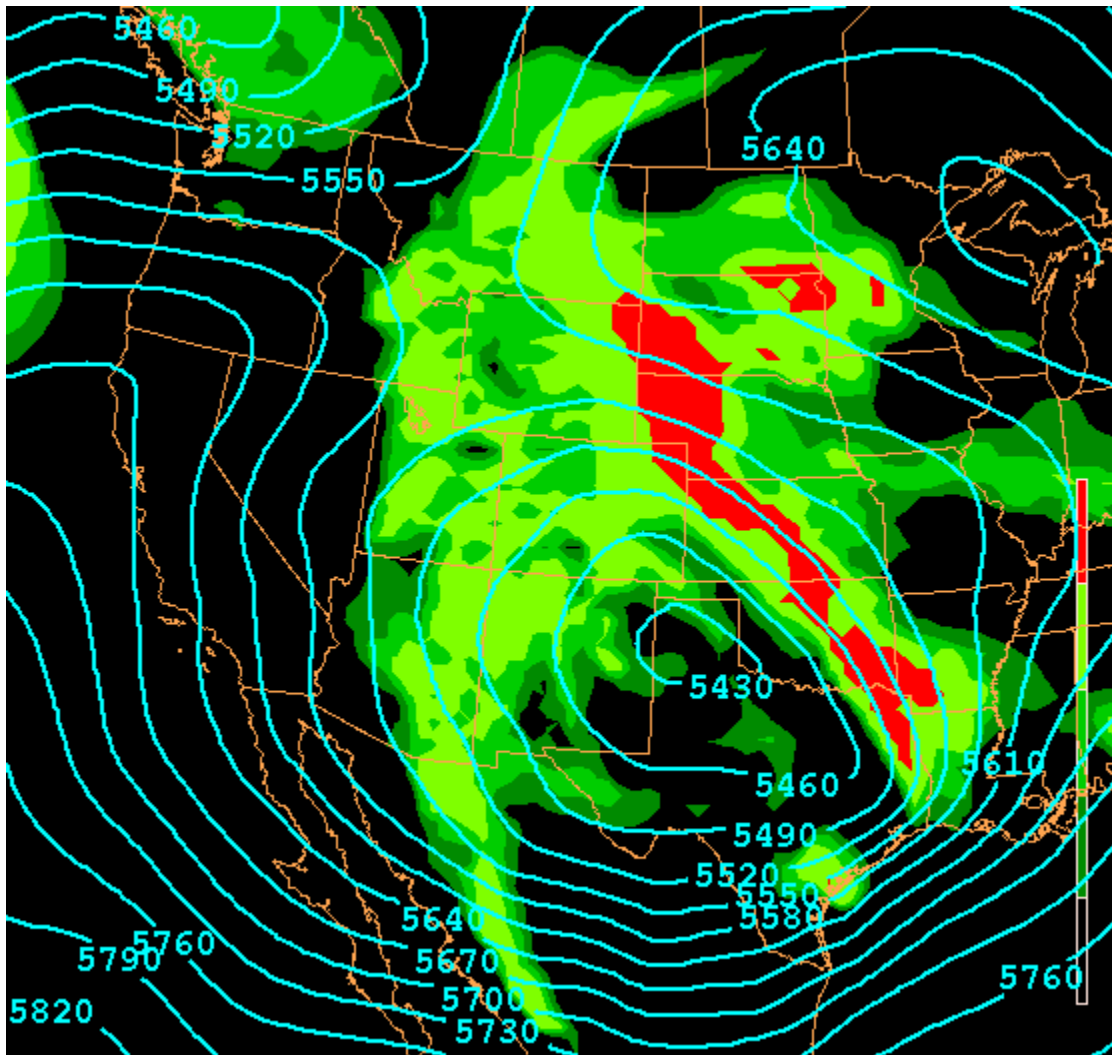


Fig. 2 500 mb heights and 700 mb RH, analyzed at 00 UTC 19 Mar. 2003. Red regions correspond to saturated conditions at 700 mb.

the central Rockies. In the northwestern portion of the cutoff system, a TROWAL-like feature set up as the occlusion matured, and this wraparound feature contributed to heavy precipitation well-removed from the cutoff center off to the southeast.

The mesoscale features of this mega-storm were of critical importance to the resulting precipitation distribution. Observationally, the role of the barrier jet in the storm in producing, first, snow instead of rain in the urban corridor, and, second, uplift strong enough to produce snowfall rates of 1-3 inches per hour for 2-3 days, cannot be overemphasized. Clearly the barrier jet was located on the cold side of a persistent rain/snow boundary that exhibited the classic characteristics of strongly diabatically-forced mesoscale dynamics, a feature documented in previous heavy springtime snowfalls in the urban

corridor (Marwitz and Toth 1993). Furthermore, the three-dimensional configuration of this barrier jet is critical to the attempt to explain the astounding snowfall and wind gradients along the urban corridor. A well-developed barrier jet was apparent by 18 March, and persisted through the 19th. Important facets of this low-level northerly flow regime over and next to the foothills:

- (a) low-level northerly zone was sloped upwards to the west, essentially modifying the obstacle encountered by upslope (easterly) flow and leading to mesoscale uplift in a saturated air mass over and just east of the jet

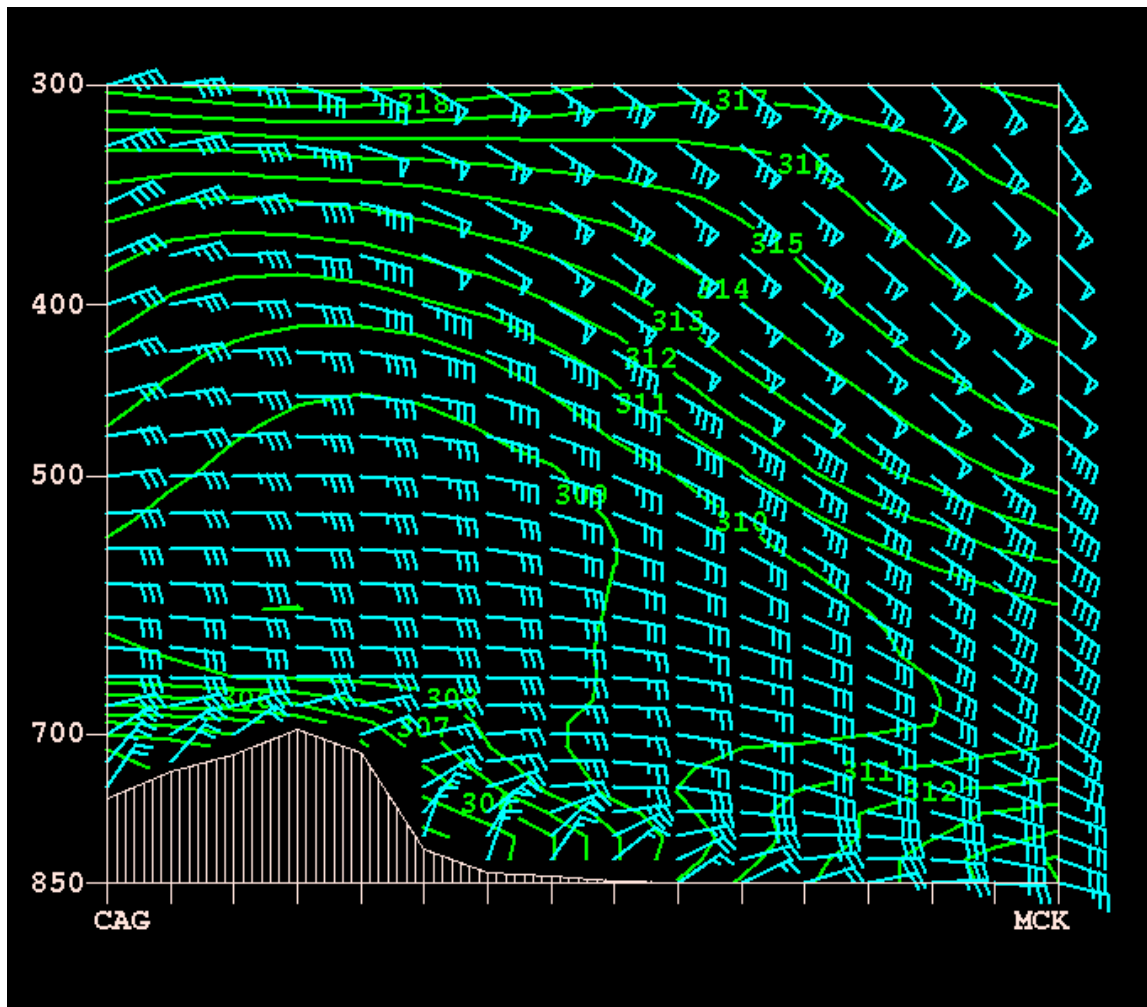


Fig. 3 Vertical cross section showing equiv. potential temp. (K) and winds (knots), 6-hr. forecast from the Eta model initialized at 18 UTC 18 Mar.

- (b) large amounts of melting in the low-levels on the east side of the barrier jet provided latent cooling, thus enhancing the blocking and barrier jet structure, similar to the March 1990 storm studied by Marwitz and Toth (1993) and others.
- (c) significant low-level cold advection from the north/northeast enhanced the stability in the air mass east of the terrain obstacles.

Note in Fig. 3 the cold air stacked up against the Front Range, and the moderate northerly flow within that cold air. Many regions just east of the foothills experienced surface wind gusts in the 30 to 40 knot range, causing extensive blowing and drifting snow. Interestingly, at this point a well-defined convergence line does not exist on the east side of the jet, and this was confirmed in surface observations. Convectively unstable conditions are noted over portions of the plains in Fig. 3.

3. Mesoscale model simulations

The MM5 was set up with a 5-grid nested configuration, the smallest domain (grid 5) centered on north-central CO and exhibiting a 1.5 km grid

spacing. Eta operational forecasts from the run initialized at 00 UTC 17 Mar. served as large-scale boundary conditions.

Fig. 4 shows the total precipitation (mm) predicted by the model through 84 hours (ending at 12 UTC 20 Mar.). Notable features are the foothills maxima in the higher terrain (but east of the Continental Divide) of Boulder and Larimer Counties (the Divide runs along the western boundaries of these two counties), with several locations predicted to have over 130 mm (more than 5 inches). Three relative minima are also very interesting:

1. northeastern Boulder Co. (less than 50 mm)
2. southeastern Larimer Co. (43.8 mm)
3. northeastern Larimer Co. (27.5 mm)

All of these regions experienced snow minima compared to observed snowfall in immediately surrounding regions of similar elevations (Fig. 1). This is best shown by examining high-resolution satellite imagery after the storm as the melting process started under sunny skies (Fig. 5).

Dataset: test RIP: rip
Fest: 84.00
Total precip. since h 0
Total precip. since h 0

Init: 0000 UTC Mon 17 Mar 03
Valid: 1200 UTC Thu 20 Mar 03 (0600 CST Thu 20 Mar 03)

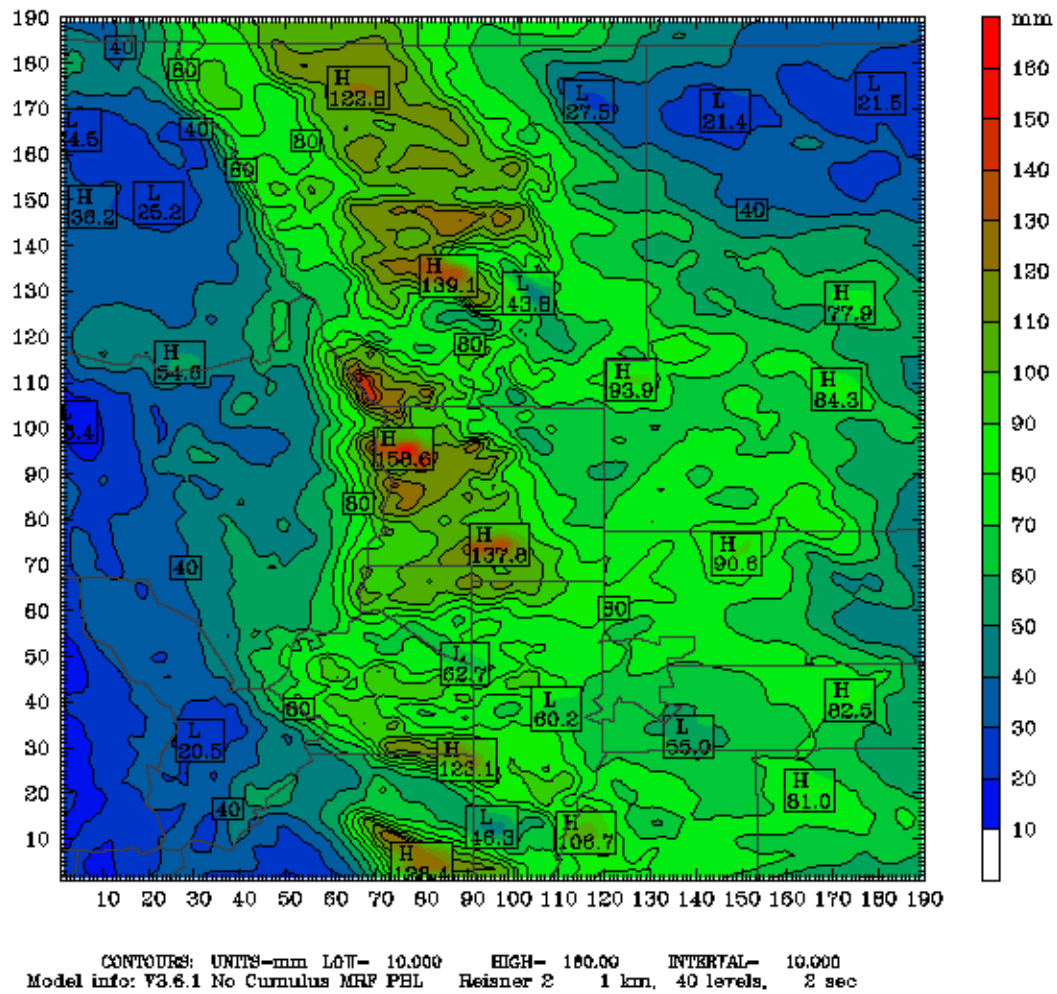


Fig. 4 MM5-predicted precipitation (mm) for 84 hours of simulations ending at 12 UTC 20 Mar..

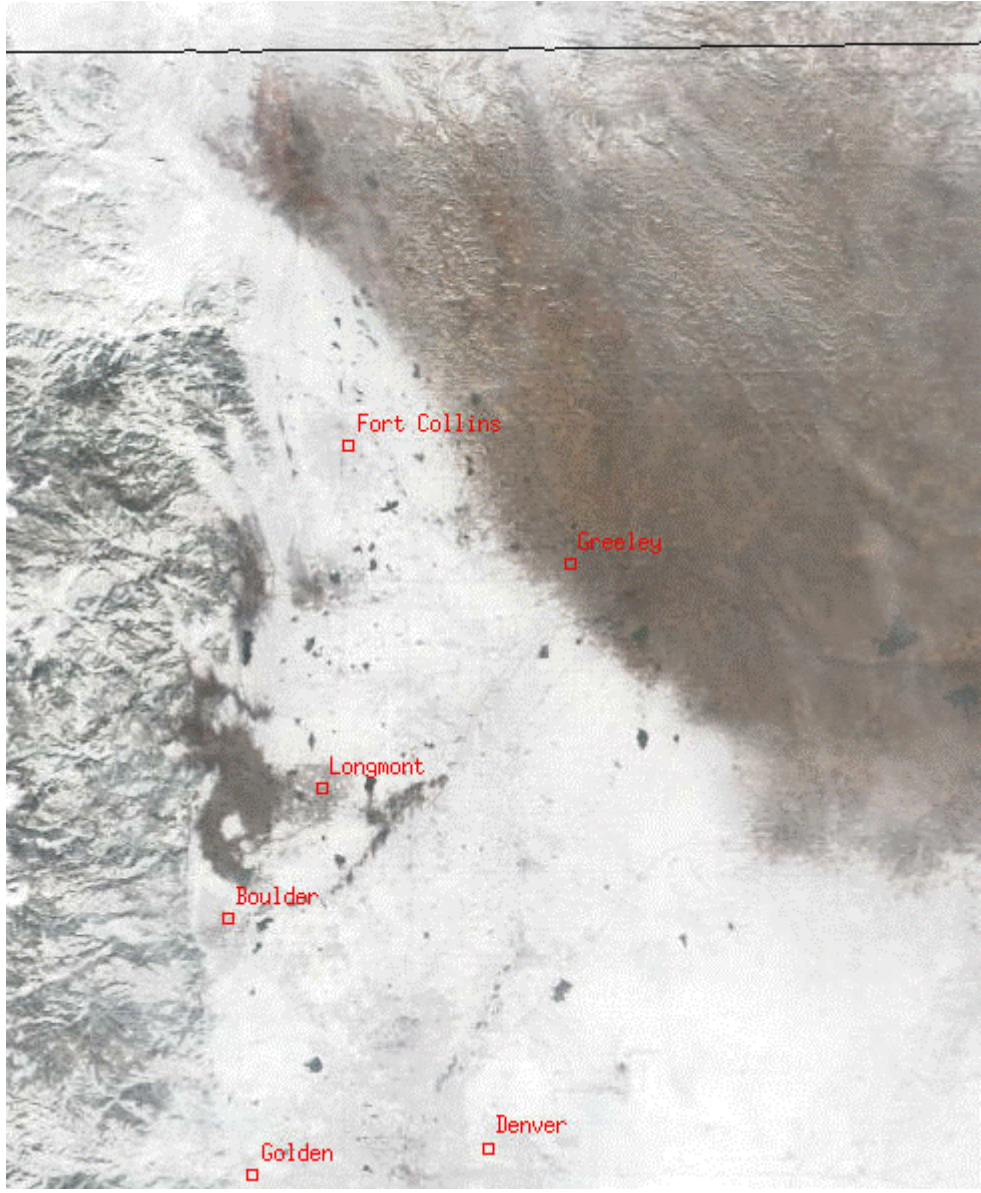


Fig. 5 High-resolution visible image (MODIS) on 22 Mar.. Complex patterns on the west side are timbered and canyon areas. Darker areas just south of the WY state line, southwest of Fort Collins and west of Longmont are areas where much less snow accumulation was observed.

Dataset: test RIP: rip
 Fcst: 42.00
 Temperature
 Temperature
 Horizontal wind vectors

Init: 0000 UTC Mon 17 Mar 03
 Valid: 1800 UTC Tue 18 Mar 03 (1200 CST Tue 18 Mar 03)
 at sigma = 0.999
 at sigma = 0.999
 at sigma = 0.999

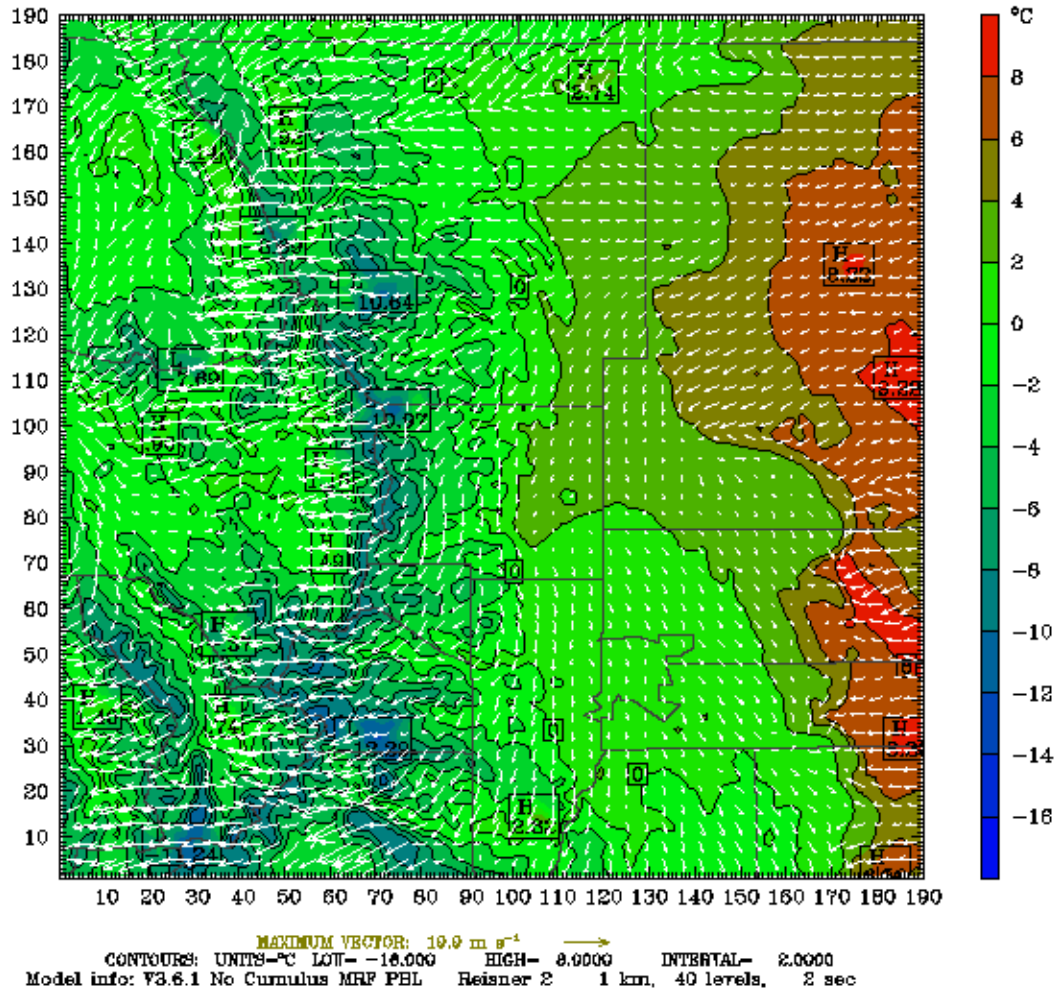


Fig. 6 MM5 42-hr. forecast of lowest level temperature (C) and winds (m/s). Note the relatively warmer areas along the foothills in southeastern Larimer Co. and northeastern Boulder Co.

Fig. 6 demonstrates several interesting aspects of the simulations. Relatively warmer conditions are predicted in general along the eastern portions of Larimer Co. and northeastern Boulder Co., in agreement with observations in two of the snow minima regions. However, in comparison with observations, these areas are predicted to be a few degrees F warmer by the model. In the urban corridor region just south of the Cheyenne Ridge, the snow minimum region discussed previously appears to be caused by lower precipitation values rather than warmer temperatures (see Wesley et al., 1995). This is often observed in storms characterized by strong north winds at the surface in this region. Also note the northerly flow over the foothills, and a strong

convergence line oriented nearly E-W along the WY border.

More results of these MM5 simulations are under investigation, including a detailed examination of the areas that experienced warmer surface conditions and less snowfall. Potential mechanisms include blocking of the barrier-jet induced cold advection by small-scale terrain features, and relatively warm air (originating over the canyons to the northwest of these locations) acting as the source region for the surface conditions over these areas.

The “workstation” Eta model was also set up non-hydrostatically, with multiple nested grid configuration and innermost grid spacing set at 2 km. Fig. 7 shows the predicted total precipitation for the 72-hr. period ending at 12 UTC 20 Mar.. Though the details in the plot do not resemble those of the MM5-predicted precipitation, especially over the eastern foothills and plains interface, note the maxima in the high terrain just east of the Continental Divide, with one elevated area in northwestern Larimer Co. exceeding 8.5". The urban corridor values are generally in the 2.25-3" range, with relatively lower values over eastern Boulder Co.. Overall, these values correlated well with observed values in a general sense, including the magnitudes of the maxima. However, some underprediction of precipitation is noted in the Fort Collins and Golden areas, and along the I-25 zone north of Denver. These issues are under further investigation, including examining the role of the diffusion processes in the Eta results.

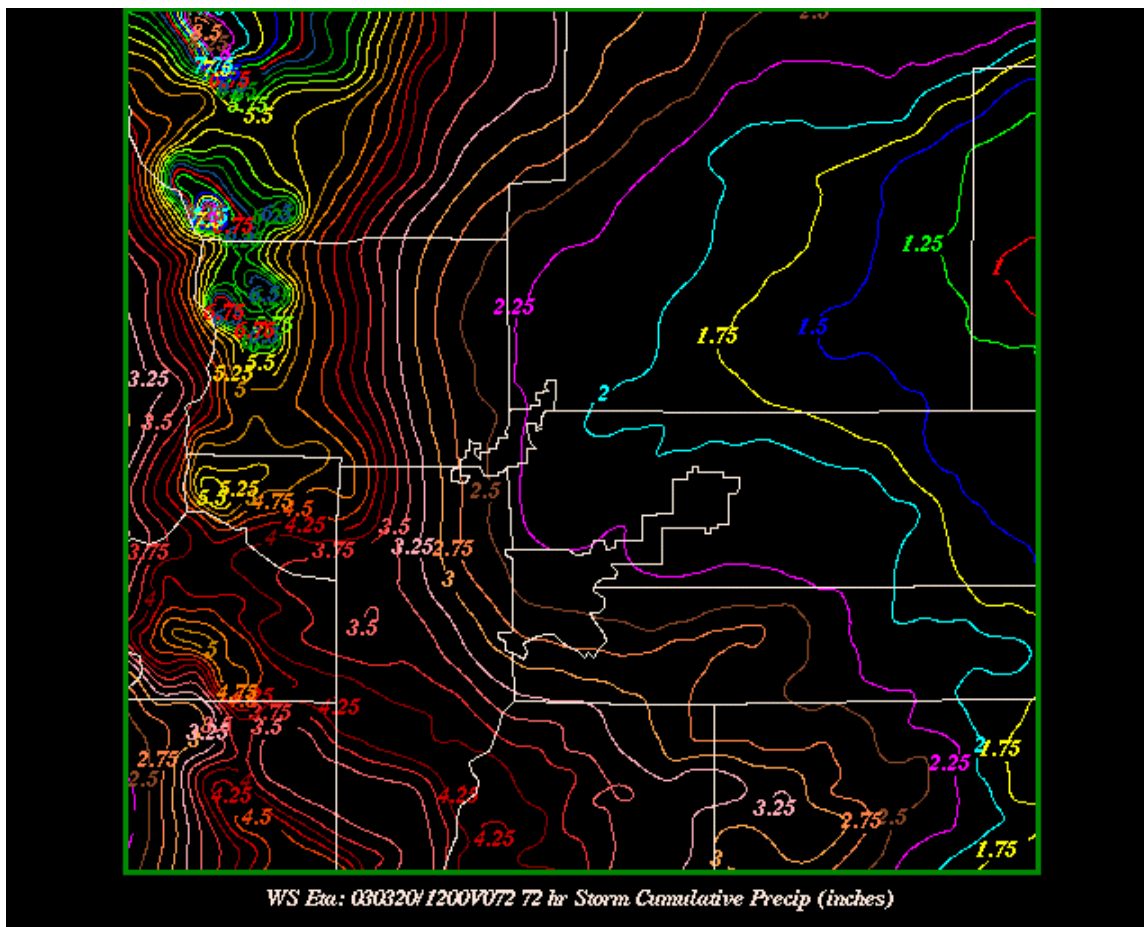


Fig. 7 High-resolution Eta predicted total precipitation (inches) for the period 12 UTC 17 Mar. through 12 UTC 20 Mar.

In regards to the precipitation type and the low-level temperature fields, the workstation Eta forecast even warmer conditions along the urban corridor than the MM5 during the storm (Fig. 8). The precipitation-type forecasts (Fig. 9) which utilize a partial-thickness approach, exhibited liquid precipitation for extreme eastern Larimer and Boulder Counties at 00 UTC 19 Mar. (at this time these areas were receiving the heaviest snowfall of the event), but do predict snowfall in some foothill/plains interface areas that were above freezing in the model through most of the storm. Note in Fig. 9 that the liquid precipitation area that extends westward over northeastern Boulder Co. has some similarity to the observed snowfall minima shown in Fig. 1. In Fig. 8, this tendency for warmer surface conditions is evident in the locations of the 2C and 3C contours over this area, especially in comparison to these locations in other areas within the urban corridor. Further examination of these thermal fields is currently underway.

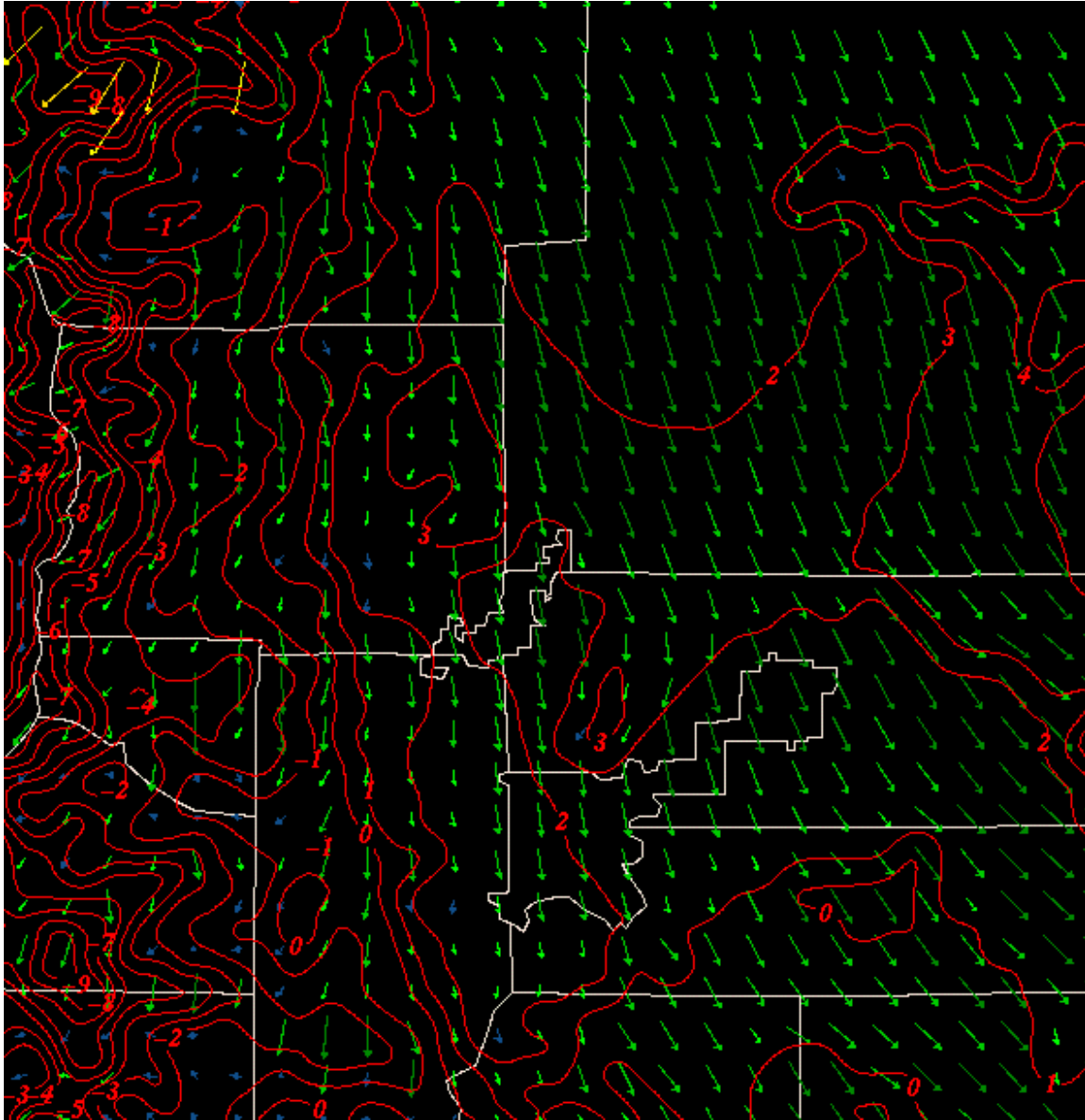


Fig. 8 High-res. Eta-forecast temperatures (C) and winds at 10m, for 00 UTC 19 Mar. The longest vector on the chart corresponds to about 25 knots.

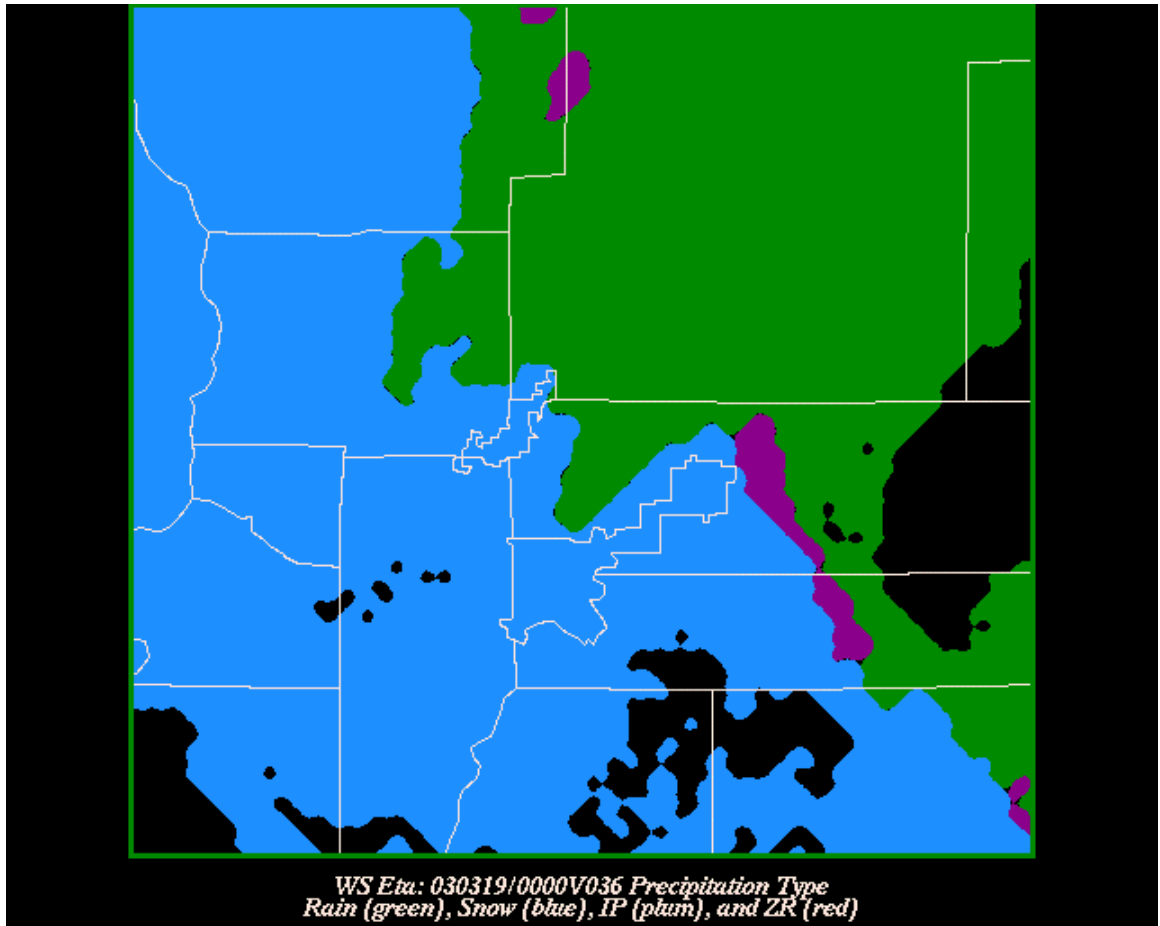


Fig. 9 Eta-forecast precipitation type, for 00 UTC 19 Mar.

The MM5 and Eta models' abilities to capture the depth and strength of the upslope flow are likely critical to the ability to predict the barrier jet regime accurately, and thus the low-level temperatures and precipitation types. This table shows a comparison of observed and predicted vertical wind speed profiles at Platteville, CO (about 25 miles north

of Denver) for the u-component at 06 UTC 19 Mar. (during the height of the storm). The "profiler" column is for the winds measured at the site. A value above 0 indicates a westerly direction.

■ Height (msl)	profiler	MM5	wEta	Eta
■-----				
■2km	+8 knots	-2	+3	~0
■3	-30	-10	-4	-8
■4	-33	-20	-22	-15
■5	-31	-32	-27	-25
■6	-40	-40	-41	-30
■7	-49	-44	-42	-40

Obviously, serious issues exist with the ability of the models to predict the upslope component accurately in the 10-15,000 (MSL) foot layer. Whether this is related to the warm biases is unclear, and at first guess is non-intuitive. Another possibility is inaccurate boundary conditions.

4. SUMMARY

This study has begun to address the applications of very high-resolution mesoscale model forecasts for a major wintertime snow event over the high plains and mountains of central/northern CO. This storm represented a situation where very strong synoptic forcing interacted with major terrain-forced processes to create snow accumulations above 40 inches in some urban areas and above 70 inches in many foothill locations during a 3-4 day period. In this research we have set up the MM5 and "workstation"-Eta models in quasi-forecast mode to investigate small-scale mechanisms for snowfall maxima and minima, precipitation type, and wind variations. Clearly the detailed precipitation and surface wind fields generated by the high-resolution models have produced insight into the physical processes involved, including blocking, melting, and barrier-jet induced uplift. Relatively high accuracy characterizes the total precipitation fields generated by the models. The three-dimensional nature of the barrier jet structure and the temporal dependence of the upslope forcing also represent important aspects of these simulations. The problem associated with the predicted vertical profiles of the upslope flow is under investigation. In addition, though the model forecasts seemed to accurately predict surface temperature gradients, the issue of forecast temperatures being too warm (by both models) in critical areas is also under further investigation. This is also the subject of a companion paper on this storm (Szoke et al., 2004).

5. ACKNOWLEDGEMENTS

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