

BLM AIR RESOURCES
TECHNICAL REPORT FOR OIL
AND GAS DEVELOPMENT IN
NEW MEXICO, OKLAHOMA,
TEXAS, AND KANSAS

2020

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1. INTRODUCTION

The purpose of this document is to present, discuss and summarize technical information on air quality, air quality related values, greenhouse gas (GHG) emissions and climate change relative to air resources with the Bureau of Land Management (BLM) New State Office Planning areas (New Mexico, Oklahoma, Texas, and Kansas). Much of the information contained in this document is directly related to air quality in the context of oil and gas development; other information is generalized air quality that can be applied to other development scenarios and assessments. This information can then be incorporated by reference into the site-specific National Environmental Policy Act (NEPA) documents (environmental assessment [EA] Application for Permit to Drill [APD], etc.) as necessary.

Since the BLM manages extensive land holdings in New Mexico, far more of its activities are centered there. The BLM has jurisdiction over mineral rights on federal lands managed by other agencies and on split estate lands in Kansas, Texas, and Oklahoma. Wherever possible, information for those states is included.

1.1 AIR RESOURCES

Air quality, GHGs, and climate are components of air resources which may be affected by BLM applications, activities, and resource management. Therefore, the BLM must consider and analyze the potential effects of BLM and BLM-authorized activities on air resources as part of the planning and decision-making process. In particular, the activities surrounding oil and gas development are likely to have impacts related to air resources.

1.2 AIR QUALITY

The Clean Air Act (CAA), as amended, is the primary authority for regulation and protection of air quality in the United States. The Federal Land Policy and Management Act also charges BLM with the responsibility to protect air and atmospheric values. Additionally, each state, tribe or local government holds additional authority for regulating air quality within their unique jurisdiction.

1.3 CLASS I, II, AND III AREAS AND THE CLEAN AIR ACT

All areas of the United States not specifically classified as Class I by the CAA are considered to be Class II for air quality. Class I areas are afforded the highest level of protection by the CAA and include all international parks, national wilderness areas and national memorial parks >5,000 acres, and national parks >6,000 acres in size which were in existence on August 7, 1977. Moderate amounts of air quality degradation are allowed in Class II areas. While the CAA allows for designation of Class III areas where greater amounts of degradation would be allowed, no areas have been successful in receiving such designation by the U.S. Environmental Protection Agency (EPA). Air quality in a given area is determined by levels and chemistry of atmospheric pollutants, dispersion meteorology, and terrain.

Regulation and enforcement of the National Ambient Air Quality Standards (NAAQS) has been delegated to the states by the EPA. New Mexico Ambient Air Quality Standards (NMAAQs) are shown in Table 1. Texas has state property line standards for sulfur dioxide (SO₂) and certain non-criteria pollutants. Other than the addition of a 30-minute SO₂ state property line standard which varies based on which county a project is located, there are no other differences between state standards and the NAAQS in Texas.

Oklahoma and Kansas do not have state standards for criteria pollutants that differ from the NAAQS (see Table 1).

The regulatory authority for air quality in Kansas is the Kansas Department of Health and Environment, Bureau of Air (KDHE BoA 2019). The regulatory authority for air quality in Oklahoma is the Oklahoma Department of Environmental Quality, Air Quality Division (ODEQ AQD 2019). The regulatory authority for air quality in Texas is the Texas Commission on Environmental Quality (TCEQ), Air Division (TCEQ 2019a).

2 CRITERIA AIR POLLUTANTS

The EPA has the primary responsibility for regulating atmospheric emissions, including six nationally regulated air pollutants defined in the CAA. These pollutants, referred to as “criteria pollutants,” include carbon monoxide (CO), nitrogen dioxide (NO₂ or NO_x), ozone (O₃), particulate matter (PM₁₀ and PM_{2.5}), SO₂ (SO₂ or SO_x), and lead (Pb). The CAA charges the EPA with establishing and periodically reviewing NAAQS for each criteria pollutant. Table 1 shows the current NAAQS for each pollutant.

Table 1. National Ambient Air Quality Standards

Pollutant	Primary Standards		Secondary Standards		NMAAQS Level (Averaging Time)
	Level	Averaging Time	Level	Averaging Time	
CO	9 parts per million (ppm) (10 milligrams/cubic meter [mg/m ³])	8-hour ⁽¹⁾	None	None	8.7 ppm
	35 ppm (40 mg/m ³)	1-hour ⁽¹⁾	None	None	13.1 ppm
Pb	0.15 micrograms/cubic meter (µg/m ³)	Rolling 3-month average ⁽²⁾	Same as primary	Same as primary	None
NO ₂ (or NO _x)	53 parts per billion (ppb) (100 µg/m ³)	Annual (arithmetic average)	Same as primary		50 ppb
	100 ppb (188 µg/m ³)	1-hour ⁽³⁾	None		100 ppb (24-hour)
PM ₁₀	150 µg/m ³	24-hour ⁽⁴⁾	Same as primary		*
PM _{2.5}	12.0 µg/m ³	Annual ⁽⁵⁾ (arithmetic average)	15.0 ug/m ³	(annual) ⁽⁵⁾ (arithmetic average)	*
	35 µg/m ³	24-hour ⁽⁶⁾	Same as primary		*

Pollutant	Primary Standards		Secondary Standards		NMAAQS Level (Averaging Time)
	Level	Averaging Time	Level	Averaging Time	
O ₃	0.070 ppm (137 µg/m ³)	8-hour ⁽⁷⁾	Same as primary		None
SO ₂ (or SO _x)	75 ppb (196 µg/m ³)	1-hour ⁽⁸⁾	0.5 ppm ⁽¹⁾ (1,300 µg/m ³)	3-hour	0.02 ppm (annual)** 0.10 ppm (24-hour)**

Source: EPA 2021a

*The New Mexico Environmental Improvement Board repealed the Total Suspended Particle NMAAQS in New Mexico Administrative Code (NMAC) 20.2.3, Ambient Air Quality Standards effective November 30, 2018.

** For additional standards of air quality related to sulfur compounds in specific areas such as Chino Mines Company smelter furnace stack at Hurley and the Pecos-Permian basin intrastate air quality control region, see NMAC 20.2.3 and Table 6 of this report.

⁽¹⁾ Not to be exceeded more than once per year.

⁽²⁾ Not to be exceeded.

⁽³⁾ To attain this standard, the 3-year average of the 98th percentile of the daily maximum 1-hour average at each monitor within an area must not exceed 100 ppb (effective January 22, 2010).

⁽⁴⁾ Not to be exceeded more than once per year on average over 3 years.

⁽⁵⁾ To attain this standard, the 3-year average of the weighted annual mean PM_{2.5} concentrations from single or multiple community-oriented monitors must not be exceeded.

⁽⁶⁾ To attain this standard, the 3-year average of the 98th percentile of 24-hour concentrations at each population-oriented monitor within an area must not exceed 35 µg/m³ (effective December 17, 2006).

⁽⁷⁾ To attain this standard, the 3-year average of the fourth-highest daily maximum 8-hour average O₃ concentrations measured at each monitor within an area over each year must not exceed 0.070 ppm.

⁽⁸⁾ To attain this standard, the 3-year average of the 99th percentile of the daily maximum 1-hour average at each monitor within an area must not exceed 75 ppb.

EPA's New Source Performance Standards (NSPS) Rules are designed to regulate criteria air pollutant and O₃ precursor emissions. The EPA NSPS regulations that are most likely to have applicability to oil and gas operations are as follows:

- NSPS Subpart JJJJ – Standards of Performance for Stationary Spark Ignition Internal Combustion Engines
- NSPS 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.
- NSPS OOOO – Standards of Performance for Crude Oil and Natural Gas Production, Transmission and Distribution
- NSPS OOOOa – Crude Oil and Natural Gas Facilities for Which Construction, Modification, or Reconstruction Commenced after September 18, 2015: NSPS – Originally this rule and its draft was promulgated to regulate VOC and GHG emissions (methane) from specific sources within

the oil and natural gas industry which would have included new, modified, and reconstructed compressors, pneumatic controllers, pneumatic pumps, storage vessels, well completions, fugitive emissions from well sites and compressor stations, and equipment leaks at natural gas processing plants. In September 2018 and August 2019, the EPA proposed changes to the rule to modify, amend and/or rescind requirements for the 2012 and 2016 NSPS for the Oil and Gas Industry, which have been incorporated into the final rule as of September 14, 2020.

2.1 OZONE AND VOLATILE ORGANIC COMPOUNDS

The current NAAQS for ozone (O_3) is the 3-year average of the annual fourth-highest daily maximum 8-hour average O_3 concentration, which for simplicity is sometimes referred to as the “design value.” Between 1997 and 2008, the NAAQS for O_3 was 0.080 ppm. To attain this standard, the design value for O_3 at any monitor in the United States could not exceed 0.084 ppm. In 2008, the NAAQS for O_3 was lowered to 0.075 ppm. In 2015, the NAAQS for O_3 was lowered to 0.070 ppm. Ground level O_3 is not emitted directly into the air but is created by chemical reactions between indicators: NO_x (oxides of nitrogen) and volatile organic compounds (VOCs), in the presence of sunlight (Figure 1). While O_3 and NO_2 are criteria air pollutants, VOCs are not. Figure 1 uses a graphical representation to show how O_3 is created in the atmosphere.

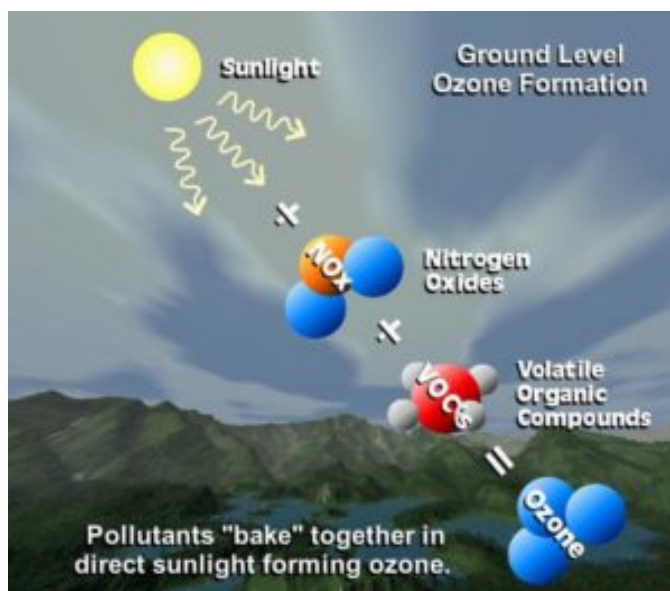


Figure 1. Ozone formation, courtesy of NASA

VOCs are components of natural gas and may be emitted from well drilling, operations and from equipment leaks, valves, pipes, and pneumatic devices. Additionally, VOCs are emitted from a variety of sources, such as refineries, oil and gas production equipment, consumer products and natural (biogenic) sources, such as trees and plants. More information on VOCs during the well completion process is discussed further in the report. NO_x emissions are discussed below under NO_2 . Emissions from industrial facilities and electric utilities, motor vehicle exhaust, gasoline vapors, and chemical solvents are some of the major sources of NO_x and VOCs (EPA 2021b).

O₃ is most likely to reach unhealthy levels on hot sunny days in urban environments but can still reach high levels during colder months. O₃ can also be transported long distances by wind, so even rural areas can experience high O₃ levels (EPA 2021b).

People most at risk from breathing air containing O₃ include people with asthma, children, older adults, and people who are active outdoors, especially outdoor workers. In addition, people with certain genetic characteristics, and people with reduced intake of certain nutrients, such as vitamins C and E, are at greater risk from O₃ exposure (EPA 2021b).

Breathing O₃ can trigger a variety of health problems including chest pain, coughing, throat irritation, and airway inflammation. It also can reduce lung function and harm lung tissue. O₃ can worsen bronchitis, emphysema, and asthma, leading to increased medical care. The environmental effects of O₃ include damaging sensitive vegetation and ecosystems, including forests, parks, wildlife refuges and wilderness areas. In particular, O₃ harms sensitive vegetation during the growing season (EPA 2021b).

2.1.1 OZONE TRENDS

Nationally, O₃ concentrations at urban and rural sites have decreased 33% from 1980 to 2020. Over the same period, emissions of the ozone precursors VOC and NO_x have decreased by 60% and 70%, respectively. Weather conditions have a significant role in the formation of O₃. O₃ is most readily formed on warm summer days when there is stagnation. Table 2 shows the ozone trends in specific cities within the BLM New Mexico State Office’s area of operations.

Table 2. Local Ozone Trends

City, ST	2000 Design value (ppm)	2010 Design Value (ppm)	2020 Design Value (ppm)	2000-2020 Trend*
Carlsbad-Artesia, NM	0.069	0.069	0.073	6%
Farmington, NM	0.08	0.064	0.066	-18%
Tulsa, OK	0.08	0.07	0.059	-26%
Oklahoma City, OK	0.081	0.071	0.066	-19%
Wichita, KS	0.081	0.075	0.059	-27%
Houston-The Woodlands-Sugar Land, TX	0.10	0.079	0.068	-32%
Longview, TX	0.099	0.078	0.055	-44%
Dallas-Fort Worth-Arlington, TX	0.09	0.076	0.069	-23%
Austin-Round Rock, TX	0.088	0.072	0.064	-27%
El Paso, TX	0.078	0.068	0.072	-8%

Source: EPA 2021c

* A positive percentage means that O₃ concentrations have increased from 2000 to 2020 while a negative percentage means that O₃ concentrations have decreased from 2000 to 2020.

NITROGEN DIOXIDE

Nitrogen dioxide (NO₂) is both a criteria pollutant and an indicator for the NO_x family of nitrogen oxide compounds that are ground-level O₃ precursors. The nitrogen oxide family of compounds includes nitric oxide (NO), NO₂, nitrous acid (HNO₂), and nitric acid (HNO₃). The primary sources of NO_x nationally are motor vehicles and fuel combustion. The excess air required for complete combustion of fuels in these processes introduces atmospheric nitrogen into the combustion reactions at high temperatures and produces nitrogen oxides. NO₂ has been shown to cause adverse respiratory impacts in both healthy people and those with asthma and is also an important contributor to the formation of ground-level O₃ (EPA 2019a).

Nationally, NO₂ concentrations have decreased substantially (61% reduction) from 1980 to 2018. In the southwest (Arizona, New Mexico, Colorado, and Utah), NO₂ concentrations have decreased 35% between 2000 and 2018; in the south (Texas, Oklahoma, Kansas, Arkansas, Louisiana, and Mississippi), NO₂ concentrations have decreased 31% between 2000 and 2018. EPA expects NO₂ concentrations will continue to decrease (EPA 2021d). EPA's national and regional rules to reduce emissions of NO₂ and NO_x will help state and local governments meet the National Ambient Air Quality Standard (NAAQS).

2.2 CARBON MONOXIDE

Carbon monoxide (CO) is produced from the incomplete burning of carbon-containing compounds such as fossil fuels; it forms when there is not enough oxygen to produce carbon dioxide (CO₂). Nationally, 86% of CO emissions come from transportation sources. CO is associated with negative health effects to human cardiovascular, central nervous, and respiratory systems (EPA 2019b).

Nationally, CO concentrations have decreased 83% from 1980 to 2018. Monitored CO concentrations in the "southwest" region (New Mexico, Arizona, Colorado, and Utah) have decreased 62% between 2000 and 2018. Monitored CO concentrations in the "south" region (Texas, Oklahoma, Kansas, Arkansas, Louisiana, and Mississippi) have decreased 69% between 2000 and 2018 (EPA 2021e).

2.3 PARTICULATE MATTER

Particulate matter, also known as particle pollution or PM, is a complex mixture of extremely small particles and liquid droplets. PM is made up of a number of components, including acids (such as nitrates and sulfates), organic chemicals, metals, and soil or dust particles. PM is measured and regulated according to particle size. PM₁₀ refers to all particles with a diameter of 10 microns or less. PM_{2.5} is made up of particles with diameters of 2.5 microns or less. Smaller particles are associated with more negative health effects, including respiratory and cardiovascular problems because they can become more deeply embedded in the lungs (EPA 2021f).

Nationally, PM_{2.5} concentrations have decreased 39% from 2000 to 2018. In that same time period, PM₁₀ concentrations decreased 31% nationally. In the Four Corners region (New Mexico, Arizona, Colorado, and Utah), PM_{2.5} concentrations have decreased 23% from 2000 to 2018, and PM₁₀ concentrations have decreased 28% in the same time period. For the southern region encompassing Texas, Oklahoma, Kansas, Arkansas, Louisiana, and Mississippi, PM_{2.5} concentrations have decreased 30% and PM₁₀ concentrations have increased 18% between 2000 and 2018 (EPA 2021g, 2021h).

2.4 SULFUR DIOXIDE

Sulfur dioxide (SO₂) is one of a group of highly reactive gases known as “oxides of sulfur,” commonly referred to as SO_x. The largest sources of SO₂ emissions nationwide are from fossil fuel combustion at power plants (73%) and other industrial facilities (20%). Smaller sources of SO₂ emissions include industrial processes, such as extracting metal from ore, and the burning of high sulfur-containing fuels by locomotives, large ships, and non-road equipment. SO₂ is linked with a number of adverse effects on the respiratory system (EPA 2019c).

Nationally, SO₂ concentrations have decreased 80% from 2000 to 2018, but substantial decreases (91% reduction) have occurred since 1980 due to implementation of federal rules requiring reductions in SO₂ emissions from power plants and other large sources of SO₂. In the Four Corners region, SO₂ concentrations decreased 23% between 2000 and 2018. In the southern region of Texas, Oklahoma, Kansas, Arkansas, Louisiana, and Mississippi, SO₂ concentrations decreased 76% between 2000 and 2018 (EPA 2021i).

2.5 LEAD

With the elimination of lead from gasoline and regulation of industrial sources, levels of lead in the atmosphere decreased 98% nationwide between 1980 and 2018. Lead concentrations decreased 93% nationally between 2000 and 2018. While still regulated as a criteria pollutant, the major sources of lead pollution are lead smelters and leaded aviation gasoline. In 2014, EPA proposed to retain the NAAQS for lead without revision (EPA 2020a).

2.6 MONITORING DATA AND DESIGN VALUES

Criteria pollutants are monitored throughout various parts of the country. Monitors measure concentrations of pollutant in the atmosphere and the results are often presented in ppm or µg/m³. EPA and states periodically analyze and review monitor locations, discontinuing monitoring at locations where pollutant concentrations have been well below the standards and adding monitors in areas where pollutant concentrations may be approaching air quality standards. *Instantaneous on-demand* monitored outdoor air quality data collected from state, local and tribal monitoring agencies can be obtained from EPA’s Air Data webpage and interactive tool (EPA 2021j).

Another type of monitoring data is *annual average concentration(s)* measured at air monitors which are then translated to annual design values to be consistent with the individual NAAQS in Table 1. A design value is a statistic that describes the air quality status of a given location relative to the level of the NAAQS. Design Values are normally updated annually and posted to the EPA’s Air Quality Design Value website. The most recent, 2020 Design Values for the measured criteria pollutants of the counties in the major oil and gas (O&G) basins of New Mexico are provided in Table 3 and Table 4.

Table 3. 2020 Design Values, Eddy and Lea Counties

Pollutant	2020 Design values	Averaging Time	NAAQS	NMAAQs ⁽⁵⁾
O ₃	0.078 ppm (Eddy County), 0.068 ppm (Lea County)	8-hour ⁽¹⁾	0.070 ppm	–

Pollutant	2020 Design values	Averaging Time	NAAQS	NMAAQS ⁽⁵⁾
NO ₂	5 ppb (Eddy County), 5 ppb (Lea County)	Annual ⁽²⁾	53 ppb	50 ppb
NO ₂	29 ppb (Eddy County), 35 ppb (Lea County)	1-hour ⁽³⁾	100 ppb	–
PM _{2.5} ⁴	7.0 micrograms per cubic meter (µg/m ³) (Lea County)	Annual ⁽⁴⁾	12 µg/m ³	–
PM _{2.5}	17 µg/m ³ (Lea County)	24-hour ⁽³⁾	35 µg/m ³	–

Source: EPA 2021k

⁽¹⁾ Annual fourth-highest daily maximum 8-hour concentration, averaged over 3 years.

⁽²⁾ Not to be exceeded during the year.

⁽³⁾ 98th percentile, averaged over 3 years.

⁽⁴⁾ Annual mean, averaged over 3 years.

⁽⁵⁾ The NMAAQS standard for total suspended particulates, which was used as a comparison for PM₁₀ and PM_{2.5}, was repealed as of November 30, 2018.

Note: While there are no NAAQS for hydrogen sulfide (H₂S), New Mexico has set 1/2-hour standards for H₂S at 0.100 ppm within Pecos-Permian AQ Control Region and 0.030 ppm, for municipal boundaries and within 5 miles of municipalities with populations greater than 20,000 in areas of the state outside of the area within 5 miles of the Pecos-Permian AQ Control Region (see Table 6).

Table 4. 2020 Design Values for Rio Arriba, Sandoval, and San Juan Counties

Pollutant	2020 Design Concentrations	Averaging Time	NAAQS	NMAAQS ⁽⁷⁾
O ₃	Rio Arriba County: 0.065 ppm Sandoval County: 0.070 ppm San Juan County: 0.069 ppm, four stations; Bloomfield at 0.066 ppm, Navajo Dam at 0.068 ppm, Shiprock at 0.069 ppm, Chaco Culture NHP 0.067 ppm	8-hour ⁽¹⁾	0.070 ppm	–
NO ₂	San Juan County: four stations; Bloomfield at 10 ppb, Navajo Dam at 6 ppb, Chaco Culture at 1 ppb and Shiprock at 3 ppb	Annual ⁽²⁾	53 ppb	50 ppb
NO ₂	San Juan County; 33 ppb, four stations; Bloomfield at 33 ppb, Navajo Dam at 23 ppb, Chaco Culture invalid, Shiprock at 24 ppb	1-hour ⁽³⁾	100 ppb	–
SO ₂	San Juan County: 1 ppb	1-hour ⁽⁵⁾	75 ppb	–
PM _{2.5}	San Juan County: No monitor data nearby	Annual ^(4,6)	60 µg/m ³	–
PM _{2.5}	San Juan County: No monitor data nearby	24-hour ^(3,6)	35 µg/m ³	–

Source: EPA 2021k

⁽¹⁾ Annual fourth highest daily maximum 8-hour concentration, averaged over 3 years.

⁽²⁾ Not to be exceeded during the year.

⁽³⁾ 98th percentile, averaged over 3 years.

⁽⁴⁾ Annual mean, average over 3 years.

⁽⁵⁾ 99th percentile of 1-hour daily maximum concentrations, averaged over 3 years.

⁽⁶⁾ PM_{2.5} monitor stations currently show installed locations in the planning area (San Juan County), however the monitor status of these stations show invalid data and cannot be used to represent design values.

⁽⁷⁾ The NMAAQs standard for total suspended particulates, which was used as a comparison with PM₁₀ and PM_{2.5}, was repealed as of November 30, 2018.

Note: While there are no NAAQS for H₂S, New Mexico has set a 1-hour standard for H₂S at 0.010 ppm for all areas of the state outside of the area within 5 miles of the Pecos-Permian AQ Control Region.

2.7 GENERAL CONFORMITY AND NONATTAINMENT

If the concentration of one or more criteria pollutants in a geographic area is found to exceed the regulated or 'threshold' level for one or more of the NAAQS, the area may be classified as a **nonattainment** area. Areas with concentrations of criteria pollutants that are below the levels established by the NAAQS are considered either **attainment** or unclassifiable areas.

To eliminate or reduce the severity and number of NAAQS violations in nonattainment areas, and to achieve expeditious attainment of the NAAQS, the EPA promulgated the Conformity Rule (40 Code of Federal Regulations [CFR] 6, 51, 93). The Conformity Rule applies to federal actions and environmental analyses in nonattainment and maintenance areas completed after March 15, 1994. This rule contains a variety of substantive and procedural requirements to show conformance with both the NAAQS and state implementation plans (SIPs).

Section 176(c) of the CAA prohibits federal agencies from taking actions in nonattainment and maintenance areas unless the emissions from the actions conform to the SIP or tribal implementation plan for the area. Federal actions must be evaluated for conformity to the local SIP if the project 1) is located within an EPA-designated nonattainment or maintenance area, 2) would result in emissions above the de minimis threshold quantities of criteria pollutants listed in 40 CFR Part 93, 3) is not a listed exempt action, and 4) has not been accounted for in an EPA-approved SIP.

EPA's conformity rule requires that all federal actions in a nonattainment area must demonstrate conformity with the SIP for the pollutant in question. If the agency can demonstrate that emissions for the action will fall below certain established levels, known as de minimis, then no further analysis is necessary. To establish de minimis, an emissions inventory for the project is required. In the case of O₃, the emissions inventory would include NO_x and VOCs. If emissions are projected to be above de minimis levels further analysis should be coordinated with the EPA and/or state agency.

Nonattainment areas in New Mexico are as follows:

- **O₃ Nonattainment Area in Dona Ana County (Sunland Park, New Mexico, located southwest of the Carlsbad Field Office [CFO] planning area, south of Las Cruces)** In 1995, the EPA declared a 42 square-mile region in the southeast corner of the County on the border of Texas and Mexico as a marginal nonattainment area for the 1-hour O₃ standard. The nonattainment area included the City of Sunland Park, Santa Teresa, and La Union, New Mexico. The 1-hour O₃ standard was revoked by EPA in 2004 with the adoption of the new 8-hour O₃ standard. Due to the revocation of the 1979 1-hour O₃ standard and based on monitoring data, Sunland Park was designated attainment for the 1997 8-hour O₃ standard (0.080 ppm).

In October 2015, the EPA again lowered the NAAQS for O₃ from 0.075 ppm to 0.070 ppm. Due to the lowering of the federal standard, the State of New Mexico recommended that a portion of Sunland Park, New Mexico be designated as nonattainment of the new 8-hour O₃ standard. On June 4, 2018, the EPA designated this area as nonattainment with an effective date of August 3, 2018 (EPA 2018).

The New Mexico Environment Department (NMED) developed new planning elements required for a marginal nonattainment classification in the Ozone NAAQS SIP revision, which was finalized on September 11, 2020. This document outlines the strategies and emissions control measures that are expected to improve air quality in the Sunland Park area that has been designated nonattainment. These strategies and emissions control measures aim to reduce the amount of nitrogen oxides and VOCs emitted to the atmosphere. They rely on current or upcoming federal rules, new or revised state rules, and other programs, such as the New Mexico Volkswagen mitigation plan projects (NMED 2020).

NMED submitted to the EPA the Baseline Emissions Inventory and Emissions Statement, which meets the Clean Air Act (CAA) section 182(a) SIP requirements for a marginal nonattainment area. In addition, the department must review its nonattainment permitting rules (NNSR) and adopt revisions, if required, to comply with federal law by August 3, 2021. The NNSR require revisions to New Mexico SIP which has undergone public review and will be subject to hearing requirements (NMED 2019a).

- **O₃ Design Value Exceedance in Eddy County (Carlsbad, New Mexico)** In May 2021 new design values for NAAQS were published by the EPA for various counties throughout the United States. The monitor at 2811 Holland St in Eddy County, show an 8-hour O₃ exceedance, 78 ppb (EPA 2021k).

NMED is required by State Statute to plan for O₃ mitigation in areas where monitors indicate O₃ levels within 95% of the O₃ standard. The area discussed above in Carlsbad, New Mexico, has not been formally declared nonattainment by the EPA through the state's recommendation.

The Ozone Attainment Initiative is a project authorized by State Statute, 74-2-5.3 New Mexico Statutes Annotated 1978. This statute directs NMED to develop plans that may include regulations more stringent than Federal rules for areas of the state in which ambient monitoring shows O₃ levels at or above 95% of the NAAQS. The 2015 8-hr primary NAAQS for O₃ is 0.070 ppm (or 70 ppb). Ninety-five percent (95%) of the O₃ NAAQS is 0.067 ppm (67 ppb). This form of the standard requires averaging of 3 years of monitoring data for the fourth highest 8-hour average, using the most recent year's data to determine the "design value". For New Mexico, seven counties show 3-year averages (2018–2020) of O₃ levels at or above 95% of the NAAQS:

- Doña Ana County (78 ppb)
 - Eddy County (78 ppb)
 - Lea County (68 ppb)
 - Rio Arriba County (65 ppb)
 - Sandoval County (70 ppb)
 - San Juan County (69 ppb)
 - Valencia County (69 ppb)
(EPA 2021k)
- **PM₁₀ Nonattainment Area in Anthony, New Mexico (located west of the CFO planning area, south of Las Cruces)** The State of New Mexico submitted the Anthony PM₁₀ SIP to the regional EPA headquarters on November 8, 1991. This area was designated nonattainment for PM₁₀ by the EPA in 1991. The nonattainment area is bounded by Anthony Quadrangle, Anthony, New Mexico-Texas. SE/4 La Mesa 15-minute Quadrangle, N32 00 - W106 30/7.5, Sections 35 and 36, Township 26 South, Range 3 East as limited by the New Mexico–Texas state line on the

south. The site is located in Doña Ana County, which submitted a Natural Events Action Plan for PM₁₀ exceedances to the EPA in December 2000 (NMED 2019b).

The NMED Air Quality Bureau developed a fugitive dust rule in conjunction with the mitigation plan to detail mandatory measures to abate certain controllable sources in Doña Ana and Luna Counties. A public hearing was held on September 28, 2018, in Las Cruces and the board adopted the rule on October 26, 2018. Mitigation plans are required by the EPA in areas where recurring natural events (in this case, high winds resulting in blowing dust) cause exceedances of the health based national standards for PM. Some of the required elements of a mitigation plan include:

- Steps to identify and study sources of dust;
 - Mandatory or voluntary control measures to abate sources of dust that cause or contribute to exceedances of the standards to better protect public health;
 - Public education and notification programs aimed at reducing individuals' exposure to unhealthy levels of PM in the air before, during, and after high wind events; and
 - Public review and periodic evaluation of the mitigation plan (NMED 2019c)
- **SO₂ Maintenance Area in Grant County (located west of the CFO planning area, at the Arizona border):** This maintenance area is located at the Phelps Dodge Chino Copper Smelter in Grant County. The maintenance area is defined as a 3.5-mile-radius region around the smelter. The maintenance area also includes high elevation areas within an 8-mile radius. The state submitted a SIP to the regional EPA headquarters in August 1978. The New Mexico Air Quality Bureau submitted a re-designation plan to the EPA in February 2003. The re-designation plan was approved by the EPA in September 2003 (NMED 2019c).
 - **TEXAS NONATTAINMENT AREAS:** There are currently 10 key nonattainment areas in Texas, one for PM₁₀ (city of El Paso), three for O₃ (Houston-Galveston-Brazoria area [eight counties], Dallas-Fort Worth area [10 counties], and Bexar County in San Antonio) and several areas not meeting the SO₂ 2010 standard and therefore designated in part as nonattainment (Freestone and Anderson Counties, Howard County, Hutchinson County, Navarro County, Rusk and Panola Counties, and Titus County) (EPA 2021).

The Houston-Galveston-Brazoria area has several counties in nonattainment status. The following counties are currently not meeting the 8-hour 2015 O₃ standard of 0.070 ppm (70 ppb): Brazoria, Chambers, Fort Bend, Galveston, Harris, and Montgomery. Each of these counties have been designated as marginal nonattainment for the 2015 O₃ nonattainment. Additionally, the six counties listed above as nonattainment with the 2015 O₃ standard are also in nonattainment with the 8-hour 2008 O₃ standard along with Liberty County, and Waller County. Each of these counties have been designated as serious nonattainment for the 2008 O₃ nonattainment. De minimis values for areas designated as serious for both NO_x and VOC are 50 tons/year (EPA 2021). The Dallas-Fort Worth area also has several counties in nonattainment status. The following counties are currently not meeting the 8-hour 2015 O₃ standard of 0.070 ppm (70 ppb): Collin, Dallas, Denton, Ellis, Johnson, Kaufman, Parker, Tarrant, and Wise) The designation of severity of the 2015 O₃ nonattainment for these counties is Marginal. Additionally, the following counties are nonattainment with respect to the 2008 8-hour O₃ standard: Collin, Dallas, Denton, Ellis, Johnson, Parker, Rockwall, Tarrant, and Wise. The designation of severity of the 2008 O₃ nonattainment for these counties is Serious. De minimis values for a Serious designation in this area is 50 tons/year for NO_x and VOC (EPA 2021).

KANSAS and OKLAHOMA

There are currently no nonattainment areas for any criteria pollutant in the states of Kansas and Oklahoma.

3 NATIONAL EMISSIONS INVENTORY DATA

The National Emissions Inventory (NEI) data presents the emissions of each criteria pollutant by national, state, county, and tribal areas for major source sectors. National emissions trends are reported in the 2017 NEI Report (EPA 2020b). The NEI data is updated every 3 years with new emission inventory data incurring a 2- to 3-year data gathering period for final use. The most recent NEI Inventory is for 2017 and the complete 2020 NEI data is expected to publicly release in the spring of 2023. Emissions data is expressed in tons per year (tpy) or total volume of pollutant released to the atmosphere. Emissions data is useful in comparing source categories to determine which industries or practices are contributing the most to the general level of pollution in an area.

Details of the anthropogenic sectors mentioned in the report are:

- (1) Electricity generation is fuel combustion from electric utilities;
- (2) Fossil fuel combustion is fuel combustion from industrial boilers, internal combustion engines, and commercial/institutional or residential use;
- (3) Industrial processes include manufacturing of chemicals, metals, and electronics, storage and transfer operations, pulp and paper production, cement manufacturing, petroleum refineries, and oil and gas production;
- (4) On-road vehicles category includes both gasoline- and diesel-powered vehicles for on-road use
- (5) Non-road equipment includes gasoline- and diesel-powered equipment for non-road use, as well as planes, trains, and ships;
- (6) Road dust includes dust from both paved and unpaved roads. Presentation of emissions data by source sector provides a better understanding of the activities that contribute to criteria pollutant emissions.

NEI data, by pollutant (CO, NO_x, PM₁₀ and PM_{2.5}, SO₂, and VOCs), for the Major Sources within New Mexico, Kansas, Oklahoma, and Texas can be found in Appendix A. Maps displaying these major sources are including in Appendix D.

3.1 2017 NATIONAL EMISSIONS INVENTORY DATA

3.1.1 VOLATILE ORGANIC COMPOUND EMISSIONS

The 2017 NEI for the entire state of New Mexico estimates VOC emissions from biogenic sources account for 63.8% of total VOCs in the state. VOCs from fire account for 7.1% and industrial processes account for 20.6% of total VOCs. According to the 2017 NEI, Texas biogenic emissions account for 55.0% of total VOC emissions, while industrial processes account for 26.0% of VOC emissions. VOC emissions from fire account for 5.8% of Texas VOC emissions, mobile sources account for 4.9% of total VOC emissions, and solvents account for 5.6% of total VOC emissions. In Oklahoma, biogenic emissions are estimated to be 44.5% of total VOC emissions, industrial processes account for 18.6%, fire accounts for

23.4% of total VOC emissions, and mobile sources account for 5.0% of total VOC emissions. In Kansas, biogenics account for 31.5% of total VOC emissions, fire accounts for 39.1% of VOC emissions, industrial processes account for 14.5% of VOC emissions, mobile sources account for 4.6% of VOC emissions, and solvents account for 6.5% of VOC emissions (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the San Juan Basin (San Juan, Rio Arriba, Sandoval, and McKinley Counties) indicates that biogenics, industrial processes, and fires account for 39.5%, 40.55.1%, and 12.9% of total VOC emissions, respectively, in the area (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the Permian Basin (Eddy, Lea, Chaves, and Roosevelt Counties) indicates that biogenics, industrial processes, and fires account for 45.9%, 48.3%, and 1.4% of total VOC emissions, respectively, in the area (EPA 2020b).

3.1.2 NITROGEN OXIDES EMISSIONS

The 2017 NEI for the state of New Mexico indicate mobile sources account for 37.4%, fuel combustion accounts for 22.5%, industrial processes account for 19.5%, biogenics account for 19.2%, and fire accounts for 1.4% of total NO_x emissions in the state. For Texas, the 2017 NEI data indicate mobile sources account for 40%, fuel combustion accounts for 18.6%, industrial processes account for 24.0%, biogenics account for 15.5%, and fire accounts for 1.4% of total NO_x emissions in the state. In Oklahoma, mobile sources account for 31.5%, fuel combustion accounts for 27.4%, industrial processes account for 19.2%, biogenics account for 15.1%, and fire accounts for 6.4% of total Oklahoma NO_x emissions. In Kansas, mobile sources account for 33.5%, fuel combustion accounts for 16.1%, industrial processes account for 19.6%, biogenics account for 20.2%, and fire accounts for 10.4% of total Kansas NO_x emissions (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the San Juan Basin (San Juan, Rio Arriba, Sandoval, and McKinley Counties) indicates that fuel combustion, industrial processes, and mobile sources account for 38.3%, 34.2%, and 18.0% of total NO_x emissions, respectively, in the area (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the Permian Basin (Eddy, Lea, Chaves, and Roosevelt Counties) indicates that fuel combustion, industrial processes, and mobile sources account for 28.5%, 39.342.6%, and 17.35.9% of total NO_x emissions, respectively, in the area (EPA 2020b).

3.1.3 CARBON MONOXIDE EMISSIONS

For New Mexico, the 2017 NEI data indicate mobile sources account for 38.2%, industrial processes account for 7.5%, fuel combustion accounts for 6.3%, fire accounts for 31.6%, and biogenics account for 15.5% of total state CO emissions. The 2017 NEI data for Texas indicate mobile sources account for 53.1%, fuel combustion accounts for 6.0%, industrial processes account for 5.8%, fire accounts for 22.1%, and biogenics account for 10.2% of total Texas CO emissions. In Oklahoma, mobile sources account for 30.5%, fuel combustion accounts for 4.9%, industrial processes account for 3.3%, fire accounts for 55.5%, and biogenics account for 4.7% of total CO emissions in the state. In Kansas, mobile sources account for 20.8%, fuel combustion accounts for 2.6%, industrial processes account for 4.8%, fire accounts for 66.9%, and biogenics account for 4.3% of total CO emissions in the state (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the San Juan Basin (San Juan, Rio Arriba, Sandoval, and McKinley Counties) indicates that fires, industrial processes, and mobile sources account for 46.3%, 16.5%, and 21.2% of total CO emissions, respectively, in the area (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the Permian Basin (Eddy, Lea, Chaves, and Roosevelt Counties) indicates that fires, industrial processes, and mobile sources account for 12.8%, 20.3%, and 33.9% of total CO emissions, respectively, in the area (EPA 2020b).

3.1.4 PARTICULATE MATTER 2.5 AND 10

According to the 2017 NEI, for the entire state of New Mexico, dust accounts for 33.4% and fire accounts for 35.5% of PM_{2.5} emissions statewide. For PM₁₀, dust accounts for 69.8% and fire accounts for 10.1% of emissions in NM. For Texas, the 2017 NEI data indicate that dust accounts for 26.2%, fire accounts for 24.2%, agriculture accounts for 24.2%, mobile sources account for 5.4%, fuel combustion accounts for 6.7% and industrial processes accounts for 6.4% of statewide PM_{2.5} emissions. Dust accounts for 53.7%, agriculture accounts for 29.3% and fire accounts for 7.0% of Texas PM₁₀ emissions. In Oklahoma, fire accounts for 48.1%, dust accounts for 19.2%, agriculture accounts for 19.3%, fuel combustion accounts for 5.7%, industrial processes account for 2.9% and mobile sources account for 2.2% of Oklahoma PM_{2.5} emissions. Dust accounts for 50.7%, agriculture accounts for 27.6% and fire accounts for 16.3% of Oklahoma PM₁₀ emissions. In Kansas, dust accounts for 19.1%, agriculture accounts for 18.8%, fire accounts for 54.1%, mobile sources account for 2.1%, industrial sources account for 2% and fuel combustion accounts for 2.3% of Kansas PM_{2.5} emissions. Agriculture accounts for 26.7%, dust accounts for 52.1% and fire accounts for 18.3% of Kansas PM₁₀ emissions (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the San Juan Basin (San Juan, Rio Arriba, Sandoval, and McKinley Counties) indicates that dust, fires, and fuel combustion account for 66.0%, 17.5%, and 6.0% of total PM₁₀ emissions, respectively, in the area (EPA 2020b). Dust, fires, and fuel combustion account for 24.8%, 48.2%, and 16.2%, respectively, of total PM_{2.5} emissions.

The 2017 NEI data for the BLM New Mexico portion of the Permian Basin (Eddy, Lea, Chaves, and Roosevelt Counties) indicates that dust, industrial processes, and agriculture account for 70.0%, 10.3%, and 13.9% of total PM₁₀ emissions, respectively, in the area (EPA 2020b). Dust, fires, and industrial processes account for 45.1%, 12.1%, and 15.93.5%, respectively, of total PM_{2.5} emissions.

3.1.5 SULFUR DIOXIDE EMISSIONS

The 2017 NEI data indicate industrial sources account for 46.7%, fuel combustion accounts for 40.5%, fires account for 10.1%, and mobile sources account for 2.4% of New Mexico SO₂ emissions. Fuel combustion accounts for 72.9% and industrial processes account for 23.0% of Texas SO₂ emissions. Fuel combustion accounts for 57.0%, industrial processes account for 29.8% and fire accounts for 12.0% of Oklahoma SO₂ emissions. Fuel combustion accounts for 32.0%, fire accounts for 51.9% and industrial processes account for 13.4% of Kansas SO₂ emissions (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the San Juan Basin (San Juan, Rio Arriba, Sandoval, and McKinley Counties) indicates that fuel combustion, industrial processes, and fires account 84.0%, 5.3%, and 9.7% of total SO₂ emissions, respectively, in the area (EPA 2020b).

The 2017 NEI data for the BLM New Mexico portion of the Permian Basin (Eddy, Lea, Chaves, and Roosevelt Counties) indicates that fuel combustion, industrial processes, and fires account for 4.0%, 94.3%, and 1.0% of total SO₂ emissions, respectively, in the area (EPA 2020b).

3.1.6 LEAD EMISSIONS

According to the 2017 NEI, aircraft account for 89.9% of the lead emissions in New Mexico. In Texas, 76.9% of the lead emissions are from aircraft. In Oklahoma, 55.2% of lead emissions are from aircraft and 14.3% are from waste disposal. In Kansas, 89.7% of lead emissions are from aircraft (EPA 2020b).

4 HAZARDOUS AIR POLLUTANTS

Currently there are 187 specific pollutants and chemical groups known as hazardous air pollutants (HAPs). The list has been modified over time. HAPs are chemicals or compounds that are known or suspected to cause cancer or other serious health effects, such as compromises to immune and reproductive systems, birth defects, developmental disorders, or adverse environmental effects and may result from either chronic (long-term) and/or acute (short-term) exposure. CAA Sections 111 and 112 establish mechanisms for controlling HAPs from stationary sources, and the EPA is required to control emissions of the 187 HAPs. The U.S. Congress amended the Federal CAA in 1990 to address a large number of air pollutants that are known to cause or may reasonably be anticipated to cause adverse effects to human health or adverse environmental effects.

Ambient air quality standards do not exist for HAPs, however the CAA requires control measures for HAPs. Mass-based emissions limits and risk-based exposure thresholds have been established as significance criteria to require maximum achievable control technologies (MACT) under the EPA promulgated National Emissions Standards for Hazardous Air Pollutants (NESHAPs) for 96 industrial source classes. NESHAPs are issued by EPA to limit the release of specified HAPs from specific industrial sectors. These standards are technology based, meaning that they represent the MACT that are economically feasible for an industrial sector.

NESHAPs for Oil and Natural Gas Production and Natural Gas Transmission and Storage were published by EPA on June 17, 1999. These NESHAPs were directed toward major sources and intended to control benzene, toluene, ethyl benzene, mixed xylenes (BTEX) and n-hexane. An additional NESHAP for Oil and Natural Gas Production Facilities directed toward area sources was published on January 3, 2007, and specifically addresses benzene emissions from triethylene glycol dehydrations units. The EPA issued a final rule revising the NESHAP rule effective October 15, 2012. The final rule includes revisions to the existing leak detection and repair requirements and established emission limits reflecting maximum achievable control technology for currently uncontrolled emission sources in oil and gas production and natural gas transmission and storage (*Federal Register* 77(159):49490–49600).

The EPA NESHAPs that are most likely to have applicability to oil and gas operations are as follows (in addition to the NESHAPs common/general provisions):

- NESHAP Subpart HH - National Emission Standard for Hazardous Air Pollutants from Oil and Natural Gas Production Facilities
- NESHAP Subpart ZZZZ - National Emission Standard for Hazardous Air Pollutants for Stationary Reciprocating Internal Combustion Engines

Note that several of the NSPS that are potentially applicable to oil and gas operations (listed in Section 2) also regulate emissions of VOCs, a component of which include HAP emissions. While the NSPS is not designed to directly regulate HAP emissions, control of VOCs results in the co-benefit of HAP reductions.

The CAA defines a major source for HAPs to be one emitting 10 tpy of any single HAP or 25 tpy of any combination of HAPs. Under state regulations, a construction or operating permit may be required for any major source though some exceptions apply. In New Mexico, these regulations are 20.2.70 and 20.2.73 New Mexico Administrative Code (NMAC), in Texas the regulation is 30 Texas Administrative Code (TAC) 122, in Kansas the regulation is Kansas Administrative Regulations (KAR) 28-19-500, and in Oklahoma the regulation is Oklahoma Register 252-100-7. Within its definition of a major source in the above referenced regulations the state of New Mexico includes the following language:

...hazardous emissions from any oil or gas exploration or production well (with its associated equipment) and hazardous emissions from any pipeline compressor or pump station shall not be aggregated with hazardous emissions from other similar units, whether or not such units are in a contiguous area or under common control, to determine whether such units or stations are major sources...

In other words, in determining a major source, each oil and gas exploration and production well must be considered singularly. Kansas, Texas, and Oklahoma regulations include similar language.

The state of New Mexico incorporates federal NESHAPs for pollutants through updates to 20.2.78 NMAC, which adopts 40 CFR Part 61, and federal NESHAPs for source categories through updates to 20.2.82 NMAC, which adopts 40 CFR Part 63. Similarly, Texas incorporates federal NESHAPs for both 40 CFR 61 and 40 CFR 63 through updates to 30 TAC 113. Kansas incorporates federal NESHAPs by adopting 40 CFR 61 through updates to KAR 28-19-735 and incorporates NESHAP source categories at 40 CFR 63 through updates to KAR 28-19-750. Oklahoma incorporates both 40 CFR 61 and 40 CFR 63 through Oklahoma Register 252-100-41-2 and Appendix Q.

4.1 NATIONAL AIR TOXICS ASSESSMENT

Every 4 years, the EPA's Office of Air Quality and Planning Standards produces a National Air Toxics Assessment (NATA). The most recent NATA for 2014 data was published August 2018. NATA is a first-pass, screening tool intended to evaluate the human-health risks posed by air toxics across the United States. Data is provided at the national, state, county, and census-tract level. NATA uses methods consistent with the general risk assessment framework used throughout EPA.

The NATA risk assessment uses emissions data compiled for a single year as inputs to air quality models, which incorporate meteorological data for the same year to estimate ambient air concentrations of certain air toxics. Modeled concentrations are then combined with census data and other information to calculate exposure concentrations of the air toxics. NATA then provides quantitative estimates of potential cancer risk and five classes of non-cancer hazards (grouped by organ/system: immunological, kidney, liver, neurological, and respiratory) associated with chronic inhalation exposure to real-world toxics.

NATA potential cancer risk values represent statistical probabilities of developing cancer over a lifetime. NATA non-cancer hazards are expressed as a ratio of an exposure concentration to a reference concentration (RfC) associated with observable adverse health effects (i.e., a hazard quotient). "For a

given air toxic, exposures at or below the RfC (i.e., hazard quotients are 1 or less) are *not* likely to be associated with adverse health effects. As exposures increase above the RfC (i.e., hazard quotients are greater than 1), the potential for adverse effects also increases” (EPA 2018b).

NATA can answer many questions including the following:

- Which air toxics pose the greatest potential risk of cancer or adverse non-cancer effects across the entire United States?
- Which air toxics pose the greatest potential risk of cancer or adverse non-cancer effects in certain areas of the United States?
- Which air toxics pose less, but still significant, potential risk of cancer or adverse non-cancer effects across the entire United States?
- When risks from long-term inhalation exposures to all outdoor air toxics are considered together, how many people could experience a lifetime cancer risk greater than levels of concern (e.g., 1-in-1 million or 100-in-1 million)?
- When considering potential adverse non-cancer effects from long-term exposures to all outdoor air toxics together for a given target organ or system, how many people could experience exposures that exceed the reference levels intended to protect against those effects (i.e., a hazard quotient greater than 1) (EPA 2018b).

NATA can be used as a tool to identify places of interest for further study, to get a starting point for local assessments, and to inform monitoring programs. For example, communities use NATA to find out what data and research is needed to better assess their local risk from air toxics. Communities have found that using NATA helps inform and empower citizens to make local decisions about their community’s health.

It is important to note that NATA focuses solely on exposures from inhalation of outdoor ambient air. The NATA framework does not address inhalation from indoor ambient air, estimate human exposure to chemicals via ingestion or through dermal contact, or account for exposures that may take place via other mechanisms. In addition, owing to the nature of the models used, the NATA Technical Support document highlights that NATA results should not be used:

- as a definitive means to pinpoint specific risk values within a census tract;
- to characterize or compare risks at local levels (such as between neighborhoods);
- to characterize or compare risks between states;
- to examine trends between or otherwise compare NATAs;
- as the sole basis for developing risk reduction plans or regulations;
- as the sole basis for determining appropriate controls on specific sources or air toxics; or
- as the sole basis to quantify benefits of reduced air toxic emissions (EPA 2018b).

In addition, although NATA reports results at the census tract level, average risk estimates are far more uncertain at this level of spatial resolution than at the county or state level. To analyze air toxics in smaller areas, such as census blocks or in suspected “hotspots,” other tools such as site-specific

monitoring and local-scale assessments should be used (EPA 2018b). NATA results are best used to focus on patterns and ranges of risks across the country.

Chronic indicators, known as RfC, are defined by EPA as the daily inhalation concentrations at which no long-term adverse health impacts are expected. Short-term (1-hour) HAPs concentrations will be compared to acute Reference Exposure Levels (RELs). RELs are defined as concentrations at or below which no adverse health effects are expected. No RELs are available for ethylbenzene and n-hexane; instead, the available Immediately Dangerous to Life or Health divided by 10 (IDLH/10) values are used. The National Institute for Occupational Safety and Health determines these IDLH values which are approximately comparable to mild effects levels for 1-hour exposures. The primary air toxics of concern for oil and gas operations are the BTEX compounds (benzene, toluene, ethylbenzene, and xylene), formaldehyde, and n-hexane.

Total cancer risk for the State of New Mexico (24.21 cases per million) was less than the United States (31.69 cases per million) (Table 5). In addition, all five non-cancer hazard quotient values were consistently lower in the State of New Mexico than the national values (immunological: 0.01; kidney: 0.005 ; liver: 0.009; neurological: 0.03; and respiratory: 0.32) than the United States (immunological: 0.02, kidney: 0.01; liver: 0.01; neurological: 0.04; and respiratory: 0.44).

At the county level, all seven counties (Chaves, Eddy, Lea, McKinley, Rio Arriba, Sandoval, and San Juan) had cancer risk values and total hazard quotients less than the United States, with all total hazard quotients reported being less than 1 (<1.0). (*However, one census tract in San Juan County, tract 35045000609, located SSW of Aztec, NM, although still less than 1 (<1.0), was greater than the U.S. total respiratory hazard quotient).

Table 5. NATA data for the United States, Texas, Oklahoma, Kansas, and New Mexico and Seven Counties in New Mexico

Location	Population	Total Cancer Risk (per million)	Total Hazard Quotients				
			Immunological	Kidney	Liver	Neurological	Respiratory
United States	312,572,412	31.6890	0.0248	0.0107	0.0149	0.0419	0.4366
Texas	25,145,472	35.0226	0.0164	0.0073	0.0120	0.0363	0.4275
Oklahoma	3,751,091	32.9178	0.0151	0.0049	0.0104	0.0272	0.4498
Kansas	2,853,116	26.8783	0.0224	0.0121	0.0176	0.0344	0.3656
New Mexico	2,059,163	24.2145	0.0108	0.0047	0.0088	0.0254	0.3162
Chaves Co.	65,645	25.8797	0.0094	0.0017	0.0070	0.0260	0.3392
Eddy Co.	53,828	27.9988	0.0117	0.0016	0.0069	0.0214	0.3479
Lea Co.	64,727	24.6598	0.0122	0.0017	0.0071	0.0193	0.2993
McKinley Co.	71,492	15.9305	0.0036	0.0008	0.0056	0.0190	0.2048
Rio Arriba Co.	40,246	16.5184	0.0039	0.0005	0.0056	0.0193	0.2002

Location	Population	Total Cancer Risk (per million)	Total Hazard Quotients				
			Immunological	Kidney	Liver	Neurological	Respiratory
Sandoval Co.	131,561	23.0073	0.0114	0.0053	0.0094	0.0259	0.2999
San Juan Co.	130,044	22.5893	0.0081	0.0020	0.0072	0.0264	0.3714

Source: EPA 2018b

Seven counties in table are: Chaves, Eddy, Lea, McKinley, Rio Arriba, Sandoval, and San Juan counties), where parcels are regularly nominated for BLM New Mexico Quarterly Oil and Gas Lease Sales.

Data for the United States, New Mexico, and seven counties in New Mexico. Total cancer risks and five total hazard quotients (immunological, kidney, liver, neurological, and respiratory) values are reported. Total Cancer Risk and Hazard Quotients at the county level in bold are those that are greater than the values for the State of New Mexico.

Detailed county and census-tract level results are reported in Appendix B, grouped according to BLM District Office/Field Office boundaries, with Pecos District counties (Chaves, Eddy, and Lea) grouped together, and Rio Puerco Field Office and Farmington Field Office (FFO) counties (McKinley, Rio Arriba, Sandoval, and San Juan) grouped together.

4.2 HYDROGEN SULFIDE

Hydrogen Sulfide (H₂S) is a colorless flammable gas with a rotten egg smell which is a naturally occurring byproduct of oil and gas development in some areas, including the New Mexico portion of the Permian Basin. H₂S is both an irritant and a chemical asphyxiant with effects on both oxygen utilization and the central nervous system. Its health effects can vary depending on the level and duration of exposure. Effects may range from eye, nose and throat irritation to dizziness, headaches, and nausea. High concentrations can cause shock, convulsions, inability to breathe, extremely rapid unconsciousness, coma, and death. Effects can occur within a few breaths, and possibly a single breath.

H₂S was originally included in the list of Toxic Air Pollutants defined by Congress in the 1990 amendments to the CAA. It was later determined that H₂S was included through a clerical error and it was removed by Congress from the list. H₂S was addressed under the accidental release provisions of the CAA. Congress also tasked EPA with assessing the hazards to public health and the environment from H₂S emissions associated with oil and gas extraction. That report was published in October 1993 (EPA 1993).

EPA found that while there was a potential for human and environmental exposure from routine emissions of H₂S from oil and gas wells, there was insufficient evidence to suggest that these exposures were a significant threat. H₂S is present in some oil and gas production zones. Flaring is used to reduce the H₂S emissions and the CFO has developed a series of standard conditions of approval for high H₂S areas in order to mitigate the risk of H₂S exposure (Lusk 2010).

H₂S was added to the Emergency Planning and Community Right-to-Know Act list of toxic chemicals in 1993. In 1994, EPA issued an administrative stay of reporting requirements for H₂S while further analysis was conducted. The administrative stay was lifted and Toxic Release Inventory reporting due in July 2013 for calendar year 2012 emissions required reporting of H₂S.

While there are no NAAQS for H₂S, a number of states, especially those with significant oil and gas production, have set standards at the state level. Table 6 summarizes these standards for states under BLM New Mexico State Office jurisdiction.

Table 6. State Ambient Air Quality Standards for H₂S

State	Standard	Averaging time	Remarks
Kansas	None	N/A	N/A
Oklahoma	200 ppb (0.2 ppm)	24 hour	N/A
New Mexico	0.010 ppm (10 ppb)	1 hour ⁽¹⁾	Statewide except Pecos-Permian Basin Intrastate Air Quality Control Region*
	0.100 ppm (100 ppb)	½ hour ⁽²⁾	Pecos-Permian Basin Intrastate Air Quality Control Region
	0.030 ppm (30 ppb)	½ hour	Within municipal boundaries and within 5 miles of municipalities with population >20,000 in Pecos-Permian Basin AQ Control Region
Texas	0.08ppm (80 ppb)	½ hour	If downwind concentration affects a property used for residential business or commercial purposes
	0.12 ppm (120 ppb)	½ hour	If downwind concentration affects only property not normally occupied by people

Source (Skrtic 2006)

* The Pecos-Permian Basin Intrastate Air Quality Control Region is composed of Quay, Curry, De Baca, Roosevelt, Chaves, Lea, and Eddy Counties in New Mexico.

⁽¹⁾ Pecos-Permian basin intrastate air quality control region has a ½ hour standard of 0.10 ppm.

⁽²⁾ Not to be exceeded more than once per year.

NMED has no routine monitors for H₂S. However, a one-time study done in 2002 (Skrtic 2006) sheds some light on the levels which can be expected near oil and gas facilities. These readings are averaged over 3-minute periods so are not comparable with the standard which has longer averaging periods. The monitoring data is presented in Table 7.

Table 7. Summary of Monitoring Data From New Mexico Study

Facility type	H ₂ S concentration measured at monitoring site (ppb)	
	Range	Average
Indian Basin hilltop, no facility	5–8	7
Indian Basin compressor station	3–9	6
Indian Basin active well drilling site	7–190	114
Indian Basin flaring, production, and tank storage site	4–1,200	203
Marathon Indian Basin refining and tank storage site	2–370	16

Facility type	H ₂ S concentration measured at monitoring site (ppb)	
	Range	Average
Carlsbad city limits, near 8 to 10 wells and tank storage sites	5–7	6
Carlsbad city limits, Tracy-A	5–8	7
Compressor station, dehydrators – Location A	4–5	4
Compressor station, dehydrators – Location B	2–15,000	1372
Huber flare/dehydrating facility	4–12	77
Snyder oil well field	2–5	4
Empire Abo gas processing plant	1–1,600	300
Navajo oil refinery	3–14	7–8

Source: Skrtic 2006

In Oklahoma, routine monitoring downwind of two refineries in Tulsa showed H₂S levels that were within state standards but above normal background levels. In Texas, which has 12 routine monitors, H₂S levels generally ranged from 0.1 to 5 ppb. One monitor at a compressor station, however, showed frequent levels in excess of the state standard of 0.8 ppm (Skrtic 2006).

5 METHODOLOGY AND ASSUMPTIONS FOR ANALYSIS OF AIR RESOURCES

Air resource impacts can be analyzed on a number of different levels. First and most basic is to compare monitored pollutant levels with NAAQS. This applies only to criteria pollutants and provides a basis for determining whether the emissions of any specific pollutant are significant in a local area. Secondly, and necessary before further analysis can be done is an estimate of actual emissions, or an emissions inventory. This may be done for all emissions in a geographic area and for a project to provide a comparison. The EPA completes an NEI (see Section 3.1) at the county level every 3 years, which provides a baseline for determining whether project emissions will cause a substantial increase in emissions or materially contribute to potential adverse cumulative air quality impacts. Finally, if impacts are anticipated to be significant, it may be necessary to apply air quality modeling to analyze the extent and geographic distribution of impacts.

Traditional air quality modeling generally falls into three categories. 1) Near-field dispersion modeling is applied to criteria pollutants, HAPs, and Air Quality Related Values (AQRVs) where a small to medium number of sources are involved to cover an area within 50 kilometers (km) of a proposed project. 2) Far-field or transport modeling is used to provide regional assessments of cumulative and incremental impacts at distances greater than 50 km. 3) Photochemical modeling is necessary for large scale projects with a large number of sources or with complex issues including O₃ and secondary particulate impacts. An Air Quality Memorandum of Understanding (MOU) signed by the U.S. Department of Agriculture, U.S. Department of the Interior (DOI), and EPA contains an appendix which describes the air quality models available and their advantages, disadvantages, and applications. The MOU and appendix are included as Appendix E of this document. As of July 25, 2019, the Air Quality MOU was terminated (see Appendix E). We retain information related to the Air Quality MOU for historical purposes.

5.1 EMISSION INVENTORIES, STUDIES, AND MODELING

An emissions inventory is a database that lists, by source, the amount of air pollutants discharged into the atmosphere during a year or other time period. Governments use emission inventories to help determine significant sources of air pollutants and to target regulatory actions. Emissions inventories are an essential input to mathematical models that estimate air quality. The effect on air quality of potential regulatory actions can be predicted by applying estimated emissions reductions to emissions inventory data in air quality models.

Emission trends over time can be established with periodic updates of the emissions inventory. Inventories also can be used to raise public awareness regarding sources of pollution. An emissions inventory includes estimates of the emissions from various pollution sources in a geographical area. It should include all pollutants associated with the air quality problems in the area. For example, an emissions inventory to support the management of ground-level O₃ should include sources of NO_x and VOCs (EPA 2020c). Emission inventories are performed to determine the current state of the atmosphere, contribute to determining a future state of the atmosphere, aid decision makers in policy and regulatory guidance, and determine emission controls. A most recent emission inventory relative to BLM New Mexico operations (Permian and San Juan basins) was completed in 2017 for base year 2014. Other emission inventories, studies and modeling years include 2005, 2007, 2009 and 2013.

5.1.1 2014 WESTERN STATES AIR RESOURCES COUNCIL–WESTERN REGIONAL AIR PARTNERSHIP EMISSIONS INVENTORY

The Western States Air Resources Council–Western Regional Air Partnership (WESTAR-WRAP) conducted an oil and gas emissions inventory report for base year 2014 to further clarify the contributions of oil and gas activities to human-caused emissions within the Permian and San Juan Basins (Ramboll Environ 2017). The inventory included the counties of Chaves, Eddy, Lea, and Roosevelt for the Permian Basin. The inventory included data from not only the New Mexico counties of McKinley, Rio Arriba, Sandoval, San Juan, and Valencia but also Archeleta and La Plata in Colorado. For purposes of our analysis, we only bring forth emissions from Eddy, Lea, Chaves, Roosevelt, Rio Arriba, San Juan, McKinley and Sandoval counties for reporting and comparison in Table 8.

Based on previous comparison of the WESTAR WRAP data versus the NEI data from the 2014 base year, there is reason to believe that there are non-point sources, including fugitive components, pneumatic devices, pumps, and well blowdown events that may not be reported through the state and federal inventories. These nonpoint sources could represent greater criteria, HAPs, and GHG emissions within these basins, in particular VOC and NO_x emissions that contribute to O₃ formation. It is therefore believed that the NEI data related to Petroleum and Related Industries is underreported in terms of VOC and NO_x emissions. Table 8 provides the 2017 NEI and WESTAR-WRAP datasets. Because the inventories are not presenting the same base-year emissions, it is not possible to make a meaningful comparison as to the magnitude of potential underestimates of emissions; however, both datasets are provided below for reference.

Table 8. 2017 NEI and 2014 WESTAR-WRAP Inventory Emissions

Basin/Counties	Emissions (metric tons)					
	NO _x	CO	VOC	PM ₁₀	PM _{2.5}	SO ₂
Permian Basin (Chavez, Eddy, Lea and Roosevelt Counties)						
2017 NEI – all sources*	33,217	78,938	172,241	39,761	7,086	7,577
2017 NEI – petroleum and related industries	13,001	–	82,793	–	–	–
WESTAR-WRAP 2014 oil and gas sources	30,351	–	121,644	–	–	–
San Juan Basin (San Juan, Rio Arriba, McKinley, Sandoval Counties)						
2017 NEI – all sources	69,967	214,889	180,956	56,049	17,238	6533
2017 NEI – petroleum and related industries	23,958	–	72,240	–	–	–
WESTAR-WRAP 2014 oil and gas sources	44,433	–	86,173	–	–	–

* Includes biogenic sources.

Sources: EPA 2020b; Ramboll Environ 2017.

Notes: Values include Tier 1 summaries for each county, including combustion, industrial, on-road/nonroad, and miscellaneous sectors. Biogenic sources are now reported in the totals.

Only precursor pollutants to O₃ formation compared in this analysis (NO_x and VOC).

The Westar WRAP data shows the following:

In the New Mexico Permian Basin, non-point and point sources of oil and gas are shown to contribute 11,790 and 18,561 tpy respectively of the 30,351 total tpy of man-made NO_x emissions. The inventory revealed that the major oil and gas sources (point and non-point sources) of NO_x emissions are attributed to: Point Source Compressor Engines 33%, Midstream Unclassified Sources 29%, Drill rigs 16%, artificial lifts 13%, Fracing 4%, non-point heaters 3% as well as other sources totaling approximately 2% of non-point and point emission sources.

In the New Mexico Permian Basin, non-point and point sources of oil and gas are shown to contribute 110,480 and 11,164 tpy respectively of the 121,644 total tpy of man-made VOC emissions. The inventory revealed that the major oil and gas sources of VOCs emissions are attributed to: Oil Tanks 58%, Midstream Unclassified Sources 7%, Venting-blowdowns 7%, pneumatic devices 7%, oil well truck loading 6% as well as other sources representing approximately 15% of non-point and point VOC emission sources.

In the New Mexico San Juan Basin, non-point sources of oil, gas and CBM wells are shown to contribute 33,435 tpy of the 44,433 total tpy of man-made NO_x emissions. These major categories of non-point NO_x emissions are attributed to nonpoint compressor engines 88%, water pump engines 6.9%, non-point heaters 2.4%, artificial lift devices 1.9%, as well as other sources representing approximately 0.8% of non-point NO_x emission sources.

In the New Mexico San Juan Basin, non-point sources of oil, gas and CBM wells are shown to contribute 79,363 tpy of the 86,173 total tpy of man-made VOC emissions. These major categories of non-point VOCs emissions are attributed to pneumatic devices 32%, non-point compressor engines 18.5%, non-

point fugitives 14.8%, pneumatic pumps 12.7%, dehydrators 9.7%, as well as other sources representing approximately 12.3% of non-point VOC emission sources (Ramboll Environ 2017).

5.1.2 2008 OZONE STUDY

In 2013, the Western Regional Air Partnership (WRAP) completed a regional technical analysis for O₃ (WestJump) that includes information about O₃ impacts and sources that contribute to the formation of O₃ for calendar year 2008. The analysis demonstrated that the largest contributor to O₃ concentrations in the western United States was international transport and stratospheric O₃. State-to-state O₃ transport was important, as well. For example, New Mexico sources significantly contribute to elevated O₃ concentrations in Texas, Arizona, and Colorado. Texas is a significant contributor to elevated O₃ in Oklahoma, Louisiana, New Mexico, Missouri, and Arizona. Kansas sources significantly contribute to elevated O₃ in Missouri and Texas, while Oklahoma sources significantly contribute to elevated O₃ in Missouri, Texas, and New Mexico (ENVIRON et al. 2013). The term significant in this context is based on the degree to which an upwind state's anthropogenic emissions contribute to a downwind state's average design value. Emission contributions from an upwind state are assumed to be significant if the contribution from the upwind state's emissions results in impacts to a nonattainment area (with an average design value above the NAAQS) in excess of 1% of the NAAQS.

The WestJump analysis also provides information about PM_{2.5} impacts and contributing sources for calendar year 2008. Interstate transport is significant for PM_{2.5}. New Mexico significantly contributes to annual PM_{2.5} exceedances in Arizona; Texas significantly contributes to exceedances in Arkansas, Missouri, Mississippi, Illinois, and Alabama; Oklahoma significantly contributes to exceedances in Arkansas and Missouri, and Kansas significantly contributes to exceedances in Iowa, Missouri, Illinois, Arkansas, and Wisconsin. For the 24-hour PM_{2.5} standard, New Mexico significantly contributes to exceedances in California, Texas and Oklahoma significantly contribute to exceedances in Iowa, and Kansas significantly contributes to exceedances in Iowa and Wisconsin (ENVIRON et al. 2013).

5.1.3 2007 EMISSIONS INVENTORY

An emissions inventory conducted for the CFO for calendar year 2007 and including Chaves, Lea, and Eddy counties (Applied Enviro Solutions 2011) shows that VOC emissions from biogenic (natural) sources are far greater than those from anthropogenic (human) sources and account for 91% of VOCs inventoried. Point source emissions (which might include such industrial sources as power plants, gas plants and oil refineries) account for 40% of anthropogenic VOC emissions in the area, solvent use accounts for 15%, and fire (including wildland, structure, and open burning) accounts for 8%. Oil and gas area sources produce only 1.4% of VOCs in the area while pipeline transport of oil and gas accounts for 1.7%.

The 2007 emissions inventory for Chaves, Eddy, and Lea Counties shows that anthropogenic sources account for 65% of CO emissions and biogenic sources 35%. Of the anthropogenic emissions 47% are from on road mobile sources, 24% from industrial point sources, 14% from non-road mobile sources, 9% from fire, and 2% each from oil and gas area sources and waste disposal burning (Applied Enviro Solutions 2011) NO_x emissions in the Carlsbad area are largely anthropogenic (88%). Of the total human-caused NO_x emissions, industrial point sources account for 84%, on-road mobile sources account for 7%, oil and gas area sources account for 5%, non-road mobile sources account for 2%, and residential heating with natural gas and propane account for 1% (Applied Enviro Solutions 2011).

A 2007 emissions inventory for Chaves, Eddy, and Lea Counties shows that the bulk of emission for both PM₁₀ and PM_{2.5} are from dust from unpaved roads (88 and 65% respectively). For PM₁₀, the next three highest categories are point sources at 2.8%, tilling and harvesting 2.6% and paved roads 2.4%. Oil and gas area sources account for only 0.1% of PM₁₀ emissions. For PM_{2.5}, the next three highest categories are point sources at 17%, fire at 4.3% and tilling and harvesting at 2.8%. Oil and gas area sources account for 0.8% of PM_{2.5} emissions in this area (Applied Enviro Solutions 2011).

The Carlsbad area, the 2007 emissions inventory does not differentiate SO₂ from SO_x but it can be assumed that the percentage of emissions by category is similar. In this region, oil and gas area sources account for 74% of all SO_x emissions with most of the remainder, 25% accounted for by industrial point (Applied Enviro Solutions 2011).

5.1.4 2005 EMISSIONS INVENTORY

An emissions inventory conducted for the Four Corners region for calendar year 2005, including counties in northwestern New Mexico, southwestern Colorado, southeastern Utah, and northeastern Arizona, estimates VOC emissions from biogenic sources account for 55% of total VOCs (Environ 2009). Oil and gas area and point sources accounted for 28% of VOCs inventoried.

The top three sources of NO_x emissions in the Farmington area in 2005 were electricity generation (72,668 tons; 33%), oil and gas (68,830 tons; 31%) and on-road mobile sources (39,340 tons; 18%) (Environ 2009).

5.1.5 2005 FOUR CORNERS AIR QUALITY TASK FORCE PHOTOCHEMICAL MODELING

In 2009, a photochemical modeling analysis was completed for the Four Corners Air Quality Task Force (FCAQTF). Potential O₃ impacts and the usefulness of certain mitigation measures were analyzed. This modeling showed that the Four Corners region would continue to meet the current O₃ standard in 2018 with continued oil and gas development and population growth. The analysis showed that emissions reductions would be required for both power plants and oil and gas sources in order to achieve measurable reductions in O₃ concentrations. The best achievable O₃ reductions from the modeling scenarios implementing control measures to reduce emissions were on the order of 5 (Environ 2009).

The modeling analysis completed for FCAQTF also used source apportionment modeling, which indicated that, in general, transport from outside the region and naturally occurring VOCs from vegetation were large contributors to 2005 O₃ levels. However, it was also shown that on days with high O₃ concentrations, oil and gas sources and electricity generation units (EGUs) both contributed significantly to the total modeled O₃ concentrations.

5.1.6 PLANNED MODELING STUDIES

An update to previous photochemical grid modeling studies is in the planning phase as part of the New Mexico Ozone Attainment Initiative, which will use 2014 Base Year Emission Inventory data and develop projections for future year 2023. The modeling study will be conducted by enhancing the WRAP/WAQS 2014 modeling platform to use a 4-km grid resolution domain covering New Mexico and surrounding areas, especially the oil and gas production regions in the Permian and San Juan Basins (Ramboll Environ and Westar 2020).

5.2 PECOS DISTRICT OFFICE ATMOSPHERIC AND PHOTOCHEMICAL GRID MODELING

An Air Resources Technical Support Document (ARTSD) by URS Corporation (URS 2013) was prepared to report the potential air quality impacts resulting from the Reasonable Foreseeable Development (RFD) scenario. This effort included atmospheric dispersion and photochemical grid modeling to predict concentrations of specific pollutants in and around the CFO (in which most of the Pecos District oil and gas activity occurs). The results of ARTSD analysis indicate that air quality impacts from the RFD scenario, while noticeable, are generally acceptable. Most predicted criteria pollutant concentrations are well below the NAAQS throughout the extensive modeling domains included in this analysis. While no exceedances of NAAQS were predicted from the modeling of federal wells associated with the RFD scenario (6,400 wells), consideration of the entire RFD scenario (16,000 wells) and other reasonably foreseeable future actions (i.e., cumulative impacts) in the ARTSD included predictions of pollutant concentrations approaching or exceeding the NAAQS (for O₃, PM_{2.5} and potentially SO₂) and indicate the need for additional ambient monitoring data, refined modeling, and consideration of additional mitigation measures. Most of the areas where NAAQS would be exceeded are out of the CFO region (URS 2013)

5.3 AIR QUALITY MODELING FOR TEXAS

Numerous reports on air quality modeling projects done by and for the TCEQ, including modeling done for the Dallas and Houston nonattainment areas can be accessed on the Air Division website (TCEQ 2019b). The TCEQ has convened advisory groups in southeastern Texas and Dallas–Fort Worth to assist the agency in addressing photochemical modeling issues.

6 CALCULATORS OIL AND GAS DEVELOPMENT

Emissions calculators were developed by air quality specialists at the BLM National Operations Center in Denver, Colorado. The calculators use an Excel spreadsheet for computation and are based on emissions factors from EPA and the American Petroleum Institute. The calculators were quality assured and improved by URS under contract with the BLM. Methodology for computing GHGs is documented in *The Climate Change Supplementary Information Report for the Montana, North Dakota, and South Dakota Bureau of Land Management* (URS 2010). More recently, Kleinfelder West, Inc. developed a calculator for a representative oil and gas well in the western United States (Kleinfelder West, Inc. 2013). Other air pollutant computations have not yet been described in a published document but are based on methods recommended in the EPA publication AP-42 *Compilation of Air Pollutant Emissions Factors* (EPA 1995, 2006).

The calculators may be considered a type of model in that they use emissions factors, mathematical algorithms, and assumptions to arrive at some approximation of reality. However, their primary purpose is to compute an emissions inventory which is a necessary ingredient to any modeling effort. The calculators account for a number of variables, including access and construction requirements, equipment and other infrastructure needs, and expected production volumes. Because the algorithms used by the calculators to quantify emissions are based on averages and numerous assumptions must be made about construction, the calculators provide an approximation of emissions levels. Actual project emissions may be greater or less than those projected by the calculators. Emissions calculators used to estimate emissions from oil and gas development assume that wells will be hydraulically fractured; if a well is not fractured, emissions will be less than calculated.

The BLM in New Mexico has modified the calculators and assumptions for use in analyzing a single well and to more closely represent oil and gas wells in the state of New Mexico; specifically, the San Juan and Permian Basins. However, it must be understood that the calculators were originally designed to make estimations of emissions at the resource management plan (RMP) level which would result in some averaging and smoothing of assumptions. At the single well level, the uncertainty in emissions projections increases substantially.

The BLM has determined that well production typically declines over time and has assumed that declining production would result in reduced emissions over time. A production history may vary from a straight line to a hyperbolic curve. The object of decline curve analysis is to model the production history. Assuming a certain abandonment pressure or gas rate, the decline curve is used to determine the productive life of the well. Well life can vary from a few years to many decades depending on the reservoir and the year it was drilled. Production is also dependent on the price of oil and gas. Since initial development in the San Juan Basin in the 1920s, all reservoirs have had significant reservoir pressure declines. Subsequent infill drilling will encounter reduced pressure reservoirs with limited well life spans compared to wells drilled earlier in the development of the field.

It should be noted that the calculations are based on recently drilled wells and tend to overestimate the average emissions over the lifetime of the well. It is not possible to estimate the lifespan of an individual well, nor is the calculator able to incorporate the decline curve into results, so we have computed one-time (construction, completion, workover, and reclamation) emissions and annual (operations and maintenance) emissions. However, the annual emissions do not take into account the declining production rates over the lifetime of the well.

6.1 ASSUMPTIONS

As mentioned above, the calculators account for numerous variables or inputs that are used to calculate the overall emissions of the different stages of oil and gas development. At the time of an APD, not all of these variables may be known. To populate the calculators with the different variables, the BLM CFO and the FFO each developed a set of assumptions pertaining to development in their respective areas. These assumptions address variables such as well depth, production, road development/maintenance, travel to and from well sites, construction times, and need for workovers. The following sections summarize the assumptions made for each field office area to populate the calculators.

6.1.1 ASSUMPTIONS – FARMINGTON FIELD OFFICE

There are several geologic formations within the FFO boundary that are known to produce natural gas. The Fruitland is the shallowest routinely produced formation at approximately 2,000 feet deep. The Dakota is the deepest formation routinely produced at approximately 6,000 feet deep. The Mancos Shale formation is approximately 2,500 feet deep. Several formations produce various amounts of water during the production phase of the well. The preferred method of disposing of the produced water is via an injection well drilled into the geologically isolated Entrada formation, which is approximately 7,500 feet deep. Although wells are not drilled to these precise depths, these generalized depths were used for the purpose of estimation in the emissions calculator.

Between 2015 and 2030, it is estimated that most of the oil and gas drilling in FFO will be within the Mancos Shale formation. The formation is thought to contain gas on the northern end (southern Colorado and northwestern New Mexico) and oil on the southern end (towards Gallup and Grants, NM).

Assumptions in the estimates account for unfavorable natural gas prices until 2019 and no gas well development in the Mancos Shale formation was expected until that time. Estimates assume that starting in 2019, approximately 100 to 200 gas wells could be drilled per year. It is likely that central collection and shipping facilities for oil will be developed, too. Oil well development in the Mancos Shale formation is expected to proceed at 100-200 wells per year, although little development is currently occurring due to unfavorable oil prices. Recent oil well drilling has used horizontal rather than vertical wells.

BLM specialists and engineers were consulted to develop a range of values to insert into the calculator to estimate the emissions from construction, completion, interim reclamation, annual operation, and final reclamation. Pad construction, interim reclamation, and final reclamation processes are generally the same across the basin. The range of values was designed to address the requirements of about 95% of the wells developed in the San Juan Basin. Unforeseen or unpredictable events may cause approximately 5% of wells to fall outside of the range.

The calculator includes construction of a “frac pond,” as future wells in the region will most likely be accomplished with hydraulic fracturing. The calculator has options for diesel-fired or natural gas-fired drill rigs. More commonly in 2014 in the region, drill rigs were diesel-fired. Many of the well pads have associated man camps where drilling personnel are housed during well drilling. Since most pads will now accommodate more than one well, the man camps allow employees to avoid commuting during the time the wells on the pad are drilled.

The ancillary activities associated with the production phase of a well such as workovers, road maintenance, and road traffic are somewhat difficult to predict. Calculations for Mancos Shale drilling recently have assumed one well workover per year. Existing gas wells in the FFO area do not require workover on a regular schedule. Three (3) to 6 years between workovers is typical, and the nature of the work required during a workover is variable.

FFO and the oil and gas industry have established a road committee to identify collector roads (main travel corridors) and have established procedures to maintain collector roads, as necessary. However, no regular maintenance schedule exists. Most new wells are drilled along existing resource roads that are not covered by the road committee and are maintained as needed. Although road maintenance within the FFO varies, a reasonable assumption is that the resource roads will be maintained once a year. The average length of new road required to drill a new well during the past 2 years has been 800 feet. Emissions are calculated based on this average assuming that an 800-foot resource road is maintained once a year and the maintenance work would require about 6 hours of work.

The majority of producing wells in the San Juan Basin utilize remote telemetry powered by solar panels to transmit well production data to centralized office locations. While the frequency of well site visits is not predictable, the need for well site visits during the production phase of the well is greatly reduced by the telemetry systems. Typically, a field technician will drive a light truck and will visit multiple wells per trip along an established service route. To estimate the miles required for each site visit, an additional 4.5 miles of travel along an existing driving route was added to the typical 800 feet of new road for a total of 5 miles. Emissions are calculated for weekly visits during the year for a light truck. For various servicing needs, heavy duty vehicles over 8,500 pounds gross vehicle weight rating (EPA 2020d) are required on-site during drilling and workovers. Heavy duty vehicles typically do not visit multiple sites per day. Emissions are calculated for driving 50 miles round trip for five trips per year.

The average San Juan Basin gas well produces at a rate of 100 mcf/d (thousand cubic feet per day). For analysis purposes, the initial production rate is assumed to be 100 mcf/d. The volume of gas and oil is normally the greatest following the completion of the well. Oil and gas production rates decline as a function of time, reservoir pressure drop, or the changing relative volumes of the produced fluids.

The FFO RMP (DOI BLM 2003) addressed air quality based on the Air Quality Modeling Analysis Technical Report prepared by Science Applications International Corporation (SAIC 2003). The 2003 FFO RMP modeling is considered here because it was used to characterize air quality for the purpose of land use planning, and this EA tiers to the 2003 FFO RMP. The 2003 SAIC modeling was based on the highest level of oil and gas development proposed based on the RFD and identified a potential for exceeding the NAAQS for NO₂. The alternative selected for the RMP proposed a lower level of development than that modeled. Lower levels of development and NO_x limits placed on engines have resulted in lower impacts than were modeled.

6.1.2 ASSUMPTIONS – CARLSBAD FIELD OFFICE

The CFO area of responsibility contains 28 different geologic zones that produce oil, natural gas, and water. The complex geology, variety of drilling techniques used (horizontal, vertical), uncertainty of production, and variation of the drilling time and equipment required makes it difficult to approximate the emissions for a proposed well. In order to provide a basis for extrapolation, the CFO selected a random sample of 70 wells out of a population of 1836 wells drilled from 2007 to 2010. Since the wells included in the sample had been recently drilled at the time the study was conducted, the production data are likely an overestimate of average annual production (and therefore emissions) as production drops with the age of the well. The sample size was selected to ensure that it was representative of 95% of the recently drilled wells.

The 70 wells were reviewed to ensure accurate production data was available and to eliminate older wells that had been re-drilled into a new formation. Sixty-eight (68) wells remained after the review. This was still a sufficient sample size to ensure statistical accuracy, so no additional wells were selected. The annual production values for oil, gas, and water, length of road constructed, well pad size and travel distances to reach the well from the nearest town were calculated for each well. The lowest, highest, and mean values were then calculated for each parameter and used to create three emissions scenarios (maximum, minimum, and average). These values represent the maximum, minimum and average emissions for 95% of the new wells in the CFO. Unforeseen or unpredictable events may cause 5% of wells to fall outside of the range. Because the minimum scenario has no production, it can be used to estimate the emissions from a salt-water disposal well.

Other values required for the calculator were conservatively estimated by BLM resource staff. It is not possible to predict the exact amount of time or equipment required for the development and operation of a well in the Permian Basin due to the varied geological formations, numerous operators, and other variables. Therefore, BLM specialists and engineers were consulted to develop a range of values to insert into the calculator to estimate the emissions from construction, completion, interim reclamation, annual operation, and final reclamation. The range was designed to include the requirements of 95% of the wells that may be developed in the Permian Basin. Where no information was available, the default values from the calculator were used. The calculator will be updated as additional information becomes available.

The ancillary activities associated with the production phase of a well such as workovers, road maintenance, and road traffic are difficult to predict. Oil and gas wells in the CFO do not require workover on a regular basis and when these activities occur, they generally are not reported to the BLM. Three (3) to 6 years between workovers is routine, and the nature of the work required during a workover is variable. It is assumed that any gas released during the completion process will be flared. The calculator assumes 100% combustion efficiency.

The emissions calculator can be used to estimate PM as a result of construction and drilling activities related to pad building and road traffic. The amount of PM emissions depends on the length, surface condition; soil types traversed, and soil moisture conditions of the road to the site. Because site visit frequencies vary and are difficult to predict, varying numbers of site visits were input into the calculator, which had almost no impact on the total tons of PM emitted. Most gas wells in the Permian Basin utilize remote telemetry powered by solar panels to transmit well production data to centralized office locations. The need for well site visits during the production phase at these wells is greatly reduced. Oil wells require site visits, and the frequency of well visits is not predictable.

While the frequency of well site visits is not predictable, the need for well site visits during the production phase of the well is greatly reduced by the telemetry systems. Typically, a field technician will drive a light truck and will visit multiple wells per trip along an established service route. It was estimated that an average trip distance consists of 2 miles three times per week. This information is used in calculating the annual operation emissions. Heavy trucks are required on site less often than light trucks for various servicing needs. Heavy trucks typically do not visit multiple sites per day. Distances to the wells were determined from the statistical sample including the total distances traveled on dirt and paved roads to reach the well from the nearest town (Carlsbad, Artesia, Hobbs, etc.). Emissions include maintenance and inspection of the well. Reclamation of the well site and road will be conducted when the well has finished producing and is plugged and abandoned. Emissions from reclamation of the well pad and road are also estimated.

County roads in the CFO have established procedures for maintenance but no regular maintenance schedule exists. Most new wells are drilled along oil and gas lease roads that are only maintained by oil and gas operators as needed. Therefore, road maintenance within the CFO is not predictable. The average length of new road required to drill a new well during the past 4 years based on the random sample has been 570 feet. Emissions are calculated based on this average assuming that a 570-foot resource road is maintained once a year.

Maximum, minimum, and average emissions for construction, completion/recompletion, workover, annual operations, annual road maintenance, and reclamation have been calculated and are presented in project APD EAs. Note that these estimates are based on hypothetical scenarios and it is unlikely that the maximum emissions scenario would ever occur.

6.1.3 ASSUMPTIONS – OKLAHOMA FIDLD OFFICE

Because development within the OFO can occur within a wide range of geographic areas and formations, a specific calculator has not been developed for this field office. However, calculators for other field offices may be used to obtain a reasonable estimate of emissions on a per well basis. Updates to assumptions may be made where warranted if there is enough detail to determine with certainty better assumptions depending on the location of a project.

6.2 VOCS AND WELL DRILLING OPERATIONS

Specifically, VOCs are emitted during well drilling and operations as exhaust from internal combustion engines. VOCs may be emitted from hydraulically fractured oil and gas wells during the fracturing and re-fracturing of the wells. In the hydraulic fracturing process, a mixture of water, chemicals and proppant is pumped into a well at extremely high pressures to fracture rock and allow oil and gas to flow from the geological formation. During one stage of well completion, fracturing fluids, water and reservoir gas come to the surface at high velocity and volume (flowback). This flowback mixture contains VOCs, methane, benzene, ethylbenzene, and n-hexane; some or all of the flowback mixture may be vented, flared or captured. The typical flowback process lasts from 3 to 10 days, so there is potential for significant VOC emissions from this stage of the well completion process. Most new oil and gas wells drilled today use the hydraulic fracturing process.

6.3 WELL COUNTS

The number of active wells can vary greatly from year-to-year as well counts are not static or logarithmic by nature. Well count data can be obtained from many sources such as state oil and gas commission databases, university and research databases, proprietary databases, as well as public federal databases. The sources reporting well counts may also differ in reporting methods. Reporting of well counts may include various types of wells such as active, new, temporarily abandoned, and inactive (shut in or temporarily abandoned). For the purposes of this report, the BLM uses the Petroleum Recovery Research Center, Automated Fluid Minerals Support System (AFMSS), and state oil and gas well count reporting.

According to data provided by the Petroleum Recovery Resource Center (PRRC), there are approximately 20,207 active wells within the San Juan Basin (McKinley, Rio Arriba, San Juan, and Sandoval counties) of which approximately 14,302 are federal. Within the Permian Basin (Chaves, Eddy, Lea, and Roosevelt) there are 41,006 active wells (primarily vertical wells) of which approximately 18,690 are federal (PRRC 2021).

7 AIR QUALITY RELATED VALUES

AQRVs are resources sensitive to air quality and can include a wide variety of atmospheric-chemistry related indicators. Monitoring and modeling of AQRVs help to provide a level of protection to sensitive areas such as Class I parks and wilderness areas. Such resources may include visibility or a specific scenic, cultural, physical, biological, ecological, or recreation resource identified for a particular area. Congress established certain national parks and wilderness areas as mandatory Class I areas where only a small amount of air quality degradation is allowed. Defined by the CAA, Class I areas include national parks greater than 6,000 acres, wilderness areas and national memorial parks greater than 5,000 acres, and international parks. These areas must have been in existence at the time the CAA was passed by Congress in August 1977.

The goal of Class I management is to protect natural conditions, rather than the conditions when first monitored. That is, if initial monitoring in a Class I area identifies human-caused changes, appropriate actions should be taken to remedy them to move toward a more natural condition. The goal of Class I management is to protect not only resources with immediate aesthetic appeal (i.e., sparkling clean streams) but also unseen ecological processes (such as natural biodiversity and gene pools) (U.S. Forest Service [USFS] et al. 2000). The Federal Land Managers' Air Quality Related Values Workgroup (FLAG)

issued a revised Phase 1 report in 2010 (USFS et al. 2010). This report was developed as a tool to provide consistent approaches to the analysis of the effects of air pollution on AQRVs. The FLAG report focuses on three areas of potential impact: visibility, aquatic and terrestrial effects of wet and dry pollutant deposition, and terrestrial effects of O₃. This report is structured to address these same three areas of potential impact.

The BLM’s goals include managing the jurisdictional field offices’ activities and development to protect and improve air quality and, within the scope of the BLM’s authority, minimize emissions that cause or contribute to violations of air quality standards or that negatively impact AQRVs (e.g., acid deposition, visibility).

7.1 VISIBILITY

Visibility is of greatest concern in Class I areas which are afforded the highest level of air quality protection by the CAA. Visibility impairment is a result of regional haze which is caused by the accumulation of pollutants from multiple sources in a region. Emissions from industrial and natural sources may undergo chemical changes in the atmosphere to form particles of a size which scatter or absorb light and result in reductions in visibility.

In 1985, the EPA initiated a network of monitoring stations to measure impacts to visibility in Class I Wilderness Areas. These monitors are known as the Interagency Monitoring for the Protection of Visual Environments (IMPROVE) monitors and exist in some, but not all, Class I wilderness areas. Table 9 shows the Class I areas in the BLM New Mexico State Office area of responsibility and whether they have an IMPROVE monitor and, if not, which monitor is considered representative for that area. There are no Class I areas in Kansas.

Table 9. Class I Areas and IMPROVE Monitors

State	Class I Area	Agency	IMPROVE
New Mexico	Bandelier	National Park Service (NPS)	Yes
	Bosque del Apache	U.S. Fish and Wildlife (USFWS)	Yes
	Carlsbad Caverns	NPS	Guadalupe Mountains
	Gila	USFS	Yes
	Pecos	USFS	Wheeler Peak
	Salt Creek	USFWS	Yes
	San Pedro Parks	USFS	Yes
	Wheeler Peak	USFS	Yes
	White Mountain	USFS	Yes
Texas	Big Bend	NPS	Yes
	Guadalupe Mountains	NPS	Yes
Oklahoma	Wichita Mountains	USFWS	Yes

Figures 2 through 4 shows visibility extinction trends for each of the IMPROVE monitors in the BLM New Mexico State Office area of responsibility. Note that peaks such as that seen for Bandelier National Monument in 2000 may be accounted for by the occurrence of large wildfires. A downward sloping line means less reduction of visibility and therefore an improvement. In most cases visibility trends have been flat or improving. Implementation of Best Available Retrofit Technology (BART) strategies as required under the federal Regional Haze Rule over the next few years should result in further improvements.

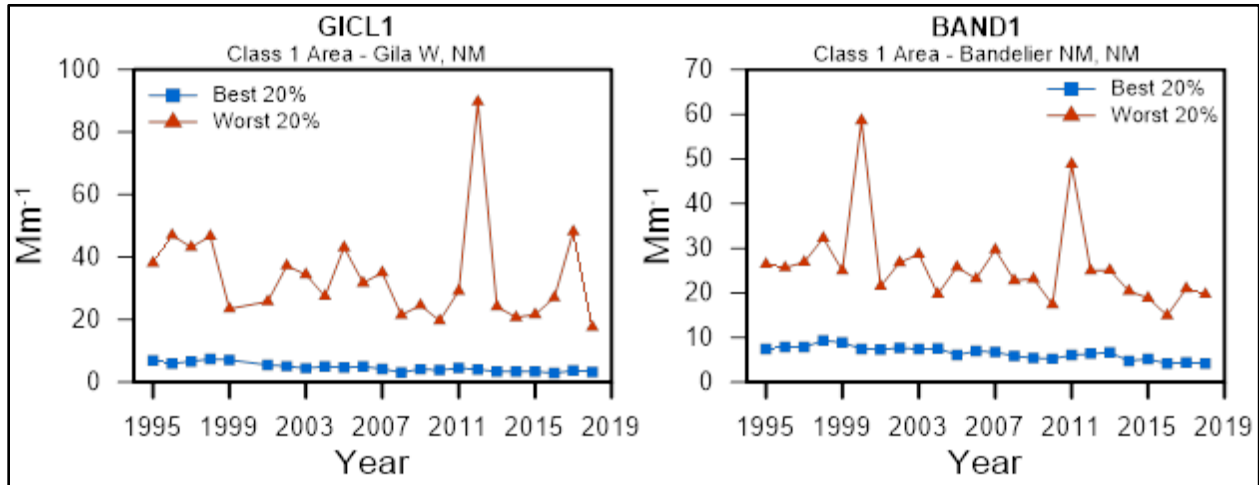


Figure 2. Visibility extinction trends for the Gila Wilderness area and Bandelier National Monument, New Mexico (data retrieved from Colorado State University, 2020).

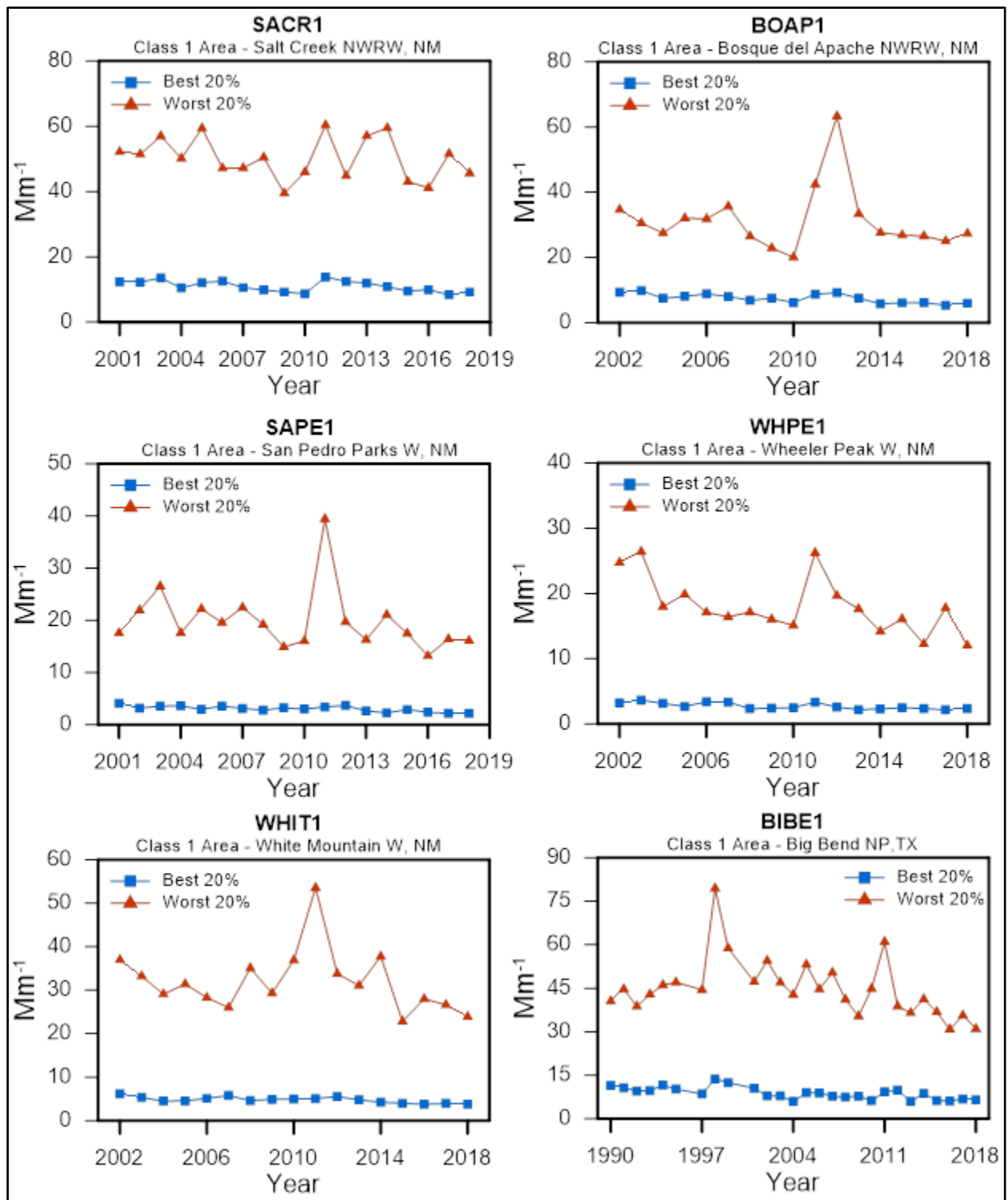


Figure 3. Visibility extinction trends for six Class 1 areas (data retrieved from Colorado State University, 2020).

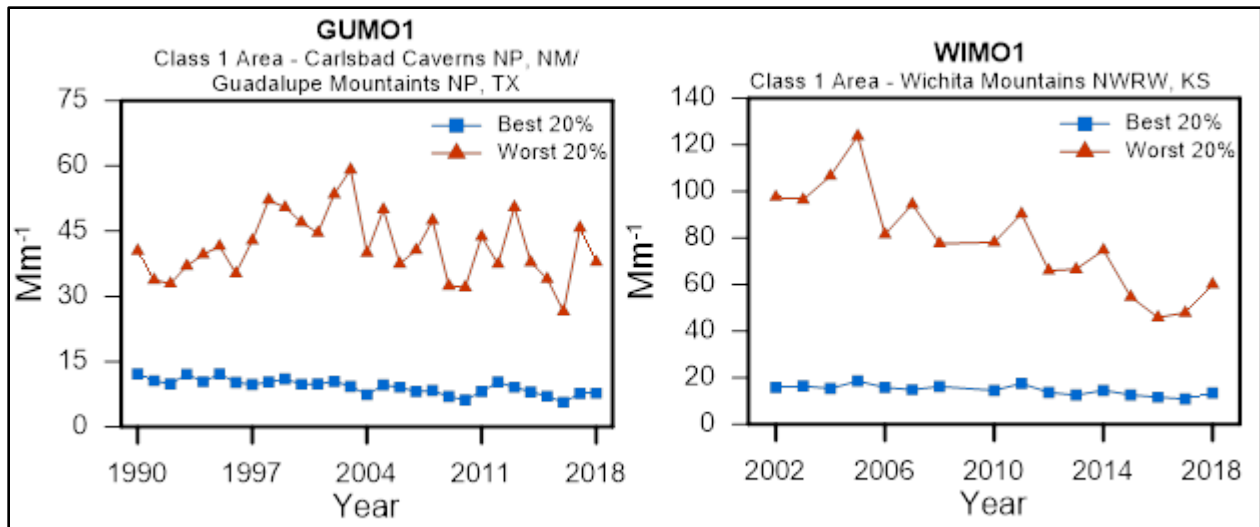


Figure 4. Visibility extinction trends for the Carlsbad Caverns National Park and Wichita Mountains National Wildlife Refuge (data retrieved from Colorado State University, 2020).

Trend lines for Class I areas affected by sources in Northwestern New Mexico (Figure 5) are similar to trend lines for Class I areas in New Mexico. While visibility on worst days at Guadalupe Mountains National Park may have diminished, a careful analysis of fire activity in the area would be necessary in order to draw conclusions about the cause of some peaks in recent years (Colorado State University 2020).

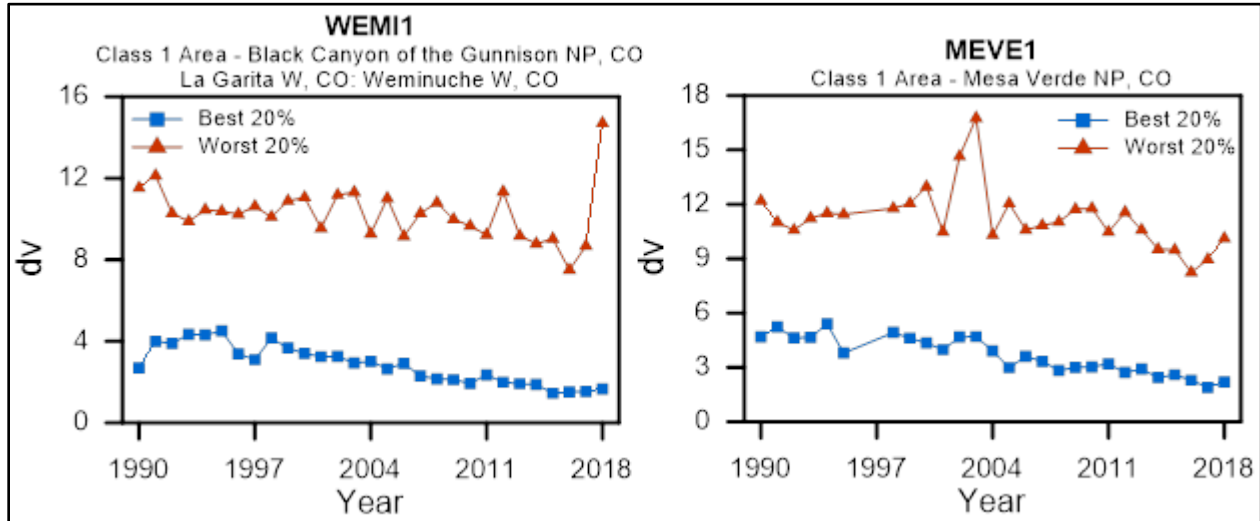


Figure 5. Visibility trends at Class I areas affected by sources in Northwestern New Mexico (data retrieved from Colorado State University, 2020)

A qualitative discussion of visibility impacts from oil and gas development in the Farmington RMP concludes that for the scenario modeled, which projected greater development than has occurred; there could potentially be significant impacts to visibility at Mesa Verde National Park, a Class I area in southwest Colorado. Occasional impacts to San Pedro Parks (northern New Mexico) and Weminuche (southern Colorado) Wilderness areas were also thought possible. However, visibility trends shown

below for San Pedro Parks, Mesa Verde, and Weminuche indicate that visibility on the best days has been flat to improving and visibility on worst days has shown little change over the period of record.

Visibility modeling was performed using the BLM CFO RFD potential oil and gas well development scenario and with mitigation using EPA's on-the-books emission controls and additional management controls. This analysis tiers to the modeling that was performed in the ARTSD (URS 2013) for the BLM CFO for results of visibility impairment, indicating that, for the Carlsbad region, visibility effects on Carlsbad Caverns National Park (CCNP) at the project level are minimal and not expected to be of concern (Engler et al. 2012, Engler and Cather 2014; URS 2013). The visibility screening analysis followed the recommendations in the FLAG Phase I Report – Revised Guidelines (USFS et al. 2010). The analysis relies on a 0.5 and 1.0 delta-deciview (change in visibility) threshold, calculated for base year 2008, base case 2017, and future RFD years. Non-project, aggregate emissions are driving the overall visibility effects. A refinement of the aggregate emissions would reduce the number of days of total visibility effects and would likely be closer to baseline and future visibility effects. Any refinement down to a smaller scope of development or project-specific level would likely reduce the number of days of total visibility effects that would be likely, closer to matching actual base and future visibility effects/baseline conditions (URS 2013). Further refinement of the URS 2013 visibility modeled results was performed to show relative effects. The results indicate that there are no days in which the threshold is exceeded at the project level for the CCNP. An additional study of Air Pollutant Emissions and Cumulative Air Impacts done for the CFO indicates that pollutants contributing to reductions in visibility are largely coming from outside the region (Applied Enviro Solutions 2011).

7.2 WET AND DRY POLLUTANT DEPOSITION

Deposition of pollutants through precipitation can result in acidification of water and soil resources in areas far removed from the source of the pollution, as well as causing harm to terrestrial and aquatic species. Some pollutants can also damage vegetation through direct or dry deposition. In general, the soils in New Mexico have a high acid neutralizing capacity and surface water is scarce, resulting in minimal impacts in this area. Also, the Acid Rain Program has resulted in greatly reduced levels of the most damaging pollutants. There are currently four wet deposition monitors in New Mexico including Gila Cliff Dwellings, Mayhill, Bandelier National Monument, and Capulin Volcano National Monument. In addition, monitors near the border at Mesa Verde and Guadalupe Mountains National Parks may shed some light on conditions in New Mexico. Data can be accessed through the National Atmospheric Deposition Network (NADP) at <http://nadp.sws.uiuc.edu/NTN/ntnData.aspx>. Wet deposition data is also available for monitoring sites in Kansas, Oklahoma, and Texas at this site.

The EPA has operated the Clean Air Status and Trends Network (CASTNET) since 1991 to provide data to assess trends in air quality, deposition, and ecological effects due to changes in air emissions. Sites are located in areas where urban influences are minimal. There are currently no CASTNET observation sites in New Mexico but there are three in Texas and one each in Oklahoma and Kansas. There is a CASTNET site at Mesa Verde National Park in the Four Corners region. National maps of pollutant concentrations can be found at <http://www.epa.gov/castnet/javaweb/airconc.html>. These maps show that New Mexico and most of the western states have much lower concentrations of all monitored pollutants than the eastern states and southern California. Nitrates are somewhat elevated in eastern Kansas and eastern Oklahoma but this is likely associated with agricultural activities rather than oil and gas development. The maps also show that the trend over the past 20 years has been for decreases in all pollutants in most areas of the country. As an example, Figures 6 and 7 show particulate nitrate and sulfate levels for 1990 and 2014. Maps of wet deposition data from NADP monitors are also available from the National

Atmospheric Deposition Program (National Atmospheric Deposition Program 2014). Total nitrogen deposition decreased by 44% from 1990 through 2014 in the eastern United States and decreased by 27% in the western United States between 1996 through 2014; however, total nitrate concentrations measured at the eastern sites were generally two to three times higher than concentrations measured at western reference sites. Total dry and wet sulfur deposition decreased by 82% from 1990 through 2014 in the eastern United States and decreased by 50% from 1996 through 2014 in the west, over 3-year mean periods. These trends in deposition levels are discussed in depth in the CASTNET annual report (AMEC Environment and Infrastructure, Inc. 2014).

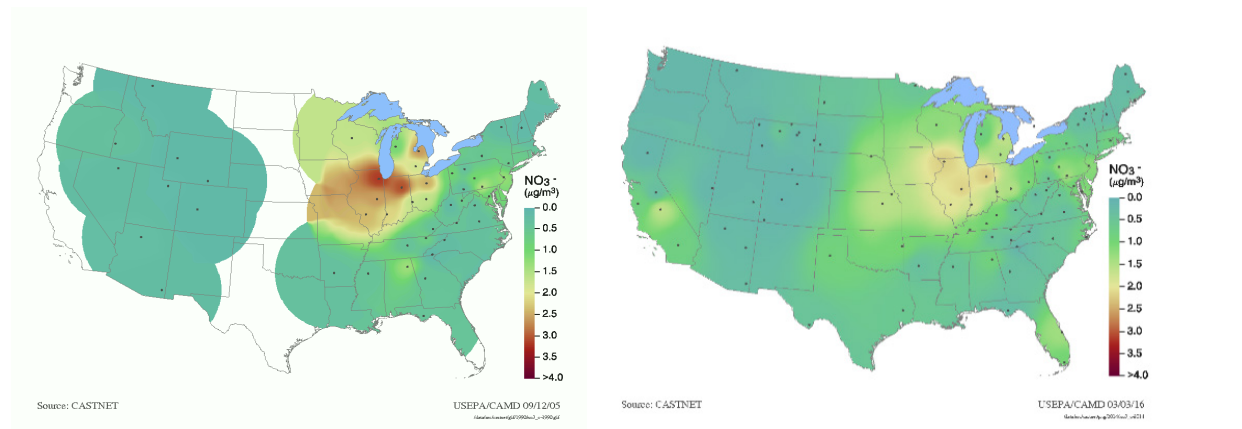


Figure 6. Particulate nitrate 1990 (left) and 2014 (right).

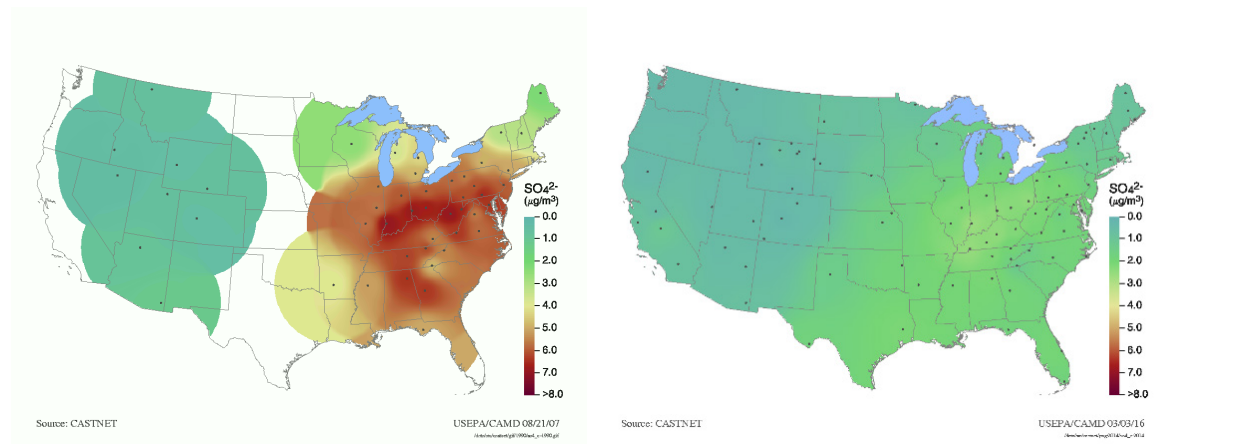


Figure 7. Particulate sulfate 1990 (left) and 2014 (right) (EPA 2015).

The WestJump analysis (ENVIRON et al. 2013) provides information about contributing sources to wet and dry deposition at IMPROVE monitoring sites. Pie charts showing the species contributing to nitrogen and sulfur deposition at Class I areas in New Mexico, Texas, Oklahoma, Kansas, Colorado, and Arizona are in Appendix G.

7.3 TERRESTRIAL EFFECTS OF OZONE

While other air pollutants may also negatively affect vegetation, ozone (O_3) is recognized as the one most likely to cause damage. Visible damage to leaf cells may be present in the form of spots or dead areas, though damage can be present long before it becomes visible. Decreased growth or altered carbon allocation may also occur. Ponderosa pine and aspen are species known to be sensitive to O_3 in the atmosphere (USFS et al. 2000)

An index has been developed to express cumulative seasonal impacts to vegetation. This is known as the W126 value. W126 is a cumulative metric that sums weighted hourly O_3 concentrations during daylight hours in the summer O_3 season. Figure 8 shows national W126 values for 2012 (AMEC Environment and Infrastructure, Inc. 2014). Higher W126 values were measured during 2012 in California, at high terrain sites in the west and at eastern sites with high daily 8-hour O_3 concentrations. At high elevations, moderate O_3 concentrations persist into the night due to lack of nighttime dry deposition and lack of fresh nitric oxide, both of which typically react with O_3 at night to reduce O_3 concentrations. The persistent, moderate O_3 concentrations at high elevation sites result in higher W126 levels, indicative of steady O_3 exposure for vegetation. In 2012, W126 values were higher than in 2011 because of higher ambient O_3 concentrations measured in 2012.

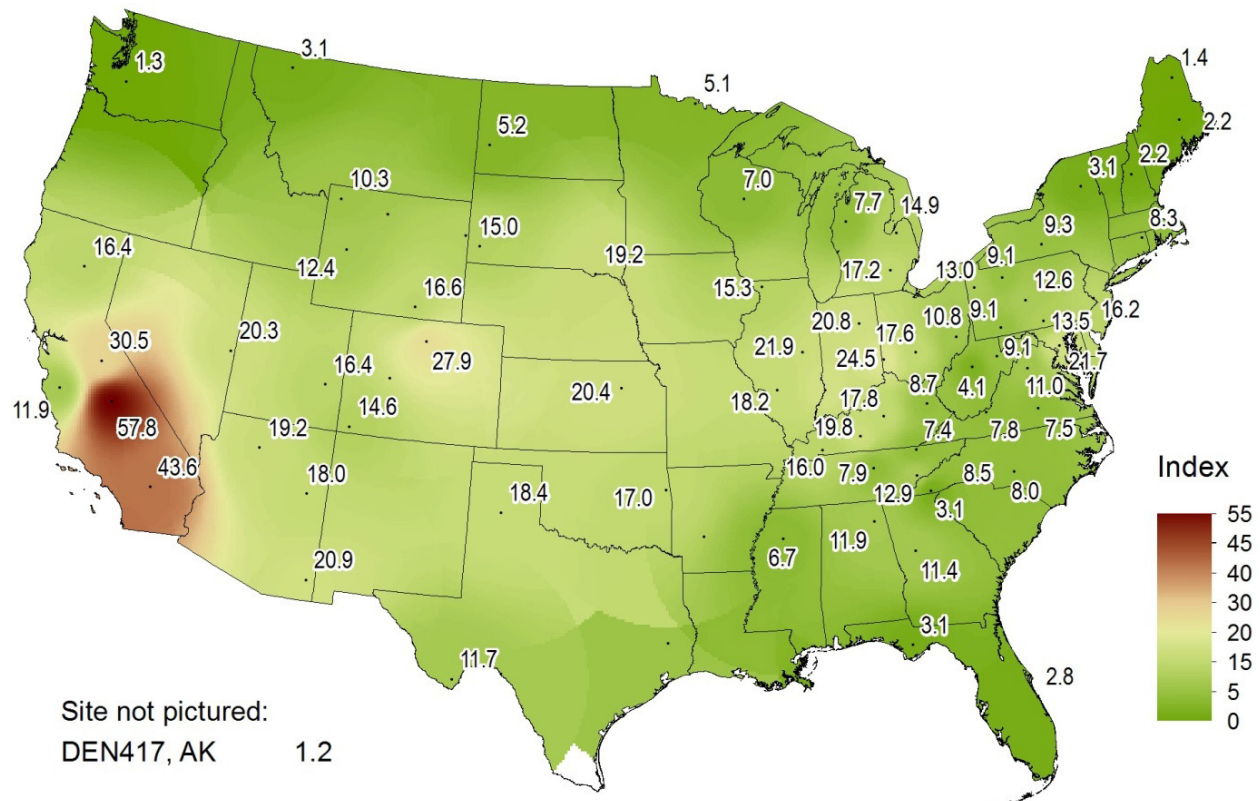


Figure 8. W126 values for 2012 in ppm-hour (AMEC Environment and Infrastructure, Inc. 2014).

7.4 VISIBILITY AND DEPOSITION MODELING

Visibility—Visibility modeling was performed using the CFO RFD potential oil and gas well development scenario and with mitigation using EPA’s on-the-books emission controls and additional management controls. This analysis tiers to the modeling that was performed in the ARTSD for the CFO for results of visibility impairment indicating that for the Carlsbad region, visibility impacts on CCNP at the project level are minimal and not expected to be of concern for the CCNP (Engler et al. 2012; URS 2013). The visibility screening analysis followed the recommendations in the FLAG Phase I Report – Revised Guidelines (USFS et al. 2010). The analysis relies on a 0.5 and 1.0 delta-deciview (change in visibility) threshold, calculated for base year 2008, base case 2017, and future RFD years. Non-project, cumulative emissions are driving the overall visibility impacts. A refinement of the cumulative emissions would reduce the number of days of total visibility impacts and would likely be closer to baseline and future visibility impacts. Any refinement down to a smaller scope of development or project-specific level would likely reduce the number of days of total visibility impacts that would be likely closer to matching actual base and future visibility impacts/baseline conditions (URS 2013). Further refinement of the URS 2013 visibility modeled results was performed to show relative impacts. The results indicate that there are no days in which the threshold is exceeded at the project level for the CCNP.

Deposition—Deposition modeling was performed using the CFO RFD potential oil and gas well development scenario and with mitigation using EPA’s on-the-books emission controls and additional management controls. This analysis tiers to the modeling that was performed in the ARTSD for results of nitrogen and sulfur deposition impairment (Engler et al. 2012; URS 2013).

To assess potential nitrogen and sulfur deposition impacts in the planning area, deposition impacts were compared to the NPS screening deposition analysis thresholds (DATs), which are defined as 0.005 kilogram per hectare per year (kg/ha/yr) in the western United States for both nitrogen and sulfur. A DAT is the additional amount of nitrogen or sulfur deposition within a Class I area, below which estimated impacts from a proposed new or modified source are considered to be insignificant. The DAT is a screening threshold that was developed primarily to assess impacts from a single stationary source (USFS et al. 2000, 2010). Modeling results showing deposition greater than a DAT do not strictly indicate the need for mitigation. If a DAT is exceeded, cumulative modeling may be required to demonstrate that cumulative deposition is below the level of concern (LOC). The LOC for the nitrogen and sulfur deposition values, defined by the NPS and USFS, is 3 kg/ha/yr for nitrogen and 5 kg/ha/yr for sulfur (Fox et al. 1989).

Results of analysis showed that the maximum annual nitrogen DAT at the project level was exceeded for CCNP but may be below the LOC at specific receptors. The CFO RFD modeling results showed that the predicted nitrogen deposition was expected to be below the LOC value of 3 kg/ha/yr for CCNP (Figure 9; URS 2013). The maximum annual sulfur DAT at the project level (CFO RFD) was below the DAT and LOC threshold for CCNP. Deposition rates that are below the LOC are believed to cause no adverse impacts. Appendix R and Appendix S of the ARTSD provide detailed nitrogen deposition results for project level and cumulative impacts, respectively (URS 2013).

Area with Greatest Predicted Impact	Maximum Modeled Project Deposition (kg/ha/yr)	DAT* (kg/ha/yr %)	Background Deposition (kg/ha/yr)	Total Project Deposition (kg/ha/yr)	LOC† (kg/ha/yr %)
Class I		<i>0.005</i>			<i>3.0</i>
Salt Creek Wilderness	0.29	5,800%	2.59	2.88	93%
Carlsbad Caverns National Park	0.19	3,800%	2.59	2.77	92%
Sensitive Class II		<i>0.005</i>			<i>3.0</i>
Bitter Lake National Wildlife Refuge	0.29	5,800%	2.59	2.88	93%
Grulla National Wildlife Refuge	0.11	2,200%	2.59	2.70	90%

Figure 9. Maximum Annual Nitrogen Deposition Source: URS 2013

* The DAT is shown in italics, while the maximum modeled deposition is provided as a percentage of the DAT.

† The LOC is shown in italics, while the maximum total deposition is shown as a percentage of the LOC.

To assess potential cumulative effects to AQRVs, the air quality assessment considers emissions and potential impacts of expected growth oil and gas development for nearby oil and gas basins as well as the Permian Basin, including the Raton Basin, San Juan Basin, Denver-Julesburg Basin, White River Field Office, Colorado River Valley Field Office, Utah Vernal Field Office, and Oklahoma, Kansas, and Texas Oil and Gas Basins (URS 2013). Cumulative scenario results were above for the Nitrogen and Sulfur LOC for CCNP. It should be noted that for a large aggregate project that includes thousands of sources (such as oil and gas development in the BLM CFO), deposition greater than the DAT as well as LOC is typical based on the uncertainty in the model parameters, and more refined modeling studies are often required to better understand potential effects. Future potential development in the region as a whole could result in degradation of air quality related to nitrogen deposition, depending on the number of sources present during development and any mitigation applied. Appropriate mitigation would be determined following further analysis at the site-specific APD stage of a project that allows for refined modeling analysis (as appropriate), which incorporates project-specific information.

In 2016, Chevron developed a Master Development Plan in which 436 oil and gas wells were projected to be developed on over 106 well pads. Although it is not anticipated that all wells will be developed concurrently during this project, similar results of AQRVs can be expected for large well development projects. The Chevron analysis extends the URS (2013) modeling that was performed and updates NO_x emissions in the project area. The results of acid deposition monitoring showed incremental exceedances of the nitrogen DAT of 0.005 kg/ha/yr in the CCNP during drilling operations but would be well below the DAT once drilling is completed (BLM 2016).

It is expected that a refined analysis may be required at the time of proposed lease development for well development that could potentially impact nitrogen deposition at the CCNP. A refined analysis of acid deposition must address the following criteria:

- Is the affected area sensitive to deposition?
- Is the affected area currently impacted by deposition?
- Have critical loads or target loads been developed for the affected area?
- Does current deposition exceed the critical load or target load?

This refined analysis should be in consultation with the NPS as prescribed in FLAG guidance (USFS et al. 2011). The Federal Land Managers will do their best to manage and protect resources at every area that they administer. Where possible, the most intrusive monitoring and instrumentation should be conducted adjacent to the Class I area if such areas adequately represent the area of concern. Federal Land Managers believe that the need to minimize potential impacts on a Class I area should be a major consideration in the best available control technology determination for a project proposed near such an area. Therefore, if a source proposes to locate near a Class I area, additional costs to minimize impacts on sensitive Class I resources may be warranted, even though such costs may be considered economically unjustified under other circumstances (USFS et al. 2010).

8 CUMULATIVE EFFECTS

More specific information about sources in New Mexico's oil and gas producing regions which have the greatest impacts on air quality and GHGs is included below. The CEQ regulations define cumulative effects as "the impact on the environment which results from the incremental impact of the action when added to other past, present, and reasonably foreseeable future actions regardless of what agency (Federal or non-Federal) or person undertakes such actions" (40 CFR 1508.7; DOI BLM 2008).

8.1 CURRENT AND REASONABLY FORESEEABLE CONTRIBUTIONS TO CUMULATIVE EFFECTS

A list of major sources (sources emitting more than 100 tons/year of CO, VOC, NO_x, SO₂, PM_{2.5}, or PM₁₀) in New Mexico, Kansas, Oklahoma, and Texas can be found in Appendix D. Any of these sources may contribute to cumulative effects within a local or regional context. All major sources in the figures represent emissions from the NEI report (EPA 2020b). Figures 10 through 15 show a map of major sources in the four-state area for pollutants of concern. These maps and their source data are also included in Appendix D as separate files available for download.

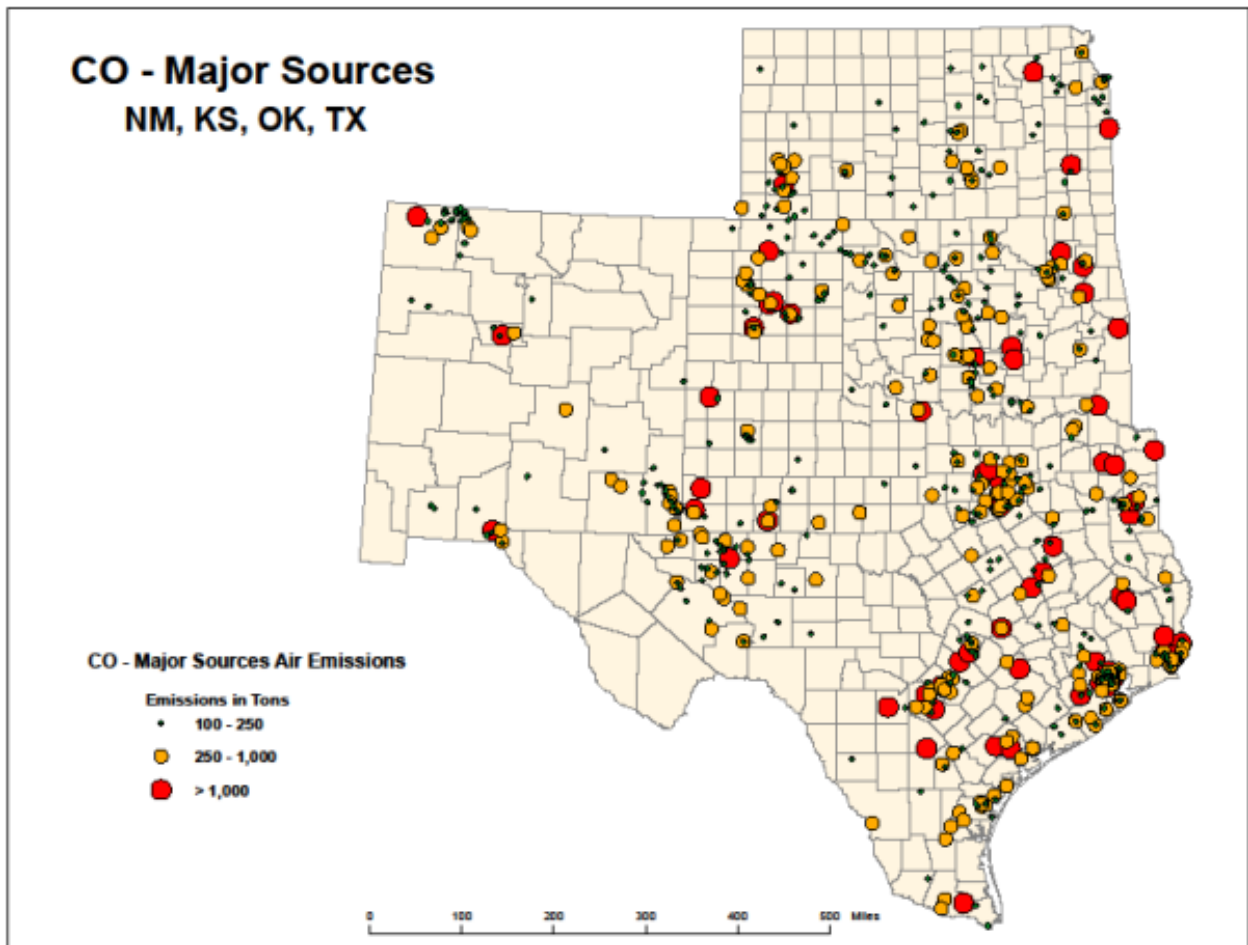


Figure 10. Major emissions sources (CO), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

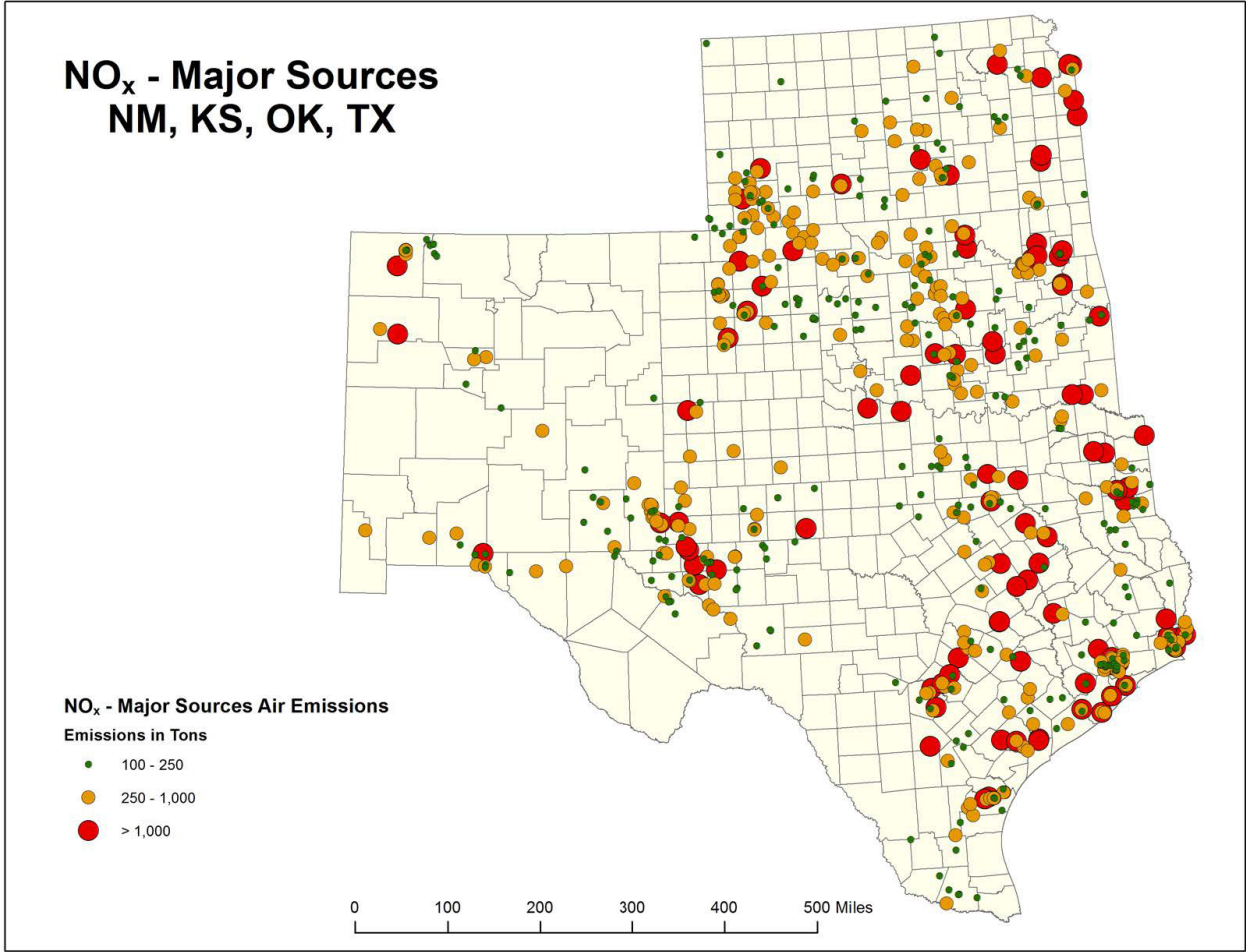


Figure 11. Major emissions sources (NO_x), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

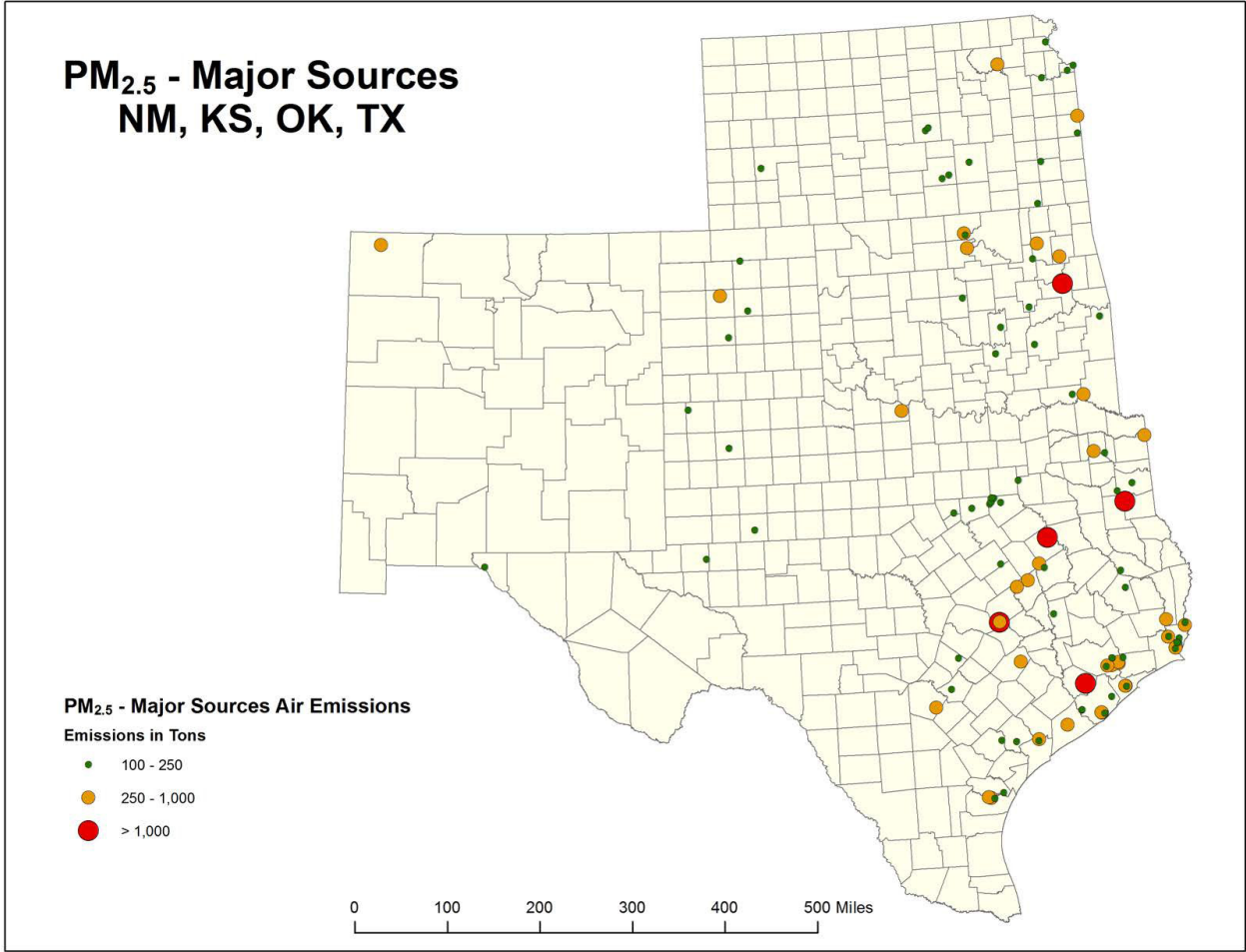


Figure 12. Major emissions sources (PM_{2.5}), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

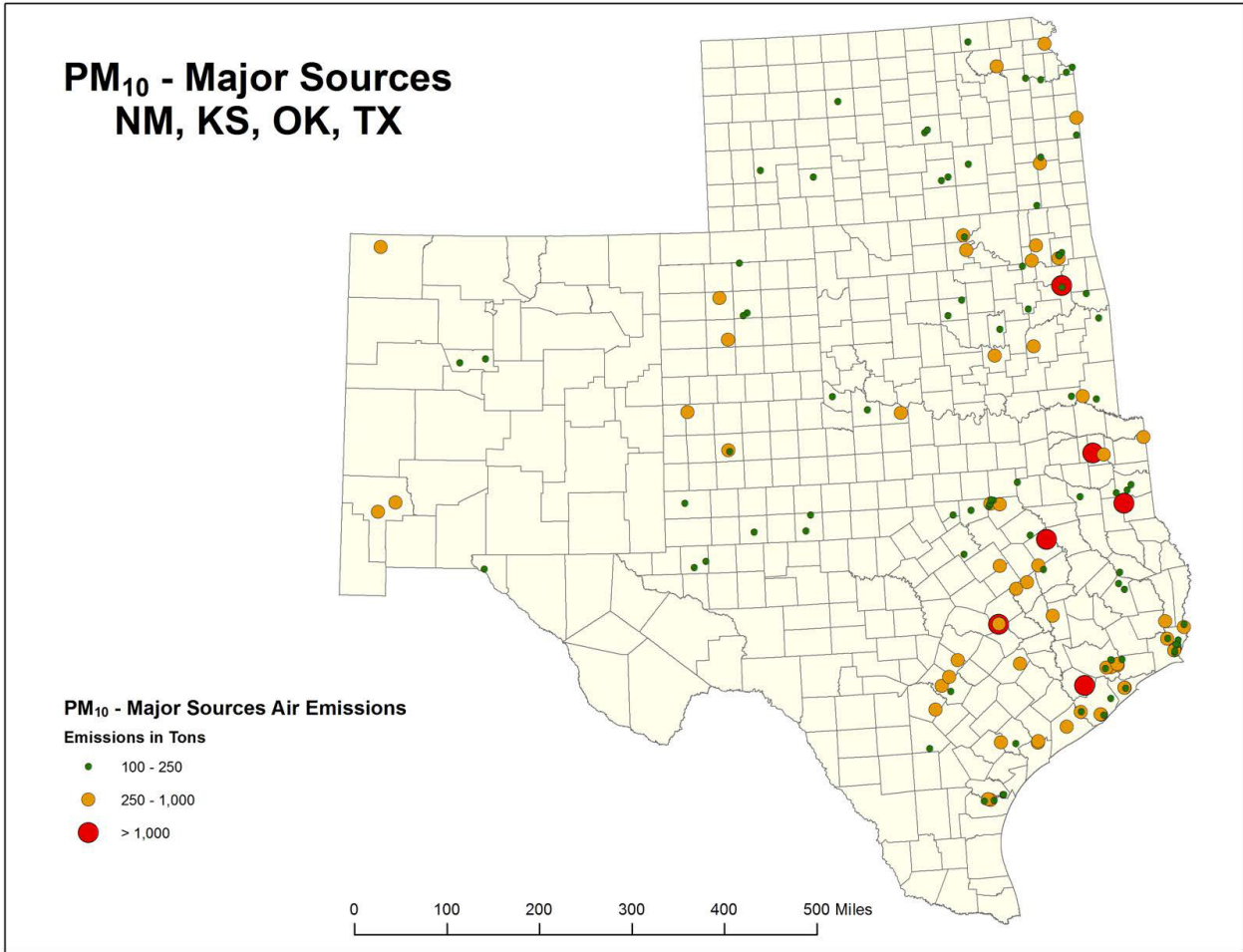


Figure 13. Major emissions sources (PM₁₀), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

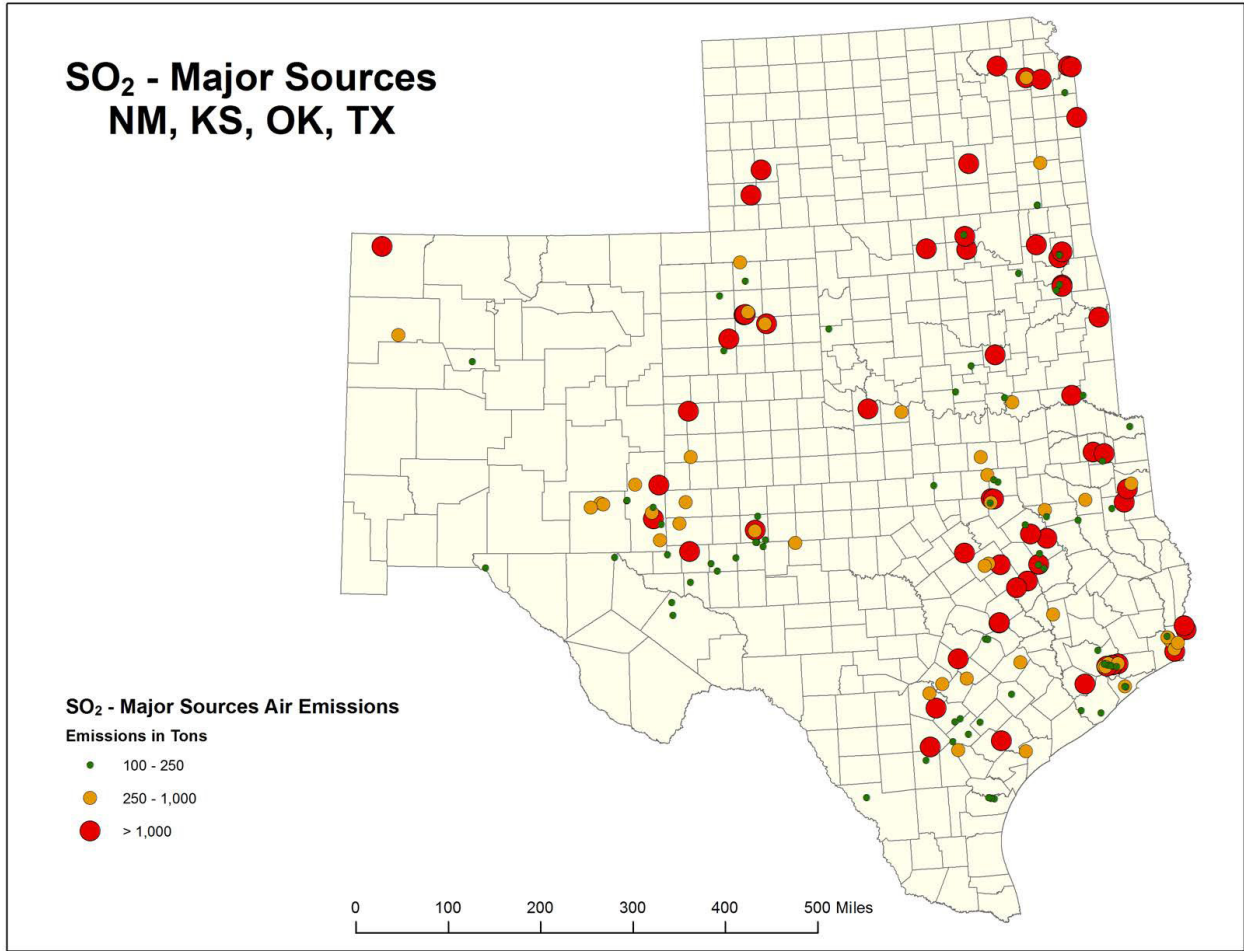


Figure 14. Major emissions sources (SO₂), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

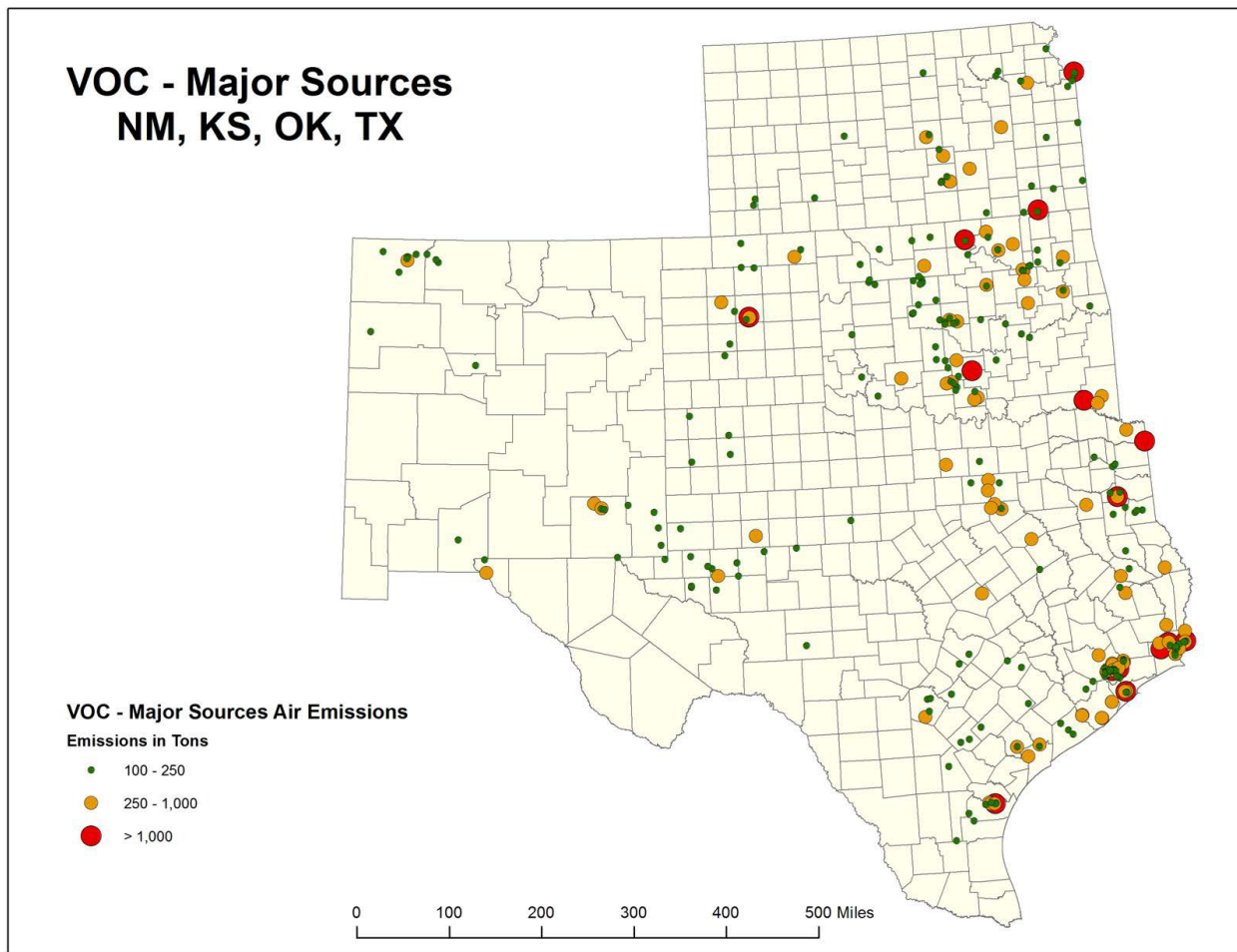


Figure 15. Major emissions sources (VOC), New Mexico, Kansas, Oklahoma, and Texas (EPA 2020b).

9 CLIMATE, CLIMATE CHANGE, AND GREENHOUSE GASES

9.1 CLIMATE

Climate is the composite of generally prevailing weather conditions of a particular region throughout the year, averaged over a series of years. Climate averages for 1981 to 2010, known as the current normal as defined by the World Meteorological Organization, are 30-year averages of temperature and precipitation for the previous 3 decades and are included in Appendix C. The next set of 30-year averages will be available again in 2021 for 1991 to 2020.

9.2 CLIMATE CHANGE

Climate change is a statistically significant and long-term change in climate patterns. The terms climate change and “global warming,” though often used interchangeably, are not the same. Climate change is any deviation from the average climate via warming or cooling and can result from both natural and human (anthropogenic) sources. Natural contributors to climate change include fluctuations in solar radiation, volcanic eruptions, and plate tectonics. Global warming refers to the apparent warming of

climate observed since the early twentieth century and is primarily attributed to human activities such as fossil fuel combustion, industrial processes, and land use changes.

Climate change may reinforce (positive feedback) or reduce (negative feedback) an expected temperatures increase. A feedback is the process by which changing one quantity results in the amplification or diminishment of another. An example of a positive feedback is the reduced albedo (reflectivity) of land surfaces from the melting of snow and ice. A warming climate is also expected to increase methane (CH₄) release from hydrates, thereby reinforcing the warming trend. There are also feedbacks related to carbon, water, and geochemical cycles. The results of most climate feedbacks are expected to amplify warming, but the exact magnitudes of these effects are difficult to quantify (IPCC 2013).

9.3 GREENHOUSE GASES

Atmospheric concentrations of naturally emitted greenhouse gases (GHGs) have varied for millennia and Earth’s climate fluctuated accordingly. However, since the beginning of the industrial revolution around 1750, human activities have significantly increased GHG concentrations and introduced man-made compounds that act as GHGs in the atmosphere. The atmospheric concentrations of carbon dioxide (CO₂), CH₄, and nitrous oxide (N₂O) have increased to levels unprecedented in at least the last 800,000 years. From pre-industrial times until today, the global average concentrations of CO₂, CH₄, and N₂O in the atmosphere have increased by around 40%, 150%, and 20%, respectively, Intergovernmental Panel on Climate Change (IPCC) 2013. Table 10 shows the average global concentrations of CO₂, CH₄, and N₂O in 1750, 2011 and 2017. Atmospheric concentrations of GHGs are reported in parts per million (ppm) and parts per billion (ppb).

Table 10. Average Global Concentrations of Greenhouse Gases in Select Years (IPCC 2007, 2013; EPA 2021n)

Greenhouse Gas	Pre-Industrial 1750	2011	2019	Increase 1750–2011
Carbon dioxide (CO ₂)	278 ppm	390.5 ppm	411 ppm ⁽¹⁾	47%
Methane (CH ₄)	722 ppb	1803 ppb	1866 ppb ⁽²⁾	167%
Nitrous oxide (N ₂ O)	270 ppb	324 ppb	331 ppb ⁽²⁾	23%

⁽¹⁾ The atmospheric CO₂ concentration is the 2019 annual average at the Mauna Loa, Hawaii, station (NOAA 2018). The concentration in 2019 at Mauna Loa was 409 ppm. The global atmospheric CO₂ concentration, computed using an average of sampling sites across the world, was 405 ppm in 2019 (EPA 2021n).

⁽²⁾ The values presented are global 2017 annual average mole fractions (EPA 2021n).

Human activities emit billions of tons of CO₂ every year. CO₂ is primarily emitted from fossil-fuel combustion but has a variety of other industrial sources. CH₄ is emitted from oil and natural gas systems, landfills, mining, agricultural activities, waste, and other industrial processes. N₂O is emitted from anthropogenic activities in the agricultural, energy-related, waste, and industrial sectors. The manufacture of refrigerants and semiconductors, electrical transmission, and metal production emit a variety of trace GHGs (including hydrofluorocarbons [HFCs], perfluorocarbons [PFCs], and sulfur hexafluoride [SF₆]). These trace gases have no natural sources and come entirely from human activities. CO₂, CH₄, N₂O, and the trace gases are considered well-mixed and long-lived GHGs. The atmospheric

lifespan for CO₂ can vary but is on the order of hundreds of years. The atmospheric lifespan of CH₄ and N₂O are 12.4 years and 121 years respectively (IPCC 2013).

9.4 OTHER GASES, ATMOSPHERIC AEROSOLS, AND PARTICULATES

Several gases do not have a direct effect on climate change, but indirectly affect the absorption of radiation by impacting the formation or destruction of GHGs. These gases include carbon monoxide (CO), oxides of nitrogen (NO_x), and non-methane volatile organic compounds (NMVOCs). Fossil fuel combustion and industrial processes account for the majority of emissions of these indirect GHGs. Unlike other GHGs, these gases are short-lived in the atmosphere.

Atmospheric aerosols, or particulate matter (PM), also contribute to climate change. Aerosols directly affect climate by scattering and absorbing radiation (aerosol-radiation interactions) and indirectly affect climate by altering cloud properties (aerosol-cloud interactions). Particles less than 10 micrometers in diameter (PM₁₀) typically originate from natural sources and settle out of the atmosphere in hours or days. Particles smaller than 2.5 micrometers in diameter (PM_{2.5}) often originate from human activities such as fossil fuel combustion. These so-called “fine” particles can exist in the atmosphere for several weeks and have local short-term impacts on climate. Aerosols can also act as cloud condensation nuclei (CCN), the particles upon which cloud droplets form.

Light-colored particles, such as sulfate aerosols, reflect and scatter incoming solar radiation, having a mild cooling effect, while dark-colored particles (often referred to as “soot” or “black carbon”) absorb radiation and have a warming effect. There is also the potential for black carbon to deposit on snow and ice, altering the surface albedo (or reflectivity), and enhancing melting. There is high confidence that aerosol effects are partially offsetting the warming effects of GHGs, but the magnitude of their effects contribute the largest uncertainty to our understanding of climate (IPCC 2013).

9.5 THE NATURAL GREENHOUSE EFFECT

The natural greenhouse effect is critical to the discussion of climate change. The greenhouse effect refers to the process by which GHGs in the atmosphere absorb heat energy radiated by Earth’s surface. Water vapor is the most abundant GHG, followed by CO₂, CH₄, N₂O, and several trace gases. Each of these GHGs exhibit a particular “heat trapping” effect, which causes additional heat retention in the atmosphere that would otherwise be radiated into space. The greenhouse effect is responsible for Earth’s warm atmosphere and temperatures suitable for life on Earth. Different GHGs can have different effects on the Earth’s warming due to their ability to absorb energy (“radiative efficiency”), and how long they stay in the atmosphere (“lifetime”). Without the natural greenhouse effect, the average surface temperature of the Earth would be about zero degrees Fahrenheit (°F). Water vapor is often excluded from the discussion of GHGs and climate change since its atmospheric concentration is largely dependent upon temperature rather than being emitted by specific sources.

9.6 GREENHOUSE GASES AND GLOBAL WARMING POTENTIALS

Common air emissions related to oil and gas activities include CO₂, CH₄, and N₂O. Other industries emit more potent GHGs including several fluorinated species of gases such as hydrofluorocarbons, perfluorocarbons, and sulfur hexafluoride. CO₂ is emitted from the combustion of fossil fuels (oil, natural gas, and coal), solid waste, trees and wood products, and as a result of other chemical reactions (e.g., manufacture of cement). The production and transport of coal, natural gas, and oil emit CH₄,

which can also be emitted from coal mining operations, naturally occurring coal CH₄ seepages, leaks from the oil and gas industry, livestock, and other agricultural practices and by the decay of organic waste in municipal solid waste landfills. Agricultural and industrial activities emit N₂O, as well as during combustion of fossil fuels and solid waste. Fluorinated gases are powerful GHGs that are emitted from a variety of industrial processes and are often used as substitutes for ozone-depleting substances (i.e., CFCs, HCFCs, and halons), but typically not from oil and gas operations. SF₆ is the most potent (highest radiative efficiency) GHG known and is typically used as an insulator in circuit breakers, gas-insulated substations, and switchgear used in the transmission system to manage the high voltages carried between power generating stations and customer load centers.

All of the different GHGs have various capacities to trap heat in the atmosphere, known as global warming potentials (GWPs). GWP is a relative measure that compares the heat absorbing ability of a certain mass of a gas relative to the same mass of CO₂. A second metric that is gaining prominence is Global Temperature change Potential (GTP). GTP is based on the change in global mean surface temperature at a chosen point in time, relative to that caused by CO₂. A number of other metrics may also be used, but no single metric accurately compares all consequences and the choice of metric is a value judgment (IPCC 2013).

Several different time horizons can express GWPs to fully account for the gases' ability to absorb infrared radiation (heat) over their atmospheric lifetime. The BLM uses the 100-year time horizon since most of the climate change impacts derived from climate models are expressed toward the end of the century. Also, in accordance with international GHG reporting standards under the United Nations Framework Convention on Climate Change (UNFCCC) and in order to maintain consistent comparisons over the years, official GHG emission estimates for the United States are reported based on the GWP values given in the Fourth Assessment Report (AR4) of the IPCC (IPCC 2007).

Updated GWPs are reported in the Fifth Assessment Report (AR5) as the level of scientific understanding increases. The atmospheric lifetimes and GWPs for the major GHGs over the 20-year and 100-year time horizons are listed below in Table 11 for comparison. CO₂ has a GWP of 1, and for the purposes of analysis a GHGs GWP is generally standardized to a carbon dioxide equivalent (CO₂e), or the equivalent amount of CO₂ mass the GHG would represent. In the AR5 report, CH₄ has a current GWP estimated to be 28 and N₂O has a GWP of 265 (IPCC 2013).

Table 11. Global Warming Potentials (100-year time horizon) (IPCC 2007, 2013)

Greenhouse Gas	GWP Values for 100-year Time Horizon	
	AR4 ⁽¹⁾	AR5
Carbon dioxide (CO ₂)	1	1
Methane (CH ₄)	25	28
Nitrous oxide (N ₂ O)	298	265
Select hydrofluorocarbons (HFCs)	124–14,800	4–12,400
Sulfur hexafluoride (SF ₆)	22,800	23,500

Greenhouse Gas	GWP Values for 20-year Time Horizon	
	AR4	AR5
Carbon dioxide (CO ₂)	1	1
Methane (CH ₄)	72	84
Nitrous oxide (N ₂ O)	299	264
Select hydrofluorocarbons (HFCs)	437–12,000	<1–10,800
Sulfur hexafluoride (SF ₆)	16,300	17,500

⁽¹⁾ For consistency with the EPA and its Inventory of Greenhouse Gas Reporting; we have represented values from AR4 of the IPCC report in this report.

9.7 CLIMATE CHANGE PROJECTIONS

Our current understanding of the climate system comes from the cumulative results of observations, experimental research, theoretical studies, and model simulations. Climate change projections are based on a hierarchy of climate models that range from simple to complex, coupled with comprehensive Earth System Models. For the Fifth Assessment Report (AR5), scientists estimated future climate impacts based on a range of Representative Concentration Pathways (RCPs) for well-mixed GHGs in model simulations carried out under the Coupled Model Intercomparison Project Phase 5 (CMIP5) of the World Climate Research Programme (IPCC 2013). The RCPs represent a range of mitigation scenarios that are dependent upon socio-economic and geopolitical factors and have different targets for radiative forcing (RF) in 2100 (2.6, 4.5, 6.0, and 8.5 W m⁻²). The scenarios are considered to be illustrative and do not have probabilities assigned to them.

AR5 uses terms to indicate the assessed likelihood of an outcome ranging from *exceptionally unlikely* (0%–1% probability) to *virtually certain* (99%–100% probability) and level of confidence ranging from *very low* to *very high*. The findings presented in AR5 indicate that warming of the climate system is unequivocal and many of the observed changes are unprecedented over decades to millennia. It is *certain* that Global Mean Surface Temperature (GMST) has increased since the late nineteenth century and *virtually certain* (99%–100% probability) that maximum and minimum temperatures over land have increased on a global scale since 1950. The globally averaged combined land and ocean surface temperature data show a warming of 0.85 degrees Celsius (°C) (1.5°F) (IPCC 2013; NOAA 2013). Human influence has been detected in warming of the atmosphere and the ocean, in changes in the global water cycle, in reductions in snow and ice, in global mean sea level rise, and in changes in some climate extremes. It is *extremely likely* (95%–100% probability) that human influence has been the dominant cause of the observed warming since the mid-twentieth century (IPCC 2013).

Additional near-term warming is inevitable due to the thermal inertia of the oceans and ongoing GHG emissions. Assuming there are no major volcanic eruptions or long-term changes in solar irradiance, global mean surface temperature increase, for the period 2016 to 2035 relative to 1986 to 2005, will likely be in the range of 0.3°C to 0.7°C (0.5°F–1.3°F). Global mean temperatures are expected to continue rising over the twenty-first century under all of the projected future RCP concentration scenarios. Global mean temperatures in 2081 to 2100 are projected to be between 0.3°C to 4.8°C (0.5°F–8.6°F) higher relative to 1986 to 2005 (IPCC 2013). The IPCC projections are consistent with reports from other organizations (NASA 2013; Joint Science Academies 2005).

Findings from AR5 and reported by other organizations (NASA 2013; NOAA 2013) also indicate that changes in the climate system are not uniform and regional differences are apparent. Some regions will experience precipitation increases, and other regions will have decreases or not much change. The contrast in precipitation between wet and dry regions and between wet and dry seasons is expected to increase. The high latitudes are *likely* (66%–100% probability) to experience greater amounts of precipitation due to the additional water carrying capacity of the warmer troposphere. Many mid-latitude arid and semi-arid regions will *likely* (66%–100% probability) experience less precipitation (IPCC 2013).

9.7.1 GENERAL CLIMATE CHANGE PREDICTIONS

Climate change is a global process that is impacted by the sum total of GHGs in the Earth's atmosphere. Currently, Global Climate Models are unable to forecast local or regional effects on resources (IPCC 2013). However, there are general projections regarding potential impacts to natural resources and plant and animal species that may be attributed to climate change from GHG emissions over time; however, these effects are likely to be varied, including those in the southwestern United States (Karl 2009). For example, if global climate change results in a warmer and drier climate, increased particulate matter impacts could occur due to increased windblown dust from drier and less stable soils. Cool season plant species' spatial ranges are predicted to move north and to higher elevations, and extinction of endemic threatened or endangered plants may be accelerated. Due to loss of habitat or competition from other species whose ranges may shift northward, the populations of some animal species may be reduced or increased. Less snow at lower elevations would likely impact the timing and quantity of snowmelt, which, in turn, could impact water resources and species dependent on historic water conditions (Karl 2009).

Climate change will impact regions differently and warming will not be equally distributed. Both observations and computer model predictions indicate that increases in temperature are likely to be greater at higher latitudes, where the temperature increase may be more than double the global average. Warming of surface air temperature over land will very likely be greater than over oceans (IPCC 2013). There is also high confidence that warming relative to the reference period will be larger in the tropics and subtropics than in mid-latitudes. Frequency of warm days and nights will increase and frequency of cold days and cold nights will decrease in most regions. Warming during the winter months is expected to be greater than during the summer and increases in daily minimum temperatures are more likely than increases in daily maximum temperatures. Models also predict increases in duration, intensity, and extent of extreme weather events. The frequency of both high and low temperature events is expected to increase. Near- and long-term changes are also projected in precipitation, atmospheric circulation, air quality, ocean temperatures and salinity, and sea ice cover.

9.7.2 REGIONAL CLIMATE CHANGE PREDICTIONS

In the region encompassing southern Colorado and New Mexico, average temperatures rose just under 0.7°F decade between 1971 and 2011, which is approximately double the global rate of temperature increase (Rahmstorf 2012). These rates of warming are unprecedented over the past 11,300 years (Marcott 2013). Climate modeling suggests that average temperatures in this region may rise by 4°F to 6°F by the end of the twenty-first century, with warming increasing from south to north. By 2080 to 2090, the southwestern United States are projected to see a 10% to 20% decline in precipitation, primarily in winter and spring, with more precipitation falling as rain rather than snow compared with historical trends (Cayan 2013).

In a recent report, the U.S. Bureau of Reclamation (U.S. Bureau of Reclamation et al. 2013) made the following projections through the end of the twenty-first century for the Upper Rio Grande Basin (southern Colorado to central southern New Mexico) based on the current and predicted future warming:

- There will be decreases in overall water availability by one quarter to one third.
- The seasonality of stream and river flows will change with summertime flows decreasing.
- Stream and river flow variability will increase. The frequency, intensity and duration of both droughts and floods will increase.

Texas, Oklahoma, and Kansas are part of the Great Plains region, which will see increases in temperatures and more frequent drought in the future. Temperature increases and precipitation decreases will stress the region's primary water supply, the Ogallala Aquifer. Seventy percent (70%) of the land in this area is used for agriculture. Threats to the region associated with climate change include:

- Pest migration as ecological zones shift northward;
- Increases in weeds; and
- Decreases in soil moisture and water availability (U.S. Bureau of Reclamation et al. 2013).

9.7.3 STATE CLIMATE CHANGE TRENDS AND PREDICTIONS

NOAA National Centers for Environmental Information released its Climate Summaries by state in 2017 with some updated 2019 information also available. The key messages bulleted below in Sections 9.7.3.1 – 9.7.3.4 represent climate summary information for each state within the New Mexico State Office (NMSO) jurisdiction. More detailed climate discussions for each state can be found through the State Climate Summaries (Revised 2019) webpage and documents (NOAA 2019).

9.7.3.1 NEW MEXICO

- Average annual temperature has increased by almost 2°F since the 1970s, and the number of hot days and warm nights has increased. Historically unprecedented future warming is likely.
- The summer monsoon rainfall, which provides much needed water for agricultural and ecological systems, varies greatly from year to year and future trends in such precipitation are highly uncertain.
- Droughts are a serious threat in this water-scarce state. Drought intensity is projected to increase and snowpack accumulation is projected to decrease, which will pose a major challenge to New Mexico's environmental, agricultural, and human systems. Wildfire frequency and severity are projected to increase in New Mexico (Frankson, Kunkel, Stevens, and Easterling 2017).

9.7.3.2 OKLAHOMA

- Average annual temperature has increased by less than 1°F since the early twentieth century. Winter warming has been characterized by the much below average occurrence of extremely cold days since 1990. Under a higher emissions pathway, historically unprecedented warming is projected by the end of the twenty-first century.

- Precipitation can vary greatly from year to year in this region of transition from humid to semi-arid conditions. Heavy precipitation events are projected to increase, which may increase the risk of flooding and associated increases in soil erosion and non-point source runoff into streams and lakes.
- The agricultural economy of Oklahoma makes the state particularly vulnerable to droughts, several of which have occurred in recent years. Higher temperatures will increase the rate of soil moisture depletion, leading to an increase in the intensity of naturally occurring future droughts (Frankson, Kunkel, Stevens, Champion, and Stewart 2017a).

9.7.3.3 KANSAS

- Average annual temperature has increased about 2°F since the early twentieth century, with greater warming in the winter and spring than in the summer and fall. The number of very cold nights has been much below average since 1990. Under a higher emissions pathway, historically unprecedented warming is projected by the end of the twenty-first century.
- Precipitation has varied greatly from year to year in this region of transition from humid conditions in the east of the state to semi-arid conditions in the west. Projected increases in winter precipitation and decreases in summer precipitation may result in both beneficial and negative impacts.
- The agricultural economy of Kansas makes the state vulnerable to droughts and heat waves, several of which occurred in the 1930s, 1950s, and in recent years. Projected increases in temperatures may increase the intensity of future droughts. The frequency of wildfire occurrence and severity is also projected to increase in Kansas (Frankson, Kunkel, Stevens, Easterling, Lin, and Shulski 2017).

9.7.3.4 TEXAS

- Mean annual temperature has increased by approximately 1°F since the first half of the twentieth century. Under a higher emissions pathway, historically unprecedented warming is projected by the end of the twenty-first century, with associated increases in extreme heat events.
- Although projected changes in annual precipitation are uncertain, increases in extreme precipitation events are projected. Higher temperatures will increase soil moisture loss during dry spells, increasing the intensity of naturally occurring droughts.
- The number of landfalling hurricanes in Texas is highly variable from year to year. As the climate warms, increases in hurricane rainfall rates, storm surge height due to sea level rise, and the intensity of the strongest hurricanes are projected (Frankson, Kunkel, Stevens, Champion, and Stewart 2017b).

9.7.4 CUMULATIVE CLIMATE CHANGE SUMMARY

Existing conditions of climate change in any given location are the result of numerous complex factors, both natural and human caused. Natural factors contributing to the current condition of air resources include existing climate resulting from long-term atmospheric weather patterns, soil types, and vegetation types. Anthropogenic factors contributing to the current condition of air resources include

long-term human habitation, growing human populations, transportation methods and patterns, recreational activities, economic patterns, the presence of power plants and other industrial sources. The presence of natural resource (primarily oil and natural gas) extraction and processing on some BLM lands also impact air quality and GHG emissions.

The IPCC concludes in AR5 that “cumulative emissions of CO₂ largely determine global mean surface warming by the late twenty-first century and beyond.” Most aspects of climate change will persist for many centuries even if emissions of CO₂ are stopped. This represents a substantial multi-century climate change commitment created by past, present and future emissions of CO₂ (IPCC 2013). Increasing concentrations may accelerate the rate of climate change in the future.

9.7.5 GLOBAL CARBON BUDGET DISCUSSION

Human activities are estimated to have caused approximately 1.0°C of global warming¹ above pre-industrial levels, with a likely range of 0.8°C to 1.2°C. Global warming is likely to reach 1.5°C between 2030 and 2052 if it continues to increase at its current rate (high confidence) (IPCC 2018).

Climate models project robust² differences in regional climate characteristics between present-day and global warming of 1.5°C, and between 1.5°C and 2.0°C. These differences include increases in

- mean temperature in most land and ocean regions (high confidence),
- hot extremes (temperatures) in most inhabited regions (high confidence),
- heavy precipitation in several regions (medium confidence), and
- drought and precipitation deficits in some regions (medium confidence) (IPCC 2018).

Limiting global warming requires limiting the total aggregate global anthropogenic emissions of CO₂ since the pre-industrial period, that is, staying within a total carbon budget (high confidence). Carbon budgeting, as defined by IPCC, refers to three concepts:

- an assessment of carbon cycle sources and sinks on a global level,
- the aggregate amount of global CO₂ emissions estimated to limit global surface temperatures to a given level above a reference period, and
- the distribution of the carbon budget defined under the regional, national, or subnational levels based on considerations of equity, cost, or efficiency (IPCC 2018).

There are at least 12 carbon budget studies with estimates that focus on limiting warming (50%, 66% probabilities, etc.) to below 1.5°C and 2.0°C (Carbon Brief 2018). Some of these studies are based on Earth System Models (ESMs), some on combined observations and ESMs, and others on integrated assessment models, all of which use varying degrees of interim physics dynamics and data methodologies to provide carbon budget estimates.

¹ Present level of global warming is defined as the average of a 30-year period centered on 2017 assuming the recent rate of warming continues.

² Robust is used here to mean that at least two-thirds of climate models show the same sign of changes at the grid point scale, and that differences in large regions are statistically significant.

Sizable uncertainties are reflected in these estimates as many different approaches are modeled into the carbon budget estimates. Some studies even show that the global carbon budget to limit warming below 1.5°C has already been expended. Attempting to show the relationship between a carbon budget and warming trends is not direct and linear and can vary drastically based on

- disagreement about the definition of “surface temperature,”
- the definition of the “pre-industrial” period,
- the observational temperature data sets that should be used,
- what happens to non-CO₂ factors that influence the climate, and
- whether Earth-system feedbacks like thawing permafrost are considered.

Drastic changes can also occur when there are net-negative emissions or depending on how quickly climate-cooling aerosols are reduced (Carbon Brief 2018). While levels of uncertainty are not uncommon to scientific projections, the challenge with carbon budgets is not the uncertainty in the budget but rather that the uncertainty is “substantial” as identified by the IPCC (2018).

Using the global mean surface air temperature, as in IPCC’s AR5, an estimate of 580 gigatonnes (Gt) CO₂ is used as the 2018 baseline for the remaining carbon budget to limit warming to 1.5°C with a 50% probability through 2100. While these estimates can be used as a baseline, “substantial” uncertainty exists (IPCC 2018). The IPCC 2018 Special Report states that the following uncertainties exist:

- the climate response from CO₂ and non-CO₂ emissions is ±400 Gt CO₂,
- the level of historic warming contributes ±250 Gt CO₂ of uncertainty,
- potential additional carbon release from future permafrost thawing and CH₄ release from wetlands would reduce budgets by up to 100 Gt CO₂ and another ±250 Gt CO₂ from future non-CO₂ mitigation efforts.

The uncertainties listed above could indicate the carbon budget could be -420 Gt CO₂ (meaning that the carbon budget to keep warming below 1.5°C may have already been expended). Similarly, the range of uncertainties listed above could result in a carbon budget that is up to 900 Gt CO₂ larger (meaning that the total carbon budget could be up to 1480 Gt to keep temperatures below 1.5°C by the end of the century). Therefore, although 580 Gt CO₂ represents a 50% chance to limit emissions to less than 1.5°C by the end of the century, the actual budget may be drastically different.

10 GREENHOUSE GAS ANALYSIS AND METHODOLOGIES

Fossil fuel extraction, construction and operation (well development), and processing and end-use production activities all contribute to air pollutants and GHG emissions in the FFO and CFO areas, especially San Juan, Northwestern Sandoval, Eddy, Lea, and Chaves counties as well as in parts of Oklahoma, Kansas and Texas. This includes midstream sources from the natural gas compressor stations and pipelines, gas plants, and petroleum refining as well as final downstream end-use by the consumer. Coal mining is also occurring in the FFO and Oklahoma Field Office (OFO) areas. Potash mining in the CFO area also contributes to air contaminant and GHG emissions.

Methodologies appropriate to GHG analysis are very different from those appropriate for air pollutant analysis. Air quality models used to predict concentrations and transport of air pollutants are not applicable to well-mixed, long-lived greenhouse gases (GHGs) which impact the atmosphere on a global scale. Global Climate Models (GCM's) cannot currently be downscaled to accurately relate GHG emissions to regional or local-scale impacts. The GHG emissions data derived from analytical tools (such as emission calculators) may be used to compare project level emissions with state, national and global emissions. However, such a comparison, while it provides context, may not always be useful information regarding impacts since project level emissions are often orders of magnitude less than national level emissions and because GHG impacts inherently cumulative. Comparisons of GHG emissions among project alternatives and an analysis of the resiliency of different project alternatives to the effects of climate change may provide more useful information. When modeling and analyzing GHG emissions, the primary sources of GHG emissions include:

- Fossil fuel combustion for construction and operation (well development) of oil and gas facilities – vehicles driving to and from production sites, engines that drive drill rigs, etc. produce CO₂ in quantities that vary depending on the age, types, and conditions of the equipment as well as the targeted formation, locations of wells with respect to processing facilities and pipelines, and other site-specific factors.

Combustion of produced oil and gas: it is expected that development will produce marketable quantities of oil and/or gas. Combustion of the oil and/or gas would release CO₂ into the atmosphere. Fossil fuel combustion is the largest source of global CO₂.

Estimated emissions for CO₂ are obtained from the calculator (Appendix H) for the drilling and operational phases of the well, as well as for other ancillary aspects of well development. These values include emissions from combustion engines used to construct and maintain the well.

- Methane (CH₄) releases from gas well development result from venting of natural gas during the well completion process, actuation of gas operated valves during well operations, and fugitive gas leaks along the infrastructure required for the production and transmission of gas. This is a major source of global CH₄ emissions. These emissions have been estimated for various aspects of the energy sector, and starting in 2011, producers were required under 40 CFR 98, to estimate and report their CH₄ emissions to the EPA (EPA 2021n). Estimated emissions for CH₄ are obtained from the calculator. These values include emissions from combustion engines used to construct and maintain the well (operations). No CH₄ emissions are predicted from ancillary construction operations.

Although incremental contributions to global GHGs from a proposed land management action cannot currently be translated into effects on climate change globally or in the area of any site-specific action, we can use GHG emission volumes as a proxy in determining impacts. In this way we can estimate emissions from a project or land management action and then compare those activities to the regional, national, or global level of GHGs or GHGs emitted by a certain industry within a region.

The BLM has jurisdiction over federal oil and gas exploration, field operations and well site-production on Federal and Indian mineral estate. Once produced oil or gas leaves the well location (via pipeline or tanker truck), the BLM no longer has jurisdiction over these products. However, it is often necessary to estimate and analyze downstream GHG emissions more completely until the product is finally combusted (end-use).

For the purpose of NEPA analysis, EPA emission factors can be used to include a qualitative and quantitative analysis of possible GHG emissions that could occur as a result of reasonably foreseeable coal, oil, gas or other development connected to federal land and resource management use. Estimates are made based on readily available data and reasonable assumptions about potential future development. More detailed emissions analysis can be qualitatively discussed and calculated at a site-specific level of analysis such as those that occur at an APD stage. Estimating direct and indirect GHG emissions attempt to provide a more complete GHG lifecycle of a well from site inspection to possible emissions through combustion.

10.1 DIRECT OIL AND GAS EMISSIONS

Direct GHG emissions from speculative future oil and gas well development and production at both the RFD and project level can be estimated based on representative well characteristics. Total carbon dioxide equivalent (CO₂e), which includes direct emissions of carbon dioxide (CO₂), CH₄, and nitrous oxide (N₂O) with applied GWPs, from an oil or gas producing well, is calculated.

10.1.1 WELL DEVELOPMENT GREENHOUSE GAS EMISSIONS

Increased GHG emissions as a can be connected to well development. The most substantial GHGs emitted by oil and gas development and production are CO₂ and CH₄. To facilitate quantification, most project-level analysis tend to assume that all wells would be developed concurrently and in the same year, though it is more likely that future potential development would not occur in this manner. Emission calculations for construction, operations, maintenance, and reclamation are included in Appendix H (Calculator) section.

Construction emissions for either an oil or gas well include well pad construction (fugitive dust), heavy equipment combustive emissions, commuting vehicles and wind erosion. Emissions from operations for an oil well include well workover operations (exhaust and fugitive dust), well site visits for inspection and repair, recompletion traffic, water and oil tank traffic, venting, compression and well pumps, dehydrators, and compression station fugitives. Operations emissions for a gas well include well workover operations (exhaust and fugitive dust), wellhead and compressor station fugitives, well site visits for inspection and repair, recompletions, compression, dehydrators, and compression station fugitives. Maintenance emissions for either an oil and gas well are for road travel, and reclamation emission activities. Interim and final activities include emissions from truck traffic, a dozer, blade, and track hoe equipment.

Emissions are anticipated to be at their highest level during the construction and completion phases of implementation (approximately 30 days in duration) because these phases require the highest degree of earth-moving activity, heavy equipment use, and truck traffic, compared with the operations and maintenance phases of implementation. Emissions are anticipated to decline during operations and maintenance as the need for earth-moving and heavy equipment declines.

Table 12 provides past well completion data and associated GHG emissions (CO₂e) based on APD activity from the BLM AFMSS I system (BLM 2021b). GHG emissions (CO₂e) are calculated for the total number of well completions using a per well emission factor based on activities during well. The emissions provide a maximum emissions scenario as the number of wells each year is multiplied by approximately 1,229 and 1,253 metric tons of CO₂e/year in the FFO and Pecos District Office (PDO)

respectively, which assumes all wells are gas wells for conservatism, since gas wells are estimated to have higher GHG emissions than oil wells.

Table 12. Well Completions and estimated GHG emissions based on APD Activity (BLM 2021b)

Farmington Field Office	2016	2017	2018	2019	2020	BLM RFD (2018–2037)	# of Wells Developed after 2018⁽³⁾
Number of BLM well completions ⁽¹⁾	19	8	61	22	0	1,980	83
Metric tons of CO ₂ e/year	23,351	9,832	74,969	27,038	0	2,433,420	102,007
Pecos District Office	2016	2017	2018	2019	2020	BLM RFD (2016–2035)	# of Wells Developed after 2016⁽³⁾
Number of BLM well completions ⁽²⁾	150	199	261	284	174	6,400	1163
Metric tons of CO ₂ e/year	187,950	249,347	327,033	355,852	218,022	8,019,200	1,457,239

Note: Methodology updated to use SNT.50 Reports from AFMSS. Wells completed reported from BLM AFMSS 1 with run date April 2021 (BLM 2021b).

⁽¹⁾ FFO number of BLM federal and non-federal wells in PDO RFD (2016–2037) is 3,200. FFO BLM wells includes completions from Farmington and Rio Puerco Field Offices.

⁽²⁾ PDO number of BLM federal and non-federal wells in PDO RFD (2016–2037) is 16,000. PDO BLM wells includes completions from Carlsbad, Hobbs, and Roswell Field Offices.

⁽³⁾ The number of wells developed after the start of the RFD for each respective field office (2018 for FFO and 2016 for PDO) are presented to disclose the current levels of development toward the total reasonably foreseeable development projections.

10.2 INDIRECT GREENHOUSE GAS EMISSIONS

Indirect GHG emissions are estimated based on speculative annual oil, gas, and/or natural gas liquids produced from oil and gas development. Indirect GHG emissions are calculated based on emission factors from 40 CFR Part 98 Subpart C which include default fuel heat values for crude oil and natural gas as well as their respective default emission factors for CO₂, CH₄, and N₂O. 40 CFR Part 98 Subpart C emission factors used in BLM GHG Indirect Greenhouse Gas Emissions are included below.

- Crude oil indirect emission factors:
 - 0.138 mmBtu/gallon
 - 74.54 kilogram CO₂/mmBtu
 - 0.003 kilogram CH₄/mmBtu
 - 0.0006 kg N₂O/mmBtu
- Gas indirect emission factors:
 - 1026 Btu/scf

- 53.06 kg CO₂/mmBtu
- 0.001 kg CH₄/mmBtu
- 0.0001 kg N₂O/mmBtu

10.2.1 OIL AND GAS PRODUCTION (DOWNSTREAM EMISSIONS [END-USE])

Estimates of production (or downstream/end use) GHG emissions are dependent on projected oil and gas production volumes. The BLM does not direct or regulate the end use of produced oil and/or gas. The challenge for estimating downstream emissions comes with understanding when and how oil and gas would be distributed and used for energy. It can be reasonably assumed the oil and gas produced on leases would be combusted primarily for electricity generation, transportation, industry, agriculture, commercial, and residential uses. From this assumption, the BLM provides potential GHG emissions estimates using currently available GHG emissions data.

Table 13 details the latest oil and gas production volumes on federal lands within the jurisdiction of the BLM New Mexico as well as the United States and state-level production as a whole.

Table 13. 2019 Oil and Gas Production (DOI ONRR 2020; EIA 2020)

Location	Oil (Mbbbl)	% U.S. Total	Gas (MMcf)	% U.S. Total
United States	4,470,528	100	40,982,458	100
New Mexico	329,483	7.37	1,852,719	4.52
Federal leases NM ⁽¹⁾	121,556	2.72	1,046,482	2.55
San Juan Basin	4,891	0.11	419,696	1.02
Permian Basin	154,266	3.45	326,451	0.79
Kansas	33,193	0.74	183,097	0.44
Federal leases KS ⁽²⁾	165	0.004	3,030	0.007
Oklahoma	211,808	4.74	3,175,008	7.74
Federal leases OK ⁽²⁾	935	0.02	13,966	0.03
Texas	1,850,715	41.40	10,355,453	25.27
Federal leases TX ⁽²⁾	396	0.009	26,166	0.06

⁽¹⁾ Federal leases in New Mexico refers to the BLM O&G activity in the following counties: Eddy, Lea, Chaves, Roosevelt, McKinley, Rio Arriba, San Juan, and Sandoval.

⁽²⁾ Federal leases in Oklahoma, Kansas, and Texas refers to BLM O&G activity in any county reporting federal leases to the Office of Natural Resources Revenue (ONRR).

10.3 UNCERTAINTIES OF GREENHOUSE GAS CALCULATIONS

Although an analysis may present a quantified estimate of potential GHG emissions associated with reasonably foreseeable oil and gas development, there is significant uncertainty in GHG emission estimates due to uncertainties with regard to eventual production volumes and variability in flaring,

construction, and transportation. A rough estimate was possible using publicly available information and using estimates from future production for reasonably foreseeable development.

Also, there is uncertainty with regard to the net effects of reasonably foreseeable oil and gas development on climate; that is, while BLM actions may contribute to the climate change phenomenon, the specific effects of those actions on global climate are speculative given the current state of the science. Inconsistencies in the results of scientific models designed to predict climate change on regional or local scales limits the ability to quantify potential future impacts of decisions made at this level and determining the significance of any discrete amount of GHG emissions is beyond the limits of existing science at the present time.

10.4 REASONABLY FORESEEABLE DEVELOPMENT SCENARIOS

10.4.1 FARMINGTON FIELD OFFICE

Table 14 provides the reasonably foreseeable future GHGs (CO₂e emissions) associated with end-use oil and gas combustion emissions for the 2018 Mancos Gallup RFD scenario from federal, state (fee) as well as Indian minerals in the planning area. Total cumulative well development will result in 3,200 new wells from 2018 to 2037, of which 1,980 are estimated to be federal. The methodology for estimating new well development as well as the volumes for oil and gas is described in the Mancos Gallup 2018 RFD (Crocker and Glover 2018).

CO₂e emissions from downstream/end-use combustion of oil and gas products are estimated annually and cumulatively for BLM development and an all development (federal and non-federal) well production development scenario (Table 14). Under the all development scenario (includes federal, Indian, state and fee minerals), cumulative emissions during the 20-year period is estimated to produce 398.4 million metric tons (MMT) of CO₂e from the end-use combustion of products from 3,200 wells. The range of annual CO₂e emissions is 15.3 MMT/year in 2024 during the development of 126 additional oil and gas wells to 28.5 MMT/year of CO₂e in 2037 when 253 annual oil and gas wells are added that year.

Table 14. Estimated Cumulative Downstream/End Use GHG Emissions Resulting from Oil and Gas Production BLM 2018 Mancos Gallup RFD Scenario (Crocker and Glover 2018)

Federal Wells in the Planning Area (Federal development only)									
Year	Number of Wells	Annual CO ₂ - (MT) Oil	Annual CH ₄ -(MT) Oil	Annual N ₂ O- (MT) Oil	Annual CO ₂ - (MT) Gas	Annual CH ₄ -(MT) Gas	Annual N ₂ O- (MT) Gas	Annual CO ₂ e (MMT) Oil and Gas	CO ₂ e % of Total RFD
2018	41	2,150,121.90	86.54	17.31	12,973,881.29	244.51	24.45	15.14	6.1%
2019	48	2,351,067.87	94.62	18.92	10,715,391.22	201.95	20.19	13.09	5.3%
2020	53	2,521,871.95	101.50	20.30	9,033,960.28	170.26	17.03	11.57	4.7%
2021	59	2,702,723.32	108.78	21.76	7,849,994.12	147.95	14.79	10.57	4.3%
2022	65	2,873,527.40	115.65	23.13	7,044,101.19	132.76	13.28	9.93	4.0%
2023	72	3,034,284.18	122.12	24.42	6,646,129.37	125.26	12.53	9.70	3.9%

Federal Wells in the Planning Area (Federal development only)									
Year	Number of Wells	Annual CO2-(MT) Oil	Annual CH4-(MT) Oil	Annual N2O-(MT) Oil	Annual CO2-(MT) Gas	Annual CH4-(MT) Gas	Annual N2O-(MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2024	78	3,215,135.55	129.40	25.88	6,397,396.98	120.57	12.06	9.63	3.9%
2025	84	3,406,034.23	137.08	27.42	6,268,056.14	118.13	11.81	9.69	3.9%
2026	90	3,606,980.20	145.17	29.03	6,297,904.03	118.69	11.87	9.92	4.0%
2027	96	3,817,973.47	153.66	30.73	6,437,194.17	121.32	12.13	10.27	4.2%
2028	103	4,049,061.34	162.96	32.59	6,685,926.55	126.01	12.60	10.76	4.3%
2029	109	4,280,149.21	172.26	34.45	7,014,253.30	132.19	13.22	11.32	4.6%
2030	112	4,461,000.58	179.54	35.91	7,342,580.05	138.38	13.84	11.83	4.8%
2031	120	4,722,230.34	190.05	38.01	7,770,399.76	146.45	14.64	12.52	5.1%
2032	127	4,983,460.11	200.57	40.11	8,238,016.65	155.26	15.53	13.25	5.4%
2033	133	5,244,689.87	211.08	42.22	8,715,582.83	164.26	16.43	13.99	5.7%
2034	139	5,515,966.93	222.00	44.40	9,232,946.19	174.01	17.40	14.78	6.0%
2035	145	5,797,291.29	233.32	46.66	9,770,208.15	184.14	18.41	15.60	6.3%
2036	150	6,078,615.66	244.65	48.93	10,347,267.28	195.01	19.50	16.46	6.7%
2037	156	6,349,892.72	255.56	51.11	10,924,326.42	205.89	20.59	17.31	7.0%
TOTAL	1980	81,182,172.73	3,267.33	653.47	165,725,414.57	3,123.36	312.34	247.36	100

All Wells in the Planning Area (including federal, Indian, state, and fee minerals)									
Year	Number of Wells	Annual CO2-(MT) Oil	Annual CH4-(MT) Oil	Annual N2O-(MT) Oil	Annual CO2-(MT) Gas	Annual CH4-(MT) Gas	Annual N2O-(MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2018	67	3,338,757.52	134.37	26.87	20,611,035.17	388.45	38.84	23.98	6.0%
2019	76	3,631,244.43	146.15	29.23	17,054,335.40	321.42	32.14	20.72	5.2%
2020	86	3,868,431.00	155.69	31.14	14,426,592.28	271.89	27.19	18.32	4.6%
2021	96	4,116,418.43	165.67	33.13	12,583,813.17	237.16	23.72	16.73	4.2%
2022	106	4,347,556.53	174.98	35.00	11,359,304.15	214.08	21.41	15.73	4.0%
2023	116	4,572,214.13	184.02	36.80	10,767,546.13	202.93	20.29	15.37	3.9%
2024	126	4,815,017.15	193.79	38.76	10,436,281.41	196.69	19.67	15.28	3.8%
2025	136	5,081,582.03	204.52	40.90	10,312,050.33	194.35	19.43	15.42	3.9%
2026	146	5,369,748.60	216.12	43.22	10,438,186.79	196.72	19.67	15.84	4.0%
2027	156	5,676,924.66	228.48	45.70	10,756,930.42	202.73	20.27	16.46	4.1%
2028	166	6,019,095.46	242.25	48.45	11,244,164.48	211.91	21.19	17.30	4.3%
2029	176	6,367,746.77	256.28	51.26	11,849,096.87	223.32	22.33	18.25	4.6%
2030	180	6,264,058.65	252.11	50.42	12,471,341.04	235.04	23.50	18.77	4.7%

All Wells in the Planning Area (including federal, Indian, state, and fee minerals)									
Year	Number of Wells	Annual CO2-(MT) Oil	Annual CH4-(MT) Oil	Annual N2O-(MT) Oil	Annual CO2-(MT) Gas	Annual CH4-(MT) Gas	Annual N2O-(MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2031	194	6,915,133.64	278.31	55.66	13,303,994.11	250.73	25.07	20.26	5.1%
2032	204	7,354,944.09	296.01	59.20	14,174,156.04	267.13	26.71	21.57	5.4%
2033	214	7,765,376.24	312.53	62.51	15,053,518.25	283.71	28.37	22.86	5.7%
2034	224	8,193,521.78	329.76	65.95	15,986,394.55	301.29	30.13	24.22	6.1%
2035	234	8,615,618.84	346.75	69.35	16,900,380.33	318.51	31.85	25.56	6.4%
2036	244	9,028,211.15	363.36	72.67	17,982,094.38	338.90	33.89	27.06	6.8%
2037	253	9,438,211.27	379.86	75.97	19,042,087.06	358.88	35.89	28.53	7.2%
TOTAL	3200	120,779,812.34	4,861.01	972.20	276,753,302.38	5,215.86	521.59	398.23	100

Over the 20-year period, cumulative federal only wells would produce 247.4 MMT of CO₂e emissions from end-use combustion of oil and gas fossil fuels from 1,980 wells. The range of annual CO₂e emissions is 9.6 MMT/year in 2024 during the development of 78 oil and gas wells to 17.3 MMT/year of CO₂e in 2037 during the development of 156 annual oil and gas wells. This would represent 0.97% to 1.75% of BLM’s 2020 annual future estimated GHG emissions from end-use combustion (Table 22). It would represent a contribution of 1.11% to 2.00% respectively to BLM’s annual 2030 future estimated downstream (end-use) GHG emissions (Table 22). It should be noted that Table 22 also includes emissions from coal which produces 50% to 60% more CO₂ emissions than natural gas.

10.4.2 PECOS DISTRICT OFFICE PLANNING AREA

Table 15 provides the reasonably foreseeable future GHGs (CO₂e emissions) associated with end-use oil and gas combustion emissions for PDO from federal, state (fee) as well as Indian minerals in the planning area. The PDO federal planning area includes oil and gas well development from CFO, Roswell Field Office as well as Hobbs Field Office. Total cumulative well development will result in 16,000 new wells from 2016 to 2035, of which 6,400 are estimated to be federal. The methodology for estimating new well development as well as the volumes for oil and gas is described in Engler et al. (2012) and Engler and Cather (2014).

CO₂e emissions from downstream/end-use combustion of oil and gas products are estimated annually and cumulatively for BLM development and for an all development (federal and non-federal) well production scenario (Table 15). Under the all development scenario (includes federal, Indian, state and fee minerals), cumulative emissions during the 20-year period is estimated to produce 5,597 MMT of CO₂e from the end-use combustion of oil and gas from 16,000 wells. The range of annual CO₂e emissions is 97.7 MMT/year in 2016 to 597.7 MMT/year of CO₂e in 2035 when additional wells are added to production.

Over the 20-year period, cumulative federal only wells could produce 1,168.5 MMT of CO₂e emissions from end-use combustion of oil and gas fossil fuels from 6,400 wells. The range of annual CO₂e emissions is 48.1 MMT/year in 2016 to 70.1 MMT/year of CO₂e in 2035. This would represent 4.86% to

7.08% respectively of BLM’s 2020 annual future estimated GHG emissions from end-use combustion (see Table 22). It would represent a contribution of 5.55% to 8.09% respectively to BLM’s 2030 annual future estimated GHG emissions (Table 22). It should be noted that Table 22 also includes emissions from coal which produces 50% to 60% more CO₂ emissions than natural gas.

Table 15. Estimated Cumulative Downstream/End Use GHG Emissions Resulting from Oil and Gas Production BLM 2018 RFD PDO Scenario (Engler et al. 2012; Engler and Cather 2014)

Federal Wells in the Planning Area (Federal development only)								
Year	Annual CO2-(MT) Oil	Annual CH4-(MT) Oil	Annual N2O-(MT) Oil	Annual CO2-(MT) Gas	Annual CH4-(MT) Gas	Annual N2O-(MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2016	32,322,159.61	1,300.87	260.17	15,640,292.47	294.77	29.48	48.09	4.1%
2017	32,965,186.72	1,326.74	265.35	15,958,669.93	300.77	30.08	49.05	4.2%
2018	33,628,308.43	1,353.43	270.69	16,277,047.38	306.77	30.68	50.04	4.3%
2019	34,301,477.44	1,380.53	276.11	16,605,374.13	312.95	31.30	51.04	4.4%
2020	34,984,693.74	1,408.02	281.60	16,933,700.88	319.14	31.91	52.06	4.5%
2021	35,688,004.64	1,436.33	287.27	17,271,976.93	325.52	32.55	53.10	4.5%
2022	36,401,362.84	1,465.04	293.01	17,620,202.27	332.08	33.21	54.16	4.6%
2023	37,124,768.34	1,494.15	298.83	17,968,427.61	338.64	33.86	55.24	4.7%
2024	37,868,268.44	1,524.08	304.82	18,326,602.25	345.39	34.54	56.34	4.8%
2025	38,621,815.84	1,554.41	310.88	18,694,726.18	352.33	35.23	57.47	4.9%
2026	39,395,457.83	1,585.54	317.11	19,072,799.41	359.46	35.95	58.62	5.0%
2027	40,189,194.42	1,617.49	323.50	19,450,872.64	366.58	36.66	59.80	5.1%
2028	40,992,978.31	1,649.84	329.97	19,838,895.16	373.90	37.39	60.99	5.2%
2029	41,806,809.49	1,682.59	336.52	20,236,866.98	381.40	38.14	62.21	5.3%
2030	42,650,782.58	1,716.56	343.31	20,644,788.09	389.08	38.91	63.46	5.4%
2031	43,494,755.66	1,750.53	350.11	21,052,709.21	396.77	39.68	64.72	5.5%
2032	44,368,870.64	1,785.71	357.14	21,470,579.62	404.65	40.46	66.01	5.6%
2033	45,253,032.92	1,821.29	364.26	21,908,348.62	412.90	41.29	67.34	5.8%
2034	46,157,289.79	1,857.69	371.54	22,346,117.62	421.15	42.11	68.68	5.9%
2035	47,081,641.26	1,894.89	378.98	22,793,835.91	429.59	42.96	70.06	6.0%
TOTAL	785,306,906.24	31,606.13	6,321.23	380,092,934.68	7,163.46	716.35	1,168.47	100

All Wells in the Planning Area (including Federal, Indian, state, and fee minerals)								
Year	Annual CO2-(MT) Oil	Annual CH4-(MT) Oil	Annual N2O-(MT) Oil	Annual CO2-(MT) Gas	Annual CH4-(MT) Gas	Annual N2O-(MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2016	65,407,913.92	2,632.46	526.49	32,056,629.99	604.16	60.42	97.72	1.7%

All Wells in the Planning Area (including Federal, Indian, state, and fee minerals)								
Year	Annual CO2- (MT) Oil	Annual CH4- (MT) Oil	Annual N2O- (MT) Oil	Annual CO2- (MT) Gas	Annual CH4- (MT) Gas	Annual N2O- (MT) Gas	Annual CO2e (MMT) Oil and Gas	CO2e % of Total RFD
2017	71,948,705.31	2,895.71	579.14	35,260,303.13	664.54	66.45	107.49	1.9%
2018	79,142,571.11	3,185.24	637.05	38,792,303.02	731.10	73.11	118.24	2.1%
2019	87,059,842.41	3,503.88	700.78	42,672,528.25	804.23	80.42	130.07	2.3%
2020	95,760,803.00	3,854.07	770.81	46,940,776.01	884.67	88.47	143.08	2.6%
2021	105,345,925.87	4,239.84	847.97	51,626,894.17	972.99	97.30	157.38	2.8%
2022	115,875,494.81	4,663.62	932.72	56,790,578.52	1,070.31	107.03	173.12	3.1%
2023	127,460,030.10	5,129.86	1,025.97	62,471,626.23	1,177.38	117.74	190.43	3.4%
2024	140,210,052.03	5,643.01	1,128.60	68,719,783.78	1,295.13	129.51	209.48	3.7%
2025	154,226,033.58	6,207.11	1,241.42	75,594,746.95	1,424.70	142.47	230.42	4.1%
2026	169,648,636.94	6,827.82	1,365.56	83,146,262.21	1,567.02	156.70	253.46	4.5%
2027	186,618,524.28	7,510.81	1,502.16	91,463,873.22	1,723.78	172.38	278.81	5.0%
2028	205,276,357.79	8,261.73	1,652.35	100,607,275.75	1,896.10	189.61	306.69	5.5%
2029	225,813,036.14	9,088.26	1,817.65	110,675,962.76	2,085.86	208.59	337.37	6.0%
2030	248,389,316.10	9,996.89	1,999.38	121,739,579.32	2,294.38	229.44	371.10	6.6%
2031	273,226,238.25	10,996.49	2,199.30	133,917,516.97	2,523.89	252.39	408.21	7.3%
2032	300,554,890.46	12,096.39	2,419.28	147,309,268.67	2,776.28	277.63	449.04	8.0%
2033	330,606,360.59	13,305.86	2,661.17	162,034,225.96	3,053.79	305.38	493.93	8.8%
2034	363,661,973.00	14,636.25	2,927.25	178,241,628.27	3,359.25	335.92	543.33	9.7%
2035	400,033,193.94	16,100.07	3,220.01	196,060,816.45	3,695.08	369.51	597.66	10.7%
TOTAL	3,746,275,946.92	150,775.80	30,155.16	1,836,122,579.62	34,604.65	3,460.46	5,597.05	100

Table 16 shows historical United States, New Mexico, and BLM New Mexico federal production in the major oil and gas basins and their associated end-use combustion GHG emissions during calendar years 2015 through 2019. Production of oil and gas on federal lands has varied over the 5-year period due to market conditions, technological advances as well as pipeline and storage infrastructure availability. In 2015, total CO₂e end-use emissions resulting from oil and gas production in the United States was 3,288.42 MMT. New Mexico, 2015 GHG emissions associated with oil and gas production end-use was 134.87 MMT, which is 4.11% of national emissions in 2015.

GHG emissions from O&G gas production in the BLM PDO planning area in 2015 was 47.18 MMT of CO₂e, which is 1.44% of national O&G GHG emissions and 35.0% of New Mexico O&G GHG emissions from production in 2015. GHG end-use emissions from the BLM PDO planning area has increased to 104.76 MMT/year of CO₂e in 2019.

GHG emissions from O&G production in the BLM FFO planning area in 2015 was 30.42 MMT of CO₂e, which is 0.93% of national O&G GHG emissions and 22.6% of New Mexico O&G GHG emissions from

production in 2015. GHG end-use emissions from the BLM PDO planning area has decreased to 24.99 MMT/year of CO₂e in 2019.

Table 16. Historical Federal Oil and Gas Production New Mexico

Oil and Gas Production	2015	2016	2017	2018	2019
U.S. oil production (Mbbbls)	3,447,970	3,239,657	3,420,545	4,001,892	4,470,528
New Mexico oil production (Mbbbls)	148,095	146,634	172,154	248,879	329,483
BLM PDO oil production (Mbbbls)	72,630	71,479	84,151	124,205	162,908
BLM FFO oil production (Mbbbls)	6,832	5,344	4,914	5,042	4,891
U.S. gas production (MMcf)	32,914,647	32,591,578	33,292,113	37,325,539	40,892,458
New Mexico gas production (MMcf)	1,296,793	1,282,666	1,323,019	1,540,984	1,852,719
BLM PDO gas production (MMcf)	287,963	308,463	352,421	478,919	626,451
BLM FFO gas production (MMcf)	503,804	470,078	440,776	438,842	419,696
GHG Emissions					
Total U.S. O&G GHG emissions (MMT CO ₂ e)	3,288.42	3,180.51	3,297.10	3,768.92	4,166.46
Total New Mexico O&G GHG emissions (MMT CO ₂ e)	134.87	133.47	146.73	191.87	243.80
Total PDO O&G GHG emissions (MMT CO ₂ e)	47.18	47.80	55.69	79.94	104.76
Total FFO O&G GHG emissions (MMT CO ₂ e)	30.42	27.93	26.15	26.10	24.99

Note: Data total for PDO, FFO includes data from both federal and mixed exploratory land classes.

GHG emissions from O&G gas production in the BLM OFO planning area in 2015 was 3.8 MMT of CO₂e, which is 0.12% of national O&G GHG emissions and 0.30% of Texas, Oklahoma, and Kansas O&G GHG emissions combined (see Table 17) from production in 2015. GHG end-use emissions from the BLM OFO planning area has decreased to 2.9 MMT/year of CO₂e in 2019.

Table 17. Historical Federal Oil and Gas Production (OFO) Oklahoma, Kansas, and Texas

Oil and Gas Production	2015	2016	2017	2018	2019
U.S. oil production (Mbbbls)	3,447,970	3,239,657	3,420,545	4,001,892	4,470,528
Oklahoma oil production (Mbbbls)	166,468	155,228	165,955	199,557	211,808
Oklahoma federal oil production (Mbbbls)	1059.219	857.562	613.381	532.066	834.788
Texas oil production (Mbbbls)	1,260,755	1,167,369	1,273,558	1,608,926	1,850,715
Texas federal oil production (Mbbbls)	270.4	260.107	421.923	448.227	314.627

Oil and Gas Production	2015	2016	2017	2018	2019
Kansas oil production (Mbbls)	45,481	37,944	35,825	34,715	33,193
Kansas federal oil production (Mbbls)	154	139	127	127	110
BLM OFO (Kansas, Oklahoma, Texas) oil production (Mbbls) ⁽¹⁾	1,484	1,257	1,163	1,107	1,259
U.S. gas production (MMcf)	32,914,647	32,591,578	33,292,113	37,325,539	40,892,458
Oklahoma gas production (MMcf)	2,499,599	2,468,312	2,513,897	2,875,787	3,175,008
Oklahoma federal gas production (MMcf)	19,673	17,862	14,757	13,710	13,966
Texas gas production (MMcf)	8,799,465	8,156,296	8,079,974	9,109,174	10,355,453
Texas federal gas production (MMcf)	34,115	33,348	30,148	27,002	26,166
Kansas gas production (MMcf)	284,184	244,795	219,639	201,391	183,097
Kansas federal gas production (MMcf)	4,418	4,137	3,865	3,743	3,030
BLM OFO (Kansas, Oklahoma, Texas) gas production (MMcf)	58,206	55,347	48,771	44,454	43,163
GHG Emissions					
Total U.S. O&G GHG emissions (MMT CO ₂ e)	3,288.42	3,180.51	3,297.10	3,768.92	4,166.46
Total Kansas, Oklahoma, Texas O&G GHG emissions (MMT CO ₂ e)	1,269.66	1,182.14	1,228.86	1,463.14	1,655.83
Total OFO (Kansas, Oklahoma, Texas) O&G GHG emissions (MMT CO ₂ e)	3.8	3.6	3.2	2.9	2.9

Note: Using emissions factors for oil and gas production from Subpart C;

⁽¹⁾ OFO includes federal oil and gas activity for Oklahoma, Kansas, and Texas.

10.5 GLOBAL, NATIONAL, AND STATE GREENHOUSE GAS EMISSIONS

It is useful to compare the relative and absolute contributions to climate change of different GHG emissions, as well as emissions from different regions/countries or sources/sectors. There are several different metrics that can be used for this comparison. A GHG emission inventory is used to identify and quantify the anthropogenic GHG emissions from different regions/countries or sources/sectors. Using the GWP concept, GHG emissions are often reported in terms of CO₂e. The World Resources Institute's Climate Analysis Indicators Tool provides data on GHG emissions from 186 countries and all 50 states. In 1990 total global GHG emissions were 34,964 MMT CO₂e; In 2017, total global GHG emissions were 49,947 MMT CO₂e, including land-use change and forestry. From 1990 to 2017, global GHG emissions have increased at an annual rate of 3.3%. Electricity generation, manufacturing/construction, and transportation account for roughly 31%, 13%, and 15% of total global GHG emissions, respectively (World Resources Institute 2017).

To meet the obligations of the UNFCCC, the EPA publishes the national GHG emissions inventory on an annual basis (EPA 2021n). The lowest GHG emissions, since reporting, 6,373 MMT of CO₂e, occurred in

1991 and the peak GHG emissions occurred about 16 years later in 2007, 7,417 MMT CO₂e. The largest source of GHG emissions from human activities in the United States is from burning of fossil fuels for electricity, heat, and transportation. Total U.S. emissions have increased by 1.8% from 1990 to 2019, down from a high of 15.6% above 1990 levels in 2007. The latest national GHG emissions are for calendar year 2019, in which total gross U.S. GHG emissions were reported at 6,558.3 MMT of CO₂e (see Figure 16). Emissions decreased from 2018 to 2019 by 1.7 MMT of CO₂e. Net emissions (including sinks) were 5,769.1 MMT CO₂e. The decline from 2018-2019 reflects the combined impacts of many long-term trends, including population, economic growth, energy market trends, technological changes including energy efficiency, and carbon intensity of energy fuel choices. Figure 16 illustrates U.S. GHG emissions (MMT/year) by gas from 1990 through 2019 (EPA 2021n).

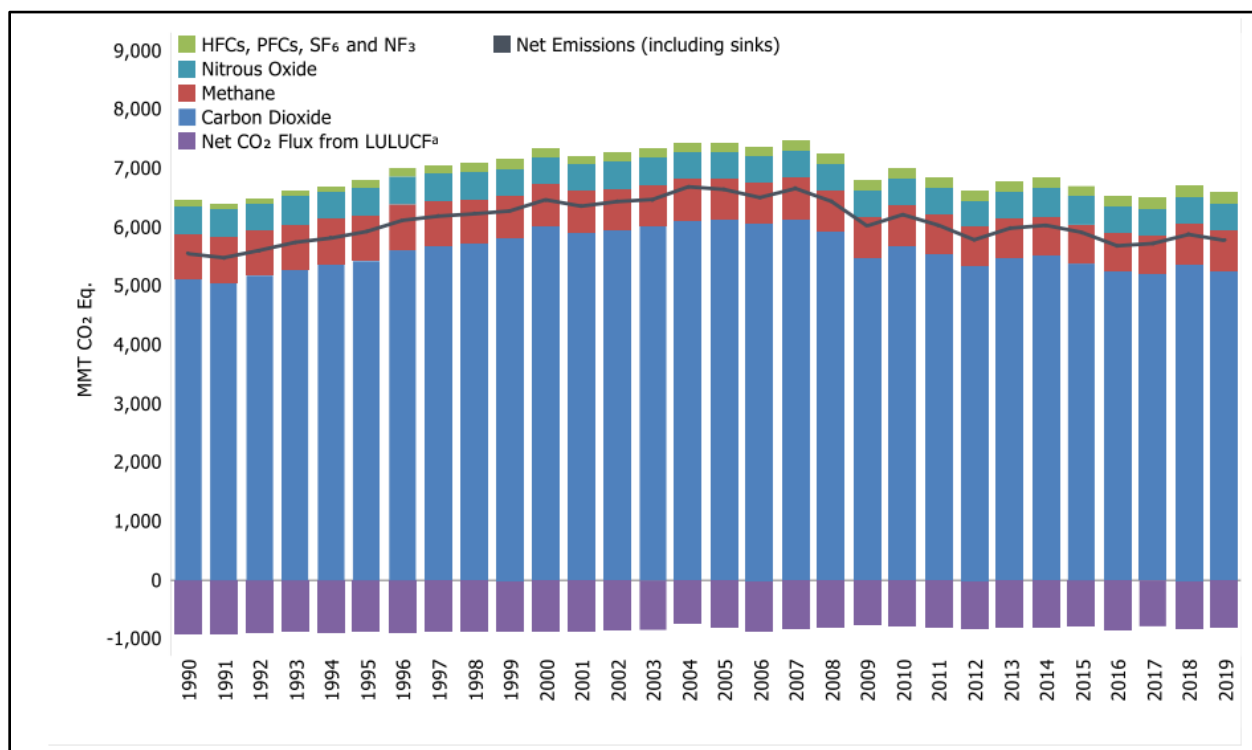


Figure 16. U.S. greenhouse gas (GHG) emissions by gas from 1990 to 2019 (EPA 2021n)

GHG emissions can be generated from a myriad of sources. The Inventory on Greenhouse Gas Emissions produced annually by the EPA categorizes these emissions by economic sector. These five major sector categories are transportation, electricity generation, industry, agriculture, and commercial and residential.

- **Transportation** – GHG emissions are a result of burning fossil fuel for use in cars, trucks ships, trains, and planes.
- **Electricity Generation** – GHG emissions in this sector are primarily from fuel methods used to generate electricity, coal, natural gas, as well as other fuels.
- **Industry** – GHG emissions from the industry sector are the result of the burning of fossil fuels for energy as well as emissions from certain chemical reactions necessary to produce goods from raw materials.

- **Commercial and Residential** – These GHG emissions are primarily from fossil fuels burned for heat and the use of certain products that contain GHGs and the handling of waste.
- **Agriculture** – GHG emissions from this sector comes from livestock such as cows, agricultural soils, and rice production.

Table 18 further breaks down GHG emissions by major source category within each major GHG and shows GHG trends from 1990 to 2019.

The gross emissions totals presented in this report for the United States exclude emissions and removals from Land Use, Land-Use Change, and Forestry (LULUCF), whereas the net emissions totals presented include emissions and removals from LULUCF (see Table 18).

Table 18. Trends in U.S. Greenhouse Gas Emissions and Sinks (MMT CO₂e)

Gas/Source	1990	2007	2015	2016	2017	2018	2019
CO₂	5,113.50	6,131.00	5,371.80	5,248.00	5,207.80	5,375.50	5,255.80
Fossil fuel combustion	4,731.50	5,754.00	5,008.30	4,911.50	4,854.50	4,991.40	4,856.70
Transportation	1,469.10	1,860.80	1,719.20	1,759.90	1,782.40	1,816.60	1,817.20
Electric power	1,820.00	2,411.10	1,900.60	1,808.90	1,732.00	1,752.90	1,606.00
Industrial	853.8	867.4	797.3	792.5	790.1	813.6	822.5
Residential	338.6	341.2	317.3	292.8	293.4	338.1	336.8
Commercial	228.3	223.1	244.6	231.6	232	245.7	249.7
U.S. Territories	21.7	50.4	29.2	26	24.6	24.6	24.6
Non-energy use of fuels	112.8	116.1	108.5	99.8	113.5	129.7	128.8
Iron and steel production and metallurgical coke production	9.7	12.5	32.4	21.8	25	37.1	47.3
Cement production	104.7	74.5	47.9	43.6	40.6	42.6	41.3
Petroleum systems	33.5	45.5	39.9	39.4	40.3	39	40.9
Natural gas systems	32	25.9	29.1	30.1	31.2	33.9	37.2
Petrochemical production	21.6	28.1	28.1	28.3	28.9	29.3	30.8
Ammonia production	13	9.1	10.6	10.2	11.1	12.2	12.3
Lime production	11.7	14.7	13.3	12.6	12.9	13.1	12.1

Gas/Source	1990	2007	2015	2016	2017	2018	2019
Incineration of waste	8.1	13	11.5	11.5	11.5	11.5	11.5
Other process uses of carbonates	6.3	8.6	12.2	11	9.9	7.5	7.5
Urea fertilization	3.8	4.9	4.6	5.1	5	5.9	6.2
Carbon dioxide consumption	2.4	3.8	4.7	4.9	5.1	5.2	5.3
Urea consumption for non-agricultural purposes	1.5	1.9	4.9	4.6	4.6	4.1	4.9
Liming	4.7	4.5	3.7	3.1	3.1	2.2	2.4
Ferroalloy production	6.8	4.3	2.8	1.3	1.2	1.5	1.9
Soda ash production	1.4	1.7	1.7	1.7	1.8	1.7	1.8
Titanium dioxide production	2.2	1.6	2	1.8	2	2.1	1.6
Aluminum production	1.2	1.9	1.6	1.7	1.7	1.5	1.5
Glass production	1.5	1.5	1.3	1.2	1.3	1.3	1.3
Zinc production	0.6	1	0.9	0.8	0.9	1	1
Phosphoric acid production	1.5	1.2	1	1	1	0.9	0.9
Lead production	0.5	0.6	0.5	0.5	0.5	0.5	0.5
Carbide production and consumption	0.4	0.2	0.2	0.2	0.2	0.2	0.2
Abandoned oil and gas wells	+	+	+	+	+	+	+
Magnesium production and processing	+	+	+	+	+	+	+
Wood biomass, ethanol, and biodiesel consumption ⁽¹⁾	219.4	244.6	317.7	316.6	312.3	319.6	316.2
International bunker fuels ⁽²⁾	103.5	115.4	110.9	116.6	120.1	122.1	116.1

Gas/Source	1990	2007	2015	2016	2017	2018	2019
CH₄⁽³⁾	776.9	693.9	651.5	642.4	648.4	655.9	659.7
Enteric fermentation	164.7	174.9	166.9	172.2	175.8	178	178.6
Natural gas systems	186.9	165.3	149.8	147.3	148.7	152.5	157.6
Landfills	176.6	127.2	111.4	108	109.4	112.1	114.5
Manure management	37.1	56	57.9	59.6	59.9	61.7	62.4
Coal mining	96.5	64.8	61.2	53.8	54.8	52.7	47.4
Petroleum systems	48.9	42.4	41.5	39.2	39.3	37.3	39.1
Wastewater treatment	20.2	19.9	18.8	18.7	18.5	18.4	18.4
Rice cultivation	16	15.9	16.2	15.8	14.9	15.6	15.1
Stationary combustion	8.6	7.7	8.5	7.9	7.6	8.5	8.7
Abandoned oil and gas wells	6.8	7.2	7.4	7.4	7.2	7.3	6.6
Abandoned underground coal mines	7.2	6.3	6.4	6.7	6.4	6.2	5.9
Mobile combustion	6.4	3.7	2.6	2.5	2.5	2.4	2.4
Composting	0.4	2	2.1	2.3	2.4	2.3	2.3
Field burning of agricultural residues	0.4	0.4	0.4	0.4	0.4	0.4	0.4
Petrochemical production	0.2	0.1	0.2	0.2	0.3	0.3	0.3
Anaerobic digestion at biogas facilities	+	0.1	0.2	0.2	0.2	0.2	0.2
Ferroalloy production	+	+	+	+	+	+	+
Carbide production and consumption	+	+	+	+	+	+	+

Gas/Source	1990	2007	2015	2016	2017	2018	2019
Iron and steel production and metallurgical coke production	+	+	+	+	+	+	+
Incineration of waste	+	+	+	+	+	+	+
International bunker fuels ⁽²⁾	0.2	0.1	0.1	0.1	0.1	0.1	0.1
N₂O ⁽³⁾	452.7	463.5	468.2	450.8	446.3	459.2	457.1
Agricultural soil management	315.9	319.5	348.5	330.1	327.6	338.2	344.6
Stationary combustion	18.7	23.6	25.4	25.9	26.4	26.1	26.4
Manure management	25.1	34.4	30.5	30	28.4	28.2	24.9
Mobile combustion	14	17.2	17.5	18.1	18.7	19.4	19.6
Adipic acid production	44.7	36.4	21.7	20.8	19.8	18.8	18
Nitric acid production	12.1	13.2	11.6	10.1	9.3	9.6	10
Wastewater treatment	15.2	10.3	4.3	7	7.4	10.3	5.3
N ₂ O from product uses	4.2	4.2	4.2	4.2	4.2	4.2	4.2
Composting	0.3	1.8	1.9	2	2.2	2	2
Caprolactam, glyoxal, and glyoxylic acid production	1.7	2.2	1.9	1.7	1.5	1.4	1.4
Incineration of waste	0.5	0.4	0.3	0.3	0.3	0.3	0.3
Electronics industry	+	0.2	0.2	0.2	0.3	0.3	0.2
Field burning of agricultural residues	0.2	0.2	0.2	0.2	0.2	0.2	0.2
Petroleum systems	+	+	+	+	+	+	+

Gas/Source	1990	2007	2015	2016	2017	2018	2019
Natural gas systems	+	+	+	+	+	+	+
International bunker fuels ⁽²⁾	0.9	1	1	1	1.1	1.1	1
HFCs	46.5	143.3	168.3	168.1	170.3	169.8	174.6
Substitution of ozone depleting substances ⁽⁴⁾	0.2	121.6	163.6	164.9	164.7	166	170.5
HCFC-22 production	46.1	21.5	4.3	2.8	5.2	3.3	3.7
Electronics industry	0.2	0.2	0.3	0.3	0.4	0.4	0.3
Magnesium production and processing	+	+	0.1	0.1	0.1	0.1	0.1
PFCs	24.3	8	5.2	4.4	4.1	4.7	4.5
Electronics industry	2.8	3.3	3.1	2.9	2.9	3	2.7
Aluminum production	21.5	4.7	2.1	1.4	1.1	1.6	1.8
Substitution of ozone depleting substances	+	+	+	+	+	0.1	0.1
SF₆	28.8	9.2	5.5	6	5.9	5.7	5.9
Electrical transmission and distribution	23.2	6.3	3.8	4.1	4.2	3.9	4.2
Magnesium production and processing	5.2	2.5	1	1.1	1	1	0.9
Electronics industry	0.5	0.5	0.7	0.8	0.7	0.8	0.8
NF₃	+	0.6	0.6	0.6	0.6	0.6	0.6
Electronics industry	+	0.6	0.6	0.6	0.6	0.6	0.6
Unspecified mix of HFCs, PFCs, SF₆, and NF₃	+	+	+	+	+	+	+
Electronics industry	+	+	+	+	+	+	+

Gas/Source	1990	2007	2015	2016	2017	2018	2019
Total emissions	6,442.70	7,449.60	6,671.10	6,520.30	6,483.30	6,671.40	6,558.30
LULUCF emissions ⁽³⁾	7.9	22.8	27.8	13.2	26	23.4	23.5
LULUCF CH ₄ emissions	5	12.8	16.6	7.7	15.3	13.8	13.8
LULUCF N ₂ O emissions	3	10	11.3	5.5	10.6	9.7	9.7
LULUCF carbon stock change ⁽⁵⁾	-908.7	-821.6	-791.7	-856	-792	-824.9	-812.7
LULUCF sector net total ⁽⁶⁾	-900.8	-798.8	-763.8	-842.8	-766.1	-801.4	-789.2

Data obtained from Table ES-2 of US Inventory of Greenhouse Gases and Sinks (EPA 2021n).

Notes: Total emissions presented without LULUCF. Net emissions presented with LULUCF. Totals may not sum due to independent rounding. Parentheses indicate negative values or sequestration.

+ Does not exceed 0.05 MMT CO₂e.

⁽¹⁾ Emissions from Wood Biomass, Ethanol, and Biodiesel Consumption are not included specifically in summing Energy sector totals. Net carbon fluxes from changes in biogenic carbon reservoirs are accounted for in the estimates for Land Use, Land-Use Change, and Forestry.

⁽²⁾ Emissions from International Bunker Fuels are not included in totals.

⁽³⁾ LULUCF emissions of CH₄ and N₂O are reported separately from gross emissions totals. LULUCF emissions include the CH₄, and N₂O emissions from Peatlands Remaining Peatlands; CH₄ and N₂O emissions reported for Non-CO₂ Emissions from Forest Fires, Non-CO₂ Emissions from Grassland Fires, and Coastal Wetlands Remaining Coastal Wetlands; CH₄ emissions from Land Converted to Coastal Wetlands; and N₂O emissions from Forest Soils and Settlement Soils.

⁽⁴⁾ Small amounts of PFC emissions also result from this source.

⁽⁵⁾ LULUCF Carbon Stock Change is the net C stock change from the following categories: Forest Land Remaining Forest Land, Land Converted to Forest Land, Cropland Remaining Cropland, Land Converted to Cropland, Grassland Remaining Grassland, Land Converted to Grassland, Wetlands Remaining Wetlands, Land Converted to Wetlands, Settlements Remaining Settlements, and Land Converted to Settlements.

⁽⁶⁾ The LULUCF Sector Net Total is the net sum of all CH₄ and N₂O emissions to the atmosphere plus net C stock changes.

Within the CO₂ pollutant category there are 28 source categories reported in the inventory; the CH₄ pollutant category has 20 source categories and N₂O pollutant category has 16 source categories reported. Other pollutant categories reported in the annual inventory report include HFCs, PFCs, SF₆ and NF₃ (see Table 18). The largest source of GHG emissions in 2018 is attributed to CO₂ emissions is from fossil fuel combustion (5,031.8 MMT of CO₂); transportation (27.3%), electric power sector (26.3%), industrial (12.5%), residential (5.1%), and commercial (3.7%) of total GHG emissions for 2018 (6,676.6 MMT of CO₂e) (EPA 2021n).

10.6 NATURAL GAS SYSTEMS AND PETROLEUM SYSTEMS

Within the fossil fuel combustion sector, in 2019, the contribution by fuel type shows that petroleum represents 45.0% of the fuel type, natural gas 34% and finally coal was 21% (Figure 17).

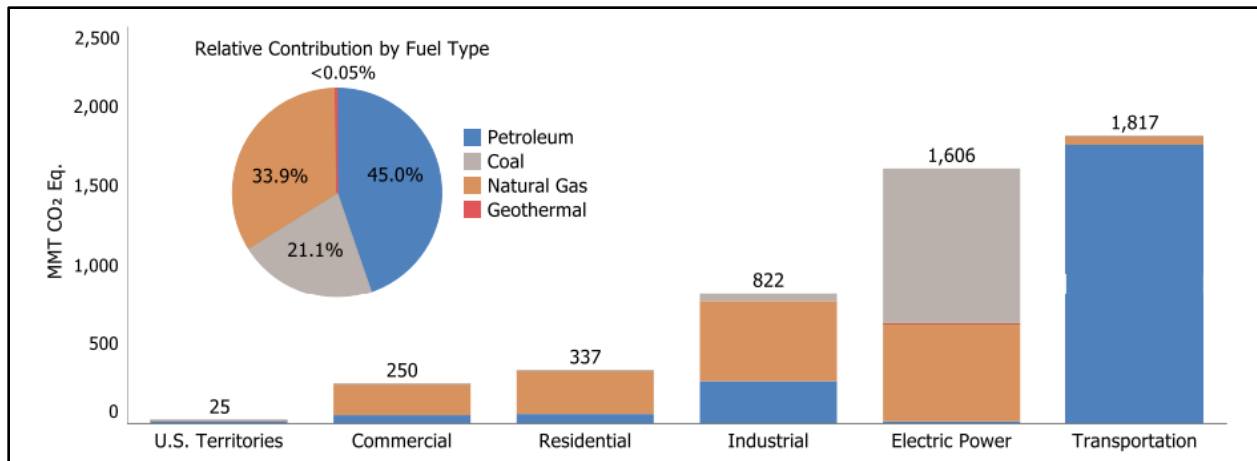


Figure 17. 2019 CO₂ emissions from fossil fuel combustion by sector and fuel type (MMT CO₂e).

The EPA’s national GHG emissions inventory describes “Natural Gas Systems” and “Petroleum Systems” as two of the major sources of U.S. GHG emissions. The inventory identifies the major contributions of natural gas and petroleum systems as total CO₂ and CH₄ emissions. Natural gas and petroleum systems do not produce noteworthy amounts of any other GHGs.

Total GHG emissions (CH₄, CO₂, and N₂O) from natural gas systems in 2019 were 194.9 MMT CO₂e, a decrease of 11% from 1990, primarily due to decreases in CH₄ emissions, and an increase of 5% from 2018, primarily due to increases in CH₄ emissions. Of the overall GHG emissions (194.9 MMT CO₂e), 81% are CH₄ emissions as expressed as CO₂e (157.6 MMT CO₂e), 19% are CO₂ emissions (37.2 MMT), and less than 0.01% are N₂O emissions as expressed as CO₂e (0.01 MMT CO₂e).

Within the category of “Natural Gas Systems”, the EPA identifies emissions occurring during distinct stages of operation, including field production, processing, transmission and storage, and distribution. Petroleum Systems” sub-activities include production field operations, crude oil transportation and crude oil refining. Within the Natural Gas Systems and Petroleum Systems the BLM has authority to regulate those field production operations that are related to oil and gas measurement and prevention of waste (via leaks, spills and unauthorized flaring and venting).

Total GHG emissions (CH₄, CO₂, and N₂O) from petroleum systems in 2019 were 86.4 MMT CO₂e, an increase of 47% from 1990, primarily due to increases in CO₂ emissions. Since 2009, total emissions increased by 64%; and since 2018, total emissions increased by 16%. Of the overall GHG emissions (86.4 MMT CO₂e), 47.3 MMT are CO₂ emissions, 39.1 MMT CO₂e are from CH₄ emissions, and 0.05 MMT CO₂e (0.24 kt N₂O), are from N₂O emissions from petroleum systems in 2018. U.S. oil production increased by 67%. In 2019, production was 129% higher than in 2009 and 12% higher than in 2018.

CH₄ emissions from petroleum systems are primarily associated with onshore and offshore crude oil production, transportation, and refining operations. During these activities, CH₄ is released to the atmosphere as emissions from leaks, venting (including emissions from operational upsets), and flaring. CO₂ emissions from petroleum systems are primarily associated with crude oil production and refining operations. Note that in the EPA’s reported data, CO₂ in the petroleum systems emissions exclude all combustion emissions (e.g., engine combustion) except for flaring CO₂ emissions. All combustion CO₂ emissions (except for flaring) are accounted for in the fossil fuel combustion category. Emissions of N₂O from petroleum systems are primarily associated with flaring.

For natural gas, extraction accounts for 55% of total life cycle CO₂e emissions, processing accounts for 27% and transmission accounts for 18% of life cycle CO₂e emissions (U.S. Department of Energy 2011). For oil, drilling and development is responsible for 8% of the total life cycle CO₂e emissions, whereas transportation of the petroleum to refineries represents about 10% of the emissions, and final consumption as transportation fuel represents fully 80% of emissions (U.S. Department of Energy 2008).

Table 19 displays GHG emissions (CO₂, CH₄ and N₂O) related to natural gas systems, petroleum systems as well as coal mining. In Table 19, CO₂ emissions listed represent CO₂ emissions which are not otherwise captured in the “fossil fuel combustion” category. The natural gas and petroleum subsectors that BLM regulates for onshore operations on federal mineral estate are highlighted in gray.

In summary, CO₂ is produced during the burning of fossil fuels to run internal combustion engines which may be used in drilling, transportation, pumping and compression. CO₂ may be a significant component of natural gas, especially coalbed CH₄, and is vented during field operations or processing. CO₂ is also used in enhanced oil production processes and may be released or escape to the atmosphere during those processes. CH₄ is the primary component of natural gas and is released to the atmosphere during both oil and gas production either intentionally during production when it cannot be captured, or accidentally through leaks and fugitive emissions.

Table 19. 2019 Greenhouse Gas Emissions for Oil and Gas subsectors and Coal Mining (EPA 2021n)

Sector	Subsector	2019 GHG Emissions (MMTCO ₂ e)				% of U.S. Total GHGs
		CO ₂	CH ₄ ⁽¹⁾	N ₂ O	Total GHGs	
Natural gas systems	Total	37.2	157.6	0.01	194.9	2.97%
	Exploration ⁽²⁾	0.2	0.5	0.0001	0.8	0.01%
	Production field operations	11.0	93.7	0.006	104.7	1.60%
	Onshore production	NE	52.0	NE	NE	NE
	Offshore production	NE	0.8	NE	NE	NE
	Gathering and boosting ⁽³⁾	NE	40.9	NE	NE	NE
	Processing	24.8	12.4	0.005	37.2	0.57%
	Transmission and storage	1.2	37.0	0.0006	38.2	0.58%
	Distribution	**	14.0	Not occurring	14.0	0.21%
Petroleum systems	Total	47.3	39.1	0.05	86.4	1.32%
	Exploration ⁽²⁾	2.1	0.3	0.0009	2.4	0.04%
	Production field operations	40.2	37.7	0.031	77.9	1.19%
	Crude oil transportation	**	0.2	NE	0.2	0.003%
	Crude refining	5.0	0.9	0.015	5.9	0.09%

Sector	Subsector	2019 GHG Emissions (MMTCO ₂ e)				% of U.S. Total GHGs
		CO ₂	CH ₄ ⁽¹⁾	N ₂ O	Total GHGs	
Coal mining	--	*+	47.4	*	47.4	0.72%
U.S. total		5,255.8	659.7	457.1	6,558.3 ***	100%

*Indicates values less than 0.1 Teragrams (Tg) CO₂e

**Indicates values that do not exceed 0.05 TgCO₂e

***Indicates that the total U.S. GHG emissions value includes U.S. emissions of three additional minor classes of GHGs not listed here.

NE=Not estimated

+ Includes data from abandoned coal mines.

Data obtained from Tables 3-30,3-33, 3-37, 3-39, 3-41, 3-62, 3-63,3-65, and 3-67 of 2018 Inventory Data (EPA 2021n)

⁽¹⁾ These values represent CH₄ emitted to the atmosphere. CH₄ that is captured, flared, or otherwise controlled (and not emitted to the atmosphere) has been calculated and removed from emission totals.

⁽²⁾ Exploration includes well drilling, testing, and completions.

⁽³⁾ Gathering and boosting includes gathering and boosting station routine vented and leak sources, gathering pipeline leaks and blowdowns, and gathering and boosting station episodic events.

10.6.1 TRENDS

Globally, emissions of CO₂ from flaring of unused gas during oil production decreased by about a quarter between 2003 and 2011; however, flaring emissions for the United States are on the rise and increased by 50% in 2011 because of the significant increase in fracking for shale oil production and the flaring of co-produced natural gas (Olivier et al. 2012). CO₂ emissions from natural gas and petroleum systems increased by 27% from 1990 to 2017, due to increases in flaring emissions.

10.7 NATIONAL GREENHOUSE GAS EMISSIONS GREENHOUSE GAS REPORTING PROGRAM (FLIGHT)

The Greenhouse Gas Reporting Program (GHGRP) is codified by regulation, (40 CFR 98) and requires reporting of GHG data and other relevant information from large GHG emission sources, fuel and industrial gas suppliers, and CO₂ injection sites in the United States. There is a total of 41 categories covered by the program. Facilities are generally required to submit annual reports under Part 98 if:

- GHG emissions from covered sources exceed 25,000 metric tons CO₂e per year.
- Supply of certain products would result in over 25,000 metric tons CO₂e of GHG emissions if those products were released, combusted, or oxidized.
- The facility receives 25,000 metric tons or more of CO₂ for underground injection.

The reported data are usually made available to the public in October of each year. It should be noted that the GHGRP does not represent total U.S. GHG emissions, but provides facility level data for large sources of direct emissions, thus representing the majority of U.S. GHG emissions. The GHGRP data collected from direct emitters represent about half of all U.S. emissions. When including GHG information reported to the GHGRP by suppliers, emissions coverage reaches approximately 85% to 90%. The *Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2019* contains information on all

GHG emissions sources and sinks in the United States. For more information, please visit the US Greenhouse Gas Reporting Program (GHGRP) <https://www.epa.gov/ghgreporting>.

10.7.1 COMPRESSOR ENGINES AND STATIONS (MIDSTREAM) REPORTED GREENHOUSE GAS EMISSIONS

Compressor engines link the natural gas pipeline infrastructure that transports natural gas from its source to points of consumption. Table 20 shows the GHG emissions from compressor stations and gas plants for each state where BLM NM has mineral estate, from the 2020 Greenhouse Gas Facility Level Information on Greenhouse Gases Tool (FLIGHT). Some gas plants and compressor stations' emissions may not be reported to FLIGHT because emissions from the plant or station do not exceed EPA's GHG reporting threshold.

Table 20. 2019 Midstream Greenhouse Gas Emissions from Gas Plants and Compressor Stations (EPA 2021o)

State	Number of Reporting Compressor Stations	Total GHG Emissions from Reporting Compressor Stations (MMT CO ₂ e)	% U.S. Total Reported Compressor Station GHG Emissions	Number of Reporting Gas Plants	Total GHG Emissions FROM Reporting Gas Plants (MMT CO ₂ e)	% U.S. Total Reported Gas Plant GHG Emissions
New Mexico	12	0.48	1.54	25	4.5	7.76
Texas	87	3.9	12.5	204	23.0	39.7
Oklahoma	19	0.60	1.94	51	3.4	5.86
Kansas	21	1.1	3.54	5	0.90	1.55

Emissions from natural gas processing, transmission/compression, transmission pipelines, and storage and distribution in the United States totaled 105 MMT of CO₂e in 2019, which was about 1.57% of total U.S. GHG emissions reported to EPA in 2019 (EPA 2021o).

10.7.2 REFINERIES (MIDSTREAM) REPORTED GREENHOUSE GAS EMISSIONS

Crude oil produced throughout the BLM-NM area is transported by pipeline and/or tanker truck to refineries where the oil is processed into various types of fuel. Table 21 shows the GHG emissions from refineries in each BLM-NM state.

Table 21. 2019 Greenhouse Gas Emissions from Refineries (EPA 2021o)

State	Number of Reporting Refineries	Total GHG Emissions from Reporting Refineries (MMT CO ₂ e)	% U.S. Total Reported Refinery GHG Emissions
New Mexico	3	1.2	0.87
Texas	29	56.0	31.46

State	Number of Reporting Refineries	Total GHG Emissions from Reporting Refineries (MMT CO ₂ e)	% U.S. Total Reported Refinery GHG Emissions
Oklahoma	5	4.5	2.52
Kansas	3	2.9	1.63

There are three refineries New Mexico, one in Jamestown (Gallup Refinery), one in Artesia, and one in Lovington. In Kansas, there are three refineries, Oklahoma has five refineries and Texas has 29 refineries. Transportation and processing of crude oil and petroleum products result in emissions of various hazardous air pollutants, criteria pollutants, and GHGs. In 2019, GHG emissions from refineries (total of 138 reporting) accounted for 178 MMT CO₂e emitted which is 2.67% of the 2019 total GHG emissions reported to EPA (EPA 2021o).

10.7.3 STATE GREENHOUSE GASES

The predicted GHG emissions are compared to the baseline statewide GHG emissions as reported in the Inventory of New Mexico Greenhouse Gas Emissions: 2000–2013 (NMED 2016). The Inventory of New Mexico Greenhouse Gas Emissions: 2000-2013 lists total statewide gross GHG emissions in 2013 as 80.9 MMT CO₂e. For 2013, New Mexico’s Environment Department reported that the primary contributors to the state’s GHG emissions were electricity generation (35%), the fossil fuel industry (26%), and transportation (17%) (NMED 2016). In 2012, World Resources Institute’s Climate Analysis Indicators Tool reports that New Mexico’s GHG emissions were 75.5 MMT CO₂e (World Resources Institute 2017). Electricity generation, transportation, and fugitive sources account for around 29%, 14%, and 11% of New Mexico’s GHG emissions in 2012 (World Resources Institute 2017).

The New Mexico Greenhouse Gas Inventory and Reference Case Projection 1990-2020 estimates that approximately 17.3 MMT of GHGs from the natural gas industry and 2.3 MMT of GHGs from the oil industry were projected in 2010 as a result of oil and natural gas production, processing, transmission and distribution. According to the New Mexico Greenhouse Gas Inventory and Reference Case Projections, 1990–2020, GHG emissions were expected to continue increasing (NMED 2006). From 1990 to 2012, New Mexico’s GHG emissions have decreased at an annual rate of 0.45% (NMED 2016).

10.7.4 OTHER MAJOR INDUSTRIES GENERATING GREENHOUSE GAS EMISSIONS

Potash mining is another major industry in the CFO area. There are two mining companies operating four potash processing plants in the CFO area. Potash production produces emissions of various hazardous air pollutants, criteria pollutants as well as GHGs. In 2015, potash mines in southeastern New Mexico emitted 97,140 metric tons of CO₂e cumulatively. This is 0.002% of total U.S. GHG emissions (EPA 2021n). In 2016 CO₂e emissions decreased significantly as some facilities discontinued reporting into the GHG emissions for valid reasons, thus in 2018 the emissions from Intrepid Potash reported only 8,408 metric tons of CO₂e; which is 0.0001% of total U.S. GHG emissions.

Coal mining is another major industry in San Juan County. Westmoreland purchased the San Juan Mine from BHP Billiton and began ownership 2016. BHP also transferred ownership of the Navajo Mine, located near Fruitland NM to the Navajo Transitional Energy Co. (NATEC) at the end of 2016. The San Juan Mine provides coal to the San Juan Generating Station and the Navajo Mine provides coal to the

Four Corners Power Plant. Coal production produces emissions of various hazardous air pollutants, criteria pollutants, and GHGs. In 2018, the San Juan Mine reported 0.57 MMT of CO₂e while data for the Navajo Mine was not available. In 2017, coal mining in the United States contributed 55.7 MMT CO₂e, from CH₄ which is 8.49% of total U.S. CH₄ emissions, and 0.87% of total U.S. GHG emissions (see Table 19).

11 CUMULATIVE GREENHOUSE GAS EMISSIONS

11.1 GOLDER ASSOCIATES REPORT (2017), GREENHOUSE GAS EMISSIONS FROM THE COAL, OIL, AND GAS LEASING PROGRAM WITHIN THE STATES OF NEW MEXICO, TEXAS, KANSAS, AND OKLAHOMA

In 2017, the BLM commissioned a climate change report with an energy focus. The report calculates GHG emissions associated with production and consumption activities related to coal, oil, natural gas, and natural gas liquids. The baseline year is 2014 and forecasts production/consumption GHG emissions for 2020 and 2030 for federal and non-federal lands on a national level and for 13 energy-producing states, not limited to New Mexico, Oklahoma, Texas, and Kansas. Inputs for the report were developed using publicly available online information from such sources as the EIA, the EPA's *Greenhouse Gas Inventory Report: 1990–2014* (EPA 2016), DOI ONRR, U.S. Extractive Industries Transparency Initiative, BLM oil and gas statistics, and others as applicable to each state. More information on the methodology and assumptions, as well as other data sources for all 13 states, is in the *Greenhouse Gas and Climate Change Report, 2017* (Golder Associates 2017), which is herein incorporated by reference.

BLM approximated national GHG emissions (CO₂e) from energy production for the baseline year 2014 and future years 2020 and 2030. Growth factors are applied as compound growth, where the exponents of each factor is raised to represent the number of years ahead of the baseline year of 2014 (Golder Associates 2017). Baseline growth or decline factors were developed based on data taken from Tables A1 and B1 of the EIA's 2016 Annual Energy Outlook (AEO). Two scenarios were developed: normal growth and high growth. Table 22 shows the 2014 baseline CO₂e emissions from fossil fuel production as well as future projections of 2020 and 2030 in the United States across federal and non-federal sectors. All projections rely on EIA's 2016 Annual Energy Outlook (AEO) growth factors. GHG emissions for future projections present only the high growth scenario, the normal growth scenario can be found in the Golder Associates report (Golder Associates 2017).

Table 22. Fossil Fuel Production and Future Year Scenarios Using AEO 2016 Outlook (Golder Associates 2017)

2014 Baseline Fossil Fuel Production in the U.S.							
	Oil Barrels (bbl)	CO₂e (MMT)	Gas (MMcf)	CO₂e (MMT)	Coal (short tons)	CO₂e (MMT)	Total CO₂e (MMT) All Fossil Fuels
U.S.	3,196,889,000	1,375	25,889,605	1,417	1,000,048,758	1900.09	4691.39
BLM national	155,424,817	67	3,399,894	186	409,345,817	777.76	1030.63
Non-BLM national	3,041,464,183	1,308	22,489,711	1,231	590,702,941.00	1122.34	3660.76
2020 Future Fossil Fuel Production in the U.S. Future, High Growth Scenarios							
	Oil Barrels (bbl)	CO₂e (MMT)	Gas (MMcf)	CO₂e (MMT)	Coal (short tons)	CO₂e (MMT)	Total CO₂e (MMT) All Fossil Fuels
U.S. high growth	3,639,277,000	1,565	30,743,208	1,682	898,459,853	1707.07	4954.17
BLM national high growth	177,967,000	77	4,062,563	222	363,786,038	691.19	990.02
Non-BLM national high growth	3,461,310,000	1,488	26,680,645	1,460	534,673,815	1015.88	3964.16
2030 Future Fossil Fuel Production in the U.S. Future, High Growth Scenarios							
	Oil Barrels (bbl)	CO₂e (MMT)	Gas (MMcf)	CO₂e (MMT)	Coal (short tons)	CO₂e (MMT)	Total CO₂e (MMT) All Fossil Fuels
U.S. high growth	3,907,285,000	1,680	37,628,912	2,059	665,345,945	1264.16	5003.27
BLM national high growth	191,073,000	82	4,972,475	272	269,398,309	511.86	866.10
Non-BLM national high growth	3,716,212,000	1,598	32,656,437	1,787	395,947,636	752.30	4137.17

BLM uses projections of the total federal and non-federal oil and gas emissions from Golder Associates (2017) to estimate expected annual future GHG emissions from energy production and consumption activity within a subnational region, including New Mexico, Oklahoma, Kansas, and Texas, which are within the administrative jurisdiction of the BLM NMSO. Assumptions of the analysis are discussed in Golder Associates (2017). The following are key assumptions:

- State-specific oil consumption is equal to state total production minus export and reserves for the state based on national averages.
- National averages for sector breakdown percentages (power, industrial, etc.) for oil, natural gas, and natural gas liquids consumptions were applied to state-specific data.
- The value of production and consumption on non-federal lands is equal to the difference of the total state or national value minus the federal lands value.

At the state level, production does not necessarily translate to 100% consumption of the fossil fuel but is representative of future energy consumption and production to show GHG emissions. New Mexico is an important supplier of electricity to the western United States. The state's power plants have historically produced more electricity than consumed in the state, and have exported significant amounts of electricity to Arizona, California, and other Western states. In 2000, for instance, New Mexico power plants produced 36% more electricity than needed for in-state use. The New Mexico electricity sector is also dominated by coal, which accounts for nearly 90% of all electricity generated in recent years. Coal-fired power plants produce as much as twice the CO₂ emissions per kilowatt-hour of electricity as natural gas-fired power plants. As a result of these factors, New Mexico power plants are the largest source of GHG emissions in the State (NMED 2016).

The development projected in the RFDs for each BLM field office under NMSO jurisdiction (such as the 2016 RFD for the PDO; see Engler et al. 2012; Engler and Cather 2014) are considered in these data. Current and future oil and gas lease sales are part of each RFD. Because the BLM NMSO administers lease sales within its jurisdictional area, this section provides a discussion of reasonably foreseeable production and consumption these states and discloses the magnitude of GHG emissions likely to result from BLM NMSO lease sale activities on an annual basis. This information is further contextualized by comparing the relative magnitude of these emission with projected national and global annual GHG emission rates.

Although quantified estimates of potential GHG emissions associated with reasonably foreseeable energy development are presented, there is uncertainty with regard to eventual production volumes and variability, flaring, construction, transportation, etc. A rough estimate was possible using publicly available information and estimates from future production for RFD. Also, there is uncertainty with regard to the net effects of reasonably foreseeable energy development on climate; that is, while BLM actions may contribute to the climate change phenomenon, the specific effects of those actions on global climate are speculative given the current state of the science. Inconsistencies in the results of scientific models designed to predict climate change on regional or local scales limits the ability to quantify potential future effects of decisions made at this level and to determine the significance of any discrete amount of GHG emissions beyond the limits of existing science.

Nonetheless, the projected emission estimates for Texas, New Mexico, Kansas, and Oklahoma contribute to global GHG emissions. This, in turn, affects the concentrations of GHG in the atmosphere which influences climate change. The relative magnitude of the contribution of the BLM NMSO GHG emissions from oil and gas as well as coal and natural gas liquids leasing activities in New Mexico, Texas,

Oklahoma, and Kansas will vary based on trends in national and global GHG emissions. However, for context, based on current global and national emission trends from the most recent available data, it is estimated that the 2014 baseline emissions for the BLM NMSO federal contribution from coal, oil, gas, and natural gas liquids leasing combined represent approximately 1.54% of national GHG emissions and 0.21% of global GHG emissions. BLM NMSO oil and gas leasing GHG emissions would be estimated to represent around 1.11% of national GHG emissions and 0.15% of global GHG emissions based on the 2014 baseline emissions.

11.1.1 NEW MEXICO COAL, OIL, AND GAS GREENHOUSE GAS EMISSIONS

BLM’s New Mexico reasonably foreseeable coal, oil, and gas production and consumption GHG emissions from federal activities are 95.09 MMT of CO₂e for the 2020 high scenario and 99.35 MMT of CO₂e for the 2030 high scenario (Table 23). These represent increases of 2.5% and 7.2%, respectively, from the 2014 baseline coal, oil, and gas GHG emissions (92.75 MMT of CO₂e). New Mexico federal coal, oil, and gas GHG emissions of 95.09 (2020 high scenario) and 99.35 (2030 high scenario) MMT CO₂e/year would represent 49% and 52% of state 2020 and 2030 high reasonably foreseeable coal, oil, and gas GHG emissions (see Table 23).

Table 23. Reasonably Foreseeable Coal, Oil and Gas Production, and GHG Consumption Emissions in New Mexico, Oklahoma, Kansas, and Texas (Golder Associates 2017)

GHG Emissions (MMT CO₂e)					
Category	New Mexico	Oklahoma	Kansas	Texas	NM, OK, KS, TX
2020 High Scenario					
Federal coal	13.89	1.25	0	0	15.14
Federal oil	25.49	0.33	0.08	0.06	25.95
Federal gas	49.60	0.96	0.29	2.40	53.25
Federal natural gas liquids	6.11	0.09	0.05	0.04	6.29
Total Federal	95.09	2.63	0.42	2.50	100.64
Federal + non-federal coal	43.12	1.87	0.13	97.46	142.58
Federal + non-federal oil	55.28	56.72	22.10	518.06	652.16
Federal + non-federal gas	83.28	152.16	18.14	694.29	947.87
Federal + non-federal natural gas liquids	12.14	20.09	3.14	84.14	119.51
Total federal and non-federal	193.82	230.84	43.51	1,393.95	1,862.12
2030 High Scenario					
Federal coal	10.14	0.91	0	0	11.05
Federal oil	25.60	0.33	0.08	0.06	26.07
Federal gas	57.44	1.11	0.34	2.78	61.67
Federal natural gas liquids	6.17	0.09	0.05	0.04	6.35

GHG Emissions (MMT CO₂e)					
Category	New Mexico	Oklahoma	Kansas	Texas	NM, OK, KS, TX
Total Federal	99.35	2.44	0.47	2.88	105.14
Federal + non-federal coal	31.52	1.37	0.1	71.12	104.11
Federal + non-federal oil	55.51	56.95	22.19	520.20	654.85
Federal + non-federal gas	96.45	176.21	21.02	804.05	1,097.72
Federal + non-federal natural gas liquids	12.25	20.27	3.17	84.88	120.57
Total federal and non-federal	195.73	254.8	46.47	1,480.25	1,977.25

Source: Golder Associates (2017).

Note: Totals may not sum exactly due to rounding.

11.1.2 OKLAHOMA COAL, OIL, AND GAS GREENHOUSE GAS EMISSIONS

BLM’s Oklahoma reasonably foreseeable coal, oil, and gas production and consumption GHG emissions from federal activities are 2.63 MMT of CO₂e for the 2020 high scenario and 2.44 MMT of CO₂e for the 2030 high scenario (see Table 23). This is a decrease of 1.9% and a decrease of 9.8%, respectively, from the 2014 baseline coal, oil, and gas GHG emissions (2.68 MMT of CO₂e). Oklahoma federal coal, oil, and gas GHG emissions of 2.63 MMT (2020 high scenario) and 2.44 (2030 high scenario) MMT CO₂e/year would represent 1.14% and 0.96%, respectively, of state 2020 and 2030 high reasonably foreseeable GHG emissions from coal, oil, and gas activities (see Table 23).

11.1.3 KANSAS COAL, OIL, AND GAS GREENHOUSE GAS EMISSIONS

BLM’s Kansas reasonably foreseeable coal, oil, and gas production and consumption GHG emissions from federal activities are 0.42 MMT CO₂e for the 2020 high scenario and 0.47 MMT CO₂e for the 2030 high scenario (see Table 23). These values represent increases of 4.8% and 14.9%, respectively, compared with the 2014 baseline coal, oil, and gas GHG emissions (0.40 MMT of CO₂e). Kansas federal coal, oil, and gas GHG emissions of 0.42 (2020 High scenario) and 0.47 (2030 High scenario) MMT CO₂e/year would represent 0.97% and 1.01%, respectively, of state 2020 and 2030 high reasonably foreseeable GHG emissions from coal, oil, and gas activities (see Table 23).

11.1.4 TEXAS COAL, OIL, AND GAS GREENHOUSE GAS EMISSIONS

BLM’s Texas reasonably foreseeable coal, oil, and gas production and consumption GHG emissions from federal activities are 2.50 MMT of CO₂e for the 2020 high scenario and 2.88 MMT of CO₂e for the 2030 high scenario (see Table 23). These are an increase of 4.0% and 16.7%, respectively, compared with the 2014 baseline coal, oil, and gas GHG emissions (2.40 MMT of CO₂e). Texas federal coal, oil, and gas GHG emissions of 2.50 (2020 high scenario) and 2.88 (2030 high scenario) MMT CO₂e/year would represent 0.18% and 0.19%, respectively, of state 2020 and 2030 high reasonably foreseeable GHG emissions from coal, oil, and gas activities (see Table 23).

11.1.5 GLOBAL CLIMATE CHANGE PROJECTIONS

Golder Associates (2017:Section 4.0) discusses future climate projections, including four RCPs as identified by the IPCC: RCP 2.6, 4.5, 6.0, and 8.5. The RCP scenarios were developed based on representative GHG emission scenarios including varying assumptions regarding levels of global GHG emissions over time. RCP 8.5 assumes increasing GHG emissions over time, with no stabilization, and is meant to be representative of scenarios leading to high GHG concentration levels. RCP 4.5 and RCP 6.0 represent scenarios where GHG emissions are reduced over time through climate policy. RCP 2.6 represents a scenario where drastic action is taken through stringent climate policy and substantial GHG emission reductions are achieved over time. The pathways are named after the radiative forcing (defined as the difference between insolation [sunlight] absorbed by the Earth and energy radiated back to space) projected to occur by 2100 (e.g., RCP 8.5 would be projected to result in 8.5 W/m² radiative forcing by 2100). The radiative forcing of the atmosphere in each pathway is driven by the concentration of GHGs accumulated in the atmosphere. The RCP characterizations and regions are further described by Golder Associates (2017:Section 4.1) climate change report.

Climate change is driven by radiative forcing, which is influenced by GHG emissions, not annual emission rates from any given subnational project. Figure 18 shows a comparison of global emissions in relation to RCPs 2.6, 4.5, and 8.5, representing low, medium, and high global emissions scenarios.

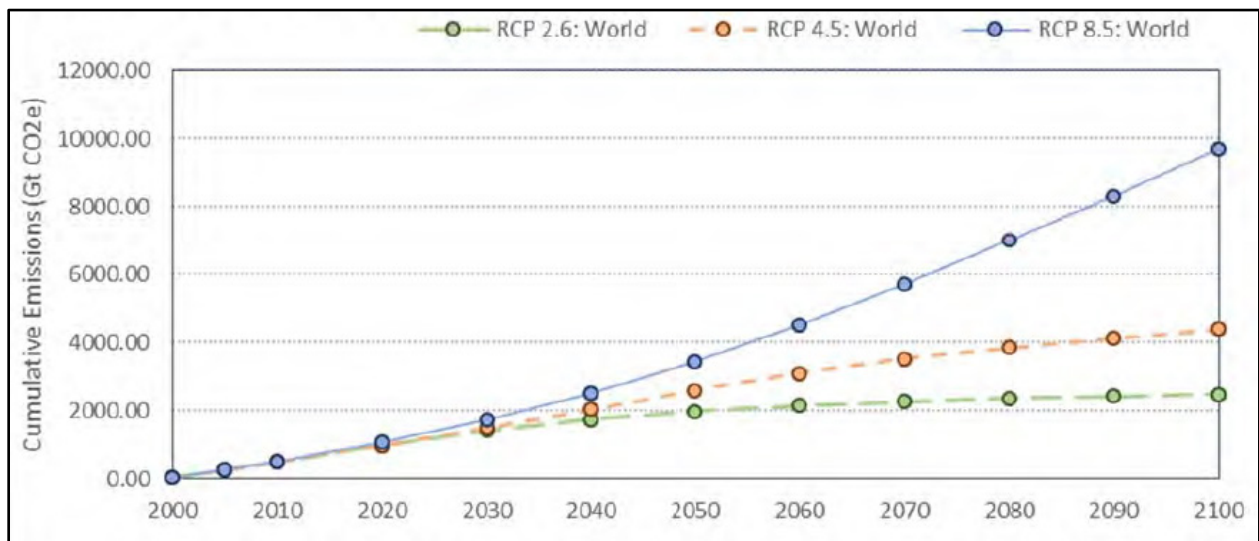


Figure 18. Comparison of RCP 2.6, RCP 4.5, and RCP 8.5 emission estimates over the twenty-first century.

When considering the emissions on a global scale, the annual emission rates of various subnational projects are one of many emission contributions. Any single contribution on a subnational scale is dwarfed by the large number of comparable national and subnational contributors on a global scale. However, the surrogate for understanding the potential effect of BLM's subnational scale emissions on climate is estimating projected annual emission rate due to BLM energy lease Sale Projects (including coal, natural gas, oil, and natural gas liquids). Golder Associates (2017) provides projections of GHG emissions from the 13 western states that regulate most of the federal fossil fuel leasing and compares these emissions with GHG emissions from other contributors. To accomplish this comparison, the Golder Associates demonstrates a comparison of the projected BLM annual emission rates derived from federal lease sale and production information from the 13 western states and compares them against the RCP

scenario emissions profile (a derived value estimating the annual GHG emission rate for each scenario). This comparison is provided in Figure 19. For additional context, 2014 baseline year federal resource production and consumption estimates for these 13 states can be compared with the 2014 baseline national energy consumption and total GHG emissions. BLM subnational emissions in these 13 states are approximately 25.97% of the total national energy consumption emissions and 19.75% of national GHG emission totals at 2014 levels. In 2014, federal mineral production and consumption emissions in these 13 states represented approximately 2.64% of the global totals from all emission sources. With the relative magnitude of these emissions in mind, climate change trends and effects are discussed below.

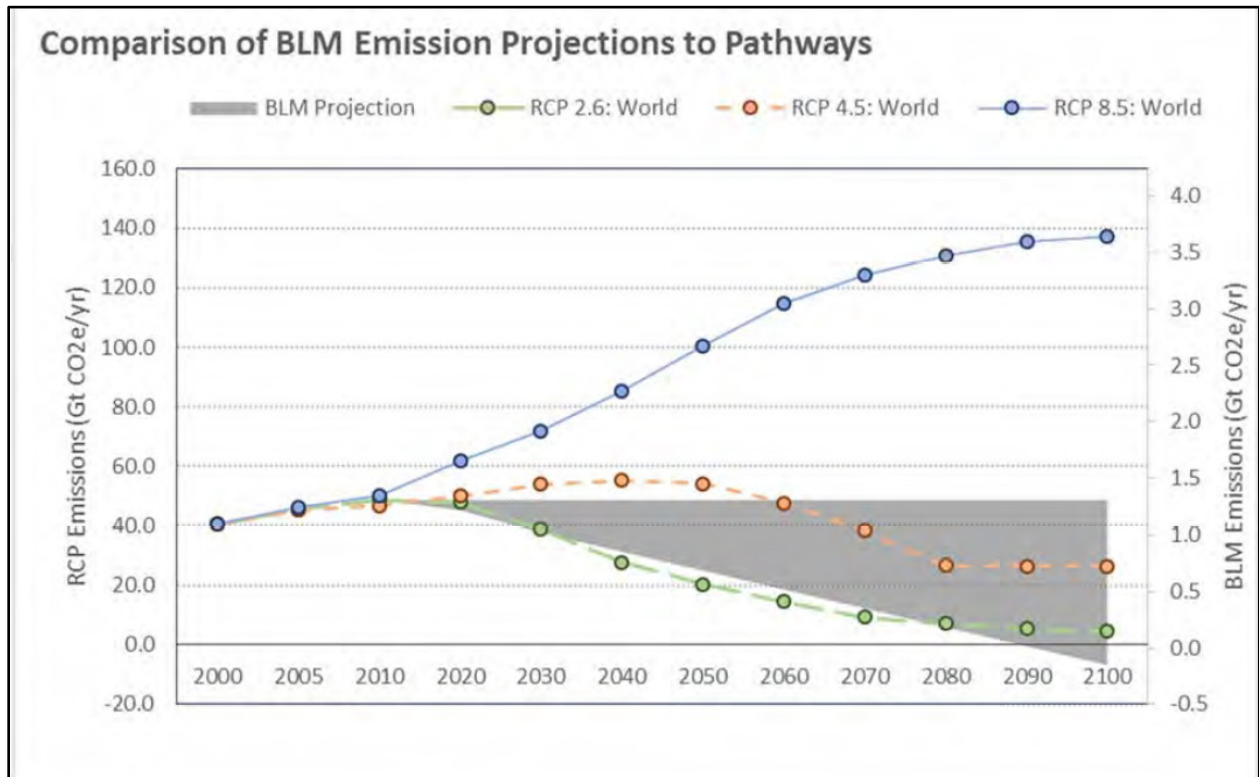


Figure 19. Comparison of BLM emission projections with RCP 2.6, RCP 4.5, and RCP 8.5.

The contribution of GHG emissions from coal, oil, natural gas, and liquefied natural gas for the 13 BLM subject states in 2020 and 2030 under both normal and high production scenarios were evaluated and compared with the GHG emissions profile (the derived annual emission rate for the three RCP scenarios shown in Figure 19). By comparing the relative emission rates of the derived ranges of BLM emissions profiles (low and high estimates) with the RCP scenarios, the BLM emissions most closely track with RCP 8.5 in 2020 and between RCP 2.6 and RCP 4.5 in 2030 (Golder Associates 2017). The reduction in BLM’s emissions profile in 2030 compared with 2020 is a result of a projected change to the federal energy resource mixture. Less coal development is projected, while a slight increase in oil, gas, and natural gas liquids are projected into 2030 relative to 2020. Because coal is the most GHG-intensive fossil fuel, the reduction in this resource development is anticipated to reduce BLM’s lease sale emissions profile (annual GHG emission rate) overall (see Figure 19).

Based on the analysis in Golder Associates (2017), BLM activities are estimated to be conducted at a level that would be in line with the level of emissions anticipated in the RCP 2.6 and RCP 4.5 through 2060. Estimates of BLM activities in future years are more uncertain and have a wider range of variability. The projections presented above are based on best available data and assumptions used to provide context to the BLM’s reasonably foreseeable environmental trends and planned actions. However, due to the levels of uncertainty, some additional information is provided below regarding BLM’s relative contribution to global emissions and, by proxy, climate change. If BLM operates under the business-as-usual scenario while all other contributors are reducing their emissions in line with RCP 2.6, the relative contribution of BLM increases as the emissions more closely resemble RCP 4.5. If BLM operates under the decreased emissions scenario, keeping their reductions in line with RCP 2.6 like all the other contributors, the relative contribution of BLM remains similar to current contributions. If BLM operates under the decreased emissions scenario while all other contributors are maintaining constant emissions (business-as-usual) or increasing emissions, the relative contribution of BLM greatly reduces. It is very unlikely that the global emissions will be strongly influenced by a single contributor at a national or subnational scale. However, the individual behavior of each contributor, through their relative contribution, has the ability to influence which RCP global emissions scenario is most closely resembled and, therefore, which climate change projections are most likely manifested toward the end of the century (Golder Associates 2017).

To understand the effects of climate change, three RCP scenario projections of global temperature and precipitation changes in both the near term (representing the period from 2021 through 2040) and far term (representing the period of 2081 through 2100) are presented in Table 24. These estimates are derived from the average of over 30 different climate change models using the inputs of each RCP scenario.

Table 24. Projected Changes in Climate under Representative Concentration Pathways

RCP Pathway	Near Term		Far Term	
	Temperature (°C)	Precipitation (%)	Temperature (°C)	Precipitation (%)
RCP 2.6	0.78	1.44	0.97	2.27
RCP 4.5	0.85	1.49	1.81	3.51
RCP 8.5	0.98	1.62	3.68	5.89

Under each RCP scenario, projected average global temperatures are expected to increase and changes in precipitation are anticipated. However, generally, the effects of climate change are least severe under the RCP 2.6 scenario and most severe under the RCP 8.5 scenario. Regardless of the specific magnitude of the effects, the effects to global climate are anticipated to include:

- long-term global temperature change;
- intensified droughts impacting agricultural, rural, and urban communities and resulting in changes in land cover and land use;
- intensified and more frequent wildfires;
- sea level rise, ocean warming, and reduced ocean oxygen, impacting global weather patterns and flora and fauna;

- intensified flooding impacting infrastructure, natural resource–based livelihoods, and cultural resources; and
- human health, such as heat-associated deaths and illnesses, chronic diseases, and other health issues associated with poor air quality (Gonzalez et al. 2018).

To understand climate change effects within the areas managed by the BLM state office, a brief discussion of potential impacts in each region (the southern Colorado/New Mexico area and the southern Great Plains Area) follows.

11.1.5.1 COLORADO/SOUTHERN NEW MEXICO

Climate modeling suggests that annual average temperatures in this region may rise by 4°F to 6°F by the end of the twenty-first century, with warming increasing from south to north. By 2080 to 2090, the southwestern United States would see a 10% to 20% decline in precipitation, primarily in winter and spring, with more precipitation falling as rain. A recent U.S. Bureau of Reclamation report (U.S. Bureau of Reclamation et al. 2013) made the following projections through the end of the twenty-first century for the Upper Rio Grande Basin (southern Colorado to central-southern New Mexico) based on the current and predicted future warming:

- There would be decreases in overall water availability by one-quarter to one-third.
- The seasonality of stream and river flows would change, with summertime flows decreasing.
- Stream and river flow variability would increase. The frequency, intensity, and duration of both droughts and floods would increase (U.S. Bureau of Reclamation et al. 2013).

The Bureau of Reclamation report also noted that reduction in water is expected to make environmental flows in the Upper Rio Grande system more difficult to maintain and reduce the shallow groundwater available to riparian vegetation. Both effects have implications for the habitat of fish and wildlife in the Upper Rio Grande Basin riparian ecosystems (U.S. Bureau of Reclamation et al. 2013). A USFS assessment of 117 species of birds, reptiles, amphibians, and mammals along the Middle Rio Grande in New Mexico (Friggens et al. 2013 as cited in U.S. Bureau of Reclamation et al. 2013) projected decreasing availability of riparian habitat, and loss of mature trees due to fire and disease that would directly and indirectly effect many species of birds and mammals. Most evaluated species were projected to experience negative effects from climate change; however, a few species that are considered generalists and highly adaptable, such as coyotes, jackrabbits, some lizards, and road runners may benefit from conversion of the bosque to a more sparsely vegetated and drier habitat (Friggens et al. 2013 as cited in U.S. Bureau of Reclamation et al. 2013).

11.1.5.2 GREAT PLAINS

Impacts anticipated in the Great Plains region (including Texas, Oklahoma, and Kansas) are used. The Great Plains region is projected to experience increases in temperatures and more frequent drought in the future. Temperature increases and precipitation decreases would stress the region’s primary water supply, the Ogallala Aquifer. Seventy percent (70%) of the land in this area is used for agriculture. Threats to the region associated with climate change include:

- pest migration as ecological zones shift northward;

- increases in weeds; and
- decreases in soil moisture and water availability (U.S. Bureau of Reclamation et al. 2013).

11.2 U.S. GEOLOGICAL SURVEY END-USE AND EXTRACTION ANALYSIS

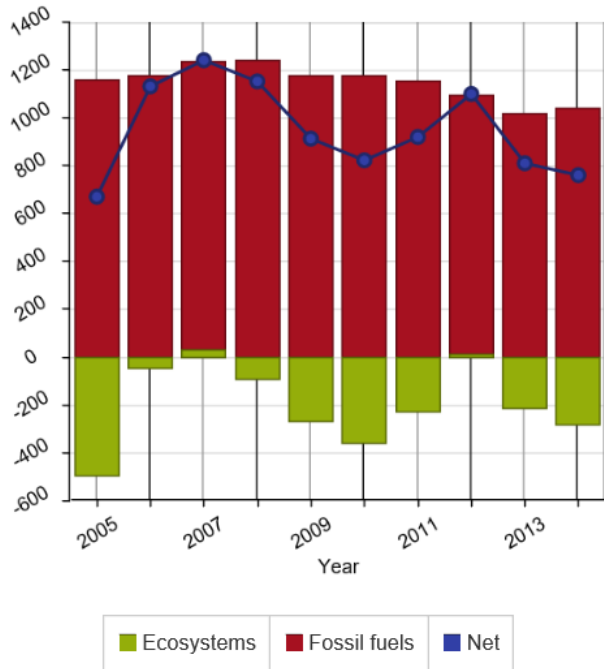
In November of 2018, the USGS published a scientific investigation report, *Federal Lands Greenhouse Gas Emissions and Sequestration in the United States: Estimates 2005-2014* (Merrill et al. 2018). The report consists of a 44-page document with four companion datasets and an interactive online mapping site in which the user can pull up data for each state (28 states included in analysis) and two offshore sites. The data itself consists of 10 years of emissions and sequestration estimates in which the emissions from combustion and extraction activities on federal lands from fossil fuels is converted into carbon dioxide equivalents (CO₂e) and measured in MMT/year of CO₂e. The estimates include the three most prominent GHGs: carbon dioxide (CO₂), methane (CH₄), and nitrous oxide (N₂O). The results are presented by state and year and the estimates are broken into categories by the sector of the economy where the combustion or extraction related emissions occurred, or the biologic process being quantified occurred. The data presents both gross and net emissions after sequestration is accounted for. For the purpose of this analysis the BLM quantifies all emissions in CO₂ equivalents (CO₂e). For context of gross and net emissions as well as sequestration activities, the BLM shows the total U.S. and New Mexico emissions from combustion and extraction activities on federal lands as well as sequestration activity (Figures 20 and 21). American Indian and Tribal lands were not included in the analysis. Additionally, the national total (gross) emissions include two offshore areas (Merrill et al. 2018).

11.2.1 GREENHOUSE GAS EMISSIONS (COMBUSTION AND EXTRACTION) FROM U.S. FEDERAL LANDS (CO₂E)

In 2014, end-use combustion and extraction (C&E) of fossil fuels produced on U.S. federal lands was 1,332 MMT of CO₂e. This reported value includes emissions from the combustion of coal, oil and natural gas from fossil fuels produced on U.S. federal lands as well as extraction emissions from activities occurring on federal lands. When compared to 2005 emissions, this results in a decrease of emissions throughout all the three prominent GHG emissions. From 2005 to 2014, GHG emissions from end-use C&E of fossil fuels produced on federal lands have resulted in an overall trend of decreased emissions (Figure 20). When compared to global and national total CO₂e emissions, 48,257 and 6,558.3 MMT respectively, from all sources (Table 25), CO₂e emissions from these activities (end-use combustion and extraction activities) of fossil fuels produced on federal lands is 2.8% and 19.4% respectively (World Resources Institute 2007; EPA 2021n). Of the 1,332 MMT CO₂e, 80.53 MMT were exported end-use combustion emissions, 752.50 MMT represented emissions from coal sources while 498.76 MMT were the result of oil and natural gas source. Figures 22 and 23 provide a graphical representation of CO₂e emissions from the fossil fuels produced on U.S. federal lands associated with end-use combustion and extraction activities.

U.S. federal lands also contribute a great deal to the sequestration of CO₂ and provide carbon storage (sinks) for CO₂ emissions. In 2014 U.S. federal lands provided 283.2 MMT of carbon storage. U.S. federal lands sequestered an average of 195 MMT of CO₂e between 2005 and 2014 offsetting approximately 15% of the CO₂ emissions resulting from the extraction of fossil fuels on federal lands and their end-use combustion (Figure 20) (Merrill et al. 2018).

National onshore CO₂ emissions and sequestration: 2005-14



Year	Total fossil fuels	Onshore fossil fuels	Onshore ecosystems	Onshore net
2005	1,361.9	1,162.3	-493.5	668.8
2006	1,378.6	1,178.1	-47.4	1,130.7
2007	1,398.3	1,207.3	32.7	1,240.0
2008	1,427.9	1,243.7	-92.3	1,151.4
2009	1,422.5	1,178.5	-266.6	911.9
2010	1,429.4	1,180.6	-359.5	821.1
2011	1,362.4	1,156.2	-237.5	918.7
2012	1,280.5	1,085.8	13.0	1,098.8
2013	1,210.5	1,022.3	-213.6	808.7
2014	1,279.0	1,042.2	-283.2	759.0

Figure 20. National CO₂ emissions and sequestration: 2005–2014

Source: Merrill et al. (2018).

All values are in million metric tons of CO₂ equivalent (MMT CO₂e) Total fossil fuels includes offshore emissions from two areas.

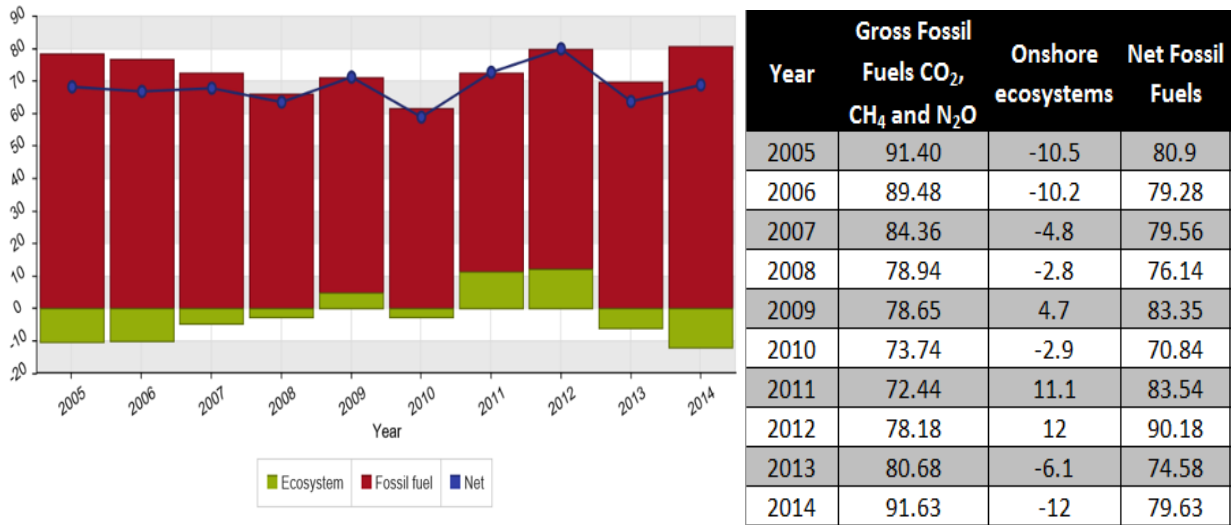


Figure 21. New Mexico CO₂ emissions and sequestration: 2005–2014

Source: Merrill et al. (2018).

While the USGS total values include GHG emissions from end-use combustion and from extraction activities of coal, oil and gas, in this section the BLM focuses on the end-use combustion emissions generated from the oil and natural gas sector, the BLM excludes coal totals (Table 25 and Figure 23).

Table 25. GHG Emissions, Combustion and Extraction, from U.S. Federal Lands (CO₂e) (World Resources Institute 2017; EPA 2021n; Merrill et al. 2018)

Level/Sector	MMT CO ₂ e
Global emissions, all sources	48,257
National emissions, all sources*	6457
End-use C&E emissions (federal lands) ^{(1),(2)}	1,332
% end-use C&E emissions (federal lands) to Global Emissions ⁽²⁾	2.76
% end-use C&E emissions (federal lands) to National Emissions ⁽²⁾	19.39
End-use combustion only emissions (federal lands) ⁽²⁾	1,201
% end-use combustion only emissions (federal lands) to global emissions ⁽²⁾	2.49
% end-use combustion only emissions (federal lands) to national emissions ⁽²⁾	17.48
Extraction only emissions (federal lands) ⁽²⁾	50.52
% of extraction only emissions (federal lands) to global emissions ⁽²⁾	0.10
% of extraction only emissions (federal lands) to national emissions ⁽²⁾	0.74
End-use C&E emissions (federal lands) O&G only ⁽³⁾	499
% end-use C&E emissions (federal lands) O&G only to global emissions ⁽³⁾	1.03
% end-use C&E emissions (federal lands) O&G only to national emissions ⁽³⁾	7.26

Level/Sector	MMT CO ₂ e
End-use combustion only emissions (federal lands) O&G only ⁽³⁾	460
% end-use combustion only emissions (federal lands) O&G only to global emissions ⁽³⁾	0.95
% end-use combustion only emissions (federal lands) O&G only to national emissions ⁽³⁾	6.70
Extraction only emissions (federal lands) O&G only ⁽³⁾	38.76
% extraction only emissions (federal lands) O&G only to global emissions ⁽³⁾	0.08
% extraction only emissions (federal lands) O&G only to national emissions ⁽³⁾	0.56

* Emissions reflect data from 2019 EPA GHG Inventory report, newer inventories may correct this value somewhat.

⁽¹⁾ Includes 80.53 MMT of exported CO₂e emissions. Emission totals are: CO₂ 1,290 MMT, CH₄ 47.6 MMT of CO₂e, N₂O 5.5 MMT of CO₂e

⁽²⁾ Includes emissions from coal, oil, and natural gas

⁽³⁾ Isolates coal from the total and only includes oil and natural gas CO₂e emissions

C&E=combustion and extraction

Global emissions represented are for 2013, national emissions and federal land emissions are for 2014.

O&G=oil and gas

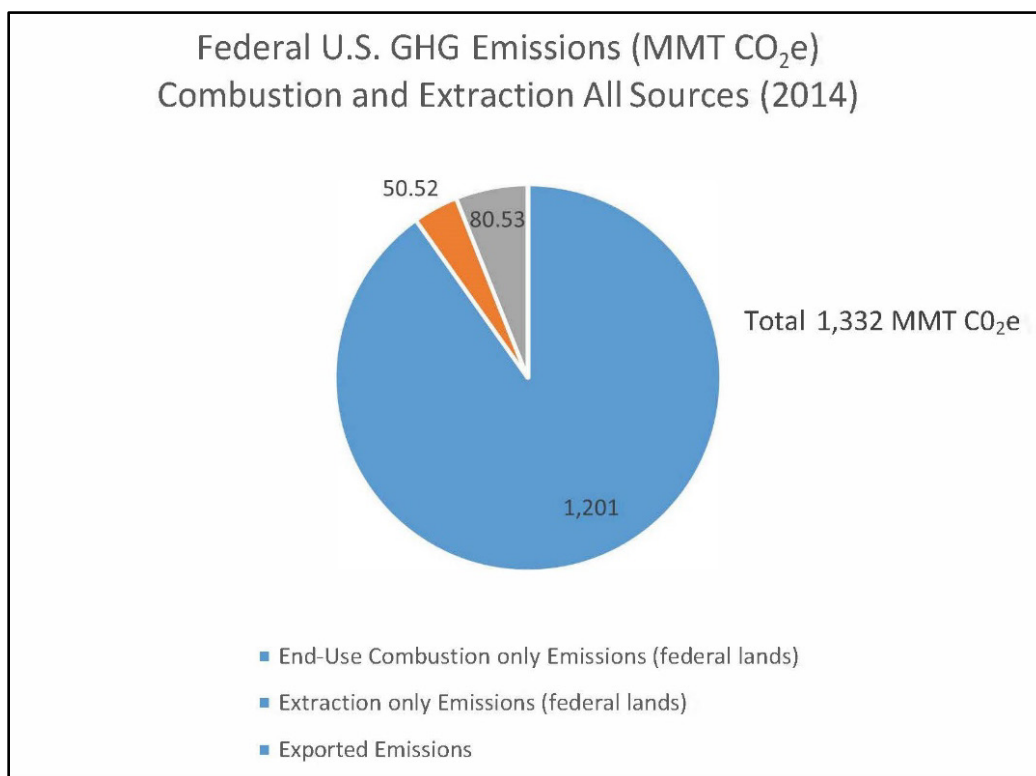


Figure 22. Federal U.S. GHG emissions (MMT CO₂e) combustion and extraction all sources (2014)

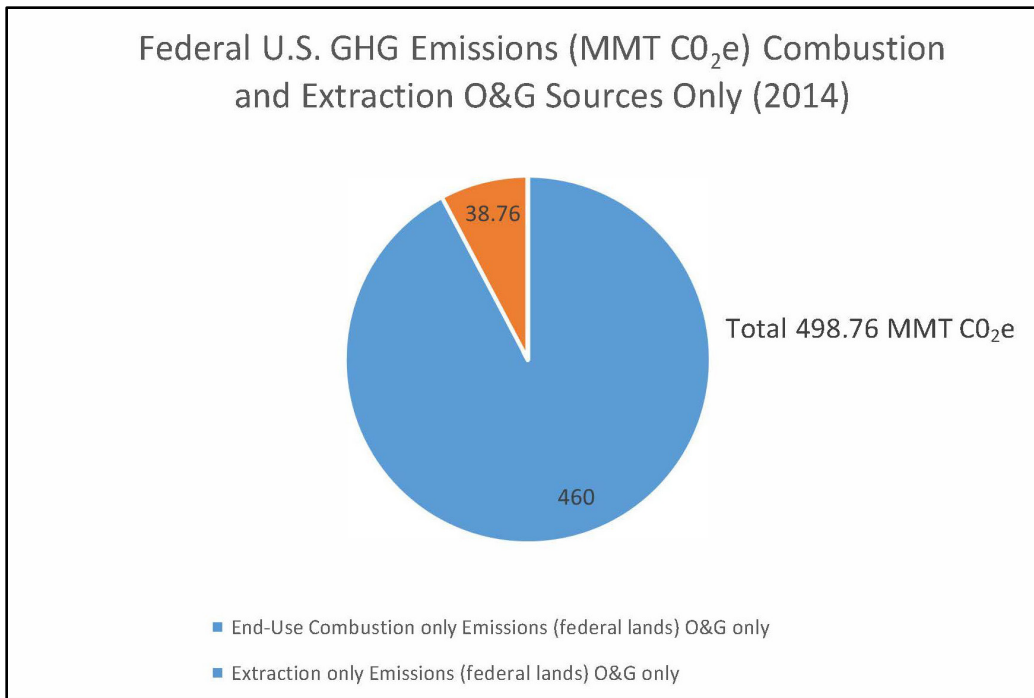


Figure 23. Federal U.S. GHG Emissions (MMT CO₂e) combustion and extraction O&G sources only (2014)

11.2.2 GREENHOUSE GAS EMISSIONS (COMBUSTION AND EXTRACTION) FROM NEW MEXICO FEDERAL LANDS (CO₂E)

In 2014, end-use combustion and extraction (C&E) of fossil fuels produced on New Mexico federal lands was 91.63 (MMT) of CO₂e. This reported value includes emissions from the combustion of coal, oil and natural gas from fossil fuels produced on federal lands as well as extraction emissions from activities occurring on federal lands. When compared to 2005 emissions this results in increased emissions throughout all the three prominent GHG emissions. From 2005 to 2014, GHG emissions from end-use C&E of fossil fuels produced on federal lands have resulted in average annual emissions of 81.95 MMT of CO₂e (Figures 20 and 21). When compared to global and national total CO₂e emissions, 48,257 and 6,558.3 MMT respectively, from all sources (Table 25), CO₂e emissions from these activities (end-use combustion and extraction activities) of fossil fuels produced on New Mexico federal lands is 0.19% and 1.33% respectively (World Resources Institute 2017; EPA 2021n).

In 2014 New Mexico federal lands provided 12 MMT of carbon storage. Federal lands sequestered an average of 9.5 MMT of CO₂e between 2005 and 2014 (Figure 21) (Merrill et al. 2018). While the USGS total values include GHG emissions from end-use combustion and from extraction activities of coal, oil, and gas, for the purposes of this analysis the BLM only focuses on the end-use combustion emissions generated from the oil and natural gas sector, the BLM exclude coal totals (Table 26 and Figure 25).

Table 26. GHG Emissions, Combustion and Extraction, from BLM New Mexico (CO₂e) (World Resources Institute 2017; EPA 2021n; Merrill et al. 2018)

Level/Sector	MMT CO ₂ e
Global emissions, all sources	48,257
National emissions, all sources*	6456.7
End-use C&E emissions (BLM NM) ^{(1), (2)}	91.63
% end-use C&E emissions (BLM NM) to global emissions ⁽²⁾	0.19
% end-use C&E emissions (BLM NM) to national emissions ⁽²⁾	1.33
End-use combustion only emissions (BLM NM) ⁽²⁾	73
% end-use combustion only emissions (BLM NM) to global emissions ⁽²⁾	0.15
% end-use combustion only emissions (BLM NM) to national emissions ⁽²⁾	1.06
Extraction only emissions (BLM NM) ⁽²⁾	12.76
% of extraction only emissions (BLM NM) to global emissions ⁽²⁾	0.03
% of extraction only emissions (BLM NM) to national emissions ⁽²⁾	0.19
End-use C&E emissions (BLM NM) O&G only ⁽³⁾	66.35
% end-use C&E emissions (BLM NM) O&G only to global emissions ⁽³⁾	0.14
% end-use C&E emissions (BLM NM) O&G only to national emissions ⁽³⁾	0.97
End-use combustion only emissions (BLM NM) O&G only ⁽³⁾	54.58
% end-use combustion only emissions (BLM NM) O&G only to Global Emissions ⁽³⁾	0.11
% end-use combustion only emissions (BLM NM) O&G only to National Emissions ⁽³⁾	0.79
Extraction only Emissions (BLM NM) O&G only ⁽³⁾	11.77
% Extraction only Emissions (BLM NM) O&G only to global emissions ⁽³⁾	0.02
% Extraction only Emissions (BLM NM) O&G only to national emissions ⁽³⁾	0.17

* Emissions reflect data from 2017 EPA GHG Inventory report, newer inventories may correct this value somewhat.

⁽¹⁾ Includes 5.86 MMT of exported CO₂e emissions. Emission totals are: CO₂ 74.78 MMT, CH₄ 10.78 MMT of CO₂e, N₂O 0.22 MMT of CO₂e

⁽²⁾ Includes emissions from coal, oil, and natural gas

⁽³⁾ Isolates coal from the total and only includes oil and natural gas CO₂e emissions

C&E=combustion and extraction

Global and state-wide emissions represented are for 2013, national emissions and federal land emissions are for 2014.

O&G=oil and gas

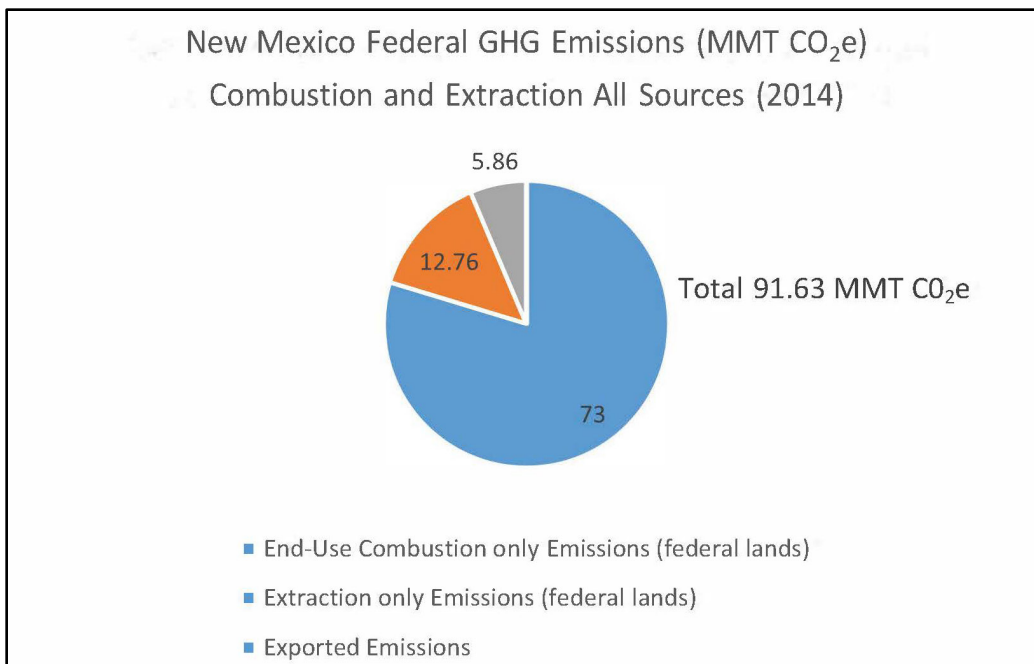


Figure 24. New Mexico Federal GHG Emissions (MMT CO₂e) combustion and extraction all sources (2014)

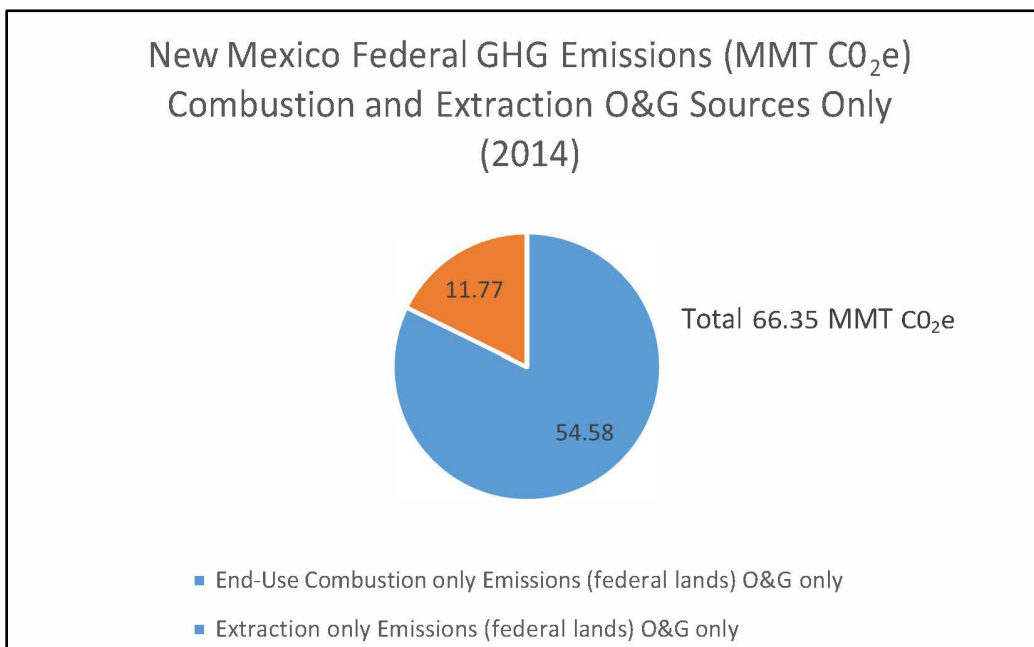


Figure 25. New Mexico Federal GHG Emissions (MMT CO₂e) combustion and extraction O&G sources only (2014)

11.3 REASONABLY FORESEEABLE FUTURE ACTIONS AFFECTING GREENHOUSE GAS EMISSIONS

Overall, total New Mexico statewide gross GHG emissions are expected to decrease (CNEE 2020). The New Mexico Greenhouse Gas Inventory and Forecast Report 2020 projects the following for year 2030 in New Mexico for emissions produced within the State (i.e., production-based emissions):

- Gross GHG emissions of 96.6 MMT of CO₂e—an increase of 22% relative to 2005 and a decrease 15% relative to 2018. New Mexico’s emissions are more than twice the national average of GHG emissions per capita. New Mexico’s high per capita emissions are largely the result of GHG - intensive oil and gas industry, which makes up a significant portion of overall GHG emissions profile.
- Top sources of GHG emissions: transportation fuel use (15.4 MMT of CO₂e,) electricity generation (12.9 MMT of CO₂e,) and oil and gas (fugitive and fuel emissions) (32.5 MMT of CO₂e,). Transportation fuel and electricity generation decreased over 2005 estimates, but oil and gas increased.
- Approximately 43 MMT of CO₂e are projected as a result of oil and natural gas production, processing, transmission and distribution. This is 44.5% of the gross New Mexico emissions (a slight decrease over the relative contribution of oil and gas production in 2018, 53.0% contribution and an increase over the relation contribution of oil and gas production in 2005, 25.0 % contribution).

All scenarios see a significant rise in emissions from 2005 to 2018, as well as a significant drop from 2018 to 2023, driven primarily by the New Source Performance Standards (NSPS) for the oil and gas sector (CNEE 2020).

12 MITIGATION

The reduction of emissions of air pollutants and GHGs from oil and gas operations has been the subject of much study and discussion in recent years. The EPA Natural Gas Star Program established in 1993 has been a leader in developing and reporting on strategies to reduce methane (CH₄) emissions (EPA 2021p). These reductions can help to control not only GHGs but also VOCs, which contribute to O₃ formation. Numerous opportunities for emissions reduction, including costs to implement, are documented on EPA’s the Natural Gas Star website.

In 2021, according to BLM AFMSS 2 Data, EPA Natural Gas Star partner companies operated 0.29% of the active federal wells in the New Mexico portion of the San Juan Basin and 12.4% of the active federal wells in the New Mexico portion of the Permian Basin. In Oklahoma, Kansas, and Texas, Natural Gas Star partner companies operate 1.76% of the active federal wells (BLM 2021a). EPA has found Natural Gas Star partners’ actions to result in measurable decreases in GHG emissions since the program’s implementation. In October 2012, EPA promulgated air quality regulations controlling VOC emissions at hydraulically fractured gas wells. These rules require air pollution mitigation measures that reduce the emissions of VOCs. These same mitigation measures have a co-benefit of reducing CH₄ emissions.

The EPA has New Source Performance Standards (NSPS) (codified in 40 CFR 60) in place to reduce VOC emissions from oil and gas sources. NSPS OOOO requires reduction of VOCs from well completion operations and storage tanks constructed after August 23, 2011. NSPS OOOOa requires reduction of VOCs from well completion operations from new or re-fractured hydraulically fractured wells and requires reduction of storage tank emissions by 95% for tanks constructed after September 18, 2015, with emissions greater than 6 tons per year of VOC (this has the co-benefit of reducing CH₄ emissions as well). NSPS OOOOa also imposes semiannual leak detection and repair requirements for the collection of fugitive emission components at well sites constructed after September 18, 2015, that produce more than 15 bbl of oil and/or gas per day. NSPS OOOOa also requires scheduled maintenance and/or emission control devices for reciprocating and centrifugal compressor venting at compressor stations and includes provisions to limit emissions from natural gas pneumatic devices and pumps. Following the 2020 amendment to OOOO and OOOOa, fugitive emissions monitoring is required only for those wells producing greater than 15 bbl per day. These provisions aim to reduce fugitive emissions of VOC at oil and gas facilities.

The NMED and EMNRD have developed rules that will regulate CH₄ emissions. The departments were charged with this task under the [Executive Order on Addressing Climate Change and Energy Waste Prevention](#) of Gov. Michelle Lujan Grisham. The order instructs NMED and the Energy, Minerals and Natural Resources Department (EMNRD) to “jointly develop a statewide, enforceable regulatory framework to secure reductions in oil and gas sector CH₄ emissions and to prevent waste from new and existing sources and enact such rules as soon as practicable” (NMED 2019d). The Natural Gas Waste Reduction Rule will reduce unnecessary natural gas venting and flaring from new and existing sources. Provisions will reduce CH₄ emissions based on stringent limitations on natural gas venting that results in non-combustible CH₄ emissions. Additionally, recovery and reuse of natural gas will reduce flaring of CO₂, CH₄, and nitrous oxide (N₂O) emissions.

A report by the U.S. Government Accountability Office (GAO) noted that opportunities exist for capturing fugitive emissions from venting and flaring of natural gas on wells under federal jurisdiction (GAO 2010). A report prepared for BLM in Montana includes an entire chapter on reduction of emissions of GHGs (URS 2010). Another report recently issued by the USFS summarizes and builds on work originally done by BLM to identify Best Management Practices for protection of air quality during oil and gas development and production (USFS 2011). Rapid development could result in an increase of criteria and HAP emissions in the planning areas. Limiting development through a phased approach could help to reduce concentrations of emissions in the air basins.

Emissions associated with the RFD, including future potential development of leases, would be offset by substantial decreases in emissions—including a 67% reduction in SO₂, 62% reduction in NO_x, 50% reduction in PM, 44% reduction in CO, and 51% reduction in VOCs—resulting from power generation due to the recent shutdown of two of the units at the San Juan Generating Station. Additionally, selective catalytic reduction technology installed on the two remaining coal-fired generators at the Four Corners Power Plant would result in additional reductions in emissions from the facility, including a 36% reduction in NO_x, 43% reduction in PM, and 24% reduction in SO₂. The San Juan Generating Station is also proposed for full closure by 2022, which would result in even further drops in future pollutant emissions for the analysis area. New Mexico will have to comply with the Federal Regional Haze Rule requirements as it develops its SIP for the second planning period. New Mexico is currently in the 2021 Regional Haze Planning Process and is in the process of updating its Regional Haze SIP. The submittal of the Proposed SIP to EPA Region 6 is expected in early 2022 (NMED 2021).

Texas proposed a 2021 Regional Haze SIP Revision that is designed to address regional haze in Big Bend and Guadalupe Mountains National Parks in Texas and Class I areas located outside of Texas that may be affected by emissions from within the state (TCEQ 2021).

As part of the process of developing Oklahoma's 2021 Regional Haze SIP, the ODEQ identified 12 facilities that are reasonably anticipated to impact visibility conditions at the Wichita Mountains Wilderness Area and identified 21 sources in neighboring states that are reasonably anticipated to impact visibility conditions at the Wichita Mountains Wilderness Area (including sources in Texas) and have asked these states to consider the potential impact of the sources identified within their states for further analysis as part of the process for developing their 2021 Regional Haze SIP (ODEQ ADQ 2021).

Cumulatively, it is expected that future levels of criteria pollutant, VOC, HAP, and GHG emissions would be lower than current levels due to the aforementioned factors despite the increases in emissions associated with reasonably foreseeable oil and gas development and future potential development of leases.

While it is beyond the scope of this report to detail the wide range of mitigation strategies available it must be understood that for the most part these strategies must be applied on a case-by-case basis at the project level.

13 OTHER TOPICS

13.1 FOUR CORNERS AIR QUALITY TASK FORCE

In 2002, NMED and local governments convened to sign an Early Action Compact for O₃ under an EPA program that required commitment for state and local action to resolving O₃ issues prior to a nonattainment designation. In 2005, the states of Colorado and New Mexico convened a group of stakeholders, then known as the FCAQTF, to address air quality issues in the Four Corners region in light of continued energy development and growth in the region and consider options for mitigating air pollution. A report detailing a wide range of mitigation options was published in November 2007 (FCAQTF 2007).

In 2008, its task complete, the group became known as the Four Corners Air Quality Group (FCAQG) and continued on as a forum for discussion of existing air quality issues and potential solutions. The FCAQG is currently comprised of more than 100 members and 150 interested parties representing a wide range of perspectives on air quality in the Four Corners region. Members include private citizens, representatives from public interest groups, universities, industry, state, tribal and local governments, and federal agencies. The BLM has been an active participant from the beginning and maintains a representative on the steering committee. The last FCAQG met Wednesday, October 23, 2019, from 9 a.m. to 4:30 p.m. at the Durango Public Library, 1900 E 3rd Ave, Durango, Colorado 81301. For more information visit the FCAQG at the NMED website <https://www.env.nm.gov/air-quality/fcagg/>.

13.2 ELECTRICAL GENERATING UNITS

There are two coal-fired electrical generation units (EGUs) in the Four Corners area: the San Juan Generating Station, located 15 miles west of Farmington, New Mexico; and the Four Corners Power Plant, located on Navajo Nation land in Fruitland, New Mexico. These EGUs are the primary source of several criteria air pollutants in the FFO area, including SO₂ (85%), NO_x (41%), and PM_{2.5} (3%)

(EPA 2020b). EGUs are responsible for 31% of New Mexico GHG emissions and 31% of U.S. GHG emissions (NMED 2016).

In 2013, NMED, Public Service Company of New Mexico and EPA agreed to meet the requirements of the federal regional haze rule through the shutdown of two units at the San Juan Generating Station by the end of 2017. The agreement also requires the installation of selective non-catalytic reduction technology on the remaining two units. This resulted in significant reductions from the previous emissions levels of many pollutants: a 67% reduction in SO₂, 62% reduction in NO_x, 50% reduction in PM, 44% reduction in CO, 51% reduction in VOC, 50% reduction in CO₂ and 50% reduction in mercury. The New Mexico Environmental Improvement Board approved a revision to the SIP containing the agreement requirements in the fall of 2013.

In December 2013, three coal-fired generators were shut down at the Four Corners Power Plant as part of a plan to meet the requirements of the federal regional haze rule. The remaining two coal-fired generators had selective catalytic reduction technology installed in 2018. These changes satisfy BART requirements from EPA. This will result in significant reductions from current emissions levels of many pollutants: a 36% reduction in NO_x, a 61% reduction in mercury, a 43% reduction in PM, a 30% reduction in CO₂, and a 24% reduction in SO₂. In Texas, NO_x emissions from EGUs in O₃ nonattainment areas (Beaumont–Port Arthur, Dallas–Fort Worth and Houston–Galveston–Brazoria) are required to limit NO_x emissions from utility boilers, auxiliary steam boilers, stationary gas turbines and duct burners under 30 TAC Chapter 117, Subchapter C. The Texas proposed regional haze SIP did not require BART-eligible EGUs to install controls because the state of Texas determined impact of each plant's emissions did not significantly degrade visibility in a Class I area, or facilities had already reduced emissions or shut down units. On December 16, 2014, EPA proposed to partially disapprove the Texas regional haze SIP and also proposed a Federal Implementation Plan to require SO₂ emissions reductions at fifteen Texas BART-eligible sources.

In Oklahoma, Tulsa Public Service Company of Oklahoma retired one coal-fired unit at Oologah in April 2016 and installed a dry sorbent injection system on a second coal-fired unit at the same time. The second unit will be shut down by December 31, 2026, to meet the requirements of the federal regional haze rule. In 2016, SO₂ emissions were reduced by 78% and NO₂ emissions were reduced by 81%. In 2011, EPA disapproved the Oklahoma SIP revision plan for controls at Oklahoma Gas and Electric's Sooner and Muskogee Units and the AEP/PSO Northeastern Units 3 and 4. EPA determined that dry scrubber control technology was needed at these units to meet federal haze rule requirements. The disapproval has been challenged by the State of Oklahoma, upheld by the courts and has now been appealed to the Supreme Court by the State of Oklahoma. Oklahoma submitted a SIP revision in 2013 that was approved by EPA in March 2014 that revises the BART determination for AEP/PSO Units 3 and 4. The revised determination included short-term compliance with emissions limits, shut down of one of the units by April 16, 2016, and shut down of the other unit by December 31, 2026.

In Kansas, emissions at four coal-fired units were significantly reduced as a result of the federal regional haze rule. At Kansas City Power and Electric's La Cygne plant, SCR was installed on both units and scrubbers were installed. This resulted in 83% reduction in NO_x emissions and 82% reduction in SO₂ emissions. At Westar's Jeffrey coal-fired units, low-NO_x burner installation and switching to natural gas combustion resulted in an 82% reduction in NO_x emissions and a 34% reduction in SO₂ emissions.

13.3 INFRARED CAMERAS

The BLM has two infrared cameras which are being used to detect leaks and fugitive emissions. BLM inspectors carry these cameras into the field and have been able to alert operators of equipment requiring repair or maintenance. At this time, the cameras are being used in an advisory rather than a regulatory role.

13.4 FOUR CORNERS METHANE HOTSPOT

In 2014, pioneering research using space-borne (satellite and aircraft) determinations of methane (CH₄) concentrations indicated anomalously large CH₄ concentrations in the Four Corners region including the northern portion of the Farmington planning area (Kort et al. 2014). A subsequent study (Schneising et al. 2014) also indicated larger anomalies over other oil and gas basins in the United States. This results from nighttime/early morning trapping of CH₄ (as well as non-methane hydrocarbons) in low-lying areas due to the San Juan Basin's topography and prevailing meteorological conditions. CH₄ is 34 times more potent at trapping GHG emissions than CO₂ when considering a time horizon of 100 years (IPCC 2013).

Sources contributing to the hotspot include fossil fuel operations, 66% to 75% of which is clearly attributed to natural gas and coalbed CH₄ operations (Pétron et al. 2020). A significant amount of CH₄ is emitted during oil and gas well completion. CH₄ is also emitted from process equipment, such as pneumatic controllers and liquid unloadings, at oil and gas production sites. Ground-based, direct source monitoring of pneumatic controllers conducted by the Center for Energy and Environmental Resources show that CH₄ emissions from controllers exhibit a wide range of emissions and a small subset of pneumatic controllers emitted more CH₄ than most (Allen, Pacsi, et al. 2014). Emissions measured in the study varied significantly by region of the United States, the application of the controller and whether the controller was continuous or intermittently venting. The Center for Energy and Environmental Resources had similar findings of variability of CH₄ emissions from liquid unloading (Allen, Sullivan, et al., 2014).

In 2016, the results from an April 2015 study were released in which researchers conducted further ground-based and space-borne studies utilizing emerging pollutant measurement technology. The NASA Jet Propulsion Laboratory conducted these studies using two JPL airborne spectrometers to identify and measure more than 250 individual sources of CH₄. The sources emitted the gas at rates ranging from a few pounds to 11,000 pounds (5,000 kilograms) per hour (NASA 2016). Overall, observed sources included gas processing facilities, storage tanks, pipeline leaks, and well pads, as well as a coal mine venting shaft. Using equipment enhancements and inferred fluxes, the CH₄ plumes showed that the top 10% of emitters contributed 49 to 66% to the inferred total point source flux of 0.23 Tg/year to 0.39 Tg/year. To understand more about the terminology and study, results are published at the Proceedings of the National Academy of Sciences in a paper titled "*Airborne methane remote measurements reveal heavy-tail flux distribution in Four Corners region.*" (Frankenburg et al. 2016).

Information on CH₄ may also be found in a new interactive mapping tool launched by NMED in 2019. This tool shows CH₄ hotspot information as well as information on CH₄ permits. The mapping tool shows elevated CH₄ levels along the northern border of San Juan County and western border of Rio Arriba County, New Mexico and along the southern borders of Montezuma County and La Plata County, Colorado. It also provides locations of NMED-permitted oil and gas wells and tank batteries for permits greater than 10 tons of CH₄ emissions per year. These sources are concentrated along State Route 550 in San Juan, Rio Arriba, and Sandoval Counties, northeast of CCNHP (NMED 2019e).

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15 APPENDICES

15.1 APPENDIX A NATIONAL EMISSIONS INVENTORY (NEI)

15.2 APPENDIX B NATIONAL AIR TOXICS ASSESSMENT (NATA)

15.3 APPENDIX C CLIMATE NORMALS

15.4 APPENDIX D MAJOR SOURCES (NEI)

15.5 APPENDIX E AIR QUALITY MEMORANDUM OF UNDERSTANDING (MOU)

15.6 APPENDIX F DEPOSITION CHARTS

15.7 APPENDIX G CALCULATORS