ANNEX 6 Additional Information

6.1. Global Warming Potential Values

Global Warming Potential (GWP) is intended as a quantified measure of the globally averaged relative radiative forcing impacts of a particular greenhouse gas. It is defined as the cumulative radiative forcing—both direct and indirect effects integrated over a specific period of time from the emission of a unit mass of gas relative to some reference gas (IPCC 2007). Carbon dioxide (CO_2) was chosen as this reference gas. Direct effects occur when the gas itself is a greenhouse gas. Indirect radiative forcing occurs when chemical transformations involving the original gas produce a gas or gases that are greenhouse gases, or when a gas influences other radiatively important processes such as the atmospheric lifetimes of other gases. The relationship between kilotons (kt) of a gas and million metric tons of CO_2 equivalents (MMT CO_2 Eq.) can be expressed as follows:

$$MMT \ CO_2 \ Eq. = (kt \ of \ gas) \times (GWP) \times \left(\frac{MMT}{1,000 \ kt}\right)$$
 where,
$$MMT \ CO_2 \ Eq. = Million \ metric \ tons \ of \ CO_2 \ equivalent$$

$$kt = kilotons \ (equivalent \ to \ a \ thousand \ metric \ tons)$$

$$GWP = Global \ warming \ potential$$

$$MMT = Million \ metric \ tons$$

$$MMT = Million \ metric \ tons$$

GWP values allow policy makers to compare the impacts of emissions and reductions of different gases. According to the IPCC, GWP values typically have an uncertainty of \$\mathbb{Z}\$35 percent, though some GWP values have larger uncertainty than others, especially those in which lifetimes have not yet been ascertained. In the following decision, the parties to the United Nations Framework Convention on Climate Change (UNFCCC) have agreed to use consistent GWP values from the IPCC Fourth Assessment Report (AR4), based upon a 100 year time horizon, although other time horizon values are available (see Table A-252). While this Inventory uses agreed-upon GWP values according to the specific reporting requirements of the UNFCCC, described below, unweighted gas emissions and sinks in kilotons (kt) are provided in the Trends chapter of this report (Table 2-2) and users of the Inventory can apply different metrics and different time horizons to compare the impacts of different greenhouse gases.

...the global warming potential values used by Parties included in Annex I to the Convention (Annex I Parties) to calculate the carbon dioxide equivalence of anthropogenic emissions by sources and removals by sinks of greenhouse gases shall be those listed in the column entitled "Global warming potential for given time horizon" in table 2.14 of the errata to the contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change, based on the effects of greenhouse gases over a 100-year time horizon...¹³⁹

Greenhouse gases with relatively long atmospheric lifetimes (e.g., CO_2 , CH_4 , N_2O , HFCs, PFCs, SF₆, and NF₃) tend to be evenly distributed throughout the atmosphere, and consequently global average concentrations can be determined. However, short-lived gases such as water vapor, carbon monoxide, tropospheric ozone, other indirect greenhouse gases (e.g., NO_x and NMVOCs), and tropospheric aerosols (e.g., SO_2 products and black carbon) vary spatially, and consequently

¹³⁹ United Nations Framework Convention on Climate Change; http://unfccc.int/resource/docs/2013/cop19/eng/10a03.pdf; 31 January 2014; Report of the Conference of the Parties at its nineteenth session; held in Warsaw from 11 to 23 November 2013; Addendum; Part two: Action taken by the Conference of the Parties at its nineteenth session; Decision 24/CP.19; Revision of the UNFCCC reporting guidelines on annual inventories for Parties included in Annex I to the Convention; p. 2. (UNFCCC 2014).

it is difficult to quantify their global radiative forcing impacts. GWP values are generally not attributed to these gases that are short-lived and spatially inhomogeneous in the atmosphere.

3 Table A-252: IPCC AR4 Global Warming Potentials (GWP) and Atmospheric Lifetimes (Years) of Gases Used in this Report

Gas	Atmospheric Lifetime	100-year GWP ^a	20-year GWP	500-year GWP
Carbon dioxide (CO ₂)	See footnote ^b	1	1	1
Methane (CH ₄) ^c	12 ^d	25	72	7.6
Nitrous oxide (N ₂ O)	114 ^d	298	289	153
HFC-23	270	14,800	12,000	12,200
HFC-32	4.9	675	2,330	205
HFC-125	29	3,500	6,350	1,100
HFC-134a	14	1,430	3,830	435
HFC-143a	52	4,470	5,890	1,590
HFC-152a	1.4	124	437	38
HFC-227ea	34.2	3,220	5,310	1,040
HFC-236fa	240	9,810	8,100	7,660
HFC-43-10mee	15.9	1,640	4,140	500
CF ₄	50,000 ^d	7,390	5,210	11,200
C_2F_6	10,000	12,200	8,630	18,200
C ₃ F ₈	2,600	8,830	6,310	12,500
C ₄ F ₆ ^e	NA	0.003	NA	NA
c-C ₅ F ₈ ^e	NA	1.97	NA	NA
C_4F_{10}	2,600	8,860	6,330	12,500
c-C ₄ F ₈	3,200	10,300	7,310	14,700
C ₅ F ₁₂	4,100	9,160	6,510	13,300
C ₆ F ₁₄	3,200	9,300	6,600	13,300
CH₃F	3.7	150	490	45
CH ₂ FCF ₃	14	1,430	3,400	420
$C_2H_2F_4$	10.6	1,000	2,900	310
SF ₆	3,200	22,800	16,300	32,600
NF ₃	740	17,200	12,300	20,700

(NA) Not Available

Source: IPCC (2007)

Table A 252 proces

Table A-253 presents direct GWP values for ozone depleting substances (ODSs). Ozone depleting substances directly absorb infrared radiation and contribute to positive radiative forcing; however, their effect as ozone-depleters also leads to a negative radiative forcing because ozone itself is a potent greenhouse gas. There is considerable uncertainty regarding this indirect effect; direct GWP values are shown, but AR4 does provide a range of net GWP values for ozone depleting substances. The IPCC Guidelines and the UNFCCC do not include reporting instructions for estimating emissions of ODSs because their use is being phased out under the Montreal Protocol (see note below Table A-253). The effects of these compounds on radiative forcing are not addressed in this report.

Table A-253: 100-year Direct Global Warming Potentials for Select Ozone Depleting Substances

Gas	Direct GWP
CFC-11	4,750
CFC-12	10.900

^a GWP values used in this report are calculated over 100 year time horizon.

^b For a given amount of CO₂ emitted, some fraction of the atmospheric increase in concentration is quickly absorbed by the oceans and terrestrial vegetation, some fraction of the atmospheric increase will only slowly decrease over a number of years, and a small portion of the increase will remain for many centuries or more.

 $^{^{}c}$ The methane GWP includes the direct effects and those indirect effects due to the production of tropospheric ozone and stratospheric water vapor. The indirect effect due to the production of CO_2 is not included.

^d Methane and N₂O have chemical feedback systems that can alter the length of the atmospheric response, in these cases, global mean atmospheric lifetime (LT) is given first, followed by perturbation time (PT), but only the perturbation time is listed here and not the atmospheric residence time.

e See Table A-1 of 40 (CFR 98).

CFC-113	6,130
HCFC-22	1,810
HCFC-123	77
HCFC-124	609
HCFC-141b	725
HCFC-142b	2,310
CH₃CCI₃	146
CCI ₄	1,400
CH₃Br	5
Halon-1211	1,890
Halon-1301	7,140

Note: Because these compounds have been shown to deplete stratospheric ozone, they are typically referred to as ODSs. However, they are also potent greenhouse gases. Recognizing the harmful effects of these compounds on the ozone layer, in 1987 many governments signed the *Montreal Protocol on Substances that Deplete the Ozone Layer* to limit the production and importation of a number of CFCs and other halogenated compounds. The United States furthered its commitment to phase-out ODSs by signing and ratifying the Copenhagen Amendments to the Montreal Protocol in 1992. Under these amendments, the United States committed to ending the production and importation of halons by 1994, and CFCs by 1996.

Source: IPCC (2007).

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The IPCC published its Fifth Assessment Report (AR5) in 2013, providing the most current and comprehensive scientific assessment of climate change (IPCC 2013). Within this report, the GWP values were revised relative to the IPCC's Fourth Assessment Report (AR4) (IPCC 2007). Although the AR4 GWP values are used throughout this Inventory report in line with UNFCCC inventory reporting guidelines, it is informative to review the changes to the 100-year GWP values and the impact they have on the total GWP-weighted emissions of the United States. All GWP values use CO2 as a reference gas; a change in the radiative efficiency of CO₂ thus impacts the GWP of all other greenhouse gases. Since the Second Assessment Report (SAR) and Third Assessment Report (TAR), the IPCC has applied an improved calculation of CO₂ radiative forcing and an improved CO₂ response function. The GWP values are drawn from IPCC (2007), with updates for those cases where new laboratory or radiative transfer results have been published. Additionally, the atmospheric lifetimes of some gases have been recalculated, and updated background concentrations were used. Table A-254 shows how the GWP values of the other gases relative to CO2 tend to be larger in AR4 and AR5 because the revised radiative forcing of CO2 is lower than in earlier assessments, taking into account revisions in lifetimes. Comparisons of GWP values are based on the 100year time horizon required for UNFCCC inventory reporting. However, there were some instances in which other variables, such as the radiative efficiency or the chemical lifetime, were altered that resulted in further increases or decreases in particular GWP values in AR5. In addition, the values for radiative forcing and lifetimes have been calculated for a variety of halocarbons. Updates in some well-mixed HFC compounds (including HFC-23, HFC-32, HFC-134a, and HFC-227ea) for AR4 result from investigation into radiative efficiencies in these compounds, with some GWP values changing by up to 40 percent; with this change, the uncertainties associated with these well-mixed HFCs are thought to be approximately 12 percent.

It should be noted that the use of IPCC AR4 GWP values for the current Inventory applies across the entire time series of the Inventory (i.e., from 1990 to 2018). As such, GWP comparisons throughout this chapter are presented relative to AR4 GWPs.

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Table A-254: Comparison of GWP values and Lifetimes Used in the SAR, AR4, and AR5

	Lif	etime (years))		GWP (10	0 year)			Diffe	rence in GW	/P (Relative	e to AR4)	
							AR5 with					AR5 with	AR5 with
							feedbacks					feedbacks	feedbacks ^b
Gas	SAR	AR4	AR5	SAR	AR4	AR5 ^a	b	SAR	SAR (%)	AR5a	AR5 (%)	b	(%)
Carbon dioxide (CO ₂)	С	d	d	1	1	1	1	NC	NC	NC	NC	NC	NC
Methane (CH ₄)e	12±3	8.7/12 ^f	12.4	21	25	28	34	(4)	-16%	3	12%	9	36%
Nitrous oxide (N₂O)	120	120/114 ^f	121	310	298	265	298	12	4%	(33)	-11%	0	0%
Hydrofluorocarbons													
HFC-23	264	270	222	11,700	14,800	12,400	13,856	(3,100)	-21%	(2,400)	-16%	(944)	-6%
HFC-32	5.6	4.9	5.2	650	675	677	817	(25)	-4%	2	+%	142	21%
HFC-125	32.6	29	28.2	2,800	3,500	3,170	3,691	(700)	-20%	(330)	-9%	191	5%
HFC-134a	14.6	14	13.4	1,300	1,430	1,300	1,549	(130)	-9%	(130)	-9%	119	8%
HFC-143a	48.3	52	47.1	3,800	4,470	4,800	5,508	(670)	-15%	330	7%	1,038	23%
HFC-152a	1.5	1.4	1.5	140	124	138	167	16	13%	14	11%	43	35%
HFC-227ea	36.5	34.2	38.9	2,900	3,220	3,350	3,860	(320)	-10%	130	4%	640	20%
HFC-236fa	209	240	242	6,300	9,810	8,060	8,998	(3,510)	-36%	(1,750)	-18%	(812)	-8%
HFC-245fa	NA	7.6	7.7	NA	1,030	858	1,032	NA	NA	(172)	-17%	2	+%
HFC-365mfc	NA	6.6	8.7	NA	794	804	966	NA	NA	10	1%	172	22%
HFC-43-10mee	17.1	15.9	16.1	1,300	1,640	1,650	1,952	(340)	-21%	10	1%	312	19%
Fully Fluorinated													
Species													
SF ₆	3,200	3,200	3,200	23,900	22,800	23,500	26,087	1,100	5%	700	3%	3,287	14%
CF ₄	50,000	50,000	50,000	6,500	7,390	6,630	7,349	(890)	-12%	(760)	-10%	(41)	-1%
C_2F_6	10,000	10,000	10,000	9,200	12,200	11,100	12,340	(3,000)	-25%	(1,100)	-9%	140	1%
C ₃ F ₈	2,600	2,600	2,600	7,000	8,830	8,900	9,878	(1,830)	-21%	70	1%	1,048	12%
C ₄ F ₁₀	2,600	2,600	2,600	7,000	8,860	9,200	10,213	(1,860)	-21%	340	4%	1,353	15%
c-C ₄ F ₈	3,200	3,200	3,200	8,700	10,300	9,540	10,592	(1,600)	-16%	(760)	-7%	292	3%
C ₅ F ₁₂	4,100	4,100	4,100	7,500	9,160	8,550	9,484	(1,660)	-18%	(610)	-7%	324	4%
C ₆ F ₁₄	3,200	3,200	3,100	7,400	9,300	7,910	8,780	(1,900)	-20%	(1,390)	-15%	(520)	-6%
NF ₃	NA	740	500	NA	17,200	16,100	17,885	NA	NA	(1,100)	-6%	685	4%

^{2 +} Does not exceed 0.05 percent.

³ NC (No Change)

⁴ NA (Not Applicable)

⁵ a The GWP values presented here are the ones most consistent with the methodology used in the AR4 report.

⁶ b The GWP values presented here from the AR5 report include climate-carbon feedbacks for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime.

^{7 °} For a given amount of CO₂ emitted, some fraction of the atmospheric increase in concentration is quickly absorbed by the oceans and terrestrial vegetation, some fraction of the atmospheric

⁸ increase will only slowly decrease over a number of years, and a small portion of the increase will remain for many centuries or more.

 $^{9^{-}d}$ No single lifetime can be determined for CO_2 (see IPCC 2007).

1 e The methane GWP includes the direct effects and those indirect effects due to the production of tropospheric ozone and stratospheric water vapor. Additionally, the AR5 rep	orted separate
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- 2 values for fossil versus biogenic methane in order to account for the CO₂ oxidation product. 3 f Methane and N2O have chemical feedback systems that can alter the length of the atmospheric response, in these cases, global mean residence time is given first, followed by perturbation
- 5 Note: Parentheses indicate negative values. Source: IPCC (2013), IPCC (2007), IPCC (1996).

The choice of GWP values between the SAR, AR4, and AR5 with or without climate-carbon feedbacks has an impact on both the overall emissions estimated by the Inventory, as well as the trend in emissions over time. To summarize, Table A-255 shows the overall trend in U.S. greenhouse gas emissions, by gas, from 1990 through 2018 using the four GWP sets. The table also presents the impact of SAR and AR5 GWP values with or without feedbacks on the total emissions for 1990 and for 2018.

Table A-255: Effects on U.S. Greenhouse Gas Emissions Using SAR, AR4, and AR5 GWP values (MMT CO₂ Eq.)

	Differen	ce in Emiss	ions Betwe	en 1990								
Gas	and 2018 (Relative to 1990)				Revisio	ns to Annu	al Emissior	n Estimates	(Relative t	o AR4)		
					SAR	AR5 ^a	AR5 ^b	SAR	AR5a	AR5 ^b		
	SAR	AR4	AR5 ^a	AR5 ^b		1990			2018			
CO ₂	300.9	300.9	300.9	300.9	NC	NC	NC	NC	NC	NC		
CH ₄	(117.5)	(139.9)	(156.7)	(190.2)	(123.9)	92.9	278.8	(101.5)	76.2	228.5		
N_2O	(0.0)	(0.0)	(0.0)	(0.0)	17.5	(48.1)	NC	17.5	(48.1)	NC		
HFCs, PFCs, SF ₆ ,												
and NF ₃	68.1	79.7	78.4	96.8	(11.9)	(9.0)	1.2	(23.6)	(10.4)	18.3		
Total	251.5	240.7	222.6	207.5	(118.3)	35.8	280.1	(107.6)	17.7	246.8		
Percent Change	4.0%	3.7%	3.4%	3.1%	-1.8%	0.6%	4.4%	-1.6%	0.3%	3.7%		

NC (No Change)

Notes: Totals may not sum due to independent rounding. Excludes sinks. Parentheses indicate negative values.

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> When the GWP values from the SAR are applied to the emission estimates presented in this report, total emissions for the year 2018 are 6,570.2 MMT CO₂ Eq., as compared to the official emission estimate of 6,677.8 MMT CO₂ Eq. using AR4 GWP values (i.e., the use of SAR GWPs results in a 1.6 percent decrease relative to emissions estimated using AR4 GWPs). Table A-256 provides a detailed summary of U.S. greenhouse gas emissions and sinks for 1990 through 2018, using the GWP values from the SAR. The percent change in emissions for a given gas resulting from using different GWPs is equal to the percent change in the GWP; however, in cases where emissions of multiple gases are combined, as with HFCs or PFCs, the percent change will be a function of the relative quantity of the individual gases. Table A-257 summarizes the resulting change in emissions from using SAR GWP values relative to emissions using AR4 values for 1990 through 2018, including the percent change for 2018.

Table A-256: Recent Trends in U.S. Greenhouse Gas Emissions and Sinks using the SAR GWP values (MMT CO₂ Eq.)

Gas/Source	1990	2005	2014	2015	2016	2017	2018
CO ₂	5,128.3	6,131.9	5,562.9	5,413.7	5,293.5	5,256.0	5,429.2
Fossil Fuel Combustion	4,740.0	5,740.7	5,185.9	5,033.0	4,942.9	4,893.9	5,033.3
Transportation	1,469.1	1,856.1	1,713.7	1,725.3	1,765.3	1,787.4	1,798.2
Electric Power Sector	1,820.0	2,400.0	2,037.1	1,900.6	1,808.9	1,732.0	1,752.8
Industrial	857.0	850.1	813.6	802.0	801.7	806.0	846.7
Residential	338.2	357.9	347.1	318.1	293.2	294.2	335.9
Commercial	228.2	226.9	233.0	245.6	232.4	232.9	258.3
U.S. Territories	27.6	49.7	41.4	41.4	41.4	41.4	41.4
Non-Energy Use of Fuels	119.5	139.7	120.0	127.0	113.7	123.1	134.5
Iron and Steel Production &							
Metallurgical Coke Production	104.7	70.1	58.2	47.9	43.6	40.8	42.7
Cement Production	33.5	46.2	39.4	39.9	39.4	40.3	40.3
Petroleum Systems	9.6	12.2	30.5	32.6	23.0	24.5	39.4
Natural Gas Systems	32.2	25.3	29.6	29.3	29.9	30.4	34.9
Petrochemical Production	21.6	27.4	26.3	28.1	28.3	28.9	29.4
Lime Production	11.7	14.6	14.2	13.3	12.9	13.1	13.9
Ammonia Production	13.0	9.2	9.4	10.6	10.8	13.2	13.5
Incineration of Waste	8.0	12.5	10.4	10.8	10.9	11.1	11.1

^a The GWP values presented here are the ones most consistent with the methodology used in the AR4 report.

b The GWP values presented here from the AR5 report include climate-carbon feedbacks for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime. Additionally, the AR5 reported separate values for fossil versus biogenic methane in order to account for the CO₂ oxidation product.

Other Process Uses of Carbonates	6.3	7.6	13.0	12.2	11.0	10.1	9.4
Urea Fertilization	2.0	3.1	3.9	4.1	4.0	4.5	4.6
Carbon Dioxide Consumption	1.5	1.4	4.5	4.5	4.5	4.5	4.5
Urea Consumption for Non-							
Agricultural Purposes	3.8	3.7	1.8	4.6	5.1	3.8	3.6
Liming	4.7	4.3	3.6	3.7	3.1	3.1	3.1
Ferroalloy Production	2.2	1.4	1.9	2.0	1.8	2.0	2.1
Soda Ash Production	1.4	1.7	1.7	1.7	1.7	1.8	1.7
Titanium Dioxide Production	1.2	1.8	1.7	1.6	1.7	1.7	1.6
Aluminum Production	6.8	4.1	2.8	2.8	1.3	1.2	1.5
Glass Production	1.5	1.9	1.3	1.3	1.2	1.3	1.3
Zinc Production	0.6	1.0	1.0	0.9	0.9	1.0	1.0
Phosphoric Acid Production	1.5	1.3	1.0	1.0	1.0	1.0	0.9
Lead Production	0.5	0.6	0.5	0.5	0.4	0.5	0.6
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 ^a	219.4	230.7	323.2	317.7	317.2	322.2	328.9
International Bunker Fuels ^b	103.5	113.1	103.4	110.9	116.6	120.1	122.1
CH ₄ ^c	650.5	570.9	536.8	536.3	527.7	529.4	533.1
Enteric Fermentation	137.9	141.8	137.9	139.9	144.3	147.3	149.2
Natural Gas Systems	153.9	132.8	118.5	119.2	117.5	116.8	117.3
Landfills	150.8	110.3	94.6	93.5	90.8	90.5	92.9
Manure Management	31.2	43.3	45.6	48.6	50.1	50.3	51.8
Coal Mining	81.1	53.9	54.2	51.4	45.2	46.0	44.3
Petroleum Systems	38.8	32.6	36.5	34.1	32.7	32.6	30.7
Wastewater Treatment	12.9	13.0	12.0	12.2	12.1	11.9	11.9
Rice Cultivation	13.4	15.1	12.9	13.6	11.3	10.7	11.2
Stationary Combustion	7.2	6.6	7.5	7.1	6.7	6.6	7.3
Abandoned Oil and Gas Wells	5.5	5.8	6.0	6.0	6.1	5.9	5.9
Abandoned Underground Coal		_					
Mines	6.0	5.5	5.3	5.4	5.6	5.4	5.2
Mobile Combustion	10.9	8.0	3.5	3.1	2.9	2.8	2.6
Composting	0.3	1.6	1.8	1.8	1.9	2.1	2.1
Field Burning of Agricultural		_					
Residues	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Petrochemical Production	0.2	0.1	0.1	0.2	0.2	0.2	0.3
Ferroalloy Production	+	+	+	+	+	+	+
Carbide Production and		_					
Consumption	+	+	+	+	+	+	+
Iron and Steel Production &		_					
Metallurgical Coke Production	+	+	+	+	+	+	+
Incineration of Waste	+	+	+	+	+	+	+
International Bunker Fuels ^b	0.1	0.1	0.1	0.1	0.1	0.1	0.1
N₂O ^c	452.1	450.1	467.4	461.9	443.5	438.2	452.1
Agricultural Soil Management	328.6	325.7	363.3	362.1	343.1	340.6	351.8
Stationary Combustion	26.1	35.7	34.3	31.8	31.3	29.8	29.6
Manure Management	14.6	17.0	18.0	18.2	18.8	19.4	20.2
Mobile Combustion	43.7	38.8	20.5	19.1	18.1	16.9	15.8
Adipic Acid Production	15.8	7.4	5.7	4.4	7.3	7.7	10.7
Nitric Acid Production	12.6	11.8	11.4	12.0	10.5	9.7	9.7
Wastewater Treatment	3.5	4.6	5.0	5.0	5.1	5.2	5.2
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N₂O from Product Uses	4.4	4.4	4.4	4.4	4.4	4.4	4.4
Composting	0.4	1.7	1.9	2.0	2.1	2.3	2.3
Caprolactam, Glyoxal, and Glyoxylic							
Acid Production	1.7	2.2	2.1	2.1	2.1	1.5	1.5
Incineration of Waste	0.5	0.4	0.3	0.3	0.3	0.3	0.3
Electronics Industry	+	0.1	0.2	0.2	0.2	0.3	0.3
Field Burning of Agricultural							
Residues	0.2	0.2	0.2	0.2	0.2	0.2	0.2
Petroleum Systems	+	+	+	+	+	+	0.1
Natural Gas Systems	+	+	+	+	+	+	+
International Bunker Fuels ^b	0.9	1.0	1.0	1.0	1.1	1.1	1.1
HFCs	36.8	111.3	141.5	144.8	144.7	146.3	145.8
Substitution of Ozone Depleting							
Substances ^d	0.3	95.3	137.2	141.1	142.1	141.8	142.8
HCFC-22 Production	36.4	15.8	4.0	3.4	2.2	4.1	2.6
Electronics Industry	0.2	0.2	0.2	0.3	0.3	0.3	0.3
Magnesium Production and							
Processing	0.0	0.0	0.1	0.1	0.1	0.1	0.1
PFCs	20.6	5.6	4.7	4.2	3.6	3.3	3.9
Electronics Industry	2.2	2.6	2.5	2.5	2.4	2.4	2.5
Aluminum Production	18.4	3.0	2.1	1.7	1.1	0.9	1.3
Substitution of Ozone Depleting							
Substances ^d	0.0	+	+	+	+	+	+
SF ₆	30.2	12.4	6.8	5.7	6.3	6.2	6.2
Electrical Transmission and							
Distribution	24.3	8.8	5.0	3.9	4.3	4.3	4.3
Magnesium Production and							
Processing	5.4	2.9	1.0	1.0	1.2	1.1	1.2
Electronics Industry	0.5	0.7	0.8	0.8	0.9	0.8	0.8
NF ₃	NA						
Electronics Industry	NA						
Unspecified Mix of HFCs, NF ₃ , PFCs,							
and SF ₆	NA						
Electronics Industry	NA						
Total	6,318.7	7,282.0	6,720.0	6,566.7	6,419.4	6,379.4	6,570.2
LULUCF Emissions ^c	6.8	15.2	15.3	25.3	11.8	24.1	24.1
LULUCF CH₄ Emissions	3.7	7.4	8.0	13.6	6.1	12.8	12.8
LULUCF N₂O Emissions	3.1	7.8	7.3	11.7	5.7	11.3	11.3
LULUCF Carbon Stock Change ^e	(860.7)	(831.0)	(739.6)	(802.9)	(801.7)	(789.9)	(799.9)
LULUCF Sector Net Totalf	(854.0)	(815.8)	(724.3)	(777.7)	(789.9)	(765.9)	(775.7)
Net Emissions (Sources and Sinks)	5,464.8	6,466.2	5,995.8	5,789.0	5,629.5	5,613.5	5,794.5
Notes: Total emissions presented withou						•	

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

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⁴ Does not exceed 0.05 MMT CO₂ Eq.

⁴ NA (Not Applicable)

^{5 °} Emissions from Wood Biomass and Biofuel Consumption are not included specifically in summing energy sector totals. Net carbon fluxes 6 from changes in biogenic carbon reservoirs are accounted for in the estimates for LULUCF.

⁷ b Emissions from International Bunker Fuels are not included in totals.

CLULUCF emissions of CH₄ and N₂O are reported separately from gross emissions totals. LULUCF emissions include the CH₄ and N₂O
 emissions reported for *Peatlands Remaining Peatlands*, Forest Fires, Drained Organic Soils, 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.Manual Settlement Soils.Small amounts of Small a

^d Small amounts of PFC emissions also result from this source.

^e 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

 $to\ Grassland,\ Wetlands\ Remaining\ Wetlands,\ Land\ Converted\ to\ Wetlands,\ Settlements\ Remaining\ Settlements,\ and\ Land\ Converted\ to\ Settlements.$

 $^{\rm f}$ The LULUCF Sector Net Total is the net sum of all CH $_{\rm 4}$ and N $_{\rm 2}$ O emissions to the atmosphere plus net carbon stock changes.

Table A-257: Change in U.S. Greenhouse Gas Emissions Using SAR GWP values relative to AR4 GWP values (MMT CO₂ Eq.)

<u></u>								Percent
Gas/Source	1990	2005	2014	2015	2016	2017	2018	Change in 2018
CO ₂	NC	NC	NC	NC	NC	NC	NC	NA
CH ₄	(123.9)	(108.7)	(102.2)	(102.2)	(100.5)	(100.8)	(101.5)	(16%)
Enteric Fermentation	(26.3)	(27.0)	(26.3)	(26.6)	(27.5)	(28.1)	(28.4)	(16%)
Natural Gas Systems	(29.3)	(25.3)	(22.6)	(22.7)	(22.4)	(22.2)	(22.3)	(16%)
Landfills	(28.7)	(21.0)	(18.0)	(17.8)	(17.3)	(17.2)	(17.7)	(16%)
Manure Management	(5.9)	(8.2)	(8.7)	(9.3)	(9.5)	(9.6)	(9.9)	(16%)
Coal Mining	(15.4)	(10.3)	(10.3)	(9.8)	(8.6)	(8.8)	(8.4)	(16%)
Petroleum Systems	(7.4)	(6.2)	(7.0)	(6.5)	(6.2)	(6.2)	(5.9)	(16%)
Wastewater Treatment	(2.5)	(2.5)	(2.3)	(2.3)	(2.3)	(2.3)	(2.3)	(16%)
Rice Cultivation	(2.6)	(2.9)	(2.5)	(2.6)	(2.2)	(2.0)	(2.1)	(16%)
Stationary Combustion	(1.4)	(1.3)	(1.4)	(1.4)	(1.3)	(1.2)	(1.4)	(16%)
Abandoned Oil and Gas Wells	(1.1)	(1.1)	(1.1)	(1.1)	(1.2)	(1.1)	(1.1)	(16%)
Abandoned Underground Coal	` /	, ,	,	,	, ,	,	, ,	(/
Mines	(1.2)	(1.1)	(1.0)	(1.0)	(1.1)	(1.0)	(1.0)	(16%)
Mobile Combustion	(2.1)	(1.5)	(0.7)	(0.6)	(0.6)	(0.5)	(0.5)	(16%)
Composting	(0.1)	(0.3)	(0.3)	(0.3)	(0.4)	(0.4)	(0.4)	(16%)
Field Burning of Agricultural	(- /	(,	(/	(/	ζ- /	(-)	(- /	(/
Residues	(0.1)	(0.1)	(0.1)	(0.1)	(0.1)	(0.1)	(0.1)	(16%)
Petrochemical Production	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
Ferroalloy Production	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
Carbide Production and	(.,	(' /	(' /	(')	(·)	(·)	(.)	(10/0)
Consumption	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
Iron and Steel Production &	(' /	(' /	(' /	(-)	(' /	(' /	(-)	(20/0)
Metallurgical Coke Production	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
Incineration of Waste	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
International Bunker Fuels ^a	(+)	(+)	(+)	(+)	(+)	(+)	(+)	(16%)
N ₂ O	17.5	17.4	18.1	17.9	17.2	17.0	17.5	4%
Agricultural Soil Management	12.7	12.6	14.1	14.0	13.3	13.2	13.6	4%
Stationary Combustion	1.0	1.4	1.3	1.2	1.2	1.2	1.1	4%
Manure Management	0.6	0.7	0.7	0.7	0.7	0.8	0.8	4%
Mobile Combustion	1.7	1.5	0.8	0.7	0.7	0.7	0.6	4%
Nitric Acid Production	0.5	0.5	0.4	0.5	0.4	0.4	0.4	4%
Adipic Acid Production	0.6	0.3	0.2	0.2	0.3	0.3	0.4	4%
Wastewater Treatment	0.1	0.2	0.2	0.2	0.2	0.2	0.2	4%
N₂O from Product Uses	0.2	0.2	0.2	0.2	0.2	0.2	0.2	4%
Composting	+	0.1	0.1	0.1	0.1	0.1	0.1	4%
Caprolactam, Glyoxal, and	·	0.1	0.1	0.1	0.1	0.1	0.1	470
Glyoxylic Acid Production	0.1	0.1	0.1	0.1	0.1	0.1	0.1	4%
Incineration of Waste	+	+	+	+	+	+	+	4%
Electronics Industry	+		+	+	+	+	+	4%
Field Burning of Agricultural	'		'	'	'	'	'	470
Residues	+	+	+	+	+	+	+	4%
Petroleum Systems	+	+	+	+	+	+	+	4% 4%
Natural Gas Systems	+	+	+	+	+	+	+	4% 4%
International Bunker Fuels ^a	+	+ +	+	+	+	+	+	4% 4%
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HFCs	(9.7)	(15.4)	(20.9)	(21.5)	(21.7)	(22.4)	(22.4)	(13%)
Substitution of Ozone Depleting								
Substances ^b	+	(11.2)	(19.8)	(20.5)	(21.0)	(21.3)	(21.7)	(13%)
HCFC-22 Production	(9.7)	(4.2)	(1.1)	(0.9)	(0.6)	(1.1)	(0.7)	(21%)
Electronics Industry	(+)	(+)	(0.1)	(0.1)	(0.1)	(0.1)	(0.1)	(21%)
Magnesium Production and								
Processing	0.0	0.0	(+)	(+)	(+)	(+)	(+)	(9%)
PFCs	(3.6)	(1.1)	(0.9)	(0.9)	(0.7)	(0.7)	(0.8)	(16%)
Electronics Industry	(0.6)	(0.7)	(0.6)	(0.5)	(0.5)	(0.5)	(0.5)	(17%)
Aluminum Production	(3.0)	(0.5)	(0.4)	(0.3)	(0.2)	(0.2)	(0.2)	(15%)
Substitution of Ozone Depleting								
Substances	0.0	(+)	(+)	(+)	(+)	(+)	(+)	(12%)
SF ₆	1.4	0.6	0.3	0.3	0.3	0.3	0.3	5%
Electrical Transmission and								
Distribution	1.1	0.4	0.2	0.2	0.2	0.2	0.2	5%
Magnesium Production and								
Processing	0.3	0.1	+	+	0.1	0.1	0.1	5%
Electronics Industry	+	+	+	+	+	+	+	5%
NF ₃	NA	NA						
Electronics Industry	NA	NA						
Unspecified Mix of HFCs, NF ₃ , PFCs,								
and SF ₆	NA	NA						
Electronics Industry	NA	NA						
Total	(118.3)	(107.8)	(106.3)	(107.0)	(106.1)	(107.3)	(107.6)	(2%)

⁺ Absolute value does not exceed 0.05 MMT CO₂ Eq.

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Table A-258 below shows a comparison of total emissions estimates by sector using both the IPCC SAR and AR4 GWP values. For most sectors, the change in emissions that result from using SAR relative to AR4 GWP values was minimal. The effect on emissions from waste was by far the greatest (14.9 percent decrease in 2018 using SAR GWP values, relative to emissions using AR4 GWP values), due the predominance of CH_4 emissions in this sector. Emissions from all other sectors were comprised of mainly CO_2 or a mix of gases, which moderated the effect of the changes.

Table A-258: Comparison of Emissions by Sector using IPCC AR4 and SAR GWP Values (MMT CO₂ Eq.)

Sector	1990	2005	2014	2015	2016	2017	2018
Energy							
AR4 GWP, Used In							
Inventory	5,338.2	6,294.4	5,705.2	5,551.3	5,426.1	5,385.4	5,551.3
SAR GWP, Updated	5,283.1	6,250.6	5,663.3	5,510.2	5,386.8	5,346.0	5,512.4
Difference (%)	(1.0%)	(0.7%)	(0.7%)	(0.7%)	(0.7%)	(0.7%)	(0.7%)
Industrial Processes and							
Product Use							
AR4 GWP, Used In							
Inventory	345.6	364.8	376.9	373.1	367.3	367.7	373.6
SAR GWP, Updated	334.9	349.3	355.7	351.3	345.4	345.1	351.0
Difference (%)	(3.1%)	(4.3%)	(5.6%)	(5.9%)	(5.9%)	(6.1%)	(6.0%)
Agriculture							
AR4 GWP, Used In							
Inventory	554.4	575.9	608.6	614.6	600.5	602.3	618.5
SAR GWP, Updated	532.8	551.0	585.8	590.7	575.3	576.5	592.4
Difference (%)	(3.9%)	(4.3%)	(3.7%)	(3.9%)	(4.2%)	(4.3%)	(4.2%)

NC (No Change)

NA (Not Applicable)

⁴ a Emissions from International Bunker Fuels are not included in totals.

^b Small amounts of PFC emissions also result from this source.

Notes: Totals may not sum due to independent rounding. Parentheses indicate negative values.

LULUCF							
AR4 GWP, Used In							
Inventory	(853.4)	(814.7)	(723.0)	(775.5)	(788.9)	(763.9)	(773.7)
SAR GWP, Updated	(854.0)	(815.8)	(724.3)	(777.7)	(789.9)	(765.9)	(775.7)
Difference (%)	0.1%	0.1%	0.2%	0.3%	0.1%	0.3%	0.3%
Waste							
AR4 GWP, Used In							
Inventory	199.0	154.7	135.6	134.7	131.6	131.4	134.4
SAR GWP, Updated	167.9	131.1	115.3	114.4	111.9	111.8	114.4
Difference (%)	(15.6%)	(15.2%)	(15.0%)	(15.0%)	(14.9%)	(14.9%)	(14.9%)
Net Emissions							
AR4 GWP, Used In							
Inventory	5,583.7	6,575.1	6,103.3	5,898.2	5,736.6	5,722.9	5,904.1
SAR GWP, Updated	5,464.8	6,466.2	5,995.8	5,789.0	5,629.5	5,613.5	5,794.5
Difference (%)	(2.1%)	(1.7%)	(1.8%)	(1.9%)	(1.9%)	(1.9%)	(1.9%)

Notes: Totals may not sum due to independent rounding. Parentheses indicate negative values.

Further, Table A-259 and Table A-260 show the comparison of emission estimates using AR5 GWP values relative to AR4 GWP values without climate-carbon feedbacks for the non-CO₂ gases, on an emissions and percent change basis. Table A-261 and Table A-262 show the comparison of emission estimates using AR5 GWP values with climate-carbon feedbacks. The use of AR5 GWP values without climate-carbon feedbacks an increase in emissions of CH₄ and SF₆ relative to AR4 GWP values, but a decrease in emissions of other gases. The use of AR5 GWP values with climate-carbon feedbacks does not impact CO2 and N2O emissions; however, it results in an increase in emissions of CH4, SF6, and NF3 relative to AR4 GWP values, and has mixed impacts on emissions of other gases. Overall, these comparisons of AR4 and AR5 GWP values do not have a significant effect on calculated U.S. emissions, resulting in an increase in emissions of less than 1 percent using AR5 GWP values, or approximately 4 percent when using AR5 GWP values with climate-carbon feedbacks. As with the comparison of SAR and AR4 GWP values presented above, the percent change in emissions is equal to the percent change in the GWP for each gas; however, in cases where multiple gases are emitted in varying amounts the percent change is variable over the years, such as with Substitution of Ozone Depleting Substances.

Table A-259: Change in U.S. Greenhouse Gas Emissions Using AR5a without Climate-Carbon Feedbacks Relative to AR4 GWP Values (MMT CO₂ Eq.)

Gas	1990	2005	2014	2015	2016	2017	2018
CO ₂	NC						
CH ₄	92.9	81.6	76.7	76.6	75.4	75.6	76.2
N_2O	(48.1)	(47.9)	(49.8)	(49.2)	(47.2)	(46.6)	(48.1)
HFCs	(7.5)	(10.9)	(9.8)	(10.0)	(9.8)	(10.2)	(10.0)
PFCs	(2.4)	(0.6)	(0.5)	(0.5)	(0.4)	(0.4)	(0.4)
SF ₆	0.9	0.4	0.2	0.2	0.2	0.2	0.2
NF ₃	(+)	(+)	(+)	(+)	(+)	(+)	(+)
Unspecified Mix of HFCs, NF ₃ ,							
PFCs, and SF ₆	NA						
Total	35.8	22.4	16.7	17.1	18.1	18.5	17.7

⁺ Absolute value does not exceed 0.05 MMT CO₂ Eq.

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^a The GWP values presented here are the ones most consistent with the methodology used in the AR4 report. The AR5 report has also calculated GWP values (shown in Table A-261) where climate-carbon feedbacks have been included for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime. Additionally, the AR5 reported separate values for fossil versus

22 biogenic methane in order to account for the CO₂ oxidation product.

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

NC (No Change)

report/ar5/wg1/WG1AR5_Chapter08_FINAL.pdf>.

Table A-260: Change in U.S. Greenhouse Gas Emissions Using AR5a without Climate-Carbon Feedbacks Relative to AR4 **GWP Values (Percent)**

Gas/Source	1990	2005	2014	2015	2016	2017	2018
CO ₂	NC						
CH ₄	12.0%	12.0%	12.0%	12.0%	12.0%	12.0%	12.0%
N ₂ O	(11%)	(11%)	(11%)	(11%)	(11%)	(11%)	(11%)
SF ₆	3.1%	3.1%	3.1%	3.1%	3.1%	3.1%	3.1%
NF ₃	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)
HFCs	(16.1%)	(8.6%)	(6.0%)	(6.0%)	(5.9%)	(6.0%)	(6.0%)
Substitution of Ozone							
Depleting Substances	11.3%	(7.2%)	(5.7%)	(5.7%)	(5.7%)	(5.7%)	(5.7%)
HCFC-22 Production ^b	(16.2%)	(16.2%)	(16.2%)	(16.2%)	(16.2%)	(16.2%)	(16.2%)
Electronics Industry ^c	(16.2%)	(16.7%)	(16.8%)	(16.4%)	(16.7%)	(16.6%)	(16.2%)
Magnesium Production and							
Processing ^d	0.0%	0.0%	(9.1%)	(9.1%)	(9.1%)	(9.1%)	(9.1%)
PFCs	(10.0%)	(9.7%)	(9.5%)	(9.5%)	(9.5%)	(9.5%)	(9.6%)
Electronics Industry ^c	(9.4%)	(9.3%)	(9.2%)	(9.2%)	(9.3%)	(9.4%)	(9.4%)
Aluminum Production ^e	(10.1%)	(10.1%)	(10.0%)	(10.0%)	(9.9%)	(9.8%)	(9.9%)
Substitution of Ozone							
Depleting Substances ^{d,f}	0.0%	(10.3%)	(10.3%)	(10.3%)	(10.3%)	(10.3%)	(10.3%)
Unspecified Mix of HFCs, NF ₃ ,							
PFCs, and SF ₆	NA						
Electronics Industry	NA						
Total	0.6%	0.3%	0.2%	0.3%	0.3%	0.3%	0.3%

NC (No Change)

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Table A-261: Change in U.S. Greenhouse Gas Emissions Using AR5 with Climate-Carbon Feedbacks^a Relative to AR4 GWP Values (MMT CO₂ Eq.)

Gas	1990	2005	2014	2015	2016	2017	2018
CO ₂	NC						
CH ₄	278.8	244.7	230.1	229.9	226.2	226.9	228.5
N_2O	NC						
HFCs	(2.9)	9.2	16.8	17.3	17.6	17.3	17.5
PFCs	(+)	+	+	+	+	+	+
SF ₆	4.2	1.7	0.9	0.8	0.9	0.9	0.9
NF ₃	+	+	+	+	+	+	+
Unspecified Mix of HFCs,							
NF ₃ , PFCs, and SF ₆	NA						
Total	280.1	255.6	247.8	248.0	244.6	245.1	246.8

¹⁷ + Absolute value does not exceed 0.05 MMT CO₂ Eq.

^a The GWP values presented here are the ones most consistent with the methodology used in the AR4 report. The AR5 report has also calculated GWP values (shown in Table A-262) where climate-carbon feedbacks have been included for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime. Additionally, the AR5 reported separate values for fossil versus biogenic methane in order to account for the CO₂ oxidation product.

^b HFC-23 emitted.

^c Emissions from HFC-23, CF₄, C₂F₆, C₃F₈, C₄F₈, SF₆, as well as other HFCs and PFCs used as heat transfer fluids.

^d Zero change in beginning of time series since emissions were zero.

¹⁰ 11 e PFC emissions from CF₄ and C₂F₆.

¹² ^f PFC emissions from CF₄.

Note: Total emissions presented without LULUCF. Parentheses indicate negative values.

¹⁸ NC (No Change)

^a The GWP values presented here from the AR5 report include climate-carbon feedbacks for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime. Additionally, the AR5 reported separate values for fossil versus biogenic methane in order to account for the CO_2 oxidation product.

Table A-262: Change in U.S. Greenhouse Gas Emissions Using AR5 with Climate-Carbon Feedbacks^a Relative to AR4 GWP Values (Percent)

Gas/Source	1990	2005	2014	2015	2016	2017	2018
CO ₂	NC						
CH ₄	36.0%	36.0%	36.0%	36.0%	36.0%	36.0%	36.0%
N ₂ O	NC						
SF ₆	14.4%	14.4%	14.4%	14.4%	14.4%	14.4%	14.4%
NF ₃	4.0%	4.0%	4.0%	4.0%	4.0%	4.0%	4.0%
HFCs	(6.2%)	7.3%	10.3%	10.4%	10.6%	10.3%	10.4%
Substitution of Ozone							
Depleting Substances	34.6%	9.9%	10.9%	10.9%	10.9%	10.8%	10.8%
HCFC-22 Production ^b	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)	(6.4%)
Electronics Industry ^c	(6.4%)	(6.9%)	(7.1%)	(6.6%)	(6.9%)	(6.8%)	(6.3%)
Magnesium Production							
and Processing ^d	0.0%	0.0%	8.3%	8.3%	8.3%	8.3%	8.3%
PFCs	(0.2%)	0.2%	0.4%	0.4%	0.5%	0.4%	0.3%
Electronics Industry ^c	0.6%	0.8%	0.8%	0.8%	0.7%	0.6%	0.5%
Aluminum Productione	(0.3%)	(0.3%)	(0.1%)	(0.1%)	0.0%	0.0%	(0.1%)
Substitution of Ozone							
Depleting Substances ^{d,f}	0.0%	(0.6%)	(0.6%)	(0.6%)	(0.6%)	(0.6%)	(0.6%)
Unspecified Mix of HFCs,							
NF ₃ , PFCs, and SF ₆	NA						
Electronics Industry	NA						
Total	4.4%	3.5%	3.6%	3.7%	3.8%	3.8%	3.7%

^{6 +} Does not exceed 0.05 percent.

NC (No Change)

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^a The GWP values presented here from the AR5 report include climate-carbon feedbacks for the non- CO_2 gases in order to be consistent with the approach used in calculating the CO_2 lifetime. Additionally, the AR5 reported separate values for fossil versus biogenic methane in order to account for the CO_2 oxidation product.

¹¹ b HFC-23 emitted.

^{12 °}Emissions from HFC-23, CF₄, C₂F₆, C₃F₈, C₄F₈, SF₆, as well as other HFCs and PFCs used as heat transfer fluids.

^d Zero change in beginning of time series since emissions were zero.

^e PFC emissions from CF₄ and C₂F₆.

¹⁵ f PFC emissions from CF₄.

Notes: Total emissions presented without LULUCF. Parentheses indicate negative values. Excludes Sinks.

Ozone is present in both the stratosphere,¹⁴¹ where it shields the earth from harmful levels of ultraviolet radiation, and at lower concentrations in the troposphere,¹⁴² where it is the main component of anthropogenic photochemical "smog." Chlorofluorocarbons (CFCs), halons, carbon tetrachloride, methyl chloroform, and hydrochlorofluorocarbons (HCFCs), along with certain other chlorine and bromine containing compounds, have been found to deplete the ozone levels in the stratosphere. These compounds are commonly referred to as ozone depleting substances (ODSs). If left unchecked, stratospheric ozone depletion could result in a dangerous increase of ultraviolet radiation reaching the earth's surface. In 1987, nations around the world signed the Montreal Protocol on Substances that Deplete the Ozone Layer. This landmark agreement created an international framework for limiting, and ultimately eliminating, the production of most ozone depleting substances. ODSs have historically been used in a variety of industrial applications, including refrigeration and air conditioning, foam blowing, fire extinguishing, sterilization, solvent cleaning, and as an aerosol propellant.

In the United States, the Clean Air Act Amendments of 1990 provide the legal instrument for implementation of the Montreal Protocol controls. The Clean Air Act classifies ozone depleting substances as either Class I or Class II, depending upon the ozone depletion potential (ODP) of the compound. The production of CFCs, halons, carbon tetrachloride, and methyl chloroform—all Class I substances—has already ended in the United States. However, large amounts of these chemicals remain in existing equipment, and stockpiles of the ODSs, as well as material recovered from equipment being decommissioned, are used for maintaining the existing equipment. As a result, emissions of Class I compounds will continue, albeit generally in decreasing amounts, for many more years. Class II designated substances, all of which are HCFCs, have been, or are being, phased out at later dates than Class I compounds because they have lower ODPs. These compounds served, and in some cases continue to serve, as interim replacements for Class I compounds in many industrial applications. The use and emissions of HCFCs in the United States is anticipated to continue for several decades as equipment that use Class II substances are retired from use. Under current Montreal Protocol controls, however, the production for domestic use of all HCFCs in the United States must end by the year 2030.

In addition to contributing to ozone depletion, CFCs, halons, carbon tetrachloride, methyl chloroform, and HCFCs are also potent greenhouse gases. However, the depletion of the ozone layer has a cooling effect on the climate that counteracts the direct warming from tropospheric emissions of ODSs. Stratospheric ozone influences the earth's radiative balance by absorption and emission of longwave radiation from the troposphere as well as absorption of shortwave radiation from the sun; overall, stratospheric ozone has a warming effect.

The IPCC has prepared both direct GWP values and net (combined direct warming and indirect cooling) GWP ranges for some of the most common ozone depleting substances (IPCC 2007). See Annex 6.1 Global Warming Potential Values, for a listing of the direct GWP values for ODS.

Although the IPCC emission inventory guidelines do not require the reporting of emissions of ozone depleting substances, the United States believes that the inventory presents a more complete picture of climate impacts when we include these compounds. Emission estimates for several ozone depleting substances are provided in Table A-263.

Table A-263: Emissions of Ozone Depleting Substances (kt)

Compound	1990	2005	2014	2015	2016	2017	2018
Class I							
CFC-11	29	12	24	25	25	25	20

¹⁴¹ The stratosphere is the layer from the top of the troposphere up to about 50 kilometers. Approximately 90 percent of atmospheric ozone is within the stratosphere. The greatest concentration of ozone occurs in the middle of the stratosphere, in a region commonly called the ozone layer.

¹⁴² The troposphere is the layer from the ground up to about 11 kilometers near the poles and 16 kilometers in equatorial regions (i.e., the lowest layer of the atmosphere, where humans live). It contains roughly 80 percent of the mass of all gases in the atmosphere and is the site for weather processes including most of the water vapor and clouds.

¹⁴³ Substances with an ozone depletion potential of 0.2 or greater are designated as Class I. All other designated substances that deplete stratospheric ozone but which have an ODP of less than 0.2 are Class II.

¹⁴⁴ Older refrigeration and air-conditioning equipment, fire extinguishing systems, and foam products blown with CFCs/HCFCs may still contain Class I ODS.

Methodology and Data Sources

Emissions of ozone depleting substances were estimated using the EPA's Vintaging Model. The model, named for its method of tracking the emissions of annual "vintages" of new equipment that enter into service, is a "bottom-up" model. It models the consumption of chemicals based on estimates of the quantity of equipment or products sold, serviced, and retired each year, and the amount of the chemical required to manufacture and/or maintain the equipment. The Vintaging Model makes use of this market information to build an inventory of the in-use stocks of the equipment in each of the end-uses. Emissions are estimated by applying annual leak rates, service emission rates, and disposal emission rates to each population of equipment. By aggregating the emission and consumption output from the different end-uses, the model produces estimates of total annual use and emissions of each chemical. Please see Annex 3.9, Methodology for Estimating HFC and PFC Emissions from Substitution of Ozone Depleting Substances, of this Inventory for a more detailed discussion of the Vintaging Model.

Uncertainties

Uncertainties exist with regard to the levels of chemical production, equipment sales, equipment characteristics, and end-use emissions profiles that are used by these models. Please see the Substitution of Ozone Depleting Substances section of this report for a more detailed description of the uncertainties that exist in the Vintaging Model.

⁺ Does not exceed 0.5 kt.

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23 24 25

Sulfur dioxide (SO₂), emitted into the atmosphere through natural and anthropogenic processes, affects the Earth's radiative budget through photochemical transformation into sulfate aerosols that can (1) scatter sunlight back to space, thereby reducing the radiation reaching the Earth's surface; (2) affect cloud formation; and (3) affect atmospheric chemical composition (e.g., stratospheric ozone, by providing surfaces for heterogeneous chemical reactions). The overall effect of SO₂-derived aerosols on radiative forcing is believed to be negative (IPCC 2007). However, because SO₂ is shortlived and unevenly distributed through the atmosphere, its radiative forcing impacts are highly uncertain. Sulfur dioxide emissions have been provided below in Table A-264.

The major source of SO₂ emissions in the United States is the burning of sulfur containing fuels, mainly coal. Metal smelting and other industrial processes also release significant quantities of SO₂. The largest contributor to U.S. emissions of SO₂ is electricity generation, accounting for 47.8 percent of total SO₂ emissions in 2018 (see Table A-265); coal combustion accounted for approximately 92.0 percent of that total. The second largest source was industrial fuel combustion, which produced 20.0 percent of 2018 SO₂ emissions (see Table A-264). Overall, SO₂ emissions in the United States decreased by 88.1 percent from 1990 to 2018. The majority of this decline came from reductions from electricity generation, primarily due to increased consumption of low sulfur coal from surface mines in western states.

Sulfur dioxide is important for reasons other than its effect on radiative forcing. It is a major contributor to the formation of urban smog and acid rain. As a contributor to urban smog, high concentrations of SO₂ can cause significant increases in acute and chronic respiratory diseases. In addition, once SO2 is emitted, it is chemically transformed in the atmosphere and returns to earth as the primary contributor to acid deposition, or acid rain. Acid rain has been found to accelerate the decay of building materials and paints, cause the acidification of lakes and streams, and damage trees. As a result of these harmful effects, the United States has regulated the emissions of SO₂ under the Clean Air Act. The EPA has also developed a strategy to control these emissions via four programs: (1) the National Ambient Air Quality Standards program, ¹⁴⁵ (2) New Source Performance Standards, ¹⁴⁶ (3) the New Source Review/Prevention of Significant Deterioration Program, ¹⁴⁷ and (4) the Sulfur Dioxide Allowance Program. ¹⁴⁸

Table A-264: SO₂ Emissions (kt)

Sector/Source	1990	2005	2014	2015	2016	2017	2018
Energy	19,628	12,364	3,742	2,844	2,187	2,050	1,983
Stationary Sources	18,407	11,541	3,532	2,635	1,978	1,841	1,774
Oil and Gas Activities	390	180	94	94	94	94	94
Mobile Sources	793	619	88	87	87	87	87
Waste Combustion	38	25	27	27	27	27	27
Industrial Processes and							
Product Use	1,307	831	497	497	497	497	497
Other Industrial Processes	362	327	151	151	151	151	151
Miscellaneous ^a	11	114	136	136	136	136	136
Chemical and Allied Product							
Manufacturing	269	228	112	112	112	112	112
Metals Processing	659	158	95	95	95	95	95
Storage and Transport	6	2	3	3	3	3	3
Solvent Use	0	+	+	+	+	+	+
Degreasing	0	0	0	0	0	0	0
Graphic Arts	0	0	0	0	0	0	0
Dry Cleaning	NA	0	0	0	0	0	0
Surface Coating	0	0	0	0	0	0	0
Other Industrial	0	+	+	+	+	+	+

¹⁴⁵ [42 U.S.C § 7409, CAA § 109]

¹⁴⁶ [42 U.S.C § 7411, CAA § 111]

¹⁴⁷ [42 U.S.C § 7473, CAA § 163]

¹⁴⁸ [42 U.S.C § 7651, CAA § 401]

Nonindustrial	NA	N	А	NA	NA	NA	NA	NA
Agriculture	NA	N	Α	NA	NA	NA	NA	NA
Agricultural Burning	NA	N	Α	NA	NA	NA	NA	NA
Waste	+		1	1	1	1	1	1
Landfills	+		1	1	1	1	1	1
Wastewater Treatment	+		0	0	0	0	0	0
Miscellaneous ^a	+		0	0	0	0	0	0
Total	20,935	13,19	6	4,240	3,342	2,685	2,548	2,481

⁺ Does not exceed 0.5 kt

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Table A-265: SO₂ Emissions from Electricity Generation (kt)

Fuel Type	1990	2005	2014	2015	2016	2017	2018
Coal	13,808	8,680	2,706	1,881	1,277	1,151	1,090
Oil	580	458	143	99	67	61	57
Gas	1	174	54	38	26	23	22
Internal Combustion	45	57	18	12	8	8	7
Other	NA	71	22	15	10	9	9
Total	14,433	9,439	2,943	2,046	1,388	1,252	1,185

NA (Not Applicable)

NA (Not Applicable)

^a Miscellaneous includes other combustion and fugitive dust categories.

² 3 4 5 6 Note: Totals may not sum due to independent rounding.

Source: Data taken from EPA (2019) and disaggregated based on EPA (2003).

Note: Totals may not sum due to independent rounding.

¹⁰ Source: Data taken from EPA (2019) and disaggregated based on EPA (2003).

6.4. Complete List of Source Categories

Chapter/Source	Gas(es)
Energy	
Fossil Fuel Combustion	CO ₂
Non-Energy Use of Fossil Fuels	CO ₂
Stationary Combustion (excluding CO ₂)	CH ₄ , N ₂ O, CO, NO _x , NMVOC
Mobile Combustion (excluding CO ₂)	CH ₄ , N ₂ O, CO, NO _x , NMVOC
Coal Mining	CH ₄
Abandoned Underground Coal Mines	CH ₄
Petroleum Systems	CH ₄ , N₂O
Natural Gas Systems	CH ₄ , N ₂ O
Abandoned Oil and Gas Wells	CO ₂ , CH ₄
Incineration of Waste	CO ₂ , CH ₄ , N ₂ O, NO _x , CO, NMVOC
Industrial Processes and Product Use	-, ,, - , ., ,
Cement Production	CO ₂
Lime Production	CO ₂
Glass Production	CO ₂
Other Process Uses of Carbonates	CO ₂
Ammonia Production	CO ₂
Urea Consumption for Non-Agricultural Purposes	CO ₂
Nitric Acid Production	N ₂ O
Adipic Acid Production	N₂O
Caprolactam, Glyoxal, and Glyoxylic Production	N ₂ O
Carbide Production and Consumption	CO ₂ , CH ₄
Titanium Dioxide Production	CO ₂ , C114 CO ₂
Soda Ash Production	CO ₂
Petrochemical Production	CO ₂ , CH ₄
HCFC-22 Production	HFC-23
Carbon Dioxide Consumption	CO ₂
Phosphoric Acid Production	CO ₂
Iron and Steel Production & Metallurgical Coke Production	CO ₂ , CH ₄
Ferroalloy Production	CO ₂ , CH ₄
Aluminum Production	CO ₂ , CF ₄ , C ₂ F ₆
Magnesium Production and Processing	CO ₂ , HFCs, SF ₆
Lead Production	CO ₂
Zinc Production	CO ₂
Electronics Industry	N ₂ O, HFCs, PFCs, ^a SF ₆ , NF ₃
Substitution of Ozone Depleting Substances	HFCs, PFCs ^b
Electrical Transmission and Distributing	SF ₆
N₂O from Product Uses	N_2O
Agriculture	
Enteric Fermentation	CH ₄
Manure Management	CH ₄ , N ₂ O
Rice Cultivation	CH ₄
Liming	CO ₂
Urea Fertilization	CO ₂
Field Burning of Agricultural Residues	CH ₄ , N ₂ O, NO _x , CO
Agricultural Soil Management	N_2O
Land Use, Land-Use Change, and Forestry ^c	
Forest Land Remaining Forest Land	CO ₂ , CH ₄ , N ₂ O, NO _x , CO
Land Converted to Forest Land	CO ₂
Cropland Remaining Cropland	CO ₂

Land Converted to Cropland CO₂

Grassland Remaining Grassland CO₂, CH₄, N₂O, NO_x, CO

Land Converted to Grassland CO₂

Wetlands Remaining Wetlands CO_2 , CH_4 , N_2O Land Converted to Wetlands CO_2 , CH_4 Settlements Remaining Settlements CO_2 , N_2O Land Converted to Settlements CO_2

Waste

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 $\begin{array}{lll} \text{Landfills} & \text{CH}_4,\,\text{NO}_x,\,\text{CO},\,\text{NMVOC} \\ \text{Wastewater Treatment} & \text{CH}_4,\,\text{N}_2\text{O},\,\text{NO}_x,\,\text{CO},\,\text{NMVOC} \\ \end{array}$

Composting CH₄, N₂O

^a Includes HFC-23, CF₄, C₂F₆, as well as a mix other HFCs and PFCs used as heat transfer fluids.

b Includes HFC-23, HFC-32, HFC-125, HFC-134a, HFC-143a, HFC-236fa, CF₄, HFC-152a, HFC-227ea, HFC-245fa, HFC-4310mee, and PFC/PFPEs.

 $[^]c$ The LULUCF Sector includes CH $_4$ and N $_2$ O emissions to the atmosphere and net carbon stock changes. The term "flux" is used to describe the net emissions of greenhouse gases accounting for both the emissions of CO $_2$ to and the removals of CO $_2$ from the atmosphere. Removal of CO $_2$ from the atmosphere is also referred to as "carbon sequestration."

Constants, Units, and Conversions 6.5.

Metric Prefixes

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Although most activity data for the United States is gathered in customary U.S. units, these units are converted into metric units per international reporting guidelines. Table A-266 provides a guide for determining the magnitude of metric units.

Table A-266: Guide to Metric Unit Prefixes

Prefix/Symbol	Factor
atto (a)	10 ⁻¹⁸
femto (f)	10 ⁻¹⁵
pico (p)	10-12
nano (n)	10 -9
micro (μ)	10-6
milli (m)	10-3
centi (c)	10-2
deci (d)	10 ⁻¹
deca (da)	10
hecto (h)	10 ²
kilo (k)	10 ³
mega (M)	10 ⁶
giga (G)	10 ⁹
tera (T)	1012
peta (P)	10 ¹⁵
exa (E)	10 ¹⁸

Unit Conversions

1 acre

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1 square mile =

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 9
      1 kilogram
                       2.205 pounds
      1 pound
                       0.454 kilograms
      1 short ton
                   = 2,000 pounds
                                            0.9072 metric tons
      1 metric ton =
                       1,000 kilograms =
                                            1.1023 short tons
10
      1 cubic meter = 35.315 cubic feet
      1 cubic foot = 0.02832 cubic meters
      1 U.S. gallon = 3.785412 liters
      1 barrel (bbl) = 0.159 cubic meters
      1 barrel (bbl) = 42 U.S. gallons
                   = 0.001 cubic meters
      1 liter
11
      1 foot
                   = 0.3048 meters
      1 meter
                   = 3.28 feet
      1 mile
                   = 1.609 kilometers
      1 kilometer
                   = 0.621 miles
12
```

Degrees Celsius = (Degrees Fahrenheit - 32)*5/9 Degrees Kelvin = Degrees Celsius + 273.15

2.589988 square kilometers

= 43,560 square feet = 0.4047 hectares = 4,047 square meters

Density Conversions¹⁴⁹

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Methane	1 cubic meter	=	0.67606 kilograms		
Carbon dioxide	1 cubic meter	=	1.85387 kilograms		
Natural gas liquids	1 metric ton	=	11.6 barrels	=	1,844.2 liters
Unfinished oils	1 metric ton	=	7.46 barrels	=	1,186.04 liters
Alcohol	1 metric ton	=	7.94 barrels	=	1,262.36 liters
Liquefied petroleum gas	1 metric ton	=	11.6 barrels	=	1,844.2 liters
Aviation gasoline	1 metric ton	=	8.9 barrels	=	1,415.0 liters
Naphtha jet fuel	1 metric ton	=	8.27 barrels	=	1,314.82 liters
Kerosene jet fuel	1 metric ton	=	7.93 barrels	=	1,260.72 liters
Motor gasoline	1 metric ton	=	8.53 barrels	=	1,356.16 liters
Kerosene	1 metric ton	=	7.73 barrels	=	1,228.97 liters
Naphtha	1 metric ton	=	8.22 barrels	=	1,306.87 liters
Distillate	1 metric ton	=	7.46 barrels	=	1,186.04 liters
Residual oil	1 metric ton	=	6.66 barrels	=	1,058.85 liters
Lubricants	1 metric ton	=	7.06 barrels	=	1,122.45 liters

1 metric ton =

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Energy Conversions

Petrochemical feedstocks

Miscellaneous products

Bitumen

Petroleum coke

Special naphtha

Waxes

Converting Various Energy Units to Joules

The common energy unit used in international reports of greenhouse gas emissions is the joule. A joule is the energy required to push with a force of one Newton for one meter. A terajoule (TJ) is one trillion (1012) joules. A British thermal unit (Btu, the customary U.S. energy unit) is the quantity of heat required to raise the temperature of one pound of water one degree Fahrenheit at or near 39.2 degrees Fahrenheit.

6.06 barrels =

7.87 barrels =

7.46 barrels

8.53 barrels

5.51 barrels =

8.00 barrels =

=

963.46 liters

876.02 liters

1,251.23 liters

1,186.04 liters

1,356.16 liters

1,271.90 liters

2.3881011 calories 23.88 metric tons of crude oil equivalent 1 TJ = 947.8 million Btus 277,800 kilowatt-hours

Converting Various Physical Units to Energy Units

Data on the production and consumption of fuels are first gathered in physical units. These units must be converted to their energy equivalents. The conversion factors in Table A-267 can be used as default factors, if local data are not available. See Appendix A of EIA's Monthly Energy Review, November 2019 (EIA 2019) for more detailed information on the energy content of various fuels.

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¹⁴⁹ Reference: EIA (2007)

1 Table A-267: Conversion Factors to Energy Units (Heat Equivalents)

Fuel Type (Units)	Factor
Solid Fuels (Million Btu/Short ton)	
Anthracite coal	22.57
Bituminous coal	23.89
Sub-bituminous coal	17.14
Lignite	12.87
Coke	21.49
Natural Gas (Btu/Cubic foot)	1,036
Liquid Fuels (Million Btu/Barrel)	
Motor gasoline	5.054
Aviation gasoline	5.048
Kerosene	5.670
Jet fuel, kerosene-type	5.670
Distillate fuel	5.825
Residual oil	6.287
Naphtha for petrochemicals	5.248
Petroleum coke	6.024
Other oil for petrochemicals	5.825
Special naphthas	5.248
Lubricants	6.065
Waxes	5.537
Asphalt	6.636
Still gas	6.000
Misc. products	5.796

Note: For petroleum and natural gas, *Monthly Energy Review, November 2019* (EIA 2019). For coal ranks, *State Energy Data Report 1992* (EIA 1993). All values are given in higher heating values (gross calorific values).

A-509

1 6.6. Abbreviations

ABS Acrylonitrile butadiene styrene

AC Air conditioner

ACC American Chemistry Council

AEDT FAA Aviation Environmental Design Tool

AEO Annual Energy Outlook
AER All-electric range

AF&PA American Forest and Paper Association

AFEAS Alternative Fluorocarbon Environmental Acceptability Study

AFOLU Agriculture, Forestry, and Other Land Use

AFV Alternative fuel vehicle
AGA American Gas Association

AGR Acid gas removal

AHEF Atmospheric and Health Effect Framework

AHRI Air-Conditioning, Heating, and Refrigeration Institute

AISI American Iron and Steel Institute

ALU Agriculture and Land Use
ANGA American Natural Gas Alliance
ANL Argonne National Laboratory
APC American Plastics Council
API American Petroleum Institute

APTA American Public Transportation Association

AR4 IPCC Fourth Assessment Report
AR5 IPCC Fifth Assessment Report
ARI Advanced Resources International
ARMA Autoregressive moving-average

ARMS Agricultural Resource Management Surveys
ASAE American Society of Agricultural Engineers

ASLRRA American Short-line and Regional Railroad Association

ASR Annual Statistical Report

ASTM American Society for Testing and Materials

AZR American Zinc Recycling

BCEF Biomass conversion and expansion factors

BEA Bureau of Economic Analysis, U.S. Department of Commerce

BIER Beverage Industry Environmental Roundtable

BLM Bureau of Land Management

BoC Bureau of Census

BOD Biological oxygen demand

BOD5 Biochemical oxygen demand over a 5-day period

BOEM Bureau of Ocean Energy Management

BOEMRE Bureau of Ocean Energy Management, Regulation and Enforcement

BOF Basic oxygen furnace
BRS Biennial Reporting System

BTS Bureau of Transportation Statistics, U.S. Department of Transportation

Btu British thermal unit

C Carbon

C&D Construction and demolition waste
C&EN Chemical and Engineering News
CAAA Clean Air Act Amendments of 1990

CaO Calcium oxide

CAPP Canadian Association of Petroleum Producers

CARB California Air Resources Board
CBI Confidential business information

C-CAP Coastal Change Analysis Program

CDAT Chemical Data Access Tool

CEAP USDA-NRCS Conservation Effects Assessment Program

CEFM Cattle Enteric Fermentation Model
CEMS Continuous emission monitoring system

CFC Chlorofluorocarbon

CFR Code of Federal Regulations
CGA Compressed Gas Association

CH₄ Methane

CHP Combined heat and power

CI Confidence interval

CIGRE International Council on Large Electric Systems

CKD Cement kiln dust
CLE Crown Light Exposure

CMA Chemical Manufacturer's Association

CMM Coal mine methane

CMOP Coalbed Methane Outreach Program

CMR Chemical Market Reporter
CNG Compressed natural gas
CO Carbon monoxide
CO₂ Carbon dioxide

COD Chemical oxygen demand

COGCC Colorado Oil and Gas Conservation Commission

CONUS Continental United States
CRF Common Reporting Format
CRM Component ratio method
CRP Conservation Reserve Program

CSRA Carbon Sequestration Rural Appraisals
CTIC Conservation Technology Information Center

CVD Chemical vapor deposition
CWNS Clean Watershed Needs Survey

d.b.h Diameter breast height DE Digestible energy

DESC Defense Energy Support Center-DoD's Defense Logistics Agency

DFAMS Defense Fuels Automated Management System
DGGS Division of Geological & Geophysical Surveys

DHS Department of Homeland Security
DLA DoD's Defense Logistics Agency

DM Dry matter

DOC Degradable organic carbon
DOC U.S. Department of Commerce
DOD U.S. Department of Defense
DOE U.S. Department of Energy
DOI U.S. Department of the Interior

DOM Dead organic matter

DOT U.S. Department of Transportation
DRE Destruction or removal efficiencies

DRI Direct Reduced Iron EAF Electric arc furnace

EDB Aircraft Engine Emissions Databank
EDF Environmental Defense Fund
EER Energy economy ratio

EF Emission factor

EFMA European Fertilizer Manufacturers Association

EJ Exajoule

EGR Exhaust gas recirculation EGU Electric generating unit

EIA Energy Information Administration, U.S. Department of Energy

EIIP Emissions Inventory Improvement Program

EOR Enhanced oil recovery

EPA U.S. Environmental Protection Agency

EREF Environment Research & Education Foundation

ERS Economic Research Service

ETMS Enhanced Traffic Management System

EV Electric vehicle

EVI Enhanced Vegetation Index
FAA Federal Aviation Administration
FAO Food and Agricultural Organization

FAOSTAT Food and Agricultural Organization database

FAS Fuels Automated System

FCCC Framework Convention on Climate Change

FEB Fiber Economics Bureau

FERC Federal Energy Regulatory Commission

FGD Flue gas desulfurization

FHWA Federal Highway Administration FIA Forest Inventory and Analysis

FIADB Forest Inventory and Analysis Database
FIPR Florida Institute of Phosphate Research

FOD First order decay

FQSV First-quarter of silicon volume

FSA Farm Service Agency
FTP Federal Test Procedure

g Gram

G&B Gathering and boosting
GaAs Gallium arsenide
GCV Gross calorific value
GDP Gross domestic product

GHG Greenhouse gas

GHGRP EPA's Greenhouse Gas Reporting Program

GIS Geographic Information Systems

GJ Gigajoule

GOADS Gulf Offshore Activity Data System

GPG Good Practice Guidance
GRI Gas Research Institute
GSAM Gas Systems Analysis Model
GTI Gas Technology Institute
GWP Global warming potential

ha Hectare

HBFC Hydrobromofluorocarbon

HC Hydrocarbon

HCFC Hydrochlorofluorocarbon Hydrochlor of luoro ole fin**HCFO HDDV** Heavy duty diesel vehicle **HDGV** Heavy duty gas vehicle **HDPE** High density polyethylene HF Hydraulically fractured HFC Hydrofluorocarbon HFO Hydrofluoroolefin **HFE** Hydrofluoroethers

HHV Higher Heating Value HMA Hot Mix Asphalt

HMIWI Hospital/medical/infectious waste incinerator

HTF Heat Transfer Fluid

HTS Harmonized Tariff Schedule HWP Harvested wood product IBF International bunker fuels

IC Integrated Circuit

ICAO International Civil Aviation Organization ICBA International Carbon Black Association

ICE Internal combustion engine
ICR Information Collection Request
IEA International Energy Agency
IFO Intermediate Fuel Oil

IGES Institute of Global Environmental Strategies

IISRP International Institute of Synthetic Rubber Products
ILENR Illinois Department of Energy and Natural Resources

IMO International Maritime Organization

IPAA Independent Petroleum Association of America
IPCC Intergovernmental Panel on Climate Change

IPPU Industrial Processes and Product Use ITC U.S. International Trade Commission

ITRS International Technology Roadmap for Semiconductors

JWR Jim Walters Resources KCA Key category analysis

kg Kilogram kt Kiloton kWh Kilowatt hour

LDPE Low density polyethylene

LDT Light-duty truck
LDV Light-duty vehicle
LEV Low emission vehicles

LFG Landfill gas

LFGTE Landfill gas-to-energy
LHV Lower Heating Value

LKD Lime kiln dust

LLDPE Linear low density polyethylene

LMOP EPA's Landfill Methane Outreach Program

LNG Liquefied natural gas

LPG Liquefied petroleum gas(es)

LTO Landing and take-off

LULUCF Land Use, Land-Use Change, and Forestry

M&R Metering and regulating

MARPOL International Convention for the Prevention of Pollution from Ships

MC Motorcycle

MCF Methane conversion factor
MCL Maximum Contaminant Levels
MCFD Thousand cubic feet per day
MDI Metered dose inhalers

MDP Management and design practices

MECS EIA Manufacturer's Energy Consumption Survey

MEMS Micro-electromechanical systems

MER Monthly Energy Review

MGO Marine gas oil

MgO Magnesium oxide

MJ Megajoule

MLRA Major Land Resource Area

mm Millimeter

MMBtu Million British thermal units

MMCF Million cubic feet

MMCFD Million cubic feet per day
MMS Minerals Management Service

MMT Million metric tons

MMTCE Million metric tons carbon equivalent

MMT CO₂ Eq. Million metric tons carbon dioxide equivalent

MODIS Moderate Resolution Imaging Spectroradiometer

MoU Memorandum of Understanding

MOVES U.S. EPA's Motor Vehicle Emission Simulator model

MPG Miles per gallon

MRLC Multi-Resolution Land Characteristics Consortium

MRV Monitoring, reporting, and verification MSHA Mine Safety and Health Administration

MSW Municipal solid waste

MT Metric ton

MTBE Methyl Tertiary Butyl Ether
MTBS Monitoring Trends in Burn Severity
MVAC Motor vehicle air conditioning

MY Model year N_2O Nitrous oxide

NA Not applicable; Not available

NACWA National Association of Clean Water Agencies

NAHMS National Animal Health Monitoring System

NAICS North American Industry Classification System

NAPAP National Acid Precipitation and Assessment Program

NARR North American Regional Reanalysis Product

NAS National Academies of Sciences, Engineering, and Medicine

NASA National Aeronautics and Space Administration
NASF National Association of State Foresters
NASS USDA's National Agriculture Statistics Service

NC No change

NCASI National Council of Air and Stream Improvement

NCV Net calorific value
NE Not estimated

NEI National Emissions Inventory

NEMA National Electrical Manufacturers Association

NEMS National Energy Modeling System

NESHAP National Emission Standards for Hazardous Air Pollutants

NEU Non-Energy Use

NEV Neighborhood Electric Vehicle

NF₃ Nitrogen trifluorideNFI National forest inventoryNGL Natural gas liquids

NIR National Inventory Report
NLA National Lime Association
NLCD National Land Cover Dataset
NMOC Non-methane organic compounds
NMVOC Non-methane volatile organic compound

NMOG Non-methane organic gas

NO Nitric oxide

NO Not occurring
NO₂ Nitrogen dioxide
NO_x Nitrogen oxides

NOAA National Oceanic and Atmospheric Administration

NOF Not on feed

NPDES National Pollutant Discharge Elimination System

NPP Net primary productivity

NPRA National Petroleum and Refiners Association

NRC National Research Council

NRCS Natural Resources Conservation Service
NREL National Renewable Energy Laboratory

NRI National Resources Inventory

NSCEP National Service Center for Environmental Publications

NSCR Non-selective catalytic reduction
NSPS New source performance standards

NWS National Weather Service
OAG Official Airline Guide

OAP EPA Office of Atmospheric Programs

OAQPS EPA Office of Air Quality Planning and Standards

ODP Ozone depleting potential
ODS Ozone depleting substances

OECD Organization of Economic Co-operation and Development

OEM Original equipment manufacturers

OGJ Oil & Gas Journal
OH Hydroxyl radical

OMS EPA Office of Mobile Sources
ORNL Oak Ridge National Laboratory

OSHA Occupational Safety and Health Administration

OTA Office of Technology Assessment

OTAQ EPA Office of Transportation and Air Quality

OVS Offset verification statement
PAH Polycyclic aromatic hydrocarbons
PCA Portland Cement Association
PCC Precipitate calcium carbonate
PDF Probability Density Function

PECVD Plasma enhanced chemical vapor deposition

PET Polyethylene terephthalate
PET Potential evapotranspiration
PEVM PFC Emissions Vintage Model

PFC Perfluorocarbon
PFPE Perfluoropolyether
PHEV Plug-in hybrid vehicles

PHMSA Pipeline and Hazardous Materials Safety Administration

PI Productivity index
PLS Pregnant liquor solution

POTW Publicly Owned Treatment Works ppbv Parts per billion (109) by volume

ppm Parts per million

ppmv Parts per million (10⁶) by volume pptv Parts per trillion (10¹²) by volume PRCI Pipeline Research Council International

PRP Pasture/Range/Paddock

PS Polystyrene

PSU Primary Sample Unit

PU Polyurethane
PVC Polyvinyl chloride
PV Photovoltaic

QA/QC Quality Assurance and Quality Control

QBtu Quadrillion Btu

R&D Research and Development
RECs Reduced Emissions Completions

RCRA Resource Conservation and Recovery Act

RFS Renewable Fuel Standard

RMA Rubber Manufacturers' Association

RPA Resources Planning Act
RTO Regression-through-the-origin
SAE Society of Automotive Engineers

SAGE System for assessing Aviation's Global Emissions

SAN Styrene Acrylonitrile

SAR IPCC Second Assessment Report SCR Selective catalytic reduction

SCSE South central and southeastern coastal

SDR Steel dust recycling

SEC Securities and Exchange Commission

SEMI Semiconductor Equipment and Materials Industry

SF₆ Sulfur hexafluoride SiC Silicon carbide

SICAS Semiconductor International Capacity Statistics SNAP Significant New Alternative Policy Program

SNG Synthetic natural gas SO₂Sulfur dioxide SOC Soil Organic Carbon SOG State of Garbage survey SOHIO Standard Oil Company of Ohio **SSURGO** Soil Survey Geographic Database STMC Scrap Tire Management Council **SULEV** Super Ultra Low Emissions Vehicle **SWANA** Solid Waste Association of North America

SWDS Solid waste disposal sites

TA Treated anaerobically (wastewater)

TAM Typical animal mass
TAME Tertiary amyl methyl ether
TAR IPCC Third Assessment Report

TBtu Trillion Btu

TDN Total digestible nutrients
TEDB Transportation Energy Data Book

TFI The Fertilizer Institute

TIGER Topologically Integrated Geographic Encoding and Referencing survey

TJ Terajoule

TLEV Traditional low emissions vehicle
TMLA Total Manufactured Layer Area

TRI Toxic Release Inventory

TSDF Hazardous waste treatment, storage, and disposal facility

TTB Tax and Trade Bureau

TVA Tennessee Valley Authority

UAN Urea ammonium nitrate

UDI Utility Data Institute

UFORE U.S. Forest Service's Urban Forest Effects model

UG Underground (coal mining)

U.S. **United States** U.S. ITC United States International Trade Commission UEP **United Egg Producers** ULEV Ultra low emission vehicle UNEP United Nations Environmental Programme UNFCCC United Nations Framework Convention on Climate Change USAA U.S. Aluminum Association USAF **United States Air Force USDA** United States Department of Agriculture USFS **United States Forest Service United States Geological Survey** USGS USITC U.S. International Trade Commission EPA's Voluntary Aluminum Industrial Partnership VAIP VAM Ventilation air methane VKT Vehicle kilometers traveled VMT Vehicle miles traveled VOCs Volatile organic compounds ٧S Volatile solids WBJ Waste Business Journal WERF Water Environment Research Federation World Fab Forecast (previously WFW, World Fab Watch) WFF WGC World Gas Conference WIP Waste in place WMO World Meteorological Organization WMS Waste management systems WTE Waste-to-energy WW Wastewater **WWTP** Wastewater treatment plant ZEVs Zero emissions vehicles

2 6.7. Chemical Formulas

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3 Table A-268: Guide to Chemical Formulas

Symbol	Name
Al	Aluminum
Al_2O_3	Aluminum oxide
Br	Bromine
С	Carbon
CH ₄	Methane
C_2H_6	Ethane
C ₃ H ₈	Propane
CF ₄	Perfluoromethane
C_2F_6	Perfluoroethane, hexafluoroethane
c-C ₃ F ₆	Perfluorocyclopropane
C ₃ F ₈	Perfluoropropane
c-C ₄ F ₈	Perfluorocyclobutane
C_4F_{10}	Perfluorobutane
C_5F_{12}	Perfluoropentane
C_6F_{14}	Perfluorohexane
CF ₃ I	Trifluoroiodomethane
CFCI ₃	Trichlorofluoromethane (CFC-11)
CF ₂ Cl ₂	Dichlorodifluoromethane (CFC-12)
CF ₃ Cl	Chlorotrifluoromethane (CFC-13)

A-517

C₂F₃Cl₃ Trichlorotrifluoroethane (CFC-113)*

CCl₃CF₃ CFC-113a*

 $C_2F_4Cl_2$ Dichlorotetrafluoroethane (CFC-114) C_2F_5Cl Chloropentafluoroethane (CFC-115)

CHCl₂F HCFC-21

CHF₂Cl Chlorodifluoromethane (HCFC-22)

 $\begin{array}{lll} C_2F_3HCl_2 & HCFC-123 \\ C_2F_4HCl & HCFC-124 \\ C_2FH_3Cl_2 & HCFC-141b \\ C_2H_3F_2Cl & HCFC-142b \\ CF_3CF_2CHCl_2 & HCFC-225ca \\ CClF_2CF_2CHClF & HCFC-225cb \\ \end{array}$

CCl₄ Carbon tetrachloride CHClCCl₂ Trichloroethylene

CCl₂CCl₂ Perchloroethylene, tetrachloroethene

 CH_3Cl Methylchloride CH_3CCl_3 Methylchloroform CH_2Cl_2 Methylenechloride

CHCl₃ Chloroform, trichloromethane

CHF₃ HFC-23 CH_2F_2 HFC-32 HFC-41 CH₃F C_2HF_5 HFC-125 $C_2H_2F_4$ HFC-134 CH₂FCF₃ HFC-134a $C_2H_3F_3$ HFC-143* $C_2H_3F_3$ HFC-143a* CH₂FCH₂F HFC-152* $C_2H_4F_2$ HFC-152a* CH₃CH₂F HFC-161 C₃HF₇ HFC-227ea CF₃CF₂CH₂F HFC-236cb $\mathsf{CF_3}\mathsf{CHFCHF_2}$ HFC-236ea HFC-236fa $C_3H_2F_6$ HFC-245ca $C_3H_3F_5$ CHF2CH2CF3 HFC-245fa CF₃CH₂CF₂CH₃ HFC-365mfc $C_5H_2F_{10}\\$ HFC-43-10mee CF₃OCHF₂ HFE-125 CF₂HOCF₂H HFE-134 CH₃OCF₃ HFE-143a CF₃CHFOCF₃ HFE-227ea CF₃CHClOCHF₂ HCFE-235da2 CF₃CHFOCHF₂ HFE-236ea2 CF₃CH₂OCF₃ HFE-236fa CF₃CF₂OCH₃ HFE-245cb2 CHF₂CH₂OCF₃ HFE-245fa1 CF₃CH₂OCHF₂ HFE-245fa2 CHF₂CF₂OCH₃ HFE-254cb2 HFE-263fb2 CF₃CH₂OCH₃ CF₃CF₂OCF₂CHF₂ HFE-329mcc2 CF₃CF₂OCH₂CF₃ HFE-338mcf2 CF₃CF₂CF₂OCH₃ HFE-347mcc3 HFE-347mcf2 CF₃CF₂OCH₂CHF₂

CF₃CHFCF₂OCH₃

HFE-356mec3

CHF₂CF₂CF₂OCH₃ HFE-356pcc3 CHF2CF2OCH2CHF2 HFE-356pcf2 HFE-356pcf3 CHF2CF2CH2OCHF2 CF₃CF₂CH₂OCH₃ HFE-365mcf3 CHF2CF2OCH2CH3 HFE-374pcf2 $C_4F_9OCH_3$ HFE-7100 $C_4F_9OC_2H_5$ HFE-7200 CH₂CFCF₃ HFO-1234yf CHFCHCF₃ HFO-1234ze(E) CF₃CHCHCF₃ HFO-1336mzz(Z) HCFO-1233zd(E) $C_3H_2CIF_3$ $\mathsf{CHF}_2\mathsf{OCF}_2\mathsf{OC}_2\mathsf{F}_4\mathsf{OCHF}_2$ H-Galden 1040x

 $\begin{array}{ll} \mathsf{CHF_2OCF_2OCHF_2} & \mathsf{HG-10} \\ \mathsf{CHF_2OCF_2CF_2OCHF_2} & \mathsf{HG-01} \end{array}$

CH₃OCH₃ Dimethyl ether
CH₂Br₂ Dibromomethane
CH₂BrCl Dibromochloromethane
CHBr₃ Tribromomethane
CHBrF₂ Bromodifluoromethane
CH₃Br Methylbromide

CF₂BrCl Bromodichloromethane (Halon 1211) CF₃Br(CBrF₃) Bromotrifluoromethane (Halon 1301)

CF₃I FIC-13I1

CO Carbon monoxide CO₂ Carbon dioxide

CaCO₃ Calcium carbonate, Limestone

CaMg(CO₃)₂ Dolomite

CaO Calcium oxide, Lime
Cl atomic Chlorine
F Fluorine
Fe Iron

 $\begin{array}{lll} \text{Fe} & & \text{Iron} \\ \text{Fe}_2\text{O}_3 & & \text{Ferric oxide} \\ \text{FeSi} & & \text{Ferrosilicon} \\ \text{GaAs} & & \text{Gallium arsenide} \end{array}$

H, H₂ atomic Hydrogen, molecular Hydrogen

H₂O Water

H₂O₂ Hydrogen peroxide

OH Hydroxyl

N, N₂ atomic Nitrogen, molecular Nitrogen

 ${
m NH_3}$ Ammonia ${
m NH_{4^+}}$ Ammonium ion ${
m HNO_3}$ Nitric acid

 $\begin{array}{ccc} \text{MgO} & \text{Magnesium oxide} \\ \text{NF}_3 & \text{Nitrogen trifluoride} \\ \text{N}_2\text{O} & \text{Nitrous oxide} \\ \text{NO} & \text{Nitric oxide} \\ \text{NO}_2 & \text{Nitrogen dioxide} \\ \text{NO}_3 & \text{Nitrate radical} \\ \text{NO}_x & \text{Nitrogen oxides} \\ \end{array}$

Na Sodium

Na₂CO₃ Sodium carbonate, soda ash

Na₃AlF₆ Synthetic cryolite

O, O₂ atomic Oxygen, molecular Oxygen

O₃ Ozone

S	atomic Sulfur
H ₂ SO ₄	Sulfuric acid
SF ₆	Sulfur hexafluoride
SF ₅ CF ₃	Trifluoromethylsulphur pentafluoride
SO ₂	Sulfur dioxide
Si	Silicon
SiC	Silicon carbide
SiO ₂	Quartz

^{*} Distinct isomers.

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References

- 4 EIA (2019) *Monthly Energy Review, November 2019*. Energy Information Administration, U.S. Department of Energy, Washington, DC. DOE/EIA-0035(2019/11). November 2019.
 - EIA (2007) Emissions of Greenhouse Gases in the United States 2006, Draft Report. Office of Integrated Analysis and Forecasting, Energy Information Administration, U.S. Department of Energy, Washington, DC. DOE-EIA-0573 (2006).
- 8 EIA (1993) *State Energy Data Report 1992*, DOE/EIA-0214(93), Energy Information Administration, U.S. Department of Energy. Washington, DC. December.
- EPA (2019) "1970-2018 Average annual emissions, all criteria pollutants in MS Excel." National Emissions Inventory (NEI)
 Air Pollutant Emissions Trends Data. Office of Air Quality Planning and Standards. Last Modified March 2018. Available online at: https://www.epa.gov/air-emissions-inventories/air-pollutant-emissions-trends-data.
 - EPA (2003) E-mail correspondence. Air pollutant data. Office of Air Pollution to the Office of Air Quality Planning and Standards, U.S. Environmental Protection Agency (EPA). December 22, 2003.
- IPCC (2013) Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment
 Report of the Intergovernmental Panel on Climate Change. [Stocker, T.F., D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen,
 J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (eds.)]. Cambridge University Press, Cambridge, United
 Kingdom and New York, NY, USA, 1535 pp.
- IPCC (2007) Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment
 Report of the Intergovernmental Panel on Climate Change. S. Solomon, D. Qin, M. Manning, Z. Chen, M. Marquis, K.B.
 Averyt, M. Tignor and H.L. Miller (eds.). Cambridge University Press. Cambridge, United Kingdom 996 pp.
- IPCC (1996) Climate Change 1995: The Science of Climate Change. Intergovernmental Panel on Climate Change,
 J.T.Houghton, L.G. Meira Filho, B.A. Callander, N. Harris, A. Kattenberg, and K. Maskell (eds.). Cambridge University
 Press. Cambridge, United Kingdom