

Wildland fire emission factors in North America: synthesis of existing data, measurement needs and management applications

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Abstract. Field and laboratory emission factors (EFs) of wildland fire emissions for 276 known air pollutants sampled across Canada and the US were compiled. An online database, the Smoke Emissions Repository Application (SERA), was created to enable analysis and summaries of existing EFs to be used in smoke management and emissions inventories. We evaluated how EFs of select pollutants (CO, CO₂, CH₄, NO_x, total particulate matter (PM), PM_{2.5} and SO₂) are influenced by combustion phase, burn type and fuel type. Of the 12 533 records in the database, over a third ($n = 5637$) are represented by 23 air pollutants, most designated as US Environmental Protection Agency criteria air pollutants, greenhouse gases, hazardous air pollutants or known air toxins. Among all pollutants in the database, including the most common pollutants PM, CO, CO₂ and CH₄, records are unevenly distributed with a bias towards flaming combustion, prescribed burning and laboratory measurements. Across all EFs, records are most common for south-eastern and western conifer forests and western shrubland types. Based on identified data gaps, we offer recommendations for future studies, including targeting underrepresented air pollutants, smouldering combustion phases and improved source characterisation of wildland fire emissions.

Additional keywords: air quality, greenhouse gas emissions, particulate matter, prescribed fire emissions, smoke management, wildfire emissions.

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Introduction

Over the past decade, field and laboratory studies have made substantial progress in characterising the chemical composition of wildland fire smoke and quantifying emission factors (EFs) in North America. EFs are defined as the mass of a gas or aerosol pollutant per dry weight unit mass of consumed fuel (Urbanski 2014). Since the seminal review of Andreae and Merlet (2001), several reviews have summarised existing EFs by vegetation type or region (Akagi *et al.* 2011; Yokelson *et al.* 2013; Urbanski 2014), and studies were compiled into a summary dataset using unpublished and published data from the 1960s to 2011 (Lincoln *et al.* 2014). These developments, including several more recent studies, represent a large advancement in emissions characterisation and justify the creation of a master repository of EFs that can be used to produce consistent values applicable for wildland

fire management, smoke modelling and emission inventories in North America.

The Smoke Emissions Repository Application (SERA) presented here is a searchable online database of existing EFs of 276 known air pollutants in standardised units (g kg^{-1}) (<https://depts.washington.edu/nwfire/sera>, accessed 26 November 2019). EF records include modified combustion efficiency (MCE, Yokelson *et al.* 1996), fuel type, study type (laboratory or field), measurement platform (ground, tower or air-based), geographic location and source reference. The database was created to enable analysis and summaries of existing EFs, and creation of average EFs to be used in decision support tools for smoke management, including Consume (Prichard 2007), FOFEM (Reinhardt *et al.* 1997) and the BlueSky Smoke Modelling Framework (Larkin *et al.* 2009), as well as future updates to national emissions

inventories (e.g. US EPA 2017). Prior to updates provided by Urbanski (2014), models that predict wildland fire emissions such as Consume, FOFEM and BlueSky used EFs published within the 2001 Smoke Management Guide (Hardy *et al.* 2001) and supported a relatively small list of pollutants, including particulate matter (PM), carbon monoxide (CO), carbon dioxide (CO₂), methane (CH₄), non-methane organic compounds (NMOCs) and sulfur dioxide (SO₂). SERA was designed to provide a synthesis of existing EFs and summary values for emissions modelling that can facilitate more rapid incorporation of EFs than was previously available through a relational database maintained by the US Forest Service Pacific Wildland Fire Sciences Laboratory (Seattle, WA).

For many air pollutants, selecting the most appropriate EFs to represent emissions from biomass burning needs to consider the combustion of fuels that burn in smouldering combustion and low-heat pyrolysis phases (e.g. coarse wood and organic soils). Modified combustion efficiency ($MCE = \Delta CO_2 / (\Delta CO_2 + \Delta CO)$) is a measure of the relative fraction of flaming and smouldering combustion, and has been demonstrated to be highly correlated with EFs for pollutant emissions (such as CO, CH₄ and NMOCs) that are more concentrated in smouldering combustion phases (Urbanski 2014). Emissions studies often include a measure of MCE to characterise source combustion phase (Yokelson *et al.* 1996; Urbanski 2014). For example, Urbanski (2013) found that Rocky Mountain wildfire emissions from prescribed fires in moderate to heavy coarse wood had a much lower MCE than that observed in studies of prescribed fires in the south-east, from where the majority of the EFs reported in the literature for NMOCs and PM_{2.5} have been derived. More recently, Sekimoto *et al.* (2018) examined relationships between high- and low-temperature pyrolysis and found that these two variables explained most of the variability in non-methane organic gas emissions and outperformed MCE. The relationship between wildland fire emissions and combustion highlights the need to develop algorithms for extrapolating EFs to broadly representative fuel types and burning conditions (e.g. flaming *v.* smouldering and prescribed underburn *v.* crown fire). Because many smoke management issues associated with wildland fire emissions are from residual long-term smouldering sources, characterising the fuels and EFs associated with these burning conditions is critical for fire management applications (Urbanski 2013; Yokelson *et al.* 2013; Hu *et al.* 2018; Peterson *et al.* 2018). Some studies also suggest that factors driving burning conditions including wind, fuel moisture and fire behaviour may more strongly influence pollutant emissions than differences in fuel type or region (Yokelson *et al.* 2013; Koss *et al.* 2018).

EFs are generally presented as best estimates, based on compiled datasets of either fire-average values (i.e. representing pollutant emissions from a combination of concurrent flaming and smouldering fuel consumption) or combustion phase (e.g. flaming, short-term smouldering and residual long-term smouldering). Flaming combustion is defined as high-temperature combustion in the presence of flames. Short-term smouldering combustion, hereafter referred to as smouldering combustion, occurs at lower temperatures without flames and is often associated with incomplete fuel combustion. Long-term residual smouldering is

generally associated with marginally combustible fuels such as coarse wood (i.e. stumps and downed logs >7.6 cm in diameter) and duff that generally have incomplete combustion and smoulder for extended periods (Ottmar 2014).

Smoke management guides have traditionally summarised EFs by major vegetation or fuel type. For example, the 2001 Smoke Management Guide (Hardy *et al.* 2001) compiled pollutant EFs from existing studies of prescribed burns, including EFs summarised from broadcast burned slash of Douglas fir, juniper and ponderosa pine forests (Ward *et al.* 1989), broadcast burned brush (Hardy *et al.* 1996) and wildfires (Hardy *et al.* 1992). Hardy *et al.* (2001) also provided EFs by flaming, short-term smouldering and long-term residual smouldering phases of combustion. Summarising EFs by combustion phase is generally useful for fire managers to inform smoke reduction techniques, such as finding optimal burn windows that maximise flaming combustion and minimise long-term smouldering (MACTEC Federal Programs 2004), which can contribute to greater overall pollutant emissions of PM and CO and nighttime intrusions into communities (Miller *et al.* 2019).

Since the original smoke management guide was published in 2001, a large number of prescribed burn studies and a relatively small number of wildfire emissions studies have been conducted. Urbanski (2014) synthesised fire-average EFs for major pollutant species by burn type (i.e. prescribed burns and wildfires) and region and/or vegetation type. Fire-average flaming and smouldering EFs and their estimated uncertainty were summarised based on six prescribed fire groupings in the US, including south-eastern (SE) conifer, south-western (SW) conifer and north-western (NW) conifer forests, western shrublands and western grasslands, and two wildfire categories including W conifer wildfires and boreal wildfires. Additional residual smouldering EFs were included for coarse wood, temperate forest organic soils and boreal forest organic soils (referred to hereafter as duff). More recently, sampled plumes from western US wildfires suggest that the quantity and composition of fuels burned in summer (May–October) wildfires may substantially increase estimates of pollutant emissions, including greenhouse gases (e.g. CO₂ and CH₄) and particulate matter (Liu *et al.* 2017).

The objectives of this study were to develop a centralised repository for evaluating and synthesising emissions factor datasets for major fuel types in North America and to provide updates to best-estimate EFs for major pollutant species. Because wildland fire emissions research is a rapidly developing field, many additional studies have been conducted since the Wildland Fire Emissions Factor Database was released in 2014 (Lincoln *et al.* 2014) and the publication of best-estimate EFs by Urbanski (2014). In the present study, we introduce SERA, an online repository of wildland fire EFs and source references, and provide revised and summarised EFs that can be used for smoke management planning for wildland fire events. The web application provides the most complete compilation of wildland EFs to date and a searchable, online database. Using the large sample size for some pollutants, we evaluated how EFs are influenced by combustion phase, burn type and fuel type. We offer recommendations for future studies including targeting under-represented air pollutants and the relative influence of fuel type, measurement platform and combustion phase on estimated pollutant emissions.

Table 1. Fields and definitions of the wildland fire emissions database (Smoke Emissions Repository Application; SERA)

Variable name	Definition	Input options or units
EPA_CAP	Critical air pollutant	True or False
EPA_HAP	Hazardous air pollutant	True or False
EPA_GHG	Greenhouse gas	True or False
EPA_TOX	Toxic air pollutant	True or False
Pollutant ID	Unique identification for each record	e.g. 275 = pyrene
Formula or acronym	Pollutant chemical formula or acronym	e.g. CO, CO ₂ , PM, PM _{2.5}
Pollutant name	Common name of pollutant	
Pollutant name 2	Secondary common name of pollutant	
Other synonyms	Synonyms (other common names)	
Fuel category	Major fuel category (e.g. shrub, herb, coarse wood, duff)	Defined in Table 3
Regional vegetation type	Major vegetation by region	e.g. SE pine, W conifer
Regional vegetation subtype	Subtype of major vegetation/fuel by region	e.g. SE pine – longleaf sandhill, W conifer – Douglas fir crown
Burn type	Type of burn including measurement platform. WF = wildfire, Rx = prescribed burn	Laboratory, WF-Ground WF-Tower, WF-Airborne, Rx-Ground, Rx-Tower, Rx-Airborne
EF	Emissions factor	g kg ⁻¹
MCE	Modified combustion efficiency	Ratio (0–1)
Combustion phase	Assigned phase based on MCE or CE or field observation. F = flaming, S = smouldering, RS = residual smouldering	F: MCE >0.89, S: MCE 0.8 – 0.89, RS: MCE <0.8
Ref ID	Unique identification for each source reference	e.g. 238 = Aurell et al. (2015)
WFEFD database ID	Cross reference ID for each reference within the Wildland Fire Emissions Factor Database (Lincoln et al. 2014)	
Lead author	Lead author of source reference	
Publication year	Publication year of source reference	
Coauthors	List of co-authors for source reference	
Location	Place name (city, town or management area)	
Country	Country (if non-US)	
State/province	State or province	
Burn name	Burn unit name (if applicable)	e.g. Rim Fire, Lolo 1
Measurement date	Date of burn	
Fuel type desc	Description of fuel type	e.g. Douglas fir crown fuels – fine branches and needles

Methods

As part of a broader study on fire behaviour and emissions funded by the Strategic Environmental Research and Development Program (SERDP), [Lincoln et al. \(2014\)](#) compiled existing emissions factor data from the 1960s through ~2010 into annotated tables, with companion references available to download (<https://www.fs.usda.gov/rds/archive/Product/RDS-2014-0012>, accessed 26 November 2019). We converted the tabular dataset into a relational database and reviewed all source references to add greater attribution and correct input errors, including location, major vegetation type, MCE and/or combustion phase and burn type (e.g. laboratory, prescribed fire or wildfire and measurement platform). Each source reference was reviewed to ensure that the EF observation was fully attributed with fields defined in [Table 1](#). Supported pollutants were assigned a molecular weight, common name, an International Union of Pure and Applied Chemistry database identification number and US Environmental Protection Agency (EPA) Substance Registry Services ID and pollutant category (SRS, https://ofmpub.epa.gov/sor_internet/registry/substreg/LandingPage.do, accessed 26 November 2019) to allow cross-referencing to other air quality and pollutant databases. Reported EF units were standardised to grams of pollutant per kilogram of dry weight

biomass consumed (g kg⁻¹). In many cases, EFs were converted from lb ton⁻¹ to g kg⁻¹ or from g kg⁻¹-C to g kg⁻¹ if the study also reported the fuel carbon content. We did not attempt to convert emission ratio data into EFs. For each study, we added individual burn data as opposed to summary data. We also confirmed that all source records were original EF data and not duplicated in other studies. For records having a measured MCE, combustion phases were assigned as flaming (MCE ≥ 0.9), smouldering (MCE = 0.8 to <0.9) and residual smouldering (MCE <0.8) (after [Akagi et al. 2011](#)). In older records in which CE was reported instead of MCE, we either used the reported phase or same thresholds to assign combustion phase. Once the existing dataset was quality assured and standardised, we added records from recent published studies (2010–18). See Supplementary Appendix 1 for the list of references augmenting the [Lincoln et al. \(2014\)](#) work. Currently, there are 12 533 records in the database from 49 source references.

The goal of SERA was to compile available EF data into a single repository with ease of access to the data. In this sense, all records are treated equally. We did not attempt to identify whether one study might be better than another study. The online, searchable database makes it straightforward to lookup the original references, allowing users to customise data for their particular application.

Statistical analysis

To evaluate the influence of combustion phase, MCE, burn type and vegetation or fuel type on estimated EFs, we concentrated on major pollutants with sufficient record counts and of high relevancy to wildland fire planners and regulators: carbon dioxide (CO₂); carbon monoxide (CO); methane (CH₄); nitrogen oxides (NO_x); particulate matter (PM_{2.5} and total PM); and sulfur dioxide (SO₂).

Explanatory variables of EFs were evaluated using Analysis of Variance tests (R statistical programming, ANOVA) (R Studio, Supplementary Appendix 2). Tukey's Honest Significant Difference testing was used to evaluate for significant differences among predictor categories (e.g. combustion phase, burn type or fuel type). Due to unequal distribution of records across predictor variables, one-way ANOVA tests were conducted for each variable and on subsets of pollutant emissions data with sufficient and comparable record counts. For example, combustion phase was evaluated as a potential predictor variable of CO₂ EFs, but because few records were available for the residual smouldering ($n < 10$), only those records recorded as either flaming or smouldering were evaluated.

We evaluated how EFs vary by burn type, which is a combination of fire type (i.e. laboratory, prescribed fire and wildfire) and measurement platform (i.e. ground-based, tower and airborne) (Fig. 1). Due to imbalances in record counts within each burn type category, pollutant datasets were subset to exclude any burn types with insufficient record counts ($n < 10$).

Because vegetation and/or fuel type are generally used to summarise and apply EFs in wildland fire management and emissions inventories, we evaluated regional vegetation type as an explanatory variable of EFs by major pollutant (Table 2). Regional vegetation types were first analysed, subsetting the dataset to exclude sparsely populated types; we also evaluated fuel category, which assigns a dominant fuel (e.g. grass, shrub, pine needles, organic soil) regardless of region for each EF record (also listed in Table 2). Fuel consumption is influenced by a range of factors, including mass of available fuel, fuel moisture, surface area-to-volume ratios, packing ratio and presence of volatile compounds (Ottmar 2014). In the absence of fuel moisture content data for each EF record, regional vegetation type may provide a surrogate for climatic differences in site moisture. In contrast, comparing EFs among fuel categories evaluates potential differences in inherent combustion properties, including differences in packing and surface area-to-volume ratios.

Sample application

To demonstrate how EFs can be used for prescribed burn planning, we used a dataset of prescribed understorey burns in SE and W conifer forests of the US that reported mean fuel consumption by major surface fuel category (Prichard *et al.* 2017). For PM_{2.5}, we had sufficient records in SERA to calculate EFs specific to flaming and smouldering consumption in addition to a fire-average short-term flaming and smouldering EF. For fuel categories that contribute to long-term smouldering, we used the residual coarse wood EF of 33 g kg⁻¹ and residual duff EF of 50 g kg⁻¹, after Urbanski (2014). Currently, there are no empirical models to estimate consumption by combustion

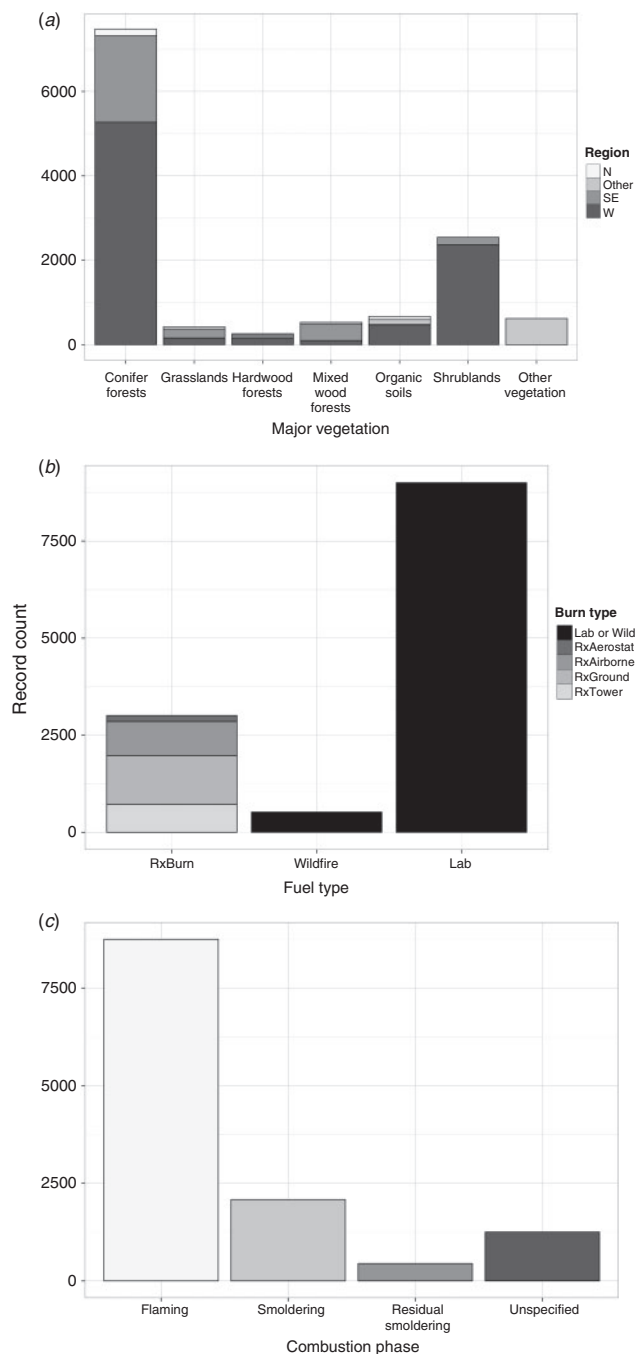


Fig. 1. Emissions factor record counts by (a) region and vegetation type, (b) burn type, including prescribed burn emissions sampled by aerostat (Rx-Aerostat), airborne (Rx-Airborne), ground (Rx-Ground) and tower (Rx-Tower) measurement platforms, laboratory burns (Laboratory) and wildfire emissions sampled using airborne platforms (Wild-Airborne) and (c) combustion phase. See Table 1 for variable definitions.

phase, and consumption was not reported by phase in Prichard *et al.* (2017). The two main operational consumption models in the US have different ways of allocating consumption into flaming v. smouldering consumption and making emissions factor assignments. Consume v 5.0 assumes a generic proportion

Table 2. Regional vegetation type and fuel category definitions

Regional vegetation type	Definition	Fuel category	Definition
<i>Northern</i>			
N conifer forests	Boreal and sub-boreal conifer forests (e.g. black spruce, white spruce forests)	Duff	Organic soils
N duff	Boreal and sub-boreal organic soil	Grass	Grassland
N grass	Boreal and sub-boreal grassland	Hardwood	Broadleaf deciduous forest (understorey fuels)
N hardwood	Boreal and sub-boreal broadleaf deciduous forest	Mixed conifer	Mixed conifer forest (understorey and canopy fuels)
N mixed wood	Boreal and sub-boreal mixed broadleaf and conifer forest	Other	
<i>South-eastern</i>			
SE duff	South-eastern organic soil	Pine needles	Pine forest with needle layer understorey
SE grass	South-eastern grassland	Pine shrub	Pine forest with shrub understorey
SE hardwood	South-eastern broadleaf deciduous and evergreen forest	Pine forest understorey	Pine forest understorey (general)
SE mixedwood	South-eastern mixedwood forests	Shrub	Shrubland
SE pine forests	South-eastern pine forests (e.g. loblolly and longleaf pine forests)	Slash	Logging slash
SE shrub	South-eastern shrublands		
<i>Western</i>			
W conifer forest	Western mixed conifer and ponderosa pine forests		
W duff	Western organic soil		
W grass	Western grassland		
W mixedwood	Western mixed broadleaf and mixed conifer forest		
W shrub	Western shrubland		
W hardwood	Western broadleaf deciduous forest		
<i>Other</i>			
Other	Other fuel types that are not regionally specific (e.g. lumber, excelsior)		
Other duff	Other organic soil types		

for each fuel category (e.g. herbs, shrubs, 1-h wood, 1000-h wood, litter or duff) to partition consumption values in flaming, smouldering and residual consumption (Prichard 2007). In contrast, FOFEM uses the BURNUP model (Albini and Reinhardt 1997) to calculate intensity of the burn with a threshold of $>15 \text{ kW m}^{-2}$ representing flaming combustion and $<15 \text{ kW m}^{-2}$ representing smouldering combustion. For these sample calculations, we used Consume's assumed proportional consumption by combustion phase to allocate the average fuel consumption reported for south-eastern forest and western ponderosa pine sites in Prichard *et al.* (2017) into flaming, smouldering and residual smouldering phases. Emissions by fuel category were then estimated by two methods – first by calculating flaming, smouldering and residual emissions by fuel category, and second by estimating total emissions using a fire-average EF. Variability in total emission estimates were also calculated using a single standard deviation of EF ($\pm 1 \text{ s.d.}$).

Results

Of the 12 533 records in the database, 45% ($n = 5637$) are represented by 23 air pollutants, most of which are designated as EPA criteria air pollutants (CAP), greenhouse gases (GHG), hazardous air pollutants (HAP) or known air toxins (TOX) (Table 3). A list of the 274 pollutants currently supported in the database, including record counts and summary statistics for EF and MCE, is available as an online appendix table (Supplementary Appendix 3). Among all pollutants in the

database, including the most common pollutants (e.g. PM, CO, CO₂ and CH₄), records are unevenly distributed with a bias towards flaming combustion, prescribed burning and laboratory measurements. Overall, EF records are most common for SE and W conifer forests and W shrubland types of the US (Fig. 1). Considering EF records by fuel category, EF records are most numerous for western mixed conifer, south-eastern and western pine understorey fuels and western shrublands.

Carbon dioxide and carbon monoxide

CO₂ and CO are well represented in the database, with a total of 578 and 640 records respectively, excluding outliers. Early emission studies did not generally record combustion phase (MCE or combustion efficiency, CE), and more than a quarter of records lack an assigned phase. Of the records with an assigned phase, around 50% of CO₂ and CO EFs were sampled during flaming combustion. Although there is a positive relationship between CO₂ and MCE, the relationship is weak ($R^2 = 0.4113$, $P < 0.0001$, Fig. 2). There are significant differences in CO₂ EFs among flaming, smouldering and residual smouldering combustion phases, with the highest EFs in flaming and lowest EFs in residual smouldering (Fig. 3).

Regional vegetation type is not a strong predictor of CO₂ EFs, with only a significant difference between SE shrub and SE pine CO₂ EFs. Comparing fuel categories, duff EFs are significantly lower than most other categories (e.g. grass, mixed conifer, mixed wood, pine forest understorey, shrub and slash).

Table 3. Pollutants with a minimum of 100 record counts in the Smoke Emissions Repository Application (SERA), including record counts (*n*) and summary statistics of reported emission factors (EFs) including mean and standard deviation (s.d.)

Pollutant categories include US Environmental Protection Agency Criteria Air Pollutant (CAP), greenhouse gas (GHG), hazardous air pollutant (HAP) and toxic air pollutant (TOX). Summary values exclude outlier and residual smouldering EFs but otherwise include all records in the database, including 'other' vegetation types and records without a regional fuel group assignment. MCE, modified combustion efficiency

Pollutant name	Category	Acronym or formula	Molecular weight	<i>n</i>	Mean EF	s.d.	Mean MCE	s.d.
Acetic acid		C ₂ H ₄ O ₂	60.052	153	2.13	2.03	0.92	0.04
Acetylene		C ₂ H ₂	26.038	291	0.83	1.27	0.93	0.04
Ammonia	CAP, TOX	NH ₃	17.031	216	1.55	1.87	0.91	0.07
Black carbon		BC		115	0.96	1.00	0.93	0.03
Carbon dioxide	GHG	CO ₂	44.009	597	1549.98	312.47	0.91	0.06
Carbon monoxide	CAP	CO	28.010	640	103.51	58.25	0.90	0.06
Ethene	TOX	C ₂ H ₄	28.054	261	1.10	2.12	0.93	0.04
Formaldehyde	HAP, TOX	H ₂ CO	30.026	204	1.59	1.13	0.92	0.04
Formic acid	TOX	HCOOH	46.025	192	0.33	0.41	0.93	0.03
Furan	TOX	C ₂ H ₄ O	68.075	179	0.34	0.36	0.92	0.04
Hydrogen cyanide	HAP, TOX	HCN	27.026	188	0.46	0.59	0.92	0.04
Methane	GHG	CH ₄	16.043	451	5.53	14.76	0.90	0.06
Methanol	HAP, TOX	CH ₃ OH	32.042	219	1.44	1.44	0.92	0.04
Nitric oxide	CAP	NO	30.006	193	2.29	2.08	0.93	0.04
Nitrogen dioxide	CAP	NO ₂	46.005	160	1.26	1.16	0.94	0.03
Nitrogen oxides	CAP	NO _x		117	3.36	3.73	0.90	0.09
Nitrous acid		HNO ₂	47.013	164	0.49	0.24	0.93	0.03
Non-methane hydrocarbons		NMHC		112	5.87	4.60	0.86	0.08
Particulate matter 2.5 µm	CAP	PM _{2.5}		337	27.87	43.09	0.90	0.07
Phenol	HAP, TOX	C ₆ H ₆ O	94.113	137	0.71	0.80	0.93	0.03
Propene	TOX	C ₃ H ₆	42.081	295	0.68	1.11	0.92	0.04
Sulfur dioxide	CAP	SO ₂	64.058	127	1.11	0.72	0.93	0.03
Total particulate matter		PM		289	23.57	25.69	0.87	0.08

Pine forest understorey EFs have a narrow range of observed values and are significantly higher than some other fuel categories (e.g. duff, slash, pine needles). When only flaming CO₂ EFs are considered, no significant differences are found among fuel categories. However, when considering only smouldering CO₂ EFs, duff EFs are significantly lower than mixed conifer and pine forest understorey fuels, and mixed conifer EFs are significantly higher than grass.

Among burn types (Laboratory, Rx-Air, Rx-Ground and WF-Air), laboratory CO₂ EF values have the widest representation and range of observed values, with no significant difference among other burn types. Rx-Tower observations range widely but are significantly lower than Rx-Ground and WF-Air. When CO₂ EFs are partitioned into flaming and smouldering combustion classes, no significant differences are found among burn types.

The CO EFs are significantly different among combustion phases, with highest and most variable CO EFs in the residual smouldering phase, moderate EFs in smouldering and lowest EFs in flaming combustion (Fig. 3). CO is negatively related to MCE ($R^2 = 0.6444$, $P < 0.0001$, Fig. 2). Regional vegetation types and fuel categories are unevenly distributed among CO EFs. CO EFs are generally not statistically different among regional vegetation, with only significant differences between SE grass and W conifer. However, there are notable differences in CO EFs by fuel category with significantly higher duff EFs than some other fuel categories (grass, pine understorey and shrub), and also differences between slash and shrubs, grass and pine understorey fuels, and between grass and shrub. Due to

significant differences by combustion phase, CO EF datasets are subset into combustion phases to evaluate the influence of burn type and fuel type. Flaming CO EFs significantly differ among tested fuel categories, with significantly lower values in grass and slash than the other types. Smouldering CO have low record counts, and comparisons can only be made among duff, mixed conifer, shrub and slash. Smouldering CO values are widely distributed but are significantly higher in mixed conifer than in shrub fuels. Overall, CO EFs are significantly higher in Rx-Tower than Laboratory, Rx-Airborne and Rx-Ground.

Methane

There are 389 records of CH₄ in the database, with 59% of observations representing flaming combustion. CH₄ is negatively correlated with MCE ($R^2 = 0.7144$, $P < 0.0001$, Fig. 2), but the linear model has a wide scatter in residuals, particularly with lower MCE values. CH₄ EFs are significantly different between flaming, smouldering and residual smouldering combustion phases, with the highest EFs in residual smouldering and lowest in flaming (Fig. 3). Only nine observations are available for the residual smouldering phase of combustion and are widely distributed. Regional vegetation is a significant predictor of CH₄ EFs. EFs for W conifer are significantly higher than SE grass, and W duff EFs are significantly higher than N conifer, SE grass, SE pine and W shrub. EFs also significantly differ by fuel category, with significantly higher EFs in coarse wood and duff than all other fuel categories except coarse wood v. duff. Mixed conifer EFs are significantly lower than coarse

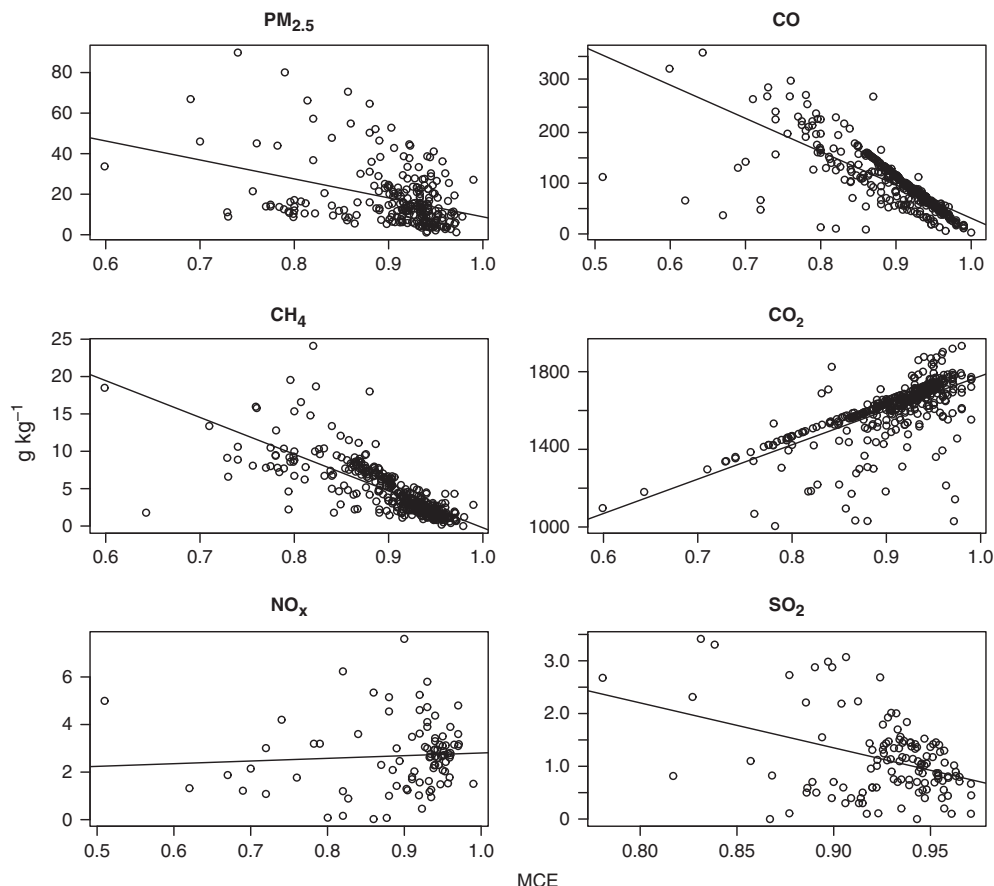


Fig. 2. Correlation plots between modified combustion efficiency and emission factor of major pollutants including $\text{PM}_{2.5}$, CO, CH_4 , CO_2 , NO_x and SO_2 .

wood and duff, but are significantly higher than grass, pine–shrub understorey, pine–understorey and shrubs. Slash is significantly higher than grass and pine–shrub understorey.

Considering only flaming combustion, mixed conifer has greater EFs than grass, pine forest understorey, shrub and slash fuel categories. However, there are no significant differences in smouldering EFs by fuel category. CH_4 EFs also significantly differ by burn type, with significantly higher EFs in Rx-Tower and WF-Air burns than in Laboratory and Rx-Ground burns. Considering only flaming combustion, WF-Air CH_4 EFs are significantly higher than other burn types. However, there are no significant differences among burn types in smouldering CH_4 EFs.

Particulate matter

Particulate matter records are recorded across a range of size fractions in the database (i.e. $<1\ \mu\text{m}$, $<2\ \mu\text{m}$, $<2.5\ \mu\text{m}$, $<3\ \mu\text{m}$, $<4\ \mu\text{m}$ and total PM). The most common records are for total PM and $\text{PM}_{2.5}$. Because the majority of PM is represented by the $<2.5\ \mu\text{m}$ size classes, we evaluated $\text{PM}_{2.5}$ EFs and also grouped all records in a generic PM category to evaluate the larger dataset.

$\text{PM}_{2.5}$ EFs are negatively related to MCE, with a wide scatter in model residuals at lower MCE values ($R^2 = 0.4735$, $P < 0.0001$, Fig. 2). Of the records with MCE values, observations are most represented by flaming combustion ($n = 177$ records

v. $n = 45$ smouldering and $n = 11$ residual smouldering) (Fig. 3). Even with a wide spread in observed values, $\text{PM}_{2.5}$ EFs significantly differ by flaming and smouldering combustion phase. Considering all $\text{PM}_{2.5}$ EFs, there are no significant differences among burn types. However, smouldering $\text{PM}_{2.5}$ EFs are significantly higher in laboratory burns than in Rx-Ground and Rx-Tower observations.

$\text{PM}_{2.5}$ EF observations are generally too unevenly distributed to evaluate statistical differences among regional vegetation types with the majority of observations in SE pine and W conifer (Fig. 1). $\text{PM}_{2.5}$ EFs are significantly higher in SE pine than W conifer. Comparisons among fuel categories are also challenged by imbalanced data with the majority of observations being in pine understorey fuels. Considering only flaming $\text{PM}_{2.5}$ emissions, pine forest understorey EFs are significantly greater than shrub EFs. Comparing smouldering $\text{PM}_{2.5}$ emissions, shrub and pine forest understorey emissions are significantly higher than slash fuels.

Total PM EFs have higher record counts ($n = 580$) but follow similar trends with $\text{PM}_{2.5}$ emissions ($n = 266$). PM EFs are weakly and negatively correlated with MCE ($R^2 = 0.1478$). Flaming PM EFs are significantly lower than smouldering and residual smouldering PM EFs. There are few significant differences among burn type; laboratory PM EFs are significantly

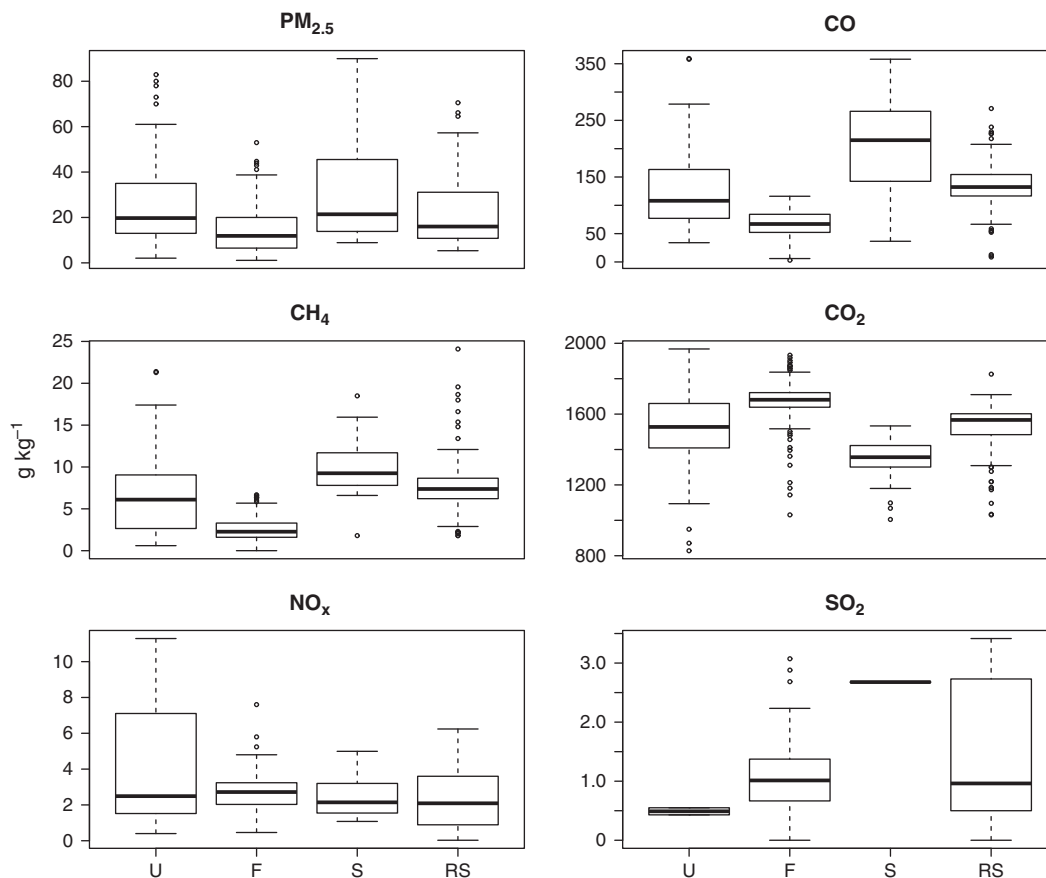


Fig. 3. Box and whisker plots of emission factor by combustion phase (U, undesigned; F, flaming; S, smouldering; RS, residual smouldering combustion phases) and major pollutant (PM_{2.5}, CO, CH₄, CO₂, NO_x and SO₂).

higher than Rx-Airborne and Rx-Tower observations, and Rx-Ground PM EFs are significantly higher than Rx-Airborne values. PM EFs are significantly higher in western shrub than in SE pine and SE shrub. Also, PM EFs are higher in shrub than in grass and pine needle fuel categories, and higher in pine understory fuels than in grass and slash. Smouldering PM values are rare, and the only possible comparison was between pine understory fuels, which were significantly higher than slash EFs.

Nitrogen oxides

Nitrogen oxides are represented as NO, NO₂ and NO_x in database records, and records are combined in this analysis to maximise the number of observations ($n = 462$). NO_x is weakly correlated with MCE ($R^2 = 0.0694$, Fig. 2). There are also no significant differences between combustion phases and particularly wide distributions of values in the residual smouldering phase (Fig. 3). Most NO_x observations were sampled in laboratory burns ($n = 427$), and although sample sizes are too unevenly distributed for statistical comparisons, Laboratory EFs are substantially higher than Rx and WF EFs. Although there are sufficient record counts to compare NO_x EFs among regional vegetation and fuel categories, only W shrub is significantly different (higher) than the SE Pine

vegetation types, and shrub fuel EFs are greater than Pine forest understory and slash fuel categories. Among flaming NO_x, only shrub EFs are significantly different (higher) than pine understory fuels. Record counts are low for smouldering NO_x observations, and there is no statistical difference among fuel categories.

Sulfur dioxide

Of the 127 SO₂ EFs, 82 were sampled during the flaming phase of combustion. SO₂ EFs are negatively but weakly correlated with MCE ($R^2 = 0.1436$, Fig. 2), with significant differences between flaming and smouldering combustion phases (Fig. 3). Almost all SO₂ values were collected in the laboratory, with an emphasis on W shrubs and W conifer vegetation types (Figs 2 and 3). Due to the uneven distribution of EFs within regional vegetation types and fuel categories, only a few statistical comparisons are possible. Of these, W conifer SO₂ EFs are statistically higher than western shrubs. Of the fuel categories, shrub EFs are significantly lower than duff and mixed conifer EFs.

Updated EFs by combustion phase

To meet land manager needs for EFs by combustion phase and vegetation type, we used SERA to produce a standardised

Table 4. Summarised emission factors (g kg^{-1}) for the south-eastern US by combustion phase for particulate matter $<2.5 \mu\text{m}$ ($\text{PM}_{2.5}$), carbon dioxide (CO_2), carbon monoxide (CO), methane (CH_4), ammonia (NH_3), sulfur dioxide (SO_2), nitrogen oxide (NO), nitrogen dioxide (NO_2) and oxides of nitrogen (NO_x)

Mean, mean emission factor (g kg^{-1}); s.d., standard deviation (g kg^{-1}); n , count of data points; fire-average F/S, combined flaming, smouldering and unspecified combustion phases. F, flaming-phase EF, S, smouldering-phase EF. Cells are missing when there were insufficient records

Pollutant	Fire-average F/S			F			S		
	Mean	s.d.	n	Mean	s.d.	n	Mean	s.d.	n
<i>Conifer forest (SE pine)</i>									
$\text{PM}_{2.5}$	25.29	16.00	102	20.91	10.95	60	29.35	18.23	18
CO_2	1576.04	248.04	86	1691.78	51.78	39	1462.00	170.14	11
CO	97.09	43.11	109	73.88	21.54	39	165.87	38.09	11
CH_4	4.12	5.02	47	2.41	1.26	38	11.36	8.06	9
NH_3	0.82	1.25	31	0.52	0.76	22	1.56	1.87	9
NO	1.86	1.80	28	1.20	0.81	18	0.78	0.94	3
NO_2	1.23	0.83	23	1.62	0.74	15	0.93		1
NO_x	3.34	2.80	27	2.43	1.10	15	1.56	0.80	3
SO_2	0.99	0.40	10	1.06	0.31	7	1.55		1
<i>Shrublands</i>									
$\text{PM}_{2.5}$	12.03	4.25	5	12.03	4.25	5			
CO_2	1707.96	192.21	15	1746.03	95.84	13	1460.5	515.48	2
CO	73.50	17.00	16	70.40	14.01	13	93.75	32.17	2
CH_4	2.38	0.87	12	2.20	0.45	10	3.30	2.12	2
NH_3	2.06	2.27	6	1.15	0.50	5	6.60		1
NO	4.71	2.45	7	3.77	2.24	5	7.05	0.78	2
NO_2	0.95	0.43	3	0.95	0.43	3			
NO_x	2.86	0.78	4	2.86	0.78	4			
SO_2	0.74	0.60	6	0.67	0.64	5	1.1		1
<i>Mixed forest</i>									
$\text{PM}_{2.5}$	14.78	0.94	4	14.78	0.94	4			
CO_2	1650.50	61.10	10	1658.33	59.24	9	1580.0		1
CO	84.19	20.30	12	76.87	12.10	9	129.50		1
CH_4	2.67	1.28	9	2.27	0.45	8	5.90		1
NH_3	2.13	1.29	5	1.74	1.09	4	3.70		1
NO	3.69	3.09	3	2.39	2.98	2	6.30		1
NO_2	1.14	0.62	2	1.14	0.62	2			
NO_x	2.18	1.31	2	2.18	1.31	2			
SO_2	0.70	0.16	4	0.77	0.12	3	0.50		1
<i>Grasslands</i>									
$\text{PM}_{2.5}$	12.080	5.24	10	12.08	5.24	10			
CO_2	1685.82	81.19	15	1696.38	72.78	14	1538.0		1
CO	68.17	27.16	16	64.6	25.27	14	119.0		1
CH_4	2.53	1.51	10	2.21	1.20	9	5.4		1
NH_3	1.30	0.70	4	1.37	0.84	3	1.1		1
NO	5.33	2.14	4	5.83	2.30	3	3.8		1
NO_2									
NO_x	3.53	0.40	3	3.53	0.40	3			
SO_2	0.83	0.34	4	0.93	0.32	3	0.5		1

dataset of summarised EFs for $\text{PM}_{2.5}$, CO, CO_2 , CH_4 , NH_3 , SO_2 and NO_x by regional fuel type (conifer forest, broadleaf deciduous forest, mixed conifer and broadleaf deciduous forest, shrubland, grassland and organic soil) for the south-eastern US (Table 4, Box 1), western US and Canada (Table 5), and northern US and Canada (Table 6). Summary EFs are listed by combustion phase, including a fire-average EF value, representing short-term flaming and smouldering emissions, in addition to separate flaming EF and smouldering EFs where available. Due to uneven record counts, EFs by combustion phase are only available for some pollutant–region

combinations, and highlight data gaps for some pollutants. Slash EFs are excluded in these summary tables, using the ‘exclude slash’ option in SERA.

Table 7 presents sample calculations of estimated $\text{PM}_{2.5}$ emissions for SE pine and W pine forests. For SE pine forests, using specific EFs to estimate flaming, smouldering and residual emissions resulted in a lower estimated total $\text{PM}_{2.5}$ emissions than the fire-average EFs. In contrast, estimates using the two approaches were markedly similar in the W pine example, reflecting the importance of coarse wood and duff on total emissions.

Box 1. Using summarised emission factors to estimate wildland fire emissions

There are two approaches to estimating PM_{2.5} emissions using summaries in Tables 4 and 5 to explore differences in estimated values using comparative approaches. We use EF values available for south-eastern conifer (Table 4) and western mixed conifer forests (Table 5) to represent flaming and smouldering emissions as well as fire average (flaming and/or smouldering; F/S) emissions. Following Urbanski (2014), we use the recommended residual coarse wood EF of 33 g kg⁻¹ and residual duff EF of 50 g kg⁻¹ to estimate residual emissions from coarse wood and duff. Using estimates of flaming, smouldering and residual smouldering consumption, based on assumed proportions in Consume v5.0 (see Eqn 1), we can multiply consumption in Mg ha⁻¹ by the recommended EF by phase (g kg⁻¹) to calculate PM_{2.5} emissions (kg ha⁻¹).

Site	Flaming EF (g kg ⁻¹)	Smouldering EF (g kg ⁻¹)	Residual Smouldering Coarse wood (g kg ⁻¹)	Residual smouldering duff (g kg ⁻¹)	Fire average EF (F/S) (g kg ⁻¹)
SE conifer	20.91	29.35	33	50	25.29
W conifer	13.5	20.05	33	50	14.32

Sample calculations

To estimate emissions from sound coarse wood (7.6 to 22.9 cm diameter) in a south-eastern US forest with hurricane damage, we can apply three separate EFs to represent flaming, smouldering and residual consumption (where EFs are expressed in kg Mg⁻¹ because 1 g kg⁻¹ = 1 kg Mg⁻¹):

$$\begin{aligned} \text{PM}_{2.5} \text{ emissions of coarse wood} = & \\ (22.4 \text{ Mg ha}^{-1} \times 0.6 \text{ Flaming} \times 20.91 \text{ kg Mg}^{-1}) + & \\ (22.4 \text{ Mg ha}^{-1} \times 0.3 \text{ Smouldering} \times 29.35 \text{ kg Mg}^{-1}) + & \\ (22.4 \text{ Mg ha}^{-1} \times 0.1 \text{ Residual} \times 33 \text{ kg Mg}^{-1}) = 552.18 \text{ kg ha}^{-1} (1) & \end{aligned}$$

For a more general estimate, fire-average EFs are available and can be used instead. For SE conifer forests, this results in a slightly higher estimation of emissions.

$$\begin{aligned} \text{PM}_{2.5} \text{ emissions of 1000-h wood} = & \\ 22.4 \text{ Mg ha}^{-1} \times 25.29 \text{ kg Mg}^{-1} = 566.5 \text{ kg ha}^{-1} (2) & \end{aligned}$$

Where fuel consumption is not available by fuel category, the fire average EF can be used to estimate consumption based on total site fuel loading. This calculation can be appropriate for some fuels such as the SE conifer forest in this example, or potentially significantly underestimate PM_{2.5} emissions in sites dominated by heavier fuels and organic soils.

For example, for an SE conifer site dominated by pine litter and shrubs that are mostly consumed in the flaming phase of combustion, the fire average EF may overestimate emissions compared with using the flaming EF:

$$\begin{aligned} \text{Fire-average EF estimate of PM}_{2.5} \text{ emissions} = & \\ 13 \text{ Mg ha}^{-1} \text{ consumed} \times 25.29 \text{ kg Mg}^{-1} = 328.77 \text{ kg ha}^{-1} (3) & \\ \text{Flaming EF estimate of PM}_{2.5} \text{ emissions} = & \\ 13 \text{ Mg ha}^{-1} \text{ consumed} \times 20.91 \text{ kg Mg}^{-1} = 271.83 \text{ kg ha}^{-1} (4) & \end{aligned}$$

For a W conifer site dominated by coarse wood, using the fire-average EF may substantially underestimate emissions:

$$\begin{aligned} \text{Fire-average EF estimate of PM}_{2.5} \text{ emissions} = & \\ 20 \text{ Mg ha}^{-1} \text{ consumed} \times 14.32 \text{ kg Mg}^{-1} = 286.40 \text{ kg ha}^{-1} (5) & \\ \text{Residual coarse wood estimate of PM}_{2.5} \text{ emissions} = & \\ 20 \text{ Mg ha}^{-1} \text{ consumed} \times 33 \text{ kg Mg}^{-1} = 660.00 \text{ kg ha}^{-1} (6) & \end{aligned}$$

Discussion

The online EF database, SERA, was designed as a clearinghouse for field and laboratory-based EFs for Canada and the US. By recording existing EFs by fuel type, MCE and burn type, the tool supports summaries of EFs for use in emissions inventories and wildland fire management. Our sample calculations for SE pine and W pine sites, presented in Table 7, demonstrate that EFs summarised by combustion phase can lead to substantially different emissions estimates than fire-average EFs in certain fuel types. For example, consumption in SE pine sites is dominated by fuels that burn in the flaming phase of combustion. Applying specific EFs for flaming, smouldering and residual smouldering

combustion produced a much lower estimate of total PM_{2.5} emissions than the fire-average EF value. In contrast, the W pine site was dominated by coarse wood and duff, and both approaches produced a comparable estimate of total PM_{2.5} emissions.

As more EFs are compiled by combustion phase and fire type (e.g. flaming v. smouldering, prescribed v. wildfire), some pollutants that are highly dependent on combustion phase or fire type (e.g. CO, CH₄ and PM_{2.5}) may be better characterised to assist with smoke management techniques that identify fuels likely to burn in short- and long-term smouldering combustion and mitigate smoke impacts. Flaming-phase EFs are more represented in SERA than are smouldering and residual smouldering EFs, but in certain vegetation and fuel types, long-term

Table 5. Summarised emission factors (g kg^{-1}) for the western US by combustion phase for particulate matter $<2.5 \mu\text{m}$ ($\text{PM}_{2.5}$), carbon dioxide (CO_2), carbon monoxide (CO), methane (CH_4), ammonia (NH_3), sulfur dioxide (SO_2), nitrogen oxide (NO), nitrogen dioxide (NO_2) and oxides of nitrogen (NO_x)

Mean, mean emission factor (g kg^{-1}); s.d., standard deviation (g kg^{-1}); n = count of data points. Fire-average F/S, combined flaming, smouldering and unspecified combustion phases; F, flaming-phase EF; S, smouldering-phase EF

Pollutant	Fire-average F/S			F			S		
	Mean	s.d.	n	Mean	s.d.	n	Mean	s.d.	n
<i>Conifer forest (western pine and mixed conifer)</i>									
$\text{PM}_{2.5}$	14.32	5.51	24	13.50	5.18	21	20.05	4.97	3
CO_2	1629.54	63.43	147	1662.92	52.08	87	1579.40	44.20	58
CO	104.01	34.93	147	79.97	20.85	87	140.30	15.57	58
CH_4	5.05	2.41	144	3.41	1.33	85	7.59	1.16	57
NH_3	0.90	0.57	54	0.83	0.53	49	1.74	0.09	3
NO	1.79	1.09	58	1.71	0.70	53	2.60	3.12	5
NO_2	1.38	0.75	56	1.40	0.71	52	1.05	1.23	4
NO_x	1.41	0.95	6	1.49	0.77	2	1.08	1.43	2
SO_2	1.38	0.61	50	1.35	0.54	47	1.73	1.44	3
<i>Shrublands</i>									
$\text{PM}_{2.5}$	7.88	2.21	14	7.12	1.22	10	9.79	3.15	4
CO_2	1587.88	141.8	98	1684.48	82.4	43	1538.81	72.42	7
CO	97.03	54.82	116	63.87	17.6	51	76.32	40.34	17
CH_4	2.16	1.37	46	1.87	1.16	38	4.14	1.15	6
NH_3	1.60	1.52	53	1.16	0.98	39	3.13	2.05	12
NO	2.42	1.29	35	2.22	0.91	32	4.53	2.84	3
NO_2	0.79	0.66	34	0.80	0.67	33	0.50		1
NO_x	3.33	1.97	35	3.11	1.21	22	2.89	2.40	10
SO_2	0.73	0.28	27	0.75	0.28	25	0.55	0.21	2
<i>Mixed forest</i>									
$\text{PM}_{2.5}$	6.83		1	6.83		1			
CO_2	1669.50	134.67	4	1669.50	134.67	4			
CO	55.26	22.07	4	55.26	22.07	4			
CH_4	2.35	1.78	4	2.35	1.78	4			
NH_3	0.60	0.34	3	0.60	0.34	3			
NO	1.65	1.39	4	1.65	1.39	4			
NO_2	1.53	1.97	4	1.53	1.97	4			
NO_x	2.55	1.40	4	2.55	1.40	4			
SO_2	0.49	0.33	3	0.49	0.33	3			
<i>Grasslands</i>									
$\text{PM}_{2.5}$									
CO_2	1421	208.48	8	1547.3	52.17	5	1210.5	200.83	3
CO	56.26	38.89	8	29.62	11.91	5	100.67	16.68	3
CH_4	2.90	1.86	5	1.63	0.75	3	4.80	0.84	2
NH_3	0.56	0.53	4	0.30	0.08	3	1.35		1
NO	2.86	1.58	7	2.77	1.50	5	3.09	2.42	2
NO_2	3.13	2.36	2	4.80		1	1.46		1
NO_x									
SO_2	2.98		1				2.98		1

smouldering can be the largest contributor to pollutant emissions and pose the greatest smoke management challenge. The SERA database also offers a standard clearinghouse for new pollutants that are just being documented in wildland smoke, including a wide range of non-methane organic gases, including volatile organic compounds and other air toxins (Urbanski 2014; Koss *et al.* 2018). It also could be expanded to support EFs in other parts of the world and adapted to incorporate other variables, such as trace indicators of low-heat and high-heat pyrolysis (Gilman *et al.* 2015; Sekimoto *et al.* 2018).

Summarising pollutants in our database by fuel type, burn type and combustion phase revealed several data gaps and a bias in the type of measurements collected to date. Across all pollutants in the database, most EFs were measured during the flaming phase of combustion. Given that some pollutants, such as $\text{PM}_{2.5}$, CO , CH_4 and a wide range of NMOCs, are more concentrated in smoke produced in short- and long-term smouldering combustion, a greater emphasis on EF characterisation during these phases is needed. For local smoke management applications including prescribed burning support, there is a particular need to

Table 6. Summarised emission factors (EFs; g kg⁻¹) for northern boreal forests and grasslands by combustion phase for particulate matter <2.5 microns (PM_{2.5}), carbon dioxide (CO₂), carbon monoxide (CO), methane (CH₄), ammonia (NH₃), sulfur dioxide (SO₂), nitrogen oxide (NO), nitrogen dioxide (NO₂) and oxides of nitrogen (NO_x)

Mean, mean emission factor (g kg⁻¹); s.d., standard deviation (g kg⁻¹); *n*, count of data points; fire-average F/S, combined flaming, smouldering and unspecified combustion phases; F, flaming-phase EF; S, smouldering-phase EF

Pollutant	Fire-average F/S			F			S		
	Mean	s.d.	<i>n</i>	Mean	s.d.	<i>n</i>	Mean	s.d.	<i>n</i>
<i>Northern forest</i>									
PM _{2.5}	10.53	0.81	3	10.53	0.81	3			
CO ₂	1614.26	116.24	11	1624.11	132.23	8	1616.00		1
CO	86.49	35.27	11	72.79	18.41	8	113.00		1
CH ₄	2.60	1.39	12	2.03	0.78	9	4.70		1
NH ₃	1.03	0.48	7	1.10	0.49	6			
NO	2.28	1.64	4	2.94	1.19	3	0.29		1
NO ₂	1.94	1.13	3	2.40	1.13	2	1.03		1
NO _x	1.28	0.32	2						
SO ₂	0.13	0.06	3	0.13	0.06	3			
<i>Northern grassland</i>									
PM _{2.5}	9.89	6.90	4	9.89	6.90	4			
CO ₂	1697.38	39.83	5	1697.38	39.83	5			
CO	64.44	16.18	5	64.44	16.18	5			
CH ₄	2.04	0.99	5	2.04	0.99	5			
NH ₃	0.77		1	0.77		1			
NO	2.27		1	2.27		1			
NO ₂									
NO _x									
SO ₂									

Table 7. Comparison of PM_{2.5} emissions (g kg⁻¹) from representative prescribed burn consumption data for south-eastern US and western US pine ecosystems using two methods

In the first method, consumption is allocated into flaming (F), smouldering (S) and residual (R) smouldering phases using Consume v5.0 defaults (Prichard 2007). In the second method, emissions by fuel category are estimated by summing the individually calculated flaming, smouldering and residual emissions. Taken from Tables 4 and 5, south-eastern conifer EF values are F = 20.91, S = 29.35, fire-average = 25.29, and western conifer EF values are F = 13.5, S = 20.05, fire average = 14.32. Smouldering and residual smouldering EFs for coarse wood and duff are assumed to be 33 and 50 g kg⁻¹, after Urbanski (2014)

Fuel category	Proportion by combustion phase			South-eastern US			Western US		
	F	S	R	Consumption Mg ha ⁻¹	Individual emissions kg ha ⁻¹	Fire average emissions kg ha ⁻¹	Consumption Mg ha ⁻¹	Individual emissions kg ha ⁻¹	Fire average emissions kg ha ⁻¹
Herb	1.00	0.00	0.00	1.00	20.91	25.29	0.00	38.22	38.66
Shrub	1.00	0.00	0.00	3.60	75.28	91.04	0.00	42.59	42.96
1h	0.95	0.05	0.00	0.80	17.07	20.23	0.00	110.76	171.60
10 h	0.90	0.10	0.00	0.60	13.05	15.17	2.70	136.08	178.20
100 h	0.85	0.10	0.05	1.10	24.17	27.82	3.00	43.88	44.39
Sound 1000 h	0.60	0.30	0.10	0.10	2.47	3.30	5.20	561.83	605.00
Rotten 1000 h	0.60	0.30	0.10	0.00	0.00	0.00	5.40	38.22	38.66
Litter	0.90	0.10	0.00	6.90	150.10	174.50	3.10	42.59	42.96
Duff	0.10	0.70	0.20	1.30	61.79	65.00	12.10	110.76	171.60
TOTAL (±1 s.d.)				15.40	364.83	422.36 ± 246.40	31.50	933.35	1080.82 ± 160.65

characterise smoke from residual smouldering combustion of coarse wood and duff (Strand *et al.* 2012; Urbanski 2013). However, smoke from residual smouldering is generally not entrained in smoke plumes and does not affect regional air quality

(Ottmar 2014; Urbanski 2014). Greater understanding of the relative contribution of flaming and short-term smouldering for lofted emissions is also needed; this is difficult to determine without source characterisation of what is actually consumed.

In many past summaries of best available EFs, pollutant EFs were summarised by major vegetation and/or region. However, the lack of relationship between some pollutant EFs and fuel type is notable, and suggests that major vegetation or fuel type is not always an important factor in emissions production. Other studies have also demonstrated a similarity in pollutant EFs across distinctly different ecosystems and fuel complexes, and have suggested that fire environment (fuel moisture, wind, fire behaviour) may be much more relevant to emissions production than fuel type. For example, Yokelson *et al.* (2013) noted that pollutant EFs from markedly different ecosystems (semiarid shrublands and western coniferous canopy fuels) had similar pollutant emissions. In a recent study of western wildfire plumes, Liu *et al.* (2017) reported much higher PM_{2.5} EFs than those summarised for prescribed burning, and suggested that differences might be attributed to greater availability of coarse wood and organic soils for consumption during summer wildfire seasons (also see Urbanski 2013). May *et al.* (2014) found that dilution and fuel moisture influenced higher organic aerosol EF values in laboratory burns *v.* field measurements during the third Fire Laboratory At Missoula Experiment (FLAME-III) experiments. Finally, Chen *et al.* (2010) investigated the effects of fuel moisture on various fuel components (e.g. duff, litter, soil, leaves, stems) in laboratory burns and reported increased emissions factors of incompletely oxidised C and N species, such as carbon monoxide (CO) and ammonia (NH₃) with higher fuel moisture content.

Although substantial advances have been made with instrumentation and characterisation of pollutant emissions in wildland smoke, the science of predicting smoke production, dispersion and chemistry is still rapidly developing. To date, providing best-estimate EFs for smoke management and emissions inventories is challenged by data gaps, inconsistencies in measurement platforms and uncertainty in how EFs should be summarised by vegetation, fuel type and other explanatory variables. In compiling and reviewing the EF literature for SERA, we were challenged by the inconsistencies and redundant reporting of many emissions studies. Many potential records either could not be used to generate best-estimate EFs or required careful parsing to resolve inconsistencies. These included: (1) inconsistency in EF units including g kg⁻¹, lb ton⁻¹, mol kg⁻¹, ppm, g kg⁻¹-C and emission ratios with a trace gas such as CO₂ or CO; (2) considerable variability in source characterisation from cursory descriptions of broad vegetation types (e.g. from airborne plume sampling of active wildfires) and detailed descriptions of vegetation and fuels, including moisture content and composition (e.g. some laboratory burns); (3) redundancy in reported datasets across papers (e.g. Hegg *et al.* (1990); Radke *et al.* (1990, 1991); Laursen *et al.* (1992)); (4) applicability of reported EFs from experimental studies that varied fuel moistures or immersed fuels in water to improve extractions of target pollutants (e.g. Chen *et al.* (2010); Grandesso *et al.* (2011)); and (5) differences in reported EFs due to instrumentation (e.g. Aurell and Gullett 2013).

To facilitate adding future studies to SERA, we recommend reporting EFs in g kg⁻¹ of dry weight biomass consumed, in addition to location, major vegetation type, fuel category, MCE, measurement platform, fuel moisture content and burn type. Additional fields that would be optimal to report are listed

in Tables 1 and 2. We also recommend reporting EF data for each burn experiment, rather than averages of multiple experiments.

Tables 4–6 present a synthesis of available EF data split by vegetation types and combustion phase, with the goal of updating the information employed in land manager tools such as Consume, FOFEM and the BlueSky smoke modelling framework. To compare our values with those previously used for emissions predictions and reported in other EF reviews, Table 8 presents comparisons of SERA summary EF values with those reported in Andreae and Merlet (2001), the 2001 Smoke Management Guide (Hardy *et al.* 2001), Yokelson *et al.* (2013) and Urbanski (2014). Across forested systems, the SERA-based analysis yielded a somewhat higher PM_{2.5} EF of 25.29 g kg⁻¹ for SE conifer than did the other studies, which ranged from 6 to 17 g kg⁻¹. This is due to higher reported EF values from Aurell and Gullett (2013), Strand *et al.* (2012) and Robertson *et al.* (2014). Otherwise, in general, the EFs range from a factor of 3 to an order of magnitude in variability across these references, with the SERA-based data falling within the bounds of previous summaries.

Although much emphasis is placed on selecting appropriate EFs for estimating pollutant emissions, it is important to place EFs in the context of their importance in estimating wildland fire emissions and dispersion. Application of best available EFs and their associated uncertainty is one of many steps in smoke prediction, including fuel characterisation, the amount and type of fuels consumed (e.g. fine wood *v.* coarse wood), the proportion of lofted smoke and factors influencing dispersion. In ensemble modelling of smoke production and dispersion, uncertainty around EF estimation is eclipsed by even larger uncertainty in fire area (factor of 4; Larkin *et al.* (2012, 2014)), fuel loading (factor of 2; Drury *et al.* (2014)), consumption estimation (23%; Drury *et al.* (2014)) and atmospheric transport (100% uncertainty; Garcia-Menendez *et al.* (2013)). The side panel calculation of emissions by fuel component and combustion phase shows a 20 to 50% difference in emissions estimation based on a single standard deviation of the EF estimate variability. This variability is easily on par with that of fuel consumption, but is still eclipsed by other factors such as the fire size estimate, fuel loading and atmospheric transport. EFs can vary from a factor of 5 to more than an order of magnitude (Figs 2 and 3). The side panel calculations illustrate that EF variability remains significant, but that having more detailed information regarding fuels and combustion constrains variability to levels less than other components in the emission calculation stream.

Conclusion

The present study revealed several important data gaps in EF records. Across all pollutants, more studies are needed for wildfires, residual smouldering and organic soils, and for specific vegetation types including grasslands, northern forests and broadleaf deciduous forests. Wildfires are a large wildland fire emission source regionally, and across the US account for ~16% of the total PM_{2.5} in the US EPA National Emissions Inventory (US Environmental Protection Agency 2018). The work of Liu *et al.* (2017) suggests that PM₁ EFs from wildfires are more than twice those of prescribed fires because wildfires

Table 8. Comparison of summary emission factors values (g kg^{-1}) in the Smoke Emissions Repository Application (SERA) and other published studies

SE, south-eastern US; SW, south-western US; W, western US; NW, north-western US. Cells with missing data have no data available

	Vegetation type	PM _{2.5}	CO ₂	CO	CH ₄	NH ₃	NO	NO ₂	NO _x	SO ₂
<i>SE conifer forests</i>										
SERA	SE conifer forest	25.29	1576	97.09	4.12	0.820	1.860	1.23	3.34	0.99
SERA	SE mixed forest	14.78	1651	84.19	2.67	2.130	3.690	1.14	2.18	0.7
Urbanski (2014)	SE conifer forest	12.58	1703	76.00	2.32	0.140	1.700			1.06
Yokelson <i>et al.</i> (2013)	Pine forest understorey	13.60	1668	72.20	3.01	0.499	0.880	2.68	2.55	1.06
Yokelson <i>et al.</i> (2013)	Conifer canopy	7.44	1670	85.30	3.27	0.936	1.740	1.01	2.4	1.06
<i>W conifer forests</i>										
SERA	W conifer forest	14.32	1630	104.01	5.05	0.900	1.790	1.38	1.41	1.38
Urbanski (2014)	SW conifer forest	14.40	1653	87.00	3.15	0.500	1.880			1.06
Urbanski (2014)	NW conifer forest	17.57	1598	105.00	4.86	1.530	2.060			1.06
Hardy <i>et al.</i> (2001)	Douglas fir/hemlock	10.90	1541	156.00	5.50					
Hardy <i>et al.</i> (2001)	Ponderosa/lodgepole pine	11.00	1601	89.00	4.10					
Hardy <i>et al.</i> (2001)	Mixed conifer	9.40	1583	100.50	6.40					
Hardy <i>et al.</i> (2001)	Juniper	9.35	1616	81.50	6.00					
Yokelson <i>et al.</i> (2013)*	Pine forest understorey	13.60	1668	72.20	3.01	0.499	0.880	2.68	2.55	1.06
Yokelson <i>et al.</i> (2013)*	Conifer canopy	7.44	1670	85.30	3.27	0.936	1.740	1.01	2.4	1.06
<i>Broadleaf/mixed forests</i>										
SERA	W mixed forest	6.83	1670	55.26	2.35	0.600	1.650	1.53	2.55	0.49
Hardy <i>et al.</i> (2001)	Hardwood forest	11.20	1536	128.00	6.60					
Andreae and Merlet (2001)	Extratropical forest	13.00	1569	107.00	4.70	1.400	3.000			1
<i>Shrublands</i>										
SERA	SE shrubland	12.03	1708	73.50	2.38	2.060	4.710	0.95	2.86	0.74
SERA	W shrubland	7.88	1588	97.03	2.16	1.600	2.420	0.79	3.33	0.73
Hardy <i>et al.</i> (2001)	Sagebrush	13.35	1563	103.00	5.95					
Hardy <i>et al.</i> (2001)	Chaparral	8.65	1629	77.00	2.85					
Yokelson <i>et al.</i> (2013)	SW shrubland	7.06	1674	73.80	3.69	1.500	0.771	2.58	2.18	0.681
Urbanski (2014)	General shrubland	7.06	1674	74.00	3.69	1.500	2.180			0.68
<i>Grasslands</i>										
SERA	SE grassland	12.08	1686	68.17	2.53	1.300	5.330		3.53	0.83
SERA	W grassland		1421	56.26	2.90	0.560	2.860	3.13		2.98
Urbanski (2014)	General grassland	8.51	1705	61.00	1.95	1.500	2.180			0.68
Andreae and Merlet (2001)	General cropland	5.40	1613	65.00	2.30	0.60–1.50	3.900			0.35

tend to burn more dead and/or down fuels and live, moist canopy fuels. Grasslands and other rangelands comprise a significant portion of the continental US, and historically have a frequent fire return interval. Fuel loadings can vary greatly depending on seasonality and drought conditions. Duff and organic soils can consume more than an order of magnitude more fuel per unit area than most other fuel types (Akagi *et al.* 2011), and can smoulder and emit for many months after ignition.

Finally, trace gases and aerosols such as non-methane organic compounds are underrepresented in the database. The detection of trace gas and aerosol species has been improving dramatically in recent years – newer studies are identifying more and more pollutant species (e.g. Akagi *et al.* 2011; Urbanski 2014; Liu *et al.* 2017; Koss *et al.* 2018), while other studies are investigating different measurement techniques for the same species (e.g. Aurell and Gullett 2013; Aurell *et al.* 2015; Koss *et al.* 2018). Accounting for this and assessing EF data in light of new technologies and measurement techniques may be necessary in the future. Furthermore, several factors greatly affect EF measurement, such as how these trace gases and aerosols react in the atmosphere (e.g. semi-volatile species condensing and

volatilising), effects of heat and dilution on the in-plume chemistry and also where measurements are made (e.g. close to the source, close to the ground, airborne, etc.). These are all items to consider when analysing EF data.

Conflicts of interest

The authors declare no conflicts of interest.

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Supplementary material for

Wildland fire emission factors in North America: synthesis of existing data, measurement needs and management applications

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Supplementary Appendices

Appendix 1. References used for emissions factor (EF) summaries in the Smoke Emissions Repository Application. The underlying database contains additional references compiled by Lincoln et al. (2014) and others that were not used in EF summaries, generally because they summarized other research and contained redundant values.

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Appendix 2: Analysis of variance tests of pollutant emissions factors (EF) categorized by combustion phase (CombPhase), regional vegetation type (regionalVegetationType) and fuel category (fuelCategory). Due to unequal sample sizes and data gaps, tests were not possible for every category by pollutant. Tables 1 and 3 provide definitions of combustion phase, regional vegetation type and fuel category codes.

Carbon dioxide (CO₂)

Combustion Phase
Record Count by Phase

Phase	Freq
F	286
R	20
S	127

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
1005	1567	1641	1613	1704.98	1932.8

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	3564305	1782152.43	136.943	0
Residuals	430	5595950	13013.84	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	-342.124	-404.179	-280.069	0
S-F	-146.196	-174.805	-117.587	0
S-R	195.928	131.384	260.472	0

Regional Vegetation Type Type
ANOVA of Regional Vegetation Type

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	4	310605.2	77651.31	2.728	0.029

Residuals	477	13575207.3	28459.55	NA	NA
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Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADL.p.value
SE pine-N conifer	-68.241	-215.613	79.130	0.711
SE shrub-N conifer	80.059	-103.312	263.430	0.754
W conifer-N conifer	-35.602	-177.765	106.561	0.960
W shrub-N conifer	-37.650	-184.319	109.019	0.956
SE shrub-SE pine	148.301	19.672	276.929	0.015
W conifer-SE pine	32.639	-23.314	88.593	0.500
W shrub-SE pine	30.591	-35.983	97.166	0.717
W conifer-SE shrub	-115.661	-238.288	6.965	0.075
W shrub-SE shrub	-117.709	-245.532	10.114	0.088
W shrub-W conifer	-2.048	-56.123	52.027	1.000

Fuel Category

ANOVA of Fuel Category

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	7	1277429	182489.87	6.577	0
Residuals	540	14983736	27747.66	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADL.p.value
Grass-Duff	191.464	35.436	347.491	0.005
MC-Duff	205.059	69.439	340.678	0.000
Mixedwood-Duff	208.000	42.993	373.007	0.003
Pine_needles-Duff	91.752	-59.908	243.412	0.592
Pine_under-Duff	240.769	99.095	382.442	0.000
Shrub-Duff	178.557	43.913	313.201	0.002
Slash-Duff	145.914	12.296	279.532	0.021
MC-Grass	13.595	-89.474	116.663	1.000
Mixedwood-Grass	16.537	-122.954	156.027	1.000
Pine_needles-Grass	-99.712	-223.127	23.703	0.216
Pine_under-Grass	49.305	-61.608	160.219	0.878
Shrub-Grass	-12.907	-114.688	88.875	1.000
Slash-Grass	-45.549	-145.969	54.870	0.866
Mixedwood-MC	2.942	-113.272	119.156	1.000
Pine_needles-MC	-113.307	-209.635	-16.978	0.009

Pine_under-MC	35.710	-43.976	115.396	0.873
Shrub-MC	-26.501	-92.890	39.887	0.927
Slash-MC	-59.144	-123.426	5.138	0.097
Pine_needles-Mixedwood	-116.249	-250.836	18.339	0.148
Pine_under-Mixedwood	32.769	-90.457	155.994	0.993
Shrub-Mixedwood	-29.443	-144.517	85.631	0.994
Slash-Mixedwood	-62.086	-175.958	51.786	0.714
Pine_under-Pine_needles	149.017	44.337	253.697	0.000
Shrub-Pine_needles	86.806	-8.144	181.755	0.102
Slash-Pine_needles	54.163	-39.327	147.652	0.646
Shrub-Pine_under	-62.212	-140.225	15.802	0.231
Slash-Pine_under	-94.855	-171.084	-18.626	0.004
Slash-Shrub	-32.643	-94.840	29.554	0.752

Carbon monoxide (CO)

Combustion Phase

Record Count by Phase

x	Freq
F	295
R	29
S	138

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
2.8	59.25	82.38	95.36	122.69	358

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	775559.8	387779.922	300.423	0
Residuals	459	592467.4	1290.779	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	135.664	119.224	152.104	0
S-F	66.845	58.132	75.557	0
S-R	-68.820	-86.076	-51.563	0

Regional Vegetation Type*ANOVA of Regional Vegetation Type*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	5	69363.67	13872.73	4.358	0.001
Residuals	547	1741068.28	3182.94	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
SE grass-N conifer	-18.560	-81.755	44.635	0.960
SE pine-N conifer	17.097	-33.824	68.018	0.930
SE shrub-N conifer	-13.232	-76.427	49.963	0.991
W conifer-N conifer	26.594	-23.045	76.234	0.644
W shrub-N conifer	9.595	-41.099	60.290	0.994
SE pine-SE grass	35.657	-7.394	78.708	0.169
SE shrub-SE grass	5.328	-51.717	62.372	1.000
W conifer-SE grass	45.154	3.627	86.682	0.024
W shrub-SE grass	28.155	-14.628	70.939	0.414
SE shrub-SE pine	-30.329	-73.380	12.722	0.335
W conifer-SE pine	9.498	-8.499	27.494	0.658
W shrub-SE pine	-7.501	-28.232	13.229	0.906
W conifer-SE shrub	39.827	-1.701	81.354	0.069
W shrub-SE shrub	22.828	-19.956	65.611	0.648
W shrub-W conifer	-16.999	-34.345	0.347	0.059

Fuel Category*ANOVA of Fuel Category*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	6	153721.5	25620.256	8.212	0
Residuals	577	1800081.9	3119.726	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
SE grass-N conifer	-18.560	-81.755	44.635	0.960
SE pine-N conifer	17.097	-33.824	68.018	0.930
SE shrub-N conifer	-13.232	-76.427	49.963	0.991
W conifer-N conifer	26.594	-23.045	76.234	0.644
W shrub-N conifer	9.595	-41.099	60.290	0.994
SE pine-SE grass	35.657	-7.394	78.708	0.169

SE shrub-SE grass	5.328	-51.717	62.372	1.000
W conifer-SE grass	45.154	3.627	86.682	0.024
W shrub-SE grass	28.155	-14.628	70.939	0.414
SE shrub-SE pine	-30.329	-73.380	12.722	0.335
W conifer-SE pine	9.498	-8.499	27.494	0.658
W shrub-SE pine	-7.501	-28.232	13.229	0.906
W conifer-SE shrub	39.827	-1.701	81.354	0.069
W shrub-SE shrub	22.828	-19.956	65.611	0.648
W shrub-W conifer	-16.999	-34.345	0.347	0.059

Methane (CH4)

Combustion Phase

Record Count by Phase

x	Freq
F	249
R	20
S	116

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
0	1.94	3.24	4.51	6.66	24.1

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	2719.959	1359.979	240.955	0
Residuals	382	2156.051	5.644	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	7.409	6.110	8.709	0
S-F	5.136	4.507	5.764	0
S-R	-2.274	-3.627	-0.921	0

Regional Vegetation Type Type

ANOVA of Regional Vegetation Type

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	6	611.234	101.872	9.249	0
Residuals	323	3557.755	11.015	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
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SE grass-N conifer	0.132	-4.539	4.803	1.000
SE pine-N conifer	2.441	-1.294	6.176	0.456
SE shrub-N conifer	-0.019	-4.514	4.476	1.000
W conifer-N conifer	2.822	-0.728	6.372	0.220
W duff-N conifer	7.733	2.120	13.347	0.001
W shrub-N conifer	-0.203	-4.009	3.603	1.000
SE pine-SE grass	2.309	-1.086	5.704	0.405
SE shrub-SE grass	-0.151	-4.367	4.066	1.000
W conifer-SE grass	2.690	-0.500	5.881	0.162
W duff-SE grass	7.602	2.208	12.995	0.001
W shrub-SE grass	-0.334	-3.807	3.139	1.000
SE shrub-SE pine	-2.460	-5.608	0.688	0.238
W conifer-SE pine	0.381	-1.139	1.902	0.990
W duff-SE pine	5.292	0.686	9.899	0.013
W shrub-SE pine	-2.644	-4.692	-0.596	0.003
W conifer-SE shrub	2.841	-0.085	5.767	0.063
W duff-SE shrub	7.752	2.511	12.994	0.000
W shrub-SE shrub	-0.184	-3.416	3.048	1.000
W duff-W conifer	4.911	0.453	9.369	0.020
W shrub-W conifer	-3.025	-4.712	-1.337	0.000
W shrub-W duff	-7.936	-12.601	-3.271	0.000

Fuel Category

ANOVA of Fuel Category

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	8	1109.206	138.651	13.726	0
Residuals	366	3697.066	10.101	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
Duff-CWD	-2.346	-7.165	2.474	0.846
Grass-CWD	-8.974	-13.094	-4.853	0.000
MC-CWD	-5.788	-9.421	-2.155	0.000
Mixedwood-CWD	-7.399	-11.494	-3.305	0.000
Pine_shrub-CWD	-9.293	-14.112	-4.473	0.000
Pine_under-CWD	-8.367	-12.108	-4.627	0.000
Shrub-CWD	-7.939	-11.665	-4.214	0.000

Slash-CWD	-5.874	-9.558	-2.190	0.000
Grass-Duff	-6.628	-10.579	-2.677	0.000
MC-Duff	-3.442	-6.882	-0.003	0.050
Mixedwood-Duff	-5.054	-8.978	-1.130	0.002
Pine_shrub-Duff	-6.947	-11.622	-2.272	0.000
Pine_under-Duff	-6.022	-9.575	-2.468	0.000
Shrub-Duff	-5.594	-9.132	-2.056	0.000
Slash-Duff	-3.528	-7.022	-0.035	0.046
MC-Grass	3.186	0.822	5.549	0.001
Mixedwood-Grass	1.574	-1.452	4.600	0.792
Pine_shrub-Grass	-0.319	-4.270	3.632	1.000
Pine_under-Grass	0.606	-1.919	3.132	0.998
Shrub-Grass	1.034	-1.470	3.538	0.934
Slash-Grass	3.099	0.658	5.541	0.003
Mixedwood-MC	-1.612	-3.930	0.706	0.428
Pine_shrub-MC	-3.505	-6.944	-0.065	0.042
Pine_under-MC	-2.579	-4.191	-0.967	0.000
Shrub-MC	-2.152	-3.729	-0.574	0.001
Slash-MC	-0.086	-1.563	1.390	1.000
Pine_shrub-Mixedwood	-1.893	-5.817	2.031	0.853
Pine_under-Mixedwood	-0.968	-3.451	1.516	0.953
Shrub-Mixedwood	-0.540	-3.001	1.921	0.999
Slash-Mixedwood	1.525	-0.872	3.923	0.555
Pine_under-Pine_shrub	0.926	-2.628	4.479	0.996
Shrub-Pine_shrub	1.353	-2.185	4.891	0.958
Slash-Pine_shrub	3.419	-0.075	6.912	0.061
Shrub-Pine_under	0.428	-1.384	2.239	0.998
Slash-Pine_under	2.493	0.769	4.217	0.000
Slash-Shrub	2.065	0.373	3.758	0.005

Nitrogen oxides (NO_x)

Combustion Phase

Record Count by Phase

<u>x</u>	<u>Freq</u>
F	63
R	11
S	17

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
0.03	1.66	2.67	2.69	3.25	7.6

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	3.291	1.646	0.816	0.446
Residuals	88	177.519	2.017	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	-0.260	-1.367	0.846	0.841
S-F	-0.477	-1.403	0.448	0.439
S-R	-0.217	-1.527	1.093	0.918

Regional Vegetation Type

ANOVA of Regional Vegetation Type

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	5	7.825	1.565	0.746	0.592
Residuals	76	159.428	2.098	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
SE grass-N conifer	0.633	-3.232	4.499	0.997
SE pine-N conifer	-0.595	-3.729	2.538	0.994
SE shrub-N conifer	-0.044	-3.711	3.623	1.000
W conifer-N conifer	-0.309	-3.589	2.971	1.000
W shrub-N conifer	0.037	-3.027	3.102	1.000
SE pine-SE grass	-1.229	-3.842	1.385	0.742
SE shrub-SE grass	-0.677	-3.911	2.557	0.990
W conifer-SE grass	-0.943	-3.730	1.845	0.920
W shrub-SE grass	-0.596	-3.127	1.935	0.983
SE shrub-SE pine	0.552	-1.759	2.862	0.982
W conifer-SE pine	0.286	-1.341	1.913	0.995
W shrub-SE pine	0.633	-0.499	1.764	0.579
W conifer-SE shrub	-0.265	-2.771	2.240	1.000
W shrub-SE shrub	0.081	-2.135	2.297	1.000
W shrub-W conifer	0.347	-1.143	1.837	0.984

Fuel Category*ANOVA of Fuel Category*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	3	76.818	25.606	7.548	0
Residuals	91	308.709	3.392	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
Pine_under-Duff	1.025	-1.085	3.136	0.583
Shrub-Duff	2.726	0.900	4.552	0.001
Slash-Duff	1.655	-0.393	3.703	0.156
Shrub-Pine_under	1.701	0.294	3.108	0.011
Slash-Pine_under	0.630	-1.056	2.315	0.762
Slash-Shrub	-1.071	-2.383	0.241	0.149

*Particulate matter (PM)***Combustion Phase***Record Count by Phase*

x	Freq
F	44
R	15
S	40

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
2.1	7.15	13.8	14.94	19.65	58.3

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	3025.839	1512.920	20.399	0
Residuals	96	7119.898	74.166	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	13.964	7.834	20.093	0.000
S-F	9.503	5.024	13.982	0.000
S-R	-4.461	-10.668	1.746	0.206

Regional Vegetation Type*ANOVA of Regional Vegetation Type*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	4	9210.836	2302.709	11.368	0
Residuals	242	49021.466	202.568	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
SE pine-SE grass	9.832	-29.595	49.259	0.959
SE shrub-SE grass	3.088	-38.403	44.578	1.000
W conifer-SE grass	4.336	-34.937	43.608	0.998
W shrub-SE grass	20.134	-19.381	59.649	0.628
SE shrub-SE pine	-6.744	-21.426	7.938	0.714
W conifer-SE pine	-5.496	-11.532	0.540	0.093
W shrub-SE pine	10.302	2.851	17.753	0.002
W conifer-SE shrub	1.248	-13.014	15.511	0.999
W shrub-SE shrub	17.046	2.130	31.963	0.016
W shrub-W conifer	15.798	9.212	22.384	0.000

Fuel Category*ANOVA of Fuel Category*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	5	8941.545	1788.309	8.822	0
Residuals	263	53311.565	202.706	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
Mixedwood-Grass	18.800	-14.573	52.173	0.588
Pine_needles-Grass	4.884	-13.346	23.114	0.972
Pine_under-Grass	13.769	-4.292	31.829	0.247
Shrub-Grass	18.040	0.482	35.598	0.040
Slash-Grass	4.861	-12.182	21.904	0.964
Pine_needles-Mixedwood	-13.916	-43.736	15.904	0.763
Pine_under-Mixedwood	-5.031	-34.748	24.685	0.997
Shrub-Mixedwood	-0.760	-30.173	28.654	1.000
Slash-Mixedwood	-13.939	-43.048	15.170	0.742
Pine_under-Pine_needles	8.885	-1.196	18.966	0.119

Shrub-Pine_needles	13.156	4.006	22.306	0.001
Slash-Pine_needles	-0.023	-8.141	8.096	1.000
Shrub-Pine_under	4.272	-4.536	13.079	0.732
Slash-Pine_under	-8.907	-16.637	-1.177	0.014
Slash-Shrub	-13.179	-19.648	-6.710	0.000

Particulate matter less than 2.5 microns (PM_{2.5})

Combustion Phase

Record Count by Phase

x	Freq
F	176
R	15
S	49

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
1.1	8.18	13.08	17.63	22.44	89.95

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	7955.672	3977.836	21.877	0
Residuals	237	43093.974	181.831	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	19.95	11.396	28.505	0.000
S-F	9.50	4.363	14.637	0.000
S-R	-10.45	-19.835	-1.065	0.025

Regional Vegetation Type

ANOVA of Regional Vegetation Type

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	4	9727.78	2431.945	9.73	0
Residuals	275	68730.90	249.931	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
SE pine-SE grass	9.449	-2.384	21.282	0.185
SE shrub-SE grass	0.940	-16.783	18.663	1.000

W conifer-SE grass	-4.214	-16.417	7.988	0.878
W shrub-SE grass	4.948	-8.070	17.966	0.835
SE shrub-SE pine	-8.509	-22.752	5.733	0.473
W conifer-SE pine	-13.664	-19.800	-7.527	0.000
W shrub-SE pine	-4.501	-12.131	3.128	0.486
W conifer-SE shrub	-5.154	-19.705	9.396	0.867
W shrub-SE shrub	4.008	-11.233	19.249	0.951
W shrub-W conifer	9.162	0.971	17.353	0.020

Fuel Category

ANOVA of Fuel Category

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	6	12412.49	2068.748	8.974	0
Residuals	301	69386.90	230.521	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
Grass-Duff	-7.296	-26.811	12.220	0.925
MC-Duff	-5.768	-28.302	16.767	0.988
Mixedwood-Duff	-9.839	-29.814	10.136	0.767
Pine_under-Duff	-0.919	-17.467	15.628	1.000
Shrub-Duff	3.961	-12.659	20.581	0.992
Slash-Duff	-12.690	-29.523	4.143	0.279
MC-Grass	1.528	-17.987	21.043	1.000
Mixedwood-Grass	-2.543	-19.037	13.950	0.999
Pine_under-Grass	6.376	-5.742	18.495	0.707
Shrub-Grass	11.257	-0.961	23.474	0.093
Slash-Grass	-5.394	-17.900	7.111	0.861
Mixedwood-MC	-4.071	-24.046	15.904	0.997
Pine_under-MC	4.848	-11.699	21.396	0.977
Shrub-MC	9.729	-6.891	26.349	0.591
Slash-MC	-6.922	-23.755	9.911	0.886
Pine_under-Mixedwood	8.920	-3.926	21.765	0.379
Shrub-Mixedwood	13.800	0.861	26.739	0.028
Slash-Mixedwood	-2.851	-16.062	10.360	0.995
Shrub-Pine_under	4.880	-1.619	11.379	0.283
Slash-Pine_under	-11.771	-18.796	-4.746	0.000

Slash-Shrub	-16.651	-23.845	-9.457	0.000
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Sulphur dioxide (SO2)

Combustion Phase

Record Count by Phase

x	Freq
F	106
R	1
S	18

Summary Statistics

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
0	0.64	1.02	1.12	1.42	3.42

ANOVA of Combustion Phase

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
CombPhase	2	5.691	2.846	5.929	0.003
Residuals	122	58.555	0.480	NA	NA

Tukey Combustion Phase

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
R-F	1.640	-0.011	3.292	0.052
S-F	0.458	0.039	0.877	0.028
S-R	-1.182	-2.871	0.507	0.225

Regional Vegetation Type

ANOVA of Regional Vegetation Type

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
regionalVegetationType	2	8.189	4.095	15.374	0
Residuals	88	23.437	0.266	NA	NA

Tukey Regional Vegetation Type

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
W conifer-SE pine	0.401	-0.024	0.825	0.068
W shrub-SE pine	-0.257	-0.710	0.196	0.371
W shrub-W conifer	-0.658	-0.945	-0.370	0.000

Fuel Category*ANOVA of Fuel Category*

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
fuelCategory	4	10.129	2.532	6.059	0
Residuals	102	42.626	0.418	NA	NA

Tukey Fuel Category

Comparison	Estimate	Conf.low	Conf.high	ADJ.p.value
Grass-Duff	-0.465	-1.435	0.504	0.672
MC-Duff	-0.393	-1.090	0.304	0.522
Pine_under-Duff	-0.575	-1.321	0.171	0.212
Shrub-Duff	-1.007	-1.714	-0.299	0.001
MC-Grass	0.072	-0.715	0.860	0.999
Pine_under-Grass	-0.109	-0.940	0.722	0.996
Shrub-Grass	-0.542	-1.338	0.255	0.331
Pine_under-MC	-0.182	-0.668	0.304	0.837
Shrub-MC	-0.614	-1.038	-0.189	0.001
Shrub-Pine_under	-0.432	-0.933	0.069	0.125

- 1 **Appendix 3:** Supported air pollutants in SERA and EF summaries, including mean, standard deviation (SD), minimum (min) and
2 maximum (max) values in g/kg, excluding outliers. Pollutant Category include: critical air pollutant (CAP), greenhouse gas (GHG),
3 hazardous air pollutant (HAX), toxic air pollutant (TOX). RefID lists source references for each pollutant.

Pollutant	Formula/ Acronym	Molecular weight	N	EF (g/kg)	SD	Min	Max	Category	Ref ID
1,1,2,2-tetrachloroethane	C2H2Cl4	167.838	1	0.01		0.01	0.01	HAP, TOX	236
1,1-dichloro-1-fluoroethane	CH3CClF2	116.944	2	3.25	2.62	1.40	5.10	TOX	230
1,1-difluoroethane	C2H4F2	66.051	2	1.75	0.92	1.10	2.40		230
1,2,3-trimethylbenzene	C9H12	120.195	7	0.04	0.03	0.02	0.09		231, 234
1,2,4-trimethylbenzene	C9H12	120.195	22	0.02	0.03	0.00	0.11	TOX	231, 234, 236, 238
1,2-dichloroethane	C2H4Cl2	98.954	2	3.10	2.83	1.10	5.10	HAP, TOX	230
1,3,5-trimethylbenzene	C9H12	120.195	18	0.01	0.01	0.00	0.05		231, 236, 238
1,3-butadiene	C4H6	54.092	59	0.17	0.15	0.00	0.74	HAP, TOX	171, 230, 231, 234, 236, 238, 243
1,3-dimethylnaphthalene	C12H12	156.228	58	0.02	0.02	0.00	0.13		242
1,4-dichlorobenzene	C6H4Cl2	146.998	1	0.00		0.00	0.00	HAP, TOX	236
1-butyne	C4H6	54.092	7	0.00	0.00	0.00	0.01	HAP, TOX	231
1-chloro-1,1-difluoroethane	C2H3ClF2	100.493	2	2.60	1.84	1.30	3.90	TOX	230
1-heptene	C7H14	98.189	7	0.04	0.03	0.01	0.10		231
1-octene	C8H16	112.216	7	0.04	0.04	0.01	0.12		231
1-pentene	C5H10	70.135	9	0.04	0.03	0.01	0.12		230, 231
2 cyclopenten 1-one	C5H6O	82.102	16	0.13	0.12	0.00	0.52		105, 231
2-(3h)furanone	C4H4O2	84.074	58	0.39	0.30	0.05	1.36		242
2(5h)-furanone	C4H4O2	84.074	7	0.10	0.06	0.01	0.22		105
2,3-dimethylbutane	C6H14	86.178	5	1.28	2.16	0.00	5.00		230, 231
2,5-dimethylfuran	C6H8O	96.129	7	0.10	0.14	0.02	0.41		231

2+3-methylpentane	C6H14		3	4.07	4.09	0.02	8.20		230, 234
2-acetylfuran	C6H6O2	110.112	8	0.04	0.02	0.01	0.06		105
2-butanone	C4H8O	72.107	22	0.10	0.06	0.01	0.29	TOX	231, 234, 236, 238
2-butyl nitrate	C4H9NO3	119.12	2	2.25	0.49	1.90	2.60		230
								HAP,	
2-butyne	C4H6	54.092	7	0.00	0.00	0.00	0.01	TOX	231
2-ethenyl benzofuran	C10H8O	144.173	57	0.02	0.01	0.00	0.06		242
2-ethylfuran	C6H8O	96.129	7	0.02	0.02	0.01	0.07		231
2-ethyltoluene	C9H12	120.195	7	0.01	0.01	0.00	0.03		231
2-furmethanol	C5H6O2	98.101	5	0.26	0.36	0.01	0.87		105
2-hexanone		100.161	10	0.00	0.00	0.00	0.01		236
2-hydroxy-2H-furan-5-one	C4H4O3	100.073	58	0.05	0.05	0.00	0.29		242
2-methyl-1-butene	C5H10	70.135	6	0.02	0.01	0.01	0.03		231
2-methyl-2-butene	C5H10	70.135	6	0.02	0.01	0.01	0.04		231
2-methylpentane	C6H14	86.178	6	0.01	0.00	0.00	0.01		231
2-methylpropanal	C4H8O	72.107	6	0.02	0.01	0.01	0.04	TOX	231
2-pentyl nitrate	C5H11NO3	133.147	2	5.15	0.07	5.10	5.20		230
2-propanol	C3H8O	60.096	3	0.00	0.00	0.00	0.00		236
3-buten-1-amine	C4H9N	71.123	56	0.00	0.00	0.00	0.01		242
3-ethyltoluene	C9H12	120.195	8	0.07	0.10	0.01	0.29		231, 234
3-furaldehyde	C5H4O2	96.085	10	0.81	0.62	0.01	1.80		105
3-methyl-1-butene	C5H10	70.135	6	0.01	0.00	0.01	0.02		231
3-methyl-2-butanone	C5H10O	86.134	7	0.03	0.04	0.01	0.12		231
3-methyl-2-butyl nitrate	C5H11NO3	133.147	2	3.90	3.96	1.10	6.70		230
3-methyl-3-buten-2-one	C5H8O	84.118	7	0.12	0.18	0.02	0.51		231
3-methylacetophenone	C9H10O	134.178	58	0.04	0.03	0.00	0.13		242
3-methylbutanal	C5H10O	86.134	6	0.03	0.01	0.02	0.05		231
3-methylpentane	C6H14	86.178	7	0.00	0.00	0.00	0.01		231
3-pentyl nitrate	C5H11NO3	133.147	2	4.40	0.00	4.40	4.40		230
4-carene	C10H16	136.238	6	0.05	0.07	0.01	0.17		231
4-ethyltoluene	C9H12	120.195	15	0.01	0.00	0.00	0.02		231, 236

4-methyl-2-pentanone	C6H12O	100.161	10	0.00	0.00	0.00	0.00	HAP, TOX	236
4-pyridone	C5H5NO	95.101	56	0.01	0.01	0.00	0.05		242
5-(hydroxymethyl)-2-furfural	C6H6O3	126.111	57	0.30	0.26	0.02	1.19		242
5-hydroxymethyl-2[3h]-furanone	C5H6O3	114.1	58	0.20	0.18	0.02	0.78		242
5-methyl-2-furaldehyde	C6H6O2	110.112	10	0.18	0.14	0.01	0.50		105
acenaphthene	C12H10	154.212	3	0.00	0.00	0.00	0.00	HAP, TOX	238
acenaphthylene	C12H8	152.196	60	0.01	0.01	0.00	0.04	HAP, TOX	238, 242
acetaldehyde	C2H4O	44.053	65	0.93	0.74	0.13	3.65	HAP, TOX	231, 242
acetamide	C2H5NO	59.068	55	0.04	0.05	0.00	0.26	HAP, TOX	242 105, 117, 118, 120, 133, 170, 171, 231, 243
acetic acid	C2H4O2	60.052	152	2.06	1.85	0.19	9.28		242
acetic anhydride	C4H6O3	102.089	58	0.09	0.07	0.01	0.39		105, 230, 231, 243
acetol	C3H6O2	74.079	84	0.79	0.60	0.00	2.63		242
acetone	C3H6O	58.08	57	0.36	0.27	0.05	1.05		230, 231, 234, 236, 238, 242
acetonitrile	C2H3N	41.053	76	0.09	0.08	0.00	0.39	HAP, TOX	63, 117, 118, 120, 137, 153, 170, 171, 172, 179, 230, 231, 234, 242, 243
acetylene	C2H2	26.038	282	0.65	0.70	0.04	4.10		
acrolein	C3H4O	56.064	80	0.70	0.49	0.07	2.49	HAP, TOX	231, 236, 238, 242
acrylonitrile	C3H3N	53.064	80	0.03	0.02	0.00	0.09	HAP, TOX	231, 236, 238, 242
allylamine	C3H7N	57.096	56	0.00	0.00	0.00	0.01		242

alpha-pinene	C10H16	136.238	22	0.05	0.03	0.00	0.12		230, 231, 234, 236, 238 63, 117, 118, 120, 133, 153, 170, 171, 172, 180, 231, 237, 243
ammonia	NH3	17.031	210	1.34	1.43	0.02	6.60	CAP, TOX	172, 180, 230
ammonium	NH4+	18.039	62	0.08	0.11	0.00	0.51		
anthracene	C14H10	178.234	3	0.00	0.00	0.00	0.00	HAP, TOX	238
benzaldehyde	C7H6O	106.124	58	0.08	0.06	0.01	0.27		242
benzene	C6H6	78.114	83	0.36	0.26	0.05	1.47	HAP, TOX	230, 231, 234, 236, 238, 242
benzo[a]anthracene	C18H12	228.294	3	0.00	0.00	0.00	0.00	HAP, TOX	238
benzo[a]pyrene	C20H12	252.316	11	0.00	0.00	0.00	0.00	HAP, TOX	178, 238
benzo[b]fluoranthene	C20H12	252.316	3	0.00	0.00	0.00	0.00	HAP, TOX	238
benzo[ghi]perylene	C22H12	276.338	3	0.00	0.00	0.00	0.00	HAP, TOX	238
benzo[k]fluoranthene	C20H12	252.316	3	0.00	0.00	0.00	0.00	HAP, TOX	238
benzofuran	C8H6O	118.135	58	0.04	0.03	0.01	0.17		242
benzoic acid	C7H6O2	122.123	3	0.08	0.07	0.02	0.16		105
benzonitrile	C7H5N	103.124	62	0.02	0.02	0.00	0.10		105, 242
beta-pinene	C10H16	136.238	10	0.19	0.27	0.01	0.72		230, 231, 234
black carbon	BC		114	0.90	0.76	0.00	3.22		236, 238, 243
brown carbon	BrC		19	1.13	0.40	0.66	2.10		236, 238
butadiene hydroxynitrates	C4O4H7N		2	0.01	0.00	0.01	0.02		230
butane	C4H10	58.124	19	0.07	0.06	0.02	0.20		63, 137, 230, 231, 234
butene hydroxynitrates	C4O4H9N		2	0.03	0.01	0.02	0.03		230

butyl acetate	C6H12O2	116.16	1	0.00		0.00	0.00		236
butyraldehyde	C4H8O	72.107	7	0.04	0.03	0.02	0.09	TOX	231
butyric acid	C4H8O2	88.106	5	0.07	0.04	0.04	0.14		105
C4-dihydroxycarbonyls	C4O3H8		2	0.05	0.04	0.02	0.08		230
C4-hydroxydicarbonyls/C5-alkenediols	C4O3H6/C5O2H10		2	0.11	0.02	0.09	0.12		230
camphene	C10H16	136.238	4	0.28	0.32	0.01	0.66		231
									4, 35, 42, 46, 63, 66, 86, 94, 100, 102, 117, 118, 120, 129, 133, 137, 140, 153, 157, 170, 171, 172, 179, 195, 201, 230, 231, 234, 237, 238, 239, 243, 244, 245
carbon dioxide	CO2	44.009	578	1586.57	171.06	828.50	1968.00	GHG HAP,	
carbon disulfide	CS2	76.131	6	0.00	0.01	0.00	0.02	TOX	236
									4, 35, 36, 37, 42, 46, 63, 66, 86, 94, 100, 102, 117, 118, 120, 129, 133, 137, 140, 153, 157, 170, 171, 172, 178, 179, 180, 195, 201, 230, 231, 234, 237, 238, 239, 243, 244, 245
carbon monoxide	CO	28.01	640	103.51	58.30	2.80	359.00	CAP	
carbon suboxide	C3O2	68.031	58	0.00	0.00	0.00	0.01		242

carbon tetrachloride	CCl4	153.811	2	0.00	0.00	0.00	0.00	GHG, HAP, TOX	236
carbonyl sulfide	COS	60.07	12	0.15	0.18	0.01	0.51	HAP, TOX	36, 230, 231, 234
chloride	Cl-	35.45	35	0.71	1.21	0.03	5.39		172, 230
chloroethane	C2H5Cl	64.512	2	0.00	0.00	0.00	0.01	HAP, TOX	236
chrysene	C18H12	228.294	3	0.00	0.00	0.00	0.00	HAP, TOX	238
cis-2-butene	C4H8	56.108	9	0.05	0.05	0.01	0.17		230, 231, 234
cis-2-pentene	C5H10	70.135	7	0.01	0.01	0.01	0.04		231
creosol	C8H10O2	138.166	66	0.27	0.26	0.01	1.36		105, 242
crotonic acid	C4H6O2	86.09	5	0.17	0.29	0.03	0.68		105
cumene	C9H12	120.195	15	0.00	0.00	0.00	0.01	HAP, TOX	231, 236, 238
cyclohexane	C6H12	84.162	3	0.00	0.00	0.00	0.00	TOX	236
decanal	C10H20O	156.269	57	0.00	0.00	0.00	0.01		242
diacetyl	C4H6O2	86.09	1	2.10		2.10	2.10		230
dibenz[a,h]anthracene	C22H14	278.354	2	0.00	0.00	0.00	0.00	HAP, TOX	238
dibromomethane	CH2Br2	173.835	2	1.80	0.28	1.60	2.00	TOX	230
dichlorodifluoromethane	CCl2F2	120.908	9	0.00	0.00	0.00	0.01		63, 137, 236
dichloromethane	CH2Cl2	84.927	8	1.05	2.30	0.00	6.50	GHG, HAP, TOX	230, 236
dihydronaphthalene	C10H10	130.19	58	0.03	0.03	0.00	0.17		242
dimethyl disulfide	C2H6S2	94.19	57	0.00	0.00	0.00	0.01		242
dimethyl sulfide	C2H6S	62.13	64	0.00	0.00	0.00	0.02		230, 231, 242
dimethyl trisulfide	C2H6S3	126.25	55	0.00	0.00	0.00	0.00		242
dimethylbenzofuran	C10H10O	146.189	58	0.04	0.03	0.00	0.16		242
d-limonene	C10H16	136.238	22	0.73	1.36	0.01	5.36		231, 236, 238

elemental carbon	EC		54	1.42	1.79	0.00	8.01		172, 180, 238
ethanal nitrate	C2O4H3N		2	2.75	0.21	2.60	2.90		230
									63, 102, 120, 137,
ethane	C2H6	30.07	72	0.39	0.19	0.07	1.03		153, 179, 230,
									231, 234
ethanol	C2H6O	46.069	76	0.04	0.04	0.00	0.23		231, 234, 236,
ethenamine	C2H5N	43.069	58	0.01	0.01	0.00	0.04		238, 242
									242
									102, 117, 118,
									120, 133, 170,
									171, 179, 230,
ethene	C2H4	28.054	258	0.94	0.80	0.11	4.70	TOX	231, 234, 242, 243
ethene hydroxynitrate	C2O4H5N		2	4.26	6.00	0.02	8.50		230
ethyl acetate	C4H8O2	88.106	1	0.00		0.00	0.00		236
ethyl nitrate	C2H5NO3	91.066	2	2.40	1.56	1.30	3.50		230
								HAP,	230, 231, 234,
ethylbenzene	C8H10	106.168	24	0.03	0.02	0.00	0.08	TOX	236, 238
ethylindene	C11H12	144.217	58	0.02	0.02	0.00	0.10		242
ethylnylpyrrole	C6H6		58	0.00	0.00	0.00	0.01		242
eugenol	C10H12O2	164.204	2	0.07	0.01	0.07	0.08		105
								HAP,	
fluoranthene	C16H10	202.256	3	0.00	0.00	0.00	0.00	TOX	238
								HAP,	
fluorene	C13H10	166.223	3	0.00	0.00	0.00	0.00	TOX	238
									117, 118, 120,
									133, 170, 171,
formaldehyde	H2CO	30.026	202	1.53	0.92	0.09	4.76	HAP,	231, 234, 242, 243
formamide	CH3NO	45.041	56	0.01	0.01	0.00	0.06	TOX	242
									105, 117, 118,
									120, 133, 170,
formic acid	HCOOH	46.025	187	0.28	0.27	0.00	1.21	TOX	171, 231, 242, 243

furan	C4H4O	68.075	178	0.33	0.34	0.02	1.60	TOX	120, 170, 171, 230, 231, 234, 242, 243
furfural	C5H4O2	96.085	62	0.69	0.45	0.01	2.70		231, 243
gamma-butyrolactone	C4H6O2	86.09	5	0.09	0.05	0.05	0.17		105
glycolaldehyde	C2H4O2	60.052	67	0.60	0.46	0.05	2.54		117, 120, 231, 243
glyoxal	C2H2O2	58.036	58	0.26	0.23	0.00	0.96		242
guaiacol	C7H8O2	124.139	66	0.38	0.38	0.00	1.92		105, 242
heptane	C7H16	100.205	24	0.01	0.01	0.00	0.04		230, 231, 234, 236, 238
hexane	C6H14	86.178	24	0.01	0.01	0.00	0.05	HAP, TOX	230, 231, 234, 236, 238
hydrogen chloride	HCl	36.458	35	0.12	0.16	0.00	0.67		170, 230, 243 118, 120, 170, 171, 230, 231, 234, 242, 243
hydrogen cyanide	HCN	27.026	185	0.40	0.40	0.01	2.12	HAP, TOX	230
hydrogen peroxide	H2O2	34.014	2	0.60	0.59	0.18	1.02		242
hydrogen sulfide	H2S	34.076	53	0.01	0.01	0.00	0.05		230
hydroperoxy acetone	C3O3H6		2	0.09	0.06	0.04	0.13		230
hydroperoxymethanol	CO3H4	64.04	2	0.19	0.20	0.05	0.33		230
indeno[1,2,3-cd]pyrene	C22H12	276.338	3	0.00	0.00	0.00	0.00	HAP, TOX	238
isoamyl cyanide	C6H11N	97.161	55	0.00	0.00	0.00	0.01		242
isobutane	C4H10	58.124	9	0.01	0.01	0.01	0.03		230, 231, 234
isobutylene	C4H8	56.108	18	0.10	0.04	0.04	0.20		230, 231, 234
isocyanic acid	HNCO	43.025	58	0.53	0.34	0.05	1.57		242
isopentane	C5H12	72.151	9	0.02	0.02	0.01	0.07		230, 231, 234 171, 230, 231, 234, 242
isoprene	C5H8	68.119	87	0.15	0.15	0.01	0.67	TOX	230
isoprene hydroperoxyaldehydes	C5O3H8		2	0.17	0.01	0.16	0.18		230
isoprene hydroxy hydroperoxides	C5O3H10		2	0.08	0.05	0.04	0.11		230
isoprene hydroxynitrates	C5O4H9N		2	0.02	0.01	0.01	0.02		230

isopropyl nitrate	C3H7NO3	105.093	2	4.55	1.77	3.30	5.80		230
m-, p-xylene	C8H10		18	0.06	0.03	0.01	0.11	HAP, TOX	230, 234, 236, 238
m,p-cresol	C7H8O		9	0.05	0.02	0.01	0.08	HAP, TOX	105
methacrolein	C4H6O	70.091	7	0.04	0.00	0.03	0.04		231, 234 35, 46, 63, 66, 94, 100, 102, 117, 118, 120, 133, 137, 153, 157, 170, 171, 172, 179, 230, 231, 234, 237, 238, 239, 243, 244, 245
methane	CH4	16.043	449	4.78	3.76	0.00	24.10	GHG	242
methanediol	CH4O2	48.041	58	0.00	0.00	0.00	0.00		242
methanethiol	CH4S	48.103	54	0.01	0.01	0.00	0.07		242
methanimine	CH3N	29.042	57	0.00	0.00	0.00	0.00		242 105, 117, 118, 120, 133, 170, 171, 230, 231, 234, 242, 243
methanol	CH3OH	32.042	217	1.37	1.25	0.09	6.42	HAP, TOX	105
methenamine	C6H12N4	140.19	3	0.18	0.11	0.06	0.28		105
methyl acetoacetate	C5H8O3	116.116	9	0.23	0.25	0.01	0.77		105
methyl benzeneacetonitrile	C9H9N	131.178	56	0.00	0.00	0.00	0.02		242
methyl benzofuran	C9H8O	132.162	57	0.05	0.04	0.01	0.18		242
methyl benzoic acid	C8H8O2	136.15	58	0.08	0.06	0.01	0.31		242
methyl bromide	CH3Br	94.939	9	0.00	0.00	0.00	0.00	GHG, HAP, TOX	230, 236
methyl chavicol	C10H12O	148.205	57	0.03	0.02	0.00	0.09		242

chemical name	chemical formula	ref. conc. (ppb)	ref. conc. (ppm)	ref. conc. (ppm)	ref. conc. (ppm)	ref. conc. (ppm)	ref. conc. (ppm)	ref. conc. (ppm)	ref. conc. (ppm)
methyl chloride	CH3Cl	50.485	22	0.02	0.01	0.01	0.05	GHG, HAP, TOX	66, 230, 234, 236, 238
methyl iodide	CH3I	141.939	2	3.70	2.55	1.90	5.50	HAP, TOX	230
methyl methacrylate	C5H8O2	100.117	7	0.00	0.00	0.00	0.01	HAP, TOX	236
methyl naphthalene	C11H10	142.201	58	0.03	0.03	0.00	0.19		242
methyl nitrate	CH3NO3	77.039	2	1.50	0.28	1.30	1.70		230
methyl propanoate	C4H8O2	88.106	58	0.08	0.07	0.01	0.30		242
methyl tert-butyl ether	C5H12O	88.15	1	0.00		0.00	0.00	HAP, TOX	236
methyl vinyl ketone	C4H6O	70.091	8	0.11	0.11	0.02	0.37		231, 234
methyl vinyl ketone/methacrolein hydroxynitrates	C4O5H7N		2	0.02	0.00	0.02	0.02		230
methyl vinyl ketone/methacrolein/crotonaldehyde	C4H6O		2	0.33	0.06	0.29	0.37		230
monoterpenes	C10H16		66	0.99	1.30	0.02	5.43		230, 231, 242
m-xylene	C8H10	106.168	6	0.06	0.03	0.01	0.11	HAP, TOX	231
myrcene	C10H16	136.238	3	0.06	0.05	0.01	0.11		231
naphthalene	C10H8	128.174	75	0.06	0.05	0.00	0.23	HAP, TOX	236, 238, 242
n-decane	C10H22	142.286	4	0.04	0.03	0.01	0.08		231
nitrate	NO3-	62.004	58	0.11	0.11	0.00	0.60		172, 180, 230, 117, 118, 142, 170, 171, 172, 195, 230, 231, 234, 237, 243
nitric oxide	NO	30.006	190	2.13	1.61	0.09	8.10		
nitrobenzene	C6H5NO2	123.111	58	0.01	0.01	0.00	0.03	HAP, TOX	242
nitroethane	C2H5NO2	75.067	58	0.00	0.00	0.00	0.00		242
nitroethene	C2H3NO2	73.051	58	0.00	0.00	0.00	0.00		242

nitrogen dioxide	NO2	46.005	157	1.15	0.87	0.05	4.80		142, 170, 171, 172, 195, 230, 231, 234, 243 37, 63, 140, 142, 153, 170, 171, 180, 201, 230, 231, 237
nitrogen oxides	Nox		115	2.96	2.03	0.03	11.29	CAP	231, 242
nitromethane	CH3NO2	61.04	65	0.06	0.04	0.00	0.15	TOX	242
nitrotoluene	C7H7NO2	137.138	55	0.01	0.01	0.00	0.03		170, 171, 231, 242, 243
nitrous acid	HNO2	47.013	164	0.49	0.24	0.04	1.28		63, 137, 153
nitrous oxide	N2O	44.013	9	0.22	0.11	0.04	0.41	GHG	
nitroxyhydroperoxide/									
nitroxyhydroxyepoxide	C5O5H9N		2	0.02	0.00	0.02	0.02		230
nonane	C9H20	128.259	14	0.01	0.01	0.00	0.03		231, 236 35, 46, 94, 157, 244, 245
nonmethane hydrocarbons	NMHC		112	5.87	4.62	0.00	22.50		231
nonmethane organic compounds	NMOC		10	18.87	14.61	6.78	57.12		231, 236
octane	C8H18	114.232	16	0.01	0.01	0.00	0.03		230, 231
organic aerosol	OA		8	13.90	9.34	5.40	30.90		172, 180, 238
organic carbon	OC		54	18.32	28.41	0.23	137.33		
orthocresol	C7H8O	108.14	9	0.03	0.01	0.00	0.05	HAP, TOX	105
o-xylene	C8H10	106.168	24	0.02	0.01	0.00	0.04	HAP, TOX	230, 231, 234, 236, 238
ozone	O3	47.997	10	2.17	4.91	-1.25	14.00	TOX	63, 137
particle-bound polycyclic aromatic hydrocarbons	PPAH		8	0.01	0.01	0.00	0.02	HAP, TOX	237
particulate matter < 1µm	PM1		3	26.03	6.15	20.40	32.60		230
particulate matter 10µm	PM10		9	8.70	3.22	3.38	12.90	CAP	140, 157 46, 66, 100, 120, 140, 141, 157,
particulate matter 2.5µm	PM2.5		321	19.83	16.19	1.10	89.95	CAP	

									171, 172, 179, 180, 236, 237, 238, 240, 244, 245
particulate matter 2µm	PM2		15	19.13	10.07	5.50	43.80		59, 137
particulate matter 3.5µm	PM3.5		13	16.14	7.86	9.70	37.40		59, 137, 153
particulate matter 48µm	PM48		22	19.97	12.13	4.40	47.90		59, 94, 137
p-cymene	C10H14	134.222	3	0.42	0.50	0.00	0.98		231
pentane	C5H12	72.151	9	0.03	0.01	0.01	0.05		230, 231, 234
peracetic acid	C2H4O3	76.051	2	0.24	0.28	0.05	0.44	TOX HAP,	230
phenanthrene	C14H10	178.234	3	0.00	0.00	0.00	0.00	TOX HAP,	238 105, 120, 231,
phenol	C6H6O	94.113	134	0.62	0.57	0.02	3.54	TOX	242, 243
phenylacetonitrile	C8H7N	117.151	56	0.01	0.01	0.00	0.05		242
phenylacetylene	C8H6	102.136	58	0.01	0.01	0.00	0.06		242
polychlorinated dibenzofurans	PCDF		14	0.00	0.00	0.00	0.00	HAP, TOX	236
polychlorinated dibenzo-p-dioxins	PCDD		13	0.00	0.00	0.00	0.00	HAP, TOX	236
potassium cation	K+	39.098	31	0.48	0.50	0.03	2.36		172
propane	C3H8	44.097	68	0.14	0.10	0.01	0.54		63, 137, 153, 179, 230, 231, 234
propanone nitrate	C3O3H5N		2	4.00	0.71	3.50	4.50		230
									63, 120, 137, 153, 170, 171, 172, 179, 230, 231, 234, 236, 238,
propene	C3H6	42.081	293	0.61	0.49	0.04	2.42	TOX	242, 243
propene hydroxynitrates	C3O4H7N		2	0.02	0.01	0.02	0.03		230
propionic acid	C3H2O2	70.047	58	0.01	0.01	0.00	0.03		242
propionitrile	C3HN	51.048	58	0.00	0.00	0.00	0.00		242

propionaldehyde	C3H6O	58.08	20	0.64	0.41	0.24	2.07	HAP, TOX	230, 231, 234, 236
propionic acid	C3H6O2	74.079	9	0.21	0.14	0.02	0.39		105
propionitrile	C3H5N	55.08	57	0.01	0.02	0.00	0.08		242
propyl nitrate	C3H7NO3	105.093	2	4.45	4.03	1.60	7.30		230
propylbenzene	C9H12	120.195	16	0.01	0.00	0.00	0.01		231, 236
propynal	C3H2O	54.048	58	0.03	0.02	0.00	0.08		242
propyne	C3H4	40.065	62	0.05	0.03	0.00	0.14		179, 230, 231, 234
p-xylene	C8H10	106.168	7	0.04	0.03	0.02	0.09	HAP, TOX	231
pyrene	C16H10	202.256	3	0.00	0.00	0.00	0.00	HAP, TOX	238
pyridine	C5H5N	79.102	5	0.05	0.08	0.01	0.20	TOX	105
pyruvaldehyde	C3H4O2	72.063	3	1.10	0.85	0.30	2.00		105
pyruvic acid	C3H4O3	88.062	58	0.01	0.01	0.00	0.03		242
quinone	C6H4O2	108.096	57	0.09	0.05	0.01	0.25	HAP, TOX	242
salicylaldehyde	C7H6O2	122.123	61	0.06	0.05	0.01	0.28		105, 242
sodium cation	Na+	22.99	15	0.06	0.08	0.00	0.25		172
styrene	C8H8	104.152	80	0.07	0.07	0.01	0.37	HAP, TOX	231, 236, 238, 242
sulfate	SO4(2-)	96.056	35	0.28	0.19	0.01	0.73		172, 230
sulfinylmethanamine	CH3NOS	77.101	57	0.00	0.00	0.00	0.00		242
sulfur dioxide	SO2	64.058	127	1.11	0.72	0.00	3.42		140, 170, 172, 230, 243
syringol	C8H10O3	154.165	56	0.04	0.03	0.00	0.17	CAP	242
terpenes			8	3.40	4.32	0.14	13.35		231
tetrahydrofuran	C4H8O	72.107	10	0.00	0.00	0.00	0.01		236
thiophene	C4H4S	84.136	58	0.01	0.01	0.00	0.05		242
toluene	C7H8	92.141	83	0.23	0.20	0.02	1.12	HAP, TOX	230, 231, 234, 236, 238, 242
total carbon	TC		5	10.06	3.87	7.10	16.00		238

total hydrocarbon	THC		56	8.36	5.30	1.06	21.80		4, 42, 129, 140 26, 35, 36, 37, 46, 53, 66, 86, 129, 147, 157, 178, 195, 229, 244, 245
total particulate matter	PM		276	19.23	15.45	2.10	77.10		230, 231, 234
trans-2-butene	C4H8	56.108	9	0.06	0.06	0.01	0.21		231
trans-2-pentene	C5H10	70.135	7	0.03	0.02	0.01	0.08		242
trimethylamine	C3H9N	59.112	56	0.00	0.00	0.00	0.01		105, 242
vanillin	C8H8O3	152.149	66	0.14	0.13	0.00	0.60		
vinyl acetate	C4H6O2	86.09	21	0.34	0.25	0.03	1.00	HAP, TOX	105, 236, 238
vinyl acetylene	C4H4	52.076	58	0.05	0.04	0.01	0.19		242
vinyl chloride	C2H3Cl	62.496	2	0.00	0.00	0.00	0.00	HAP, TOX	236
vinyl guaiacol	C9H10O2	150.177	58	0.15	0.16	0.01	0.84		242
vinylpyridine	C7H7N	105.14	58	0.00	0.00	0.00	0.02		242
water	H2O	18.015	55	0.66	0.39	0.09	1.63		242
xylene	C8H10		10	0.08	0.03	0.04	0.13	HAP, TOX	236