



**City of Philadelphia  
Department of Public Health  
Air Management Services**

# **Philadelphia's Air Quality Report 2019**



## Executive Summary<sup>1</sup>

This report focuses on the air quality of the City of Philadelphia, as presented by the Philadelphia Department of Public Health, Air Management Services, the local air pollution control agency for the City of Philadelphia. As an urban area, Philadelphia faces many of the same pollution challenges as other densely populated areas, such as emissions from vehicles and industries. The information contained in this report reviews Philadelphia's air quality for the calendar year 2019 and reports how the City's air compared with the National Ambient Air Quality Standards. This report covers the following criteria pollutants: **ozone, carbon monoxide, nitrogen dioxide, sulfur dioxide, particulate matter, and lead**. It also provides an overview of **hazardous air pollutants**, also referred to as **air toxics**.

In general, trends show many air pollutants in Philadelphia to be decreasing. In 2019, Philadelphia attained the NAAQS for all pollutants, except for ozone. There were 244 good days, 115 moderate days, and 6 unhealthy days (6 from ozone and 0 from PM<sub>2.5</sub>) in Philadelphia.

In 2019, AMS continued the Philadelphia Air Quality Survey. This project set up 50 street level, neighborhood-oriented air sampling sites throughout the City to sample the ambient air for PM<sub>2.5</sub>, PM<sub>2.5</sub> speciation, NO<sub>2</sub>, SO<sub>2</sub>, and O<sub>3</sub>.

For further information, please visit the Air Management Services website at: <https://www.phila.gov/departments/departments-of-public-health/about-us/contact-us/>

or contact us at:  
215-685-7580

James Kenney, Mayor  
Thomas A. Farley, Health Commissioner

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<sup>1</sup> Cover photo by [Leo Serrat](#) on [Unsplash](#)

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## Introduction

The Philadelphia Department of Public Health, Air Management Services (AMS) is responsible for the prevention, abatement, and control of air pollution and air pollution nuisances; achieving and maintain the National Ambient Air Quality Standards (NAAQS) in Philadelphia; and protecting the health and quality of life of the Philadelphia community from the adverse effects of air contaminants and noise.

AMS implements the environmental protection mandates contained in city, state, and federal regulations; reviews construction and operating permits for compliance with air regulations, standards, and guidelines; develops emission inventories for large stationary sources; operates and maintains a citywide air sampling network to continuously monitor Philadelphia's air; routinely inspects pollution sources; services citywide complaints of air pollution, asbestos, and noise; issues violations; conducts enforcement actions; and advances voluntary emissions reductions.

## Air Monitoring Network

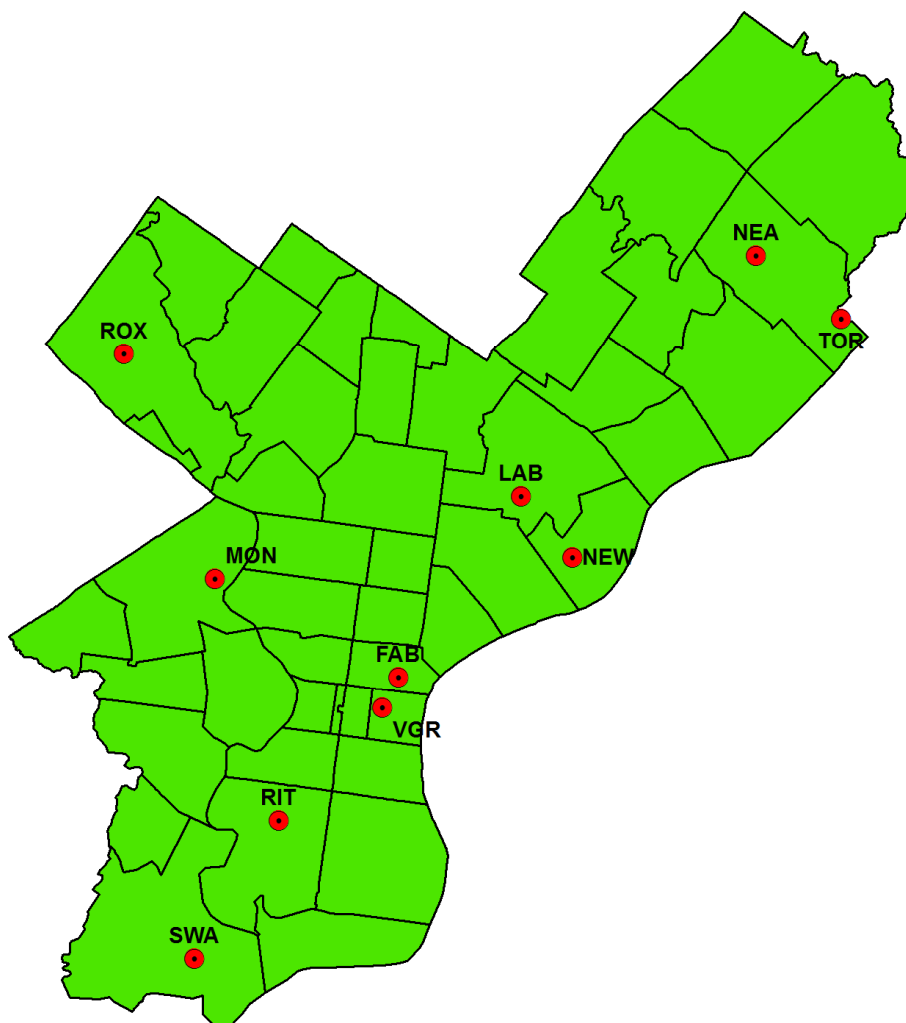
The City of Philadelphia is served by a network of ten air monitoring sites located throughout the City that measure the criteria pollutants: ozone (O<sub>3</sub>), carbon monoxide (CO), nitrogen dioxide (NO<sub>2</sub>), sulfur dioxide (SO<sub>2</sub>), particulate matter (PM<sub>10</sub> and PM<sub>2.5</sub>), and lead (Pb). Four of the sites also measure toxics, such as 1,3-butadiene, benzene, and carbon tetrachloride. Many of the measurements are made in "real time", meaning that the measurements show pollution levels as they occur, instead of after the fact. The map on page 6 shows the location of air monitors and the pollutants measured at each monitoring location. AMS measures air quality for several reasons:

- To ensure that long-term goals and targets to reduce levels of air pollution are being met.
- To provide information to the public as to how good or bad the air quality is in Philadelphia.
- To ensure attainment with standards set forth by the United States Environmental Protection Agency (EPA).

An air monitoring network plan (AMNP) has been made available to the public annually starting in the year 2007. The most recent AMNP is located on the AMS reports and documents website:

<https://www.phila.gov/documents/air-management-reports-and-documents/>.

Figure 1 – 2019 Philadelphia Air Monitoring Network



			Parameter																
AQS Site Code	AMS Site	Address	CO	SO <sub>2</sub>	Ozone	NO <sub>2</sub>	NOy/NO	PM <sub>10</sub>	PM <sub>2.5</sub>	Speciated PM <sub>2.5</sub>	PM Coarse	Black Carbon / Ultrafine PM	Carbonyls	PAMS VOC	BaP	TSP Metals (Be, Cr, Mn, Ni, As, Cd, Pb)	Toxics TO15	MET	AMS Site
421010004	LAB	1501 E Lycoming St			X														LAB
421010014	ROX	Eva & Dearnley Sts											X				X		ROX
421010024	NEA	Grant Ave & Ashton Rd			X														NEA
421010048	NEW	2861 Lewis St	X	X	X	X	X	X	X	X	X		X	X			X	X	NEW
421010055	RIT	24th & Ritner Sts		X					X	X			X			X	X		RIT
421010057	FAB	3rd & Spring Garden Sts							X										FAB
421010063	SWA	8200 Enterprise Ave											X				X		SWA
421010075	TOR	4901 Grant Ave & James St	X			X			X									X	TOR
421010076	MON	I-76 & Montgomery Drive	X			X			X			X			X	X		X	MON
	VGR	6th & Arch Sts			X				X									X	VGR

**Table 1 – Site Summary Table**

<b>AMS Site</b>	<b>Address</b>	<b>Statement of Purpose</b>
<b>LAB</b>	1501 E. Lycoming St	Built in 1964, this monitor assesses the City's impact on ozone precursors and is a designated Photochemical Assessment Monitoring Station (PAMS) site. New monitoring methods are often evaluated on this site.
<b>ROX</b>	Eva St & Dearnley St	As a periphery site, this site is used for measuring Air Toxics and Carbonyls.
<b>NEA</b>	Grant Ave & Ashton Rd	As a periphery site, this site is best for measuring ozone in the City, because as a secondary pollutant, ozone requires some time to form (longer time periods allow precursor emissions to distribute more uniformly across a region, and thus allow ozone concentrations to develop more uniformly across subregions and even large-scale regions). We tend to see fewer "hot spots" as ozone is not directly emitted from combustion activities as other pollutants are.
<b>NEW</b>	2861 Lewis St	This site was one few sites that was originally established to measure the impact of specific industrial facilities which are now closed. Today, the monitors conduct continuous particulate monitoring and provide information about the nearby wastewater treatment plant. As of October 2, 2013, the NCore site has been moved to this site from Baxter water treatment plant (BAX).
<b>RIT</b>	24 <sup>th</sup> St & Ritner St	This site was selected to help assess the impact of the petroleum refinery on the local community. The area was identified by air quality modeling.
<b>FAB</b>	3 <sup>rd</sup> St & Spring Garden St	This site was established to monitor high levels of fine particulates in the City based on EPA Region III's air quality modeling of air toxics in Philadelphia. It shows high levels of fine particulate created by vehicle traffic.
<b>SWA</b>	8200 Enterprise Ave	This site was established to measure toxics, carbonyls, and metals. Fine particulates may also be monitored. EPA Region III modeling analysis has shown that areas near the airport have high levels of aldehydes.
<b>TOR</b>	4901 Grant Ave & James St.	This site was established as the 1 <sup>st</sup> near-road NO <sub>2</sub> monitor in the Philadelphia-Camden-Wilmington, PA-NJ-DE-MD Metropolitan Statistical Area .
<b>MON</b>	I-76 & Montgomery Drive	This site was established as the 2nd near-road monitor in the Philadelphia-Camden-Wilmington, PA-NJ-DE-MD Metropolitan Statistical Area.
<b>VGR</b>	6 <sup>th</sup> St & Arch St	EPA's Village Green Air Monitoring Station. Utilizes solar and wind turbine power as energy sources. Sited to increase community awareness of environmental conditions.



## Quality Assurance

The AMS Air Monitoring Laboratory's main responsibility is to provide accurate data on the quality of the City's air. Pollutants in the atmosphere are measured to answer a number of questions such as:

- Are the NAAQS being met in Philadelphia?
- How close or far away are we from meeting these standards?
- Which pollutants are getting worse (increasing in concentration) or improving?

Many of our measurements require detecting very small amounts of a pollutant, often expressed as parts per million (ppm) or parts per billion (ppb). An illustration: imagine a million yellow balls all the same size with several red balls in the middle of them; we would need to find those red balls and then be able to count them. The instruments used to measure air pollutants need to be reliable in identifying the pollutant and accurate in making the measurement every time. The main way we check to see if our instruments are giving accurate measurements is to send a sample of air which has a known amount (concentration) of a pollutant and compare what the instrument says is the concentration to what we know is the right concentration. Then adjustments (calibration) to an instrument can be made to give a better measurement. If the equipment is off by a significant margin, the instrument needs to be repaired or replaced. The EPA and our Laboratory have standard operating procedures on how accurate and reliable measurements need to be to answer the questions being asked. The instruments being used now are much more reliable than those available years ago. Steps to assure good data quality include:

- Automated calibration.
- Manual calibration conducted by chemists.
- Review of the data by an experienced engineer or scientist.

The system is geared towards public safety; for example, a few measurements can be enough to identify a problem in meeting the NAAQS, but many good measurements over a period of time (often three years) as well as additional types of analysis are needed to “demonstrate compliance” with the corresponding pollutant’s standard on the NAAQS.

The EPA’s regulations on establishing an air monitoring network are found in Title 40 – Protection of Environment in the Code of Federal Regulations Part 58 – Ambient Air Quality Surveillance<sup>2</sup>. A comprehensive technical systems audit (TSA) by EPA occurs at least once every three years to evaluate an ambient air quality monitoring program. AMS's TSA occurred August 2019. The TSA included on-site interviews with program personnel, evaluations of ambient air monitoring sites and laboratories operated by the agency, and a review of quality assurance and data processing procedures. AMS follows EPA's

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<sup>2</sup> [https://www.ecfr.gov/cgi-bin/text-idx?SID=af0f262c08e4f8aad56a91c70e0ce294&mc=true&tpl=/ecfrbrowse/Title40/40cfr58\\_main\\_02.tpl](https://www.ecfr.gov/cgi-bin/text-idx?SID=af0f262c08e4f8aad56a91c70e0ce294&mc=true&tpl=/ecfrbrowse/Title40/40cfr58_main_02.tpl)



guidance in the Quality Assurance Handbook for Air Pollution Measurement Systems, Volume II<sup>3</sup>, dedicated to the Ambient Air Quality Surveillance Program and the data collection activities inherent to that program. Additionally, workshops are provided annually to staff as an opportunity to interact with and learn from nearby state and local monitoring agency program representatives as well as expert speakers. Participants share experiences meeting challenges with new air quality monitoring technology, updated data analysis and quality assurance methods, and revised regulatory requirements. The 2019 Mid-Atlantic Regional Air Management Association (MARAMA) Air Monitoring Workshop occurred in November 2019<sup>4</sup>.

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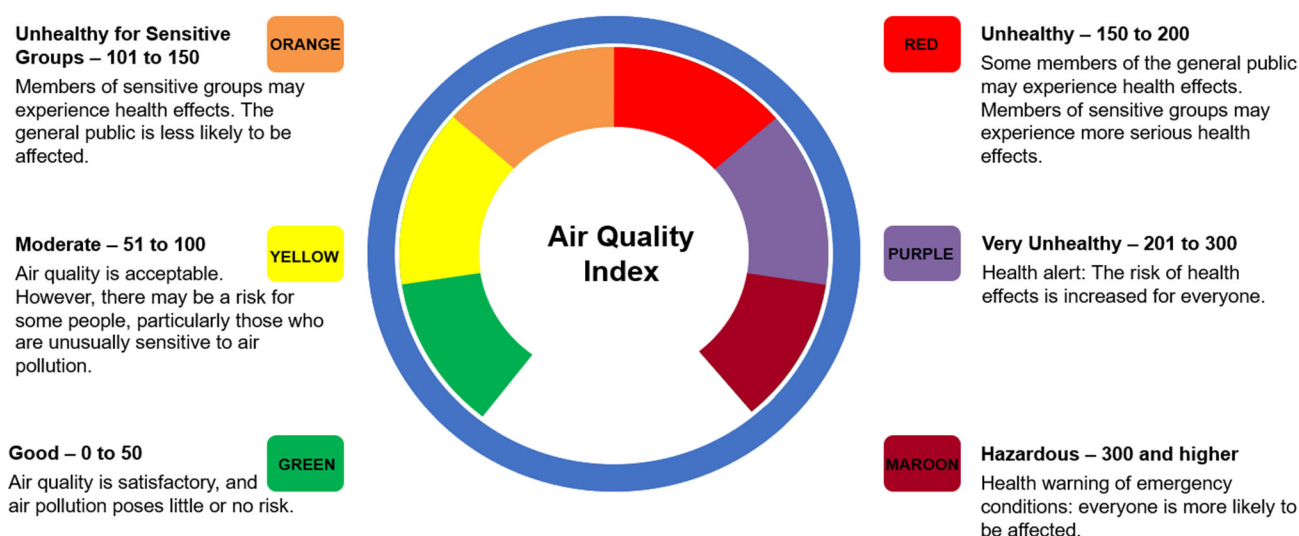
<sup>3</sup> [https://www3.epa.gov/ttn/amtic/files/ambient/pm25/qa/Final%20Handbook%20Document%201\\_17.pdf](https://www3.epa.gov/ttn/amtic/files/ambient/pm25/qa/Final%20Handbook%20Document%201_17.pdf)

<sup>4</sup> <https://marama.org/events/2019-monitoring-committee-workshop/>

## Air Quality Index

The Air Quality Index (AQI) is a color-coding system for air quality used by government agencies across the United States. Daily pollution levels for five pollutants (ground-level ozone, particulate matter, carbon monoxide, sulfur dioxide, and nitrogen dioxide) are converted into a score ranging from 1 to 500. A level of 101 generally corresponds to the National Ambient Air Quality Standard for each pollutant, and an “Action Day” occurs when the AQI for any pollutant exceeds 100. The color coded AQI is shown in Figure 2.

Figure 2 – Color Coded AQI



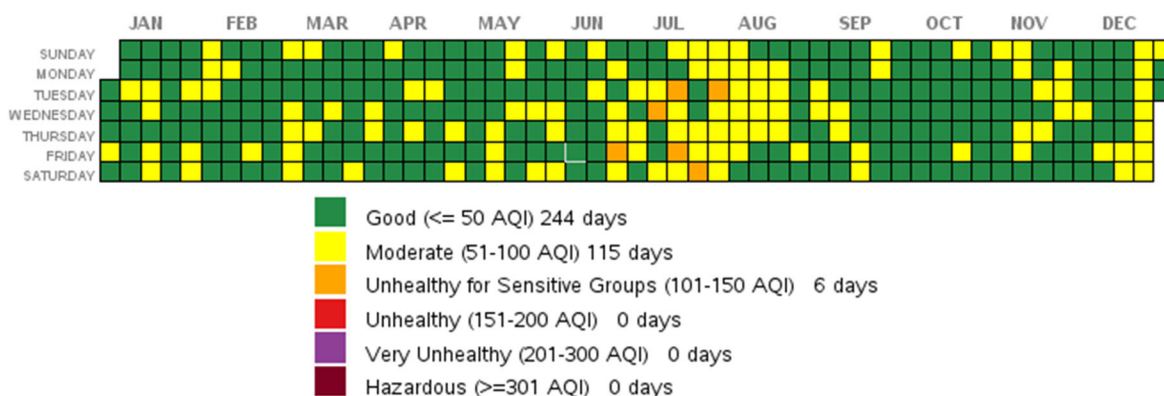
Action days are more likely to occur in the summer months but can happen any time of the year. On these days, the public is advised to do their part to reduce pollution and take precautions to protect themselves and their families from health effects. For example, on an Orange day, children, seniors, and those with respiratory ailments are advised to minimize prolonged outdoor exposure. On a Red or Purple day, all residents are advised to limit outdoor activity. Red and purple days are uncommon. The highest of the five pollutant scores is reported as the overall air quality rating for Philadelphia for a given day. The daily AQI values for 2019 are shown in Figure 3.<sup>5</sup>

Action Days are reported through print, radio and television media, online, and by local and regional air agencies. Current air quality conditions in Philadelphia can be found on the website: <https://www.phila.gov/services/mental-physical-health/environmental-health-hazards/air-quality/>.

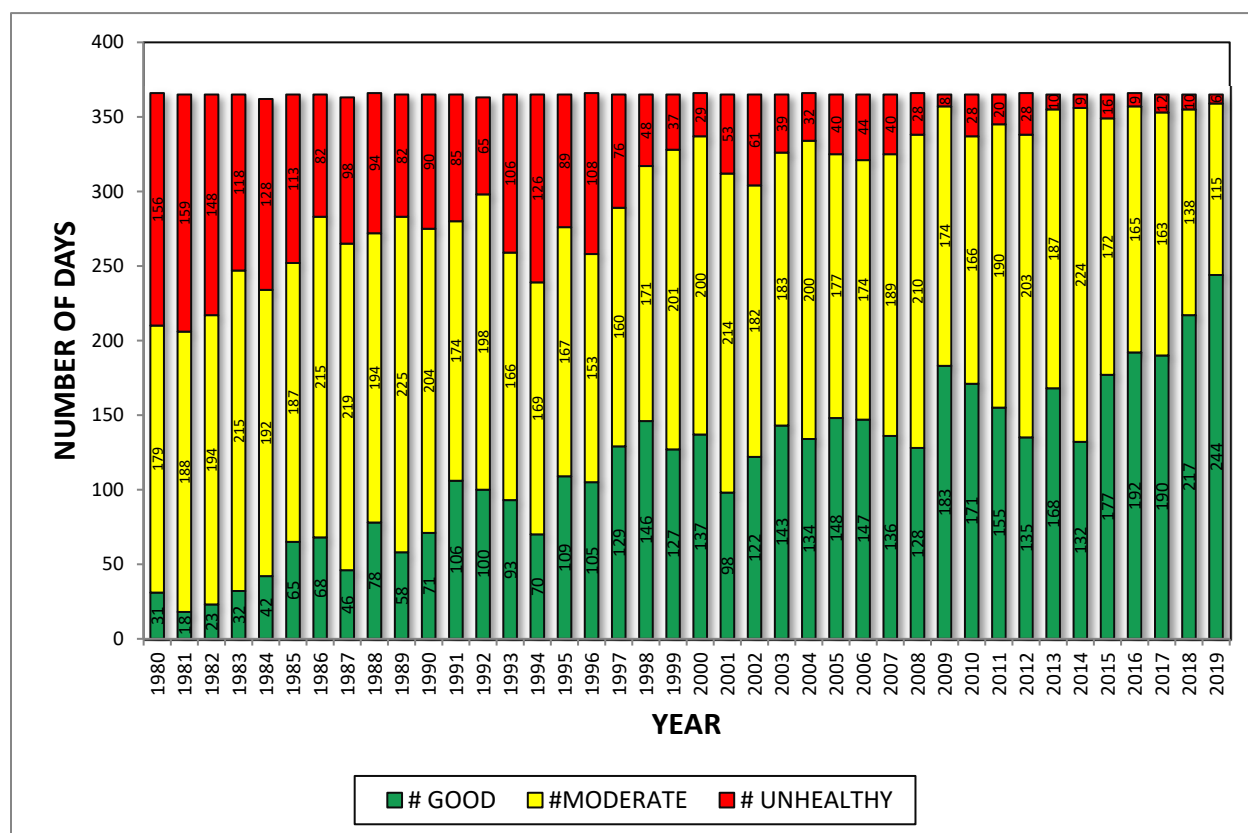
<sup>5</sup> <https://www.epa.gov/outdoor-air-quality-data/air-data-tile-plot>. Downloaded August 19, 2020.

Figure 3.2 shows the annual summary of the number of good, moderate, and unhealthy air quality days in Philadelphia based on monitoring conducted by AMS since 1980<sup>6</sup>. The chart has been standardized with the current EPA AQI breakpoints or pollutant concentration cut-offs and are consistent with the 2015 ozone standards.

**Figure 3.1 – Daily AQI Values for 2019**



**Figure 3.2 – Philadelphia Annual AQI Summary**



<sup>6</sup> <https://www.epa.gov/outdoor-air-quality-data/air-quality-index-report>. Downloaded August 4, 2020.

## National Ambient Air Quality Standards

The Clean Air Act (CAA), which was last amended in 1990, requires EPA to set NAAQS for pollutants considered harmful to public health and the environment. The CAA identifies two types of national ambient air quality standards. Primary standards provide public health protection, including protecting the health of "sensitive" populations such as asthmatics, children, and the elderly. Secondary standards provide public welfare protection, including protection against decreased visibility and damage to animals, crops, vegetation, and buildings.

The EPA has set NAAQS for six criteria air pollutants: carbon monoxide, sulfur dioxide, nitrogen dioxide, lead, particulate matter ("dust" or "soot"), and ozone. Periodically, the standards are reviewed and may be revised. The current standards are listed here: <https://www.epa.gov/criteria-air-pollutants/naaqs-table>. A history of the standard for each criteria pollutant is shown in Appendix C. Units of measure for the standards are parts per million (ppm) by volume, parts per billion (ppb) by volume, and micrograms per cubic meter of air ( $\mu\text{g}/\text{m}^3$ ).

In 2019, Philadelphia was in attainment for all pollutants, except for ozone.

## The Pollutants We Measure

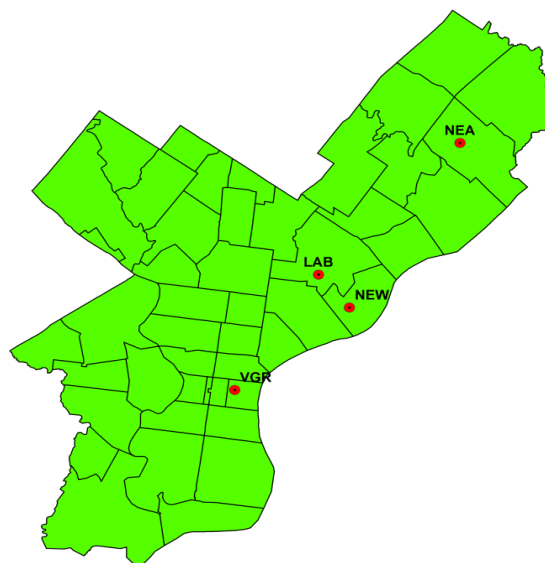
The following pages provide information on the health effects, sources, and trends of pollutants measured in Philadelphia. Included are the six pollutants, commonly called criteria pollutants, for which EPA has established NAAQS, as well as pollutants identified as being toxic or hazardous. Each of the criteria pollutants are graphed to show the historical trends compared with national standards. The graphs identify the range of values ("MIN / MAX RANGE") and with a solid black line, the mean of all recorded levels. It is important to note the mean, as it factors out extreme levels, and thereby provides a better indication of general air quality levels. In addition, Appendix B provides tables of historical information downloaded from EPA's Air Quality System data mart.

## Ozone (O<sub>3</sub>)

### NAAQS:

- The 3-year average of the annual fourth-highest daily maximum 8-hour average O<sub>3</sub> concentration is less than or equal to 0.070 ppm

Figure 4.1 – Ozone Monitoring Map



Ground level ozone (the primary constituent of smog) is the pollutant most often responsible for unhealthy air quality in the Philadelphia region. Ozone is not emitted into the atmosphere directly but is formed by chemical reactions between other pollutants. Specifically, Volatile Organic Compounds (VOCs) and Nitrogen Oxides (NO<sub>x</sub>) react to create ozone in the presence of heat and sunlight. Ozone levels are consistently higher during the summer months.



VOCs are organic (i.e. carbon-containing) compounds that evaporate readily, such as gasoline vapors and paint fumes. NO<sub>x</sub> stands for two compounds, nitric oxide (NO) and nitrogen dioxide (NO<sub>2</sub>). Some VOCs are considerably more reactive in the atmosphere than others, and the reactivity of a VOC influences how quickly ozone forms. A compound that reacts in a few minutes to produce ozone will have a much greater impact near its source than one that reacts more slowly. Thus, ozone can form at various distances downwind of a VOC source due to the speed of these chemical reactions.

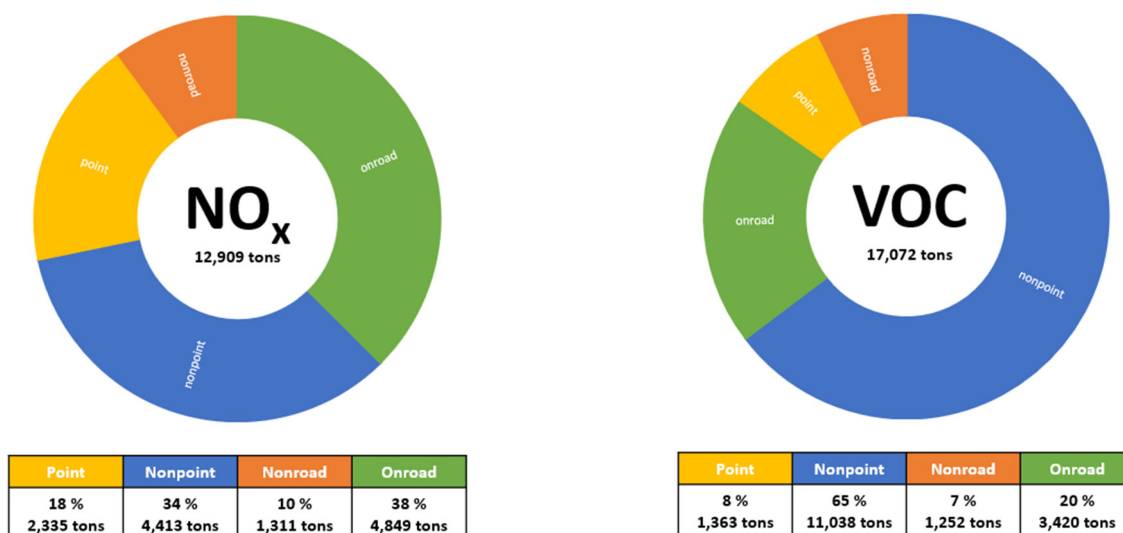
There are four categories of emission sources from human activity that produce VOC and NO<sub>x</sub>:

- Point sources – The largest utilities, industries, and other operations.

- Nonpoint sources – Commercial, solvent use, waste disposal, and other smaller categories.
- Nonroad sources – Construction and agricultural equipment, recreational boats, lawnmowers and other sources.
- Onroad sources – Cars, trucks, buses, and motorcycles.

Figure 4.2 shows NO<sub>x</sub> and VOC emissions from all source categories for Philadelphia County for 2017. This is the most recent data from EPA's National Emission Inventory.

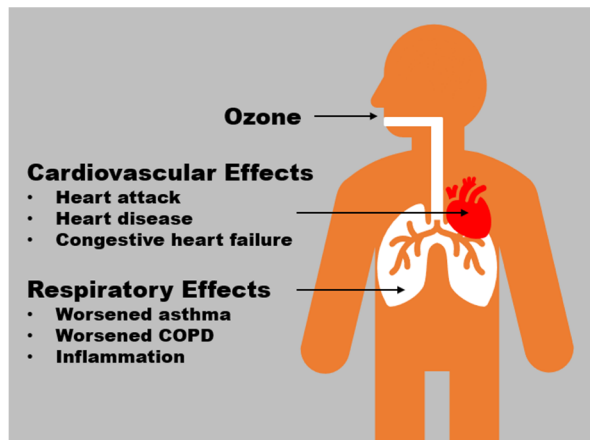
**Figure 4.2 – 2017 NO<sub>x</sub> and VOC Emissions From All Sources<sup>7</sup>**



Emissions of VOC and NO<sub>x</sub> may be carried by wind currents while reacting to produce high ozone levels hundreds of miles from their sources. In the eastern United States during the summer months, ground level ozone is frequently high over wide areas containing several states. This phenomenon is caused by ozone and its precursors traveling via wind currents across great distances.

In any discussion of ozone, it is important to distinguish between the effects of ozone at the ground and ozone high in the atmosphere. An advertisement might use the slogan “good up high, bad nearby,” to describe ozone. Up high, in what’s called the ozone layer, ozone is essential to the health of nearly every living thing, since it protects the Earth from harmful ultraviolet (UV) light. Near the ground, ozone reacts with buildings, plants, animals, and people, and is one of the most irritating, harmful components of smog. Smog refers to the whole mixture of air pollution in an area, and may include ozone, a whole host of other gases, as well as fine particles and the hazy conditions they cause.

<sup>7</sup> Data from <https://www.epa.gov/air-emissions-inventories/2017-national-emissions-inventory-nei-data>.



People who are very young, very old, or have chronic lung problems such as asthma are particularly sensitive to ground level ozone. EPA concluded that ozone pollution causes respiratory harm such as worsened asthma, worsened COPD, and inflammation; may cause reproductive and developmental harm as well as harm to the central nervous system; and likely to cause cardiovascular harm such as heart attacks, heart disease, and congestive heart failure.<sup>8</sup>

In 2019, as seen in Figure 4.1, there were four ozone monitoring sites: LAB, NEA, NEW, and VGR. The ozone monitor at VGR is part of EPA's Village Green Project to demonstrate the capabilities of new real-time monitoring technology using solar power for residents and citizen scientists to learn about local air quality. The real-time data is available here: <http://villagegreen.airnowtech.org/welcome?siteID=24292>. Data from the VGR monitor is not used for comparison to the NAAQS or AQI.

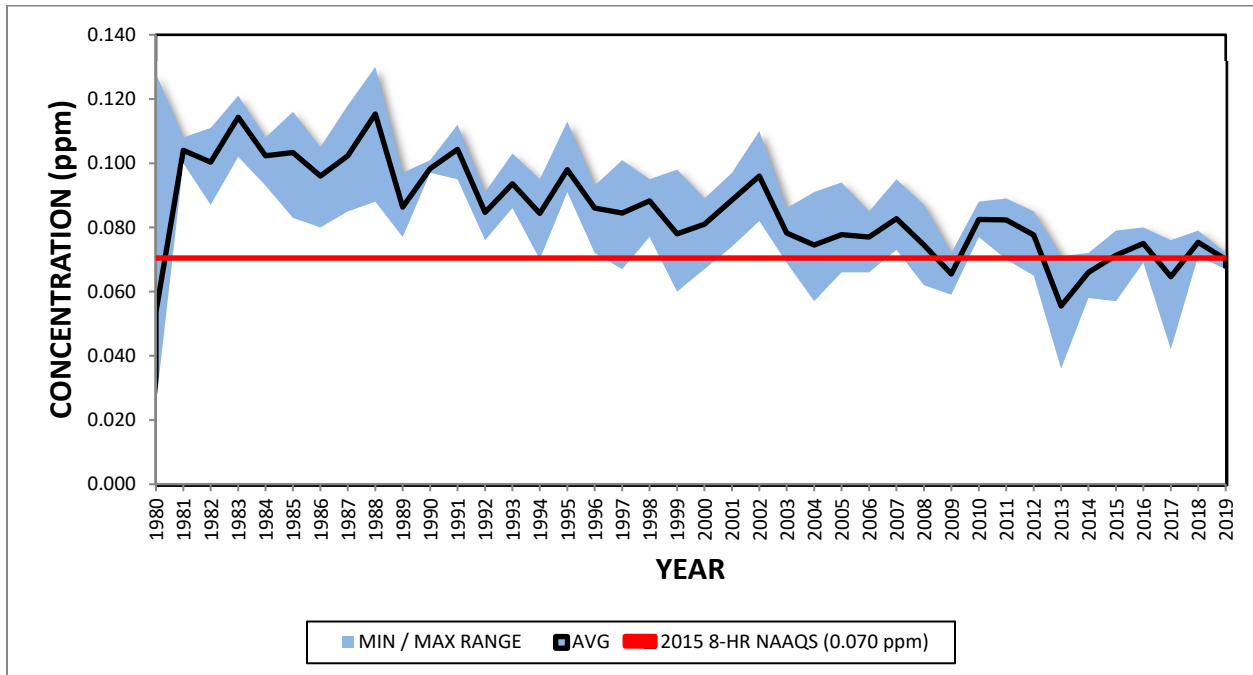
Figures 4.3 and 4.4 show the trends for the 2015 ozone 8-hour concentration in Philadelphia and the 3-year design value at Northeast Airport (NEA), respectively. The NEA monitoring site has the highest ozone design value in Philadelphia.

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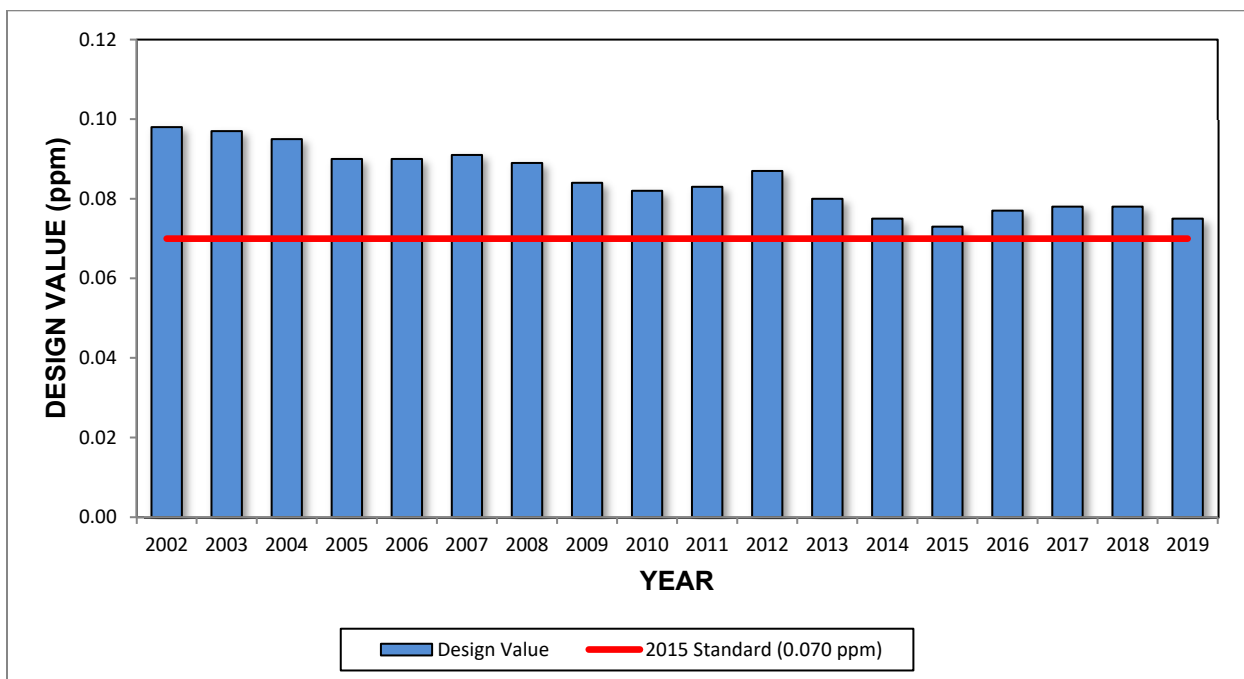
<sup>8</sup> U.S. Environmental Protection Agency, Integrated Science Assessment for Ozone and Related Photochemical Oxidants, 2013. EPA/600/R-10/076F.



**Figure 4.3 – Ozone Trends for the 4<sup>th</sup> Highest Daily Maximum 8-Hour Concentration From All Sites**



**Figure 4.4 – 3-Year Design Value at NEA Monitoring Site (AQS ID 421010024)**

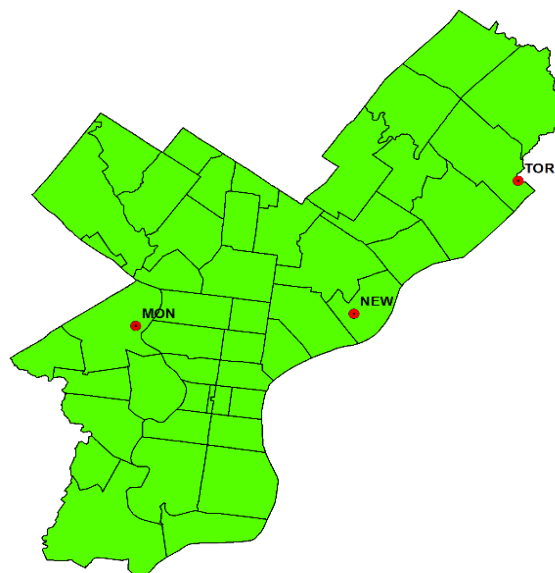


## Carbon Monoxide (CO)

### NAAQS:

- 9 parts per million for an 8-hour average concentration (2nd highest) not to be exceeded more than once per year.
- 35 parts per million for a 1-hour average concentration not to be exceeded more than once per year.

Figure 5.1 – CO Monitoring Map



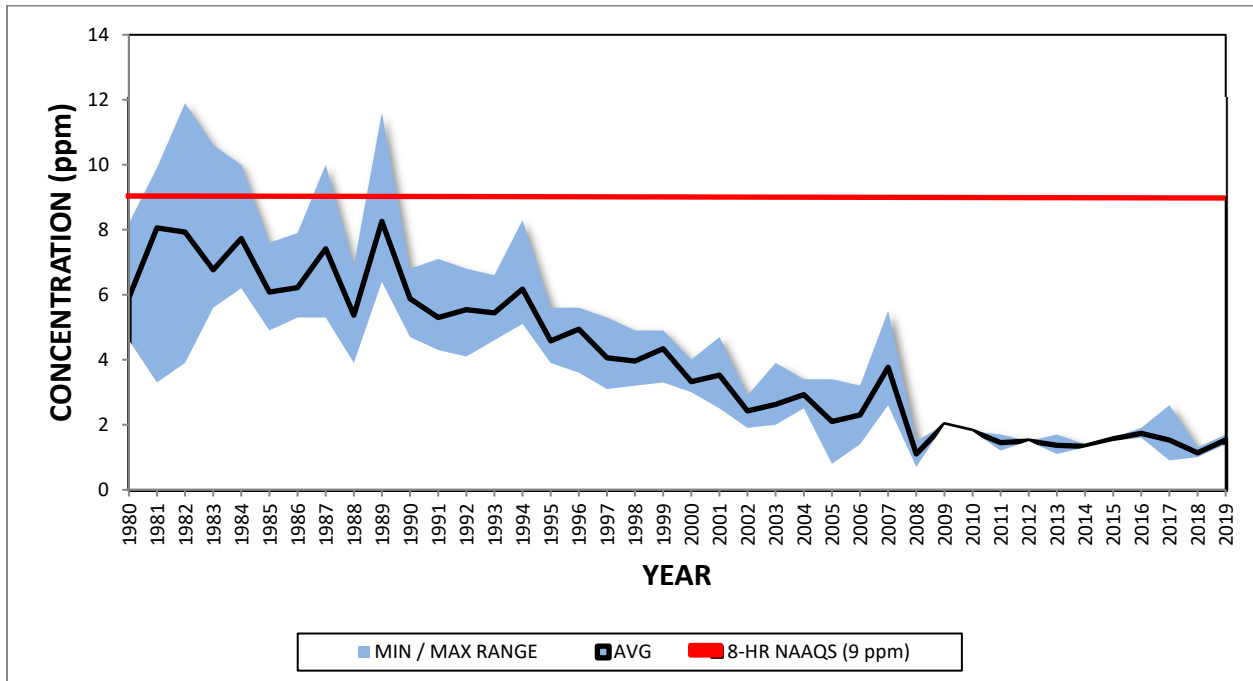
Carbon monoxide (CO) is colorless, odorless, and at high concentrations a poisonous gas. It is formed when carbon in fuels are not burned completely. The major source of CO is motor vehicle emissions. Other sources of CO include residential, industrial, and natural processes. Weather greatly affects CO levels, and peak CO concentrations typically occur during the colder months of the year.

Carbon monoxide enters the bloodstream and reduces oxygen delivery to the body's organs and tissues. The health threat from carbon monoxide is most serious for those who suffer from cardiovascular disease. Exposure to elevated CO levels is associated with impairment of vision, reduced work capacity, reduced manual dexterity, poor learning ability, and difficulty in performing complex tasks. At very high levels, carbon monoxide can be fatal.

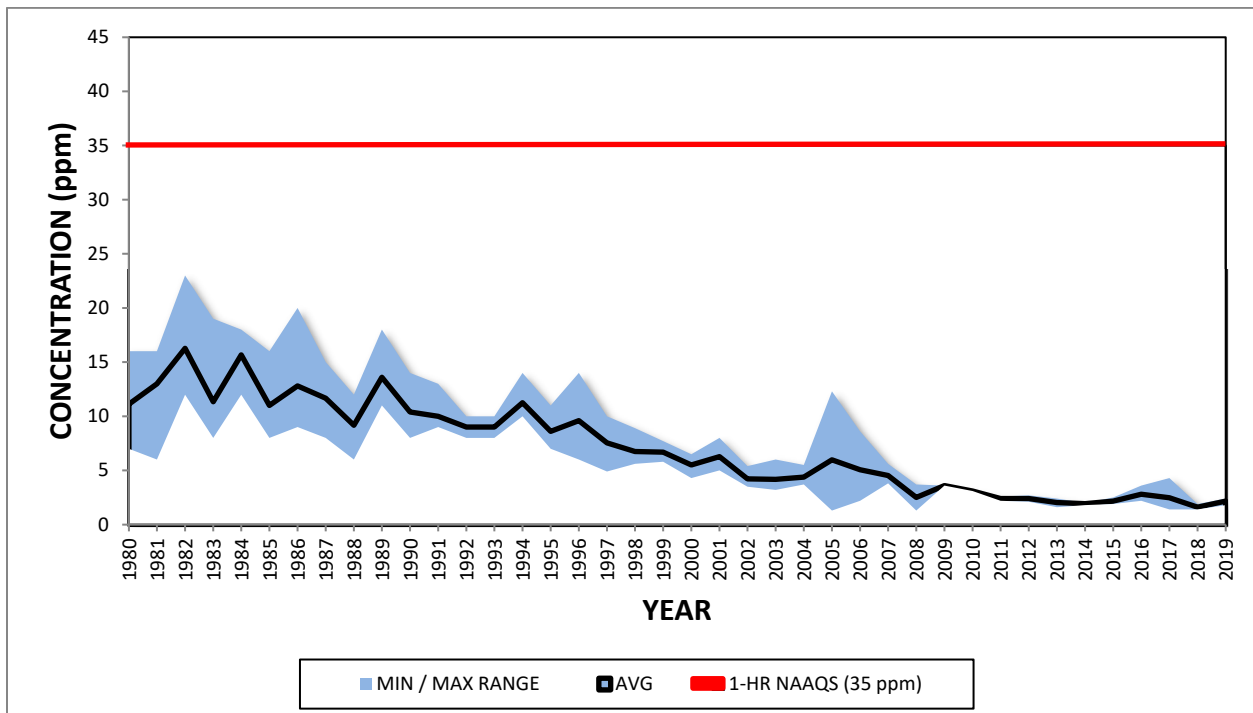
Over a thirty year period, there has been a continued reduction in carbon monoxide levels. This is mainly the result of federal requirements for cleaner automobiles and fuel and state inspection/maintenance programs.

Figures 5.2 and 5.3 on the following page show the trends for the CO 8-hour concentration and 1-hour concentration, respectively, in Philadelphia.

**Figure 5.2 – CO Trends for the 2<sup>nd</sup> Highest 8-Hour Average Concentration**



**Figure 5.3 – CO Trends for the Highest 1-Hour Average Concentration**

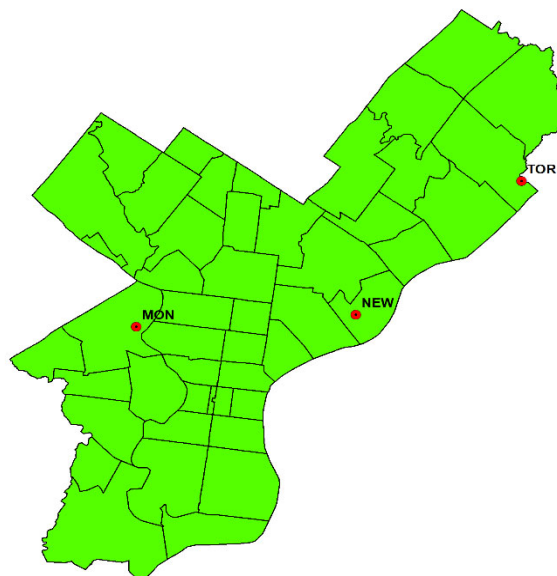


## Nitrogen Dioxide (NO<sub>2</sub>)

### NAAQS:

- The annual average concentration is less than or equal to 53 ppb.
- The 3-year average of the annual 98th percentile of the daily maximum 1-hour average concentration is less than or equal to 100 ppb.

Figure 6.1 – NO<sub>2</sub> Monitoring Map



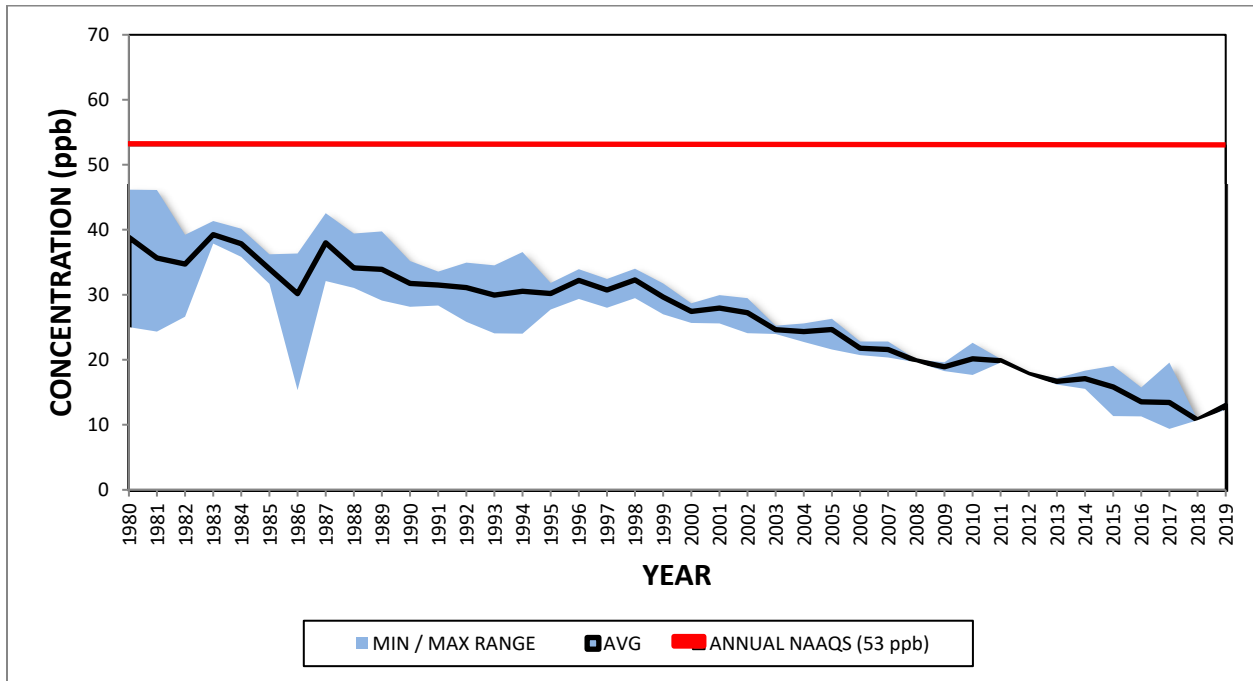
Nitrogen dioxide is a light brown gas that is an important component of urban haze. The compound is created primarily from fuel combustion in motor vehicles, utilities, and industrial sources.

Nitrogen dioxide can irritate the lungs and lower resistance to respiratory infections such as influenza. Nitrogen oxides (NO<sub>x</sub>) are an important precursor to both ozone and acid rain and can affect both land and water ecosystems. They contribute to the formation of fine particulate matter, haze and reductions in visibility.

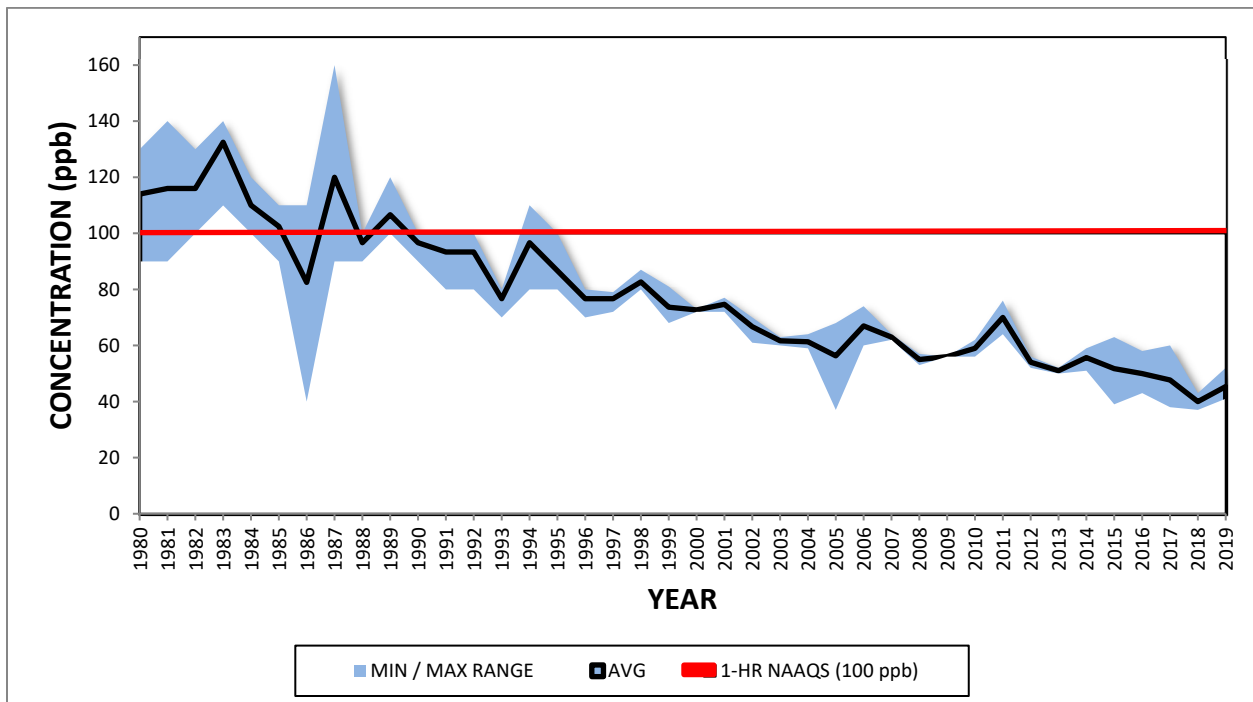
Ambient levels of nitrogen dioxide in Philadelphia are significantly below the NAAQS, showing a sustained downward trend over time.

Figures 6.2 and 6.3 show the NO<sub>2</sub> trends for annual average and daily maximum of one-hour concentrations, respectively.

**Figure 6.2 – NO<sub>2</sub> Trends for Annual Average Concentration**



**Figure 6.3 – NO<sub>2</sub> Trends for 98th Percentile Daily Maximum 1-Hour Concentration**

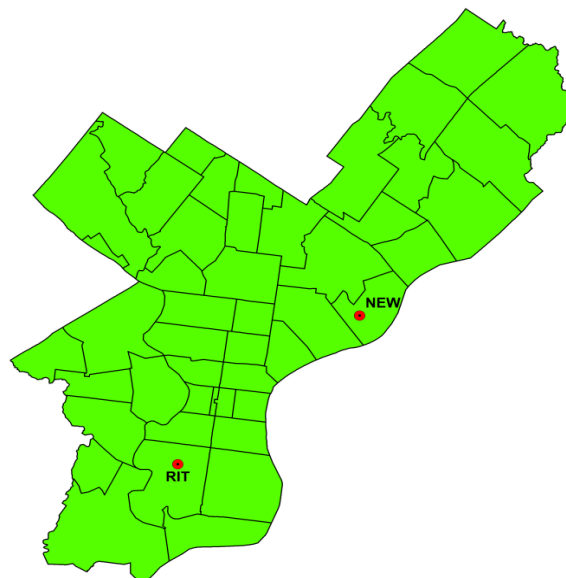


## Sulfur Dioxide (SO<sub>2</sub>)

### NAAQS:

- The 3-year average of the annual 99th percentile of the daily maximum 1-hour average concentrations is less than or equal to 75 ppb.

Figure 7.1 – SO<sub>2</sub> Monitoring Map



Sulfur dioxide is emitted from the burning of fuels that contain sulfur. Industrial grade fuel oils are the primary source in Philadelphia.

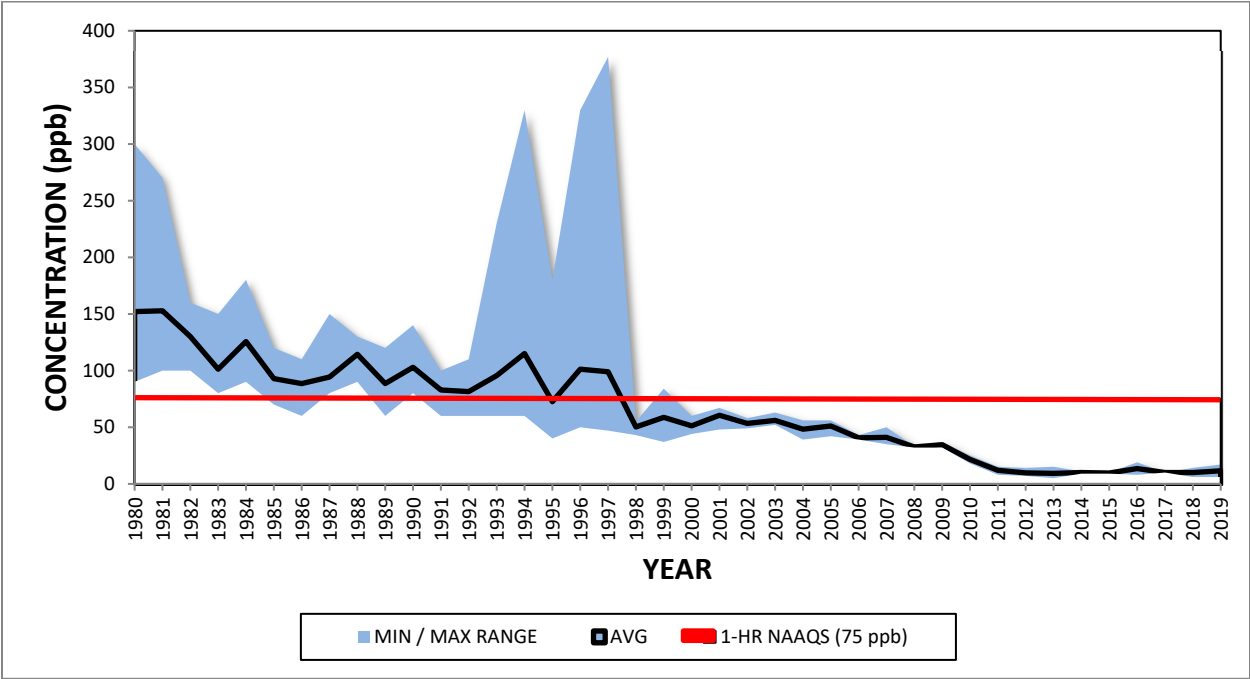
The major health concerns associated with exposure to high concentrations of SO<sub>2</sub> include effects on breathing, respiratory illness, alterations in the lungs' defenses, and aggravation of existing respiratory and cardiovascular disease. Together, SO<sub>2</sub> and NO<sub>x</sub> are the major ingredients of acid rain. SO<sub>2</sub> also plays a significant role in the formation of fine particulate matter.

SO<sub>2</sub> levels are well within air quality standards and show a slow, continued improvement over time. This is mainly due to industry, businesses, and homes changing to fuels with lower sulfur content such as natural gas.

In 2019, the NEW and RIT sites were operating as the monitoring sites for SO<sub>2</sub> as seen in Figure 7.1.

The following graph, Figure 7.2, shows the trends for the one-hour SO<sub>2</sub> concentration for Philadelphia.

Figure 7.2 – SO<sub>2</sub> Trends for 99<sup>th</sup> Percentile Daily Maximum 1-Hour Average Concentration





## Lead (Pb)

### **NAAQS:**

- The maximum arithmetic rolling 3-month mean concentration for a 3-year period is less than or equal to 0.15  $\mu\text{g}/\text{m}^3$  micrograms per cubic meter.

The processing of metals is the major source of lead emissions to the atmosphere. Lead does not travel over great distances in the air and so concentrations vary, with the highest levels near specific industrial sites.

Lead is a metal that is highly toxic when inhaled or ingested. Lead accumulates in the blood, bone, and soft tissue and may affect the kidneys, liver, nervous system and other organs. It also can cause learning difficulties in children.

Ambient lead levels have been decreasing throughout the city due to the elimination of leaded gasoline and greater control of emissions from companies that produce or process lead compounds.

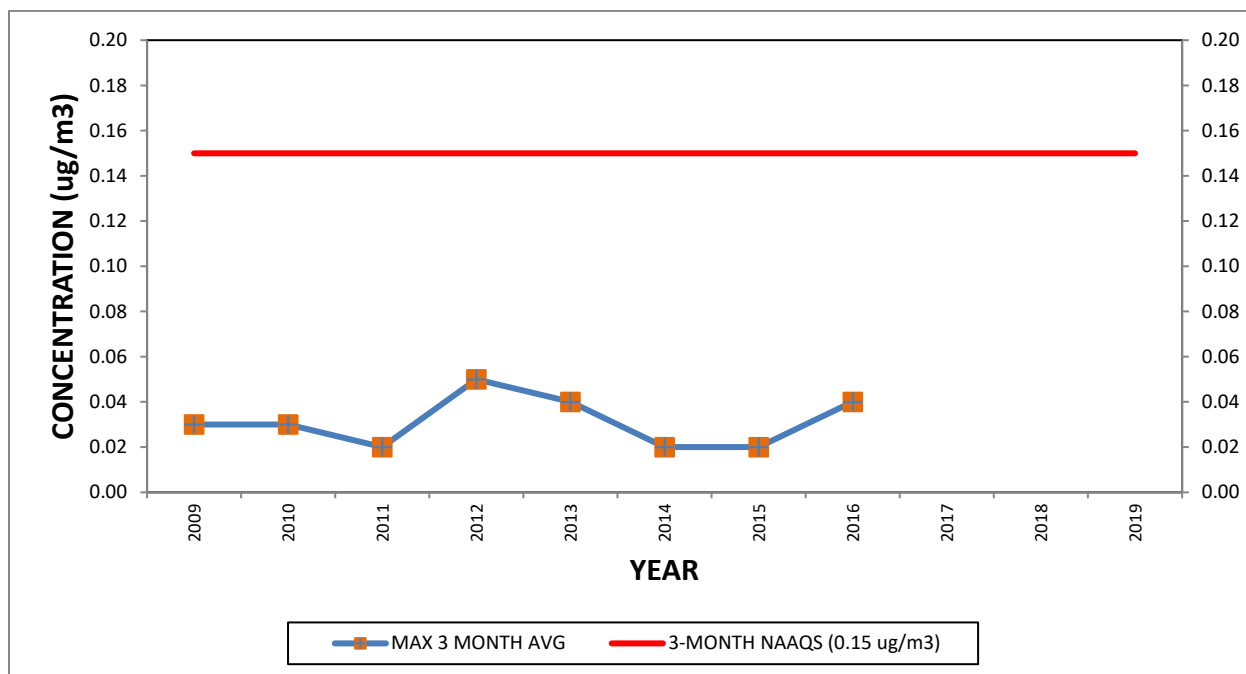
Prior to 1998, lead levels in certain parts of the city were once extremely high due to the concentration of particular industries in the areas near Castor and Delaware Avenues. The levels of lead in these areas have drastically improved and are now comparable to the rest of the city.

On October 15, 2008, the EPA strengthened its regulation for lead. The standard was revised from the 1978 standard of 1.5  $\mu\text{g}/\text{m}^3$  to a level that is 10 times more stringent, 0.15  $\mu\text{g}/\text{m}^3$ , with a different averaging time. For the previous standard, the averaging time used a quarterly average while the new standard uses a rolling 3-month average. The revision is based on more than 6000 studies performed since 1990 on the health effects of high lead concentrations in the bloodstream. The studies show that adverse effects

from lead in the blood occur at a much lower level than previously thought. Figure 8.3 shows the trends for the 2008 lead standard.

As of January 1, 2017, Total Suspended Particulate Lead monitor was shutdown at NEW site. Philadelphia has no sources that emit 0.5 or more tons of Pb per year. On 4/28/17, EPA approved Philadelphia AMS' waiver of the requirement for a source oriented Lead-TSP monitor in Philadelphia effective 1/1/17.

**Figure 8.1 – Lead Trends (Maximum) Rolling 3-Month Average**



## Particulate Matter (PM<sub>10</sub>, PM<sub>2.5</sub>)

Particulate matter is the general term used for a mixture of solid particles and liquid droplets found in the air. These particles come in a wide range of sizes and originate from stationary, mobile, and natural sources.

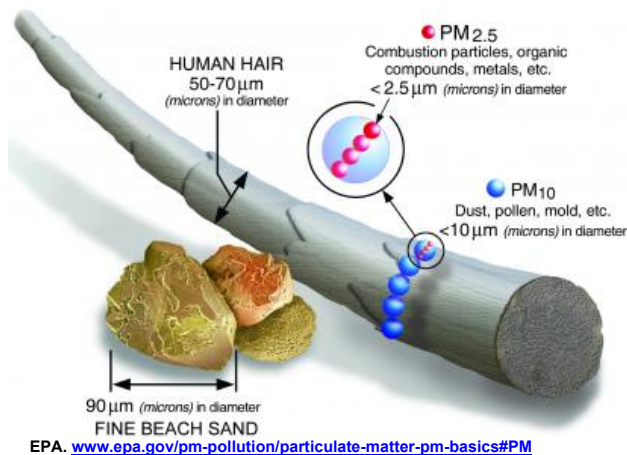
PM<sub>10</sub> and PM<sub>2.5</sub> refer to small particulates that measure less than 10 micrometers (0.00001 meters) and 2.5 micrometers (0.0000025 meters) in diameter, respectively. In addition to health problems, particulate matter can cause reduced visibility, soiling, and damage to materials such as buildings. Particles of this size remain airborne for long periods of time and disperse in uniform concentrations across wide areas, crossing geographic boundaries.

In 1997, the EPA set a separate standard for PM<sub>2.5</sub>. Particles in the PM<sub>2.5</sub> size range are able to travel deeply into the respiratory tract, reaching the lungs. Exposure to fine particles can cause short-term health effects such as eye, nose, throat and lung irritation, coughing, sneezing, runny nose and shortness of breath. Exposure to fine particles can also affect lung function and worsen medical conditions such as asthma and heart disease. Scientific studies have linked increases in daily PM<sub>2.5</sub> exposure with increased respiratory and cardiovascular hospital admissions, emergency department visits and deaths. Recent studies suggest that long term exposure to particulate matter may be associated with increased rates of bronchitis and reduced lung function.

Particles come in a wide variety of shapes and sizes, which affect their impacts on the environment and human health. Bigger particles, such as dust, are easier to see and can cause problems, but smaller particles are likely to be worse for our health.

Fine particles are treated as though they are a single pollutant, but fine particles come from many different sources and are composed of thousands of different compounds. Fortunately, these compounds fall into a few dominant categories: sulfates, nitrates, ammonium compounds, soil, organic carbon compounds, and elemental carbon. Water is nearly always an important and variable part of PM, and sea salt is often significant near the coast. Given the complex composition of PM, it is no surprise that its chemistry is also complex. Particles may be dry or wet. When the wind blows hard enough, soil, silt, and sand can be lifted from the surface. Human activities such as mining, construction, plowing, and driving on unpaved roads also lift particles into the air. Soot, also referred to as black carbon or elemental carbon, is emitted directly by diesel engines and forest fires, among other sources. Most individual particles are likely mixtures of different substances, the products of growing by collisions with other particles and by taking on gases.

**Figure 9 – Size Comparisons for PM Particles**

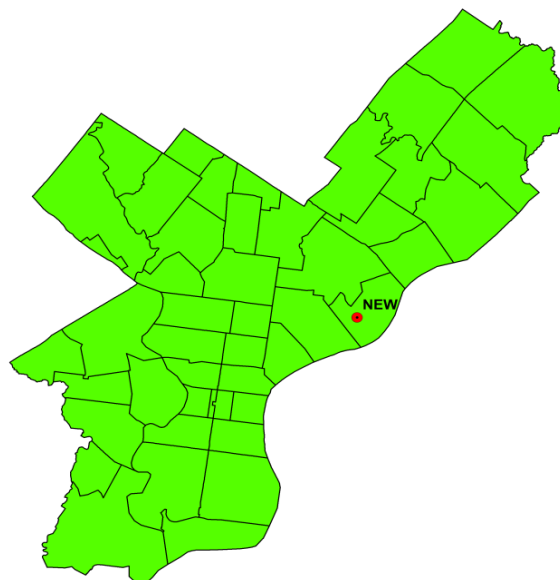


## PM<sub>10</sub>

### NAAQS:

- 150 µg/m<sup>3</sup> for a daily 24-hour average concentration not to be exceeded more than once per year on average over a 3-year period.

Figure 9.1 – PM<sub>10</sub> Monitoring Map



Particulate matter levels have been decreasing due to regulations limiting the amount of emissions allowed and the change to cleaner fuels, for example, switching from oil to natural gas by industry, businesses and homes.

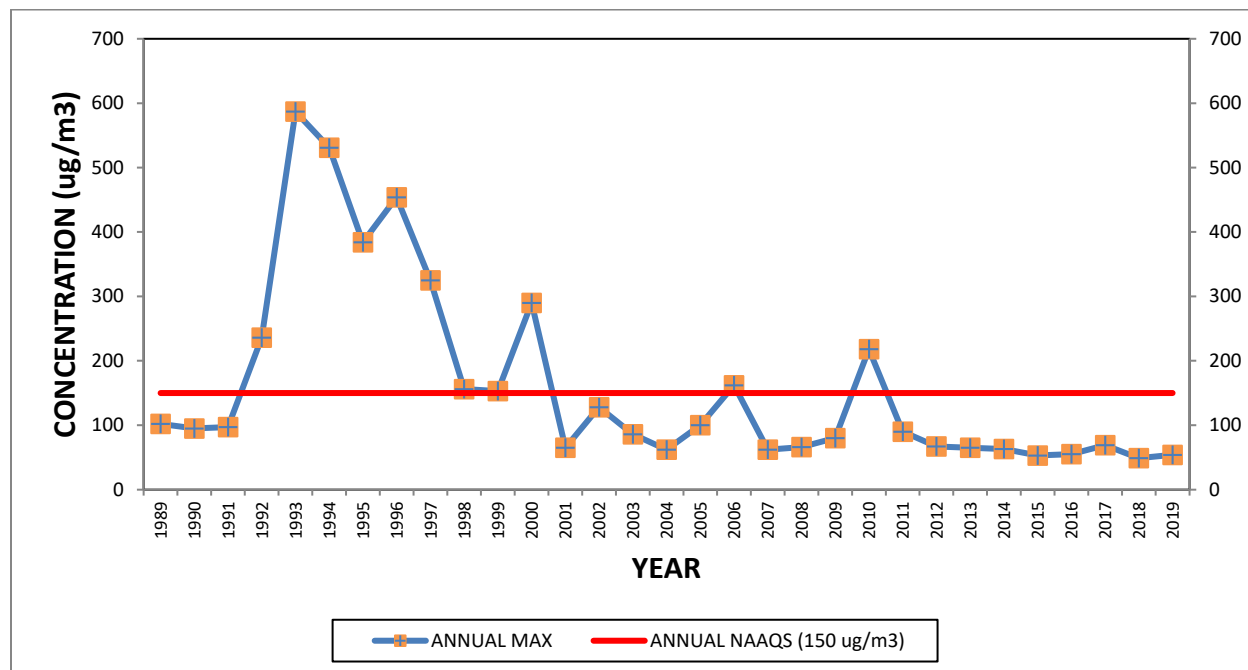
During the mid-1990s, particulate emissions from several sources in the area of Castor and Delaware Avenues caused extremely high-localized measurements and the levels were many times higher than those measured at other city locations. Specific actions to abate these sources have resulted in air quality that now meets the national standards and are now comparable to levels in the rest of the city.

The EPA revoked the annual standard for PM<sub>10</sub> on December 17, 2006, due to a lack of evidence linking health problems to long-term exposure to coarse particle pollution.

As seen in Figure 9.1, there is one PM<sub>10</sub> monitoring site, NEW.

Figure 9.2 shows the trends for PM<sub>10</sub> for the maximum 24-hour average concentration from all monitoring sites.

**Figure 9.2 – PM<sub>10</sub> Trends for the Highest 24-Hour Average Concentration for All Monitoring Sites**

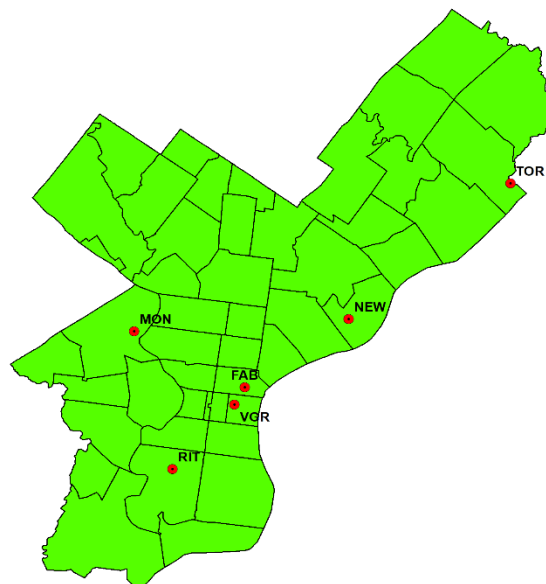


## PM<sub>2.5</sub>

### NAAQS:

- The 3-year average of the annual arithmetic mean concentration is less than or equal to 12.0 µg/m<sup>3</sup>.
- The 3-year average of the 98th percentile 24-hour concentration is less than or equal to 35 µg/m<sup>3</sup>.

Figure 10.2 – PM<sub>2.5</sub> Monitoring Map



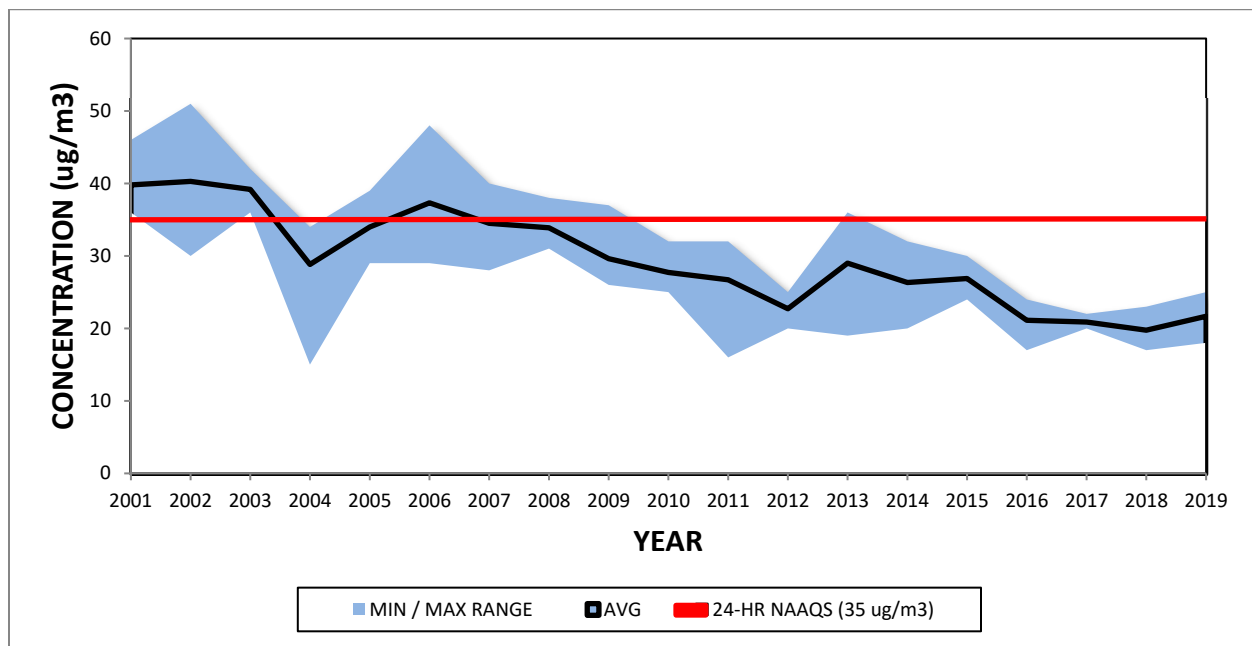
PM<sub>2.5</sub> consists of those particles that are less than 2.5 micrometers in diameter. They are also referred to as "fine" particles. Fine particles result from fuel combustion from motor vehicles, power generation, and industrial facilities, as well as from residential fireplaces and wood stoves. A significant amount of fine particles are also formed in the atmosphere by the transformation of gaseous emissions such as SO<sub>2</sub>, NO<sub>x</sub>, VOCs, and ammonia.

Fine particles can accumulate in the respiratory system and are associated with numerous health effects such as premature death, respiratory symptoms and disease, and decreased lung function. Sensitive groups that appear to be at the greatest risk for such effects include children, seniors, and individuals with cardiopulmonary disease or respiratory ailments such as asthma.

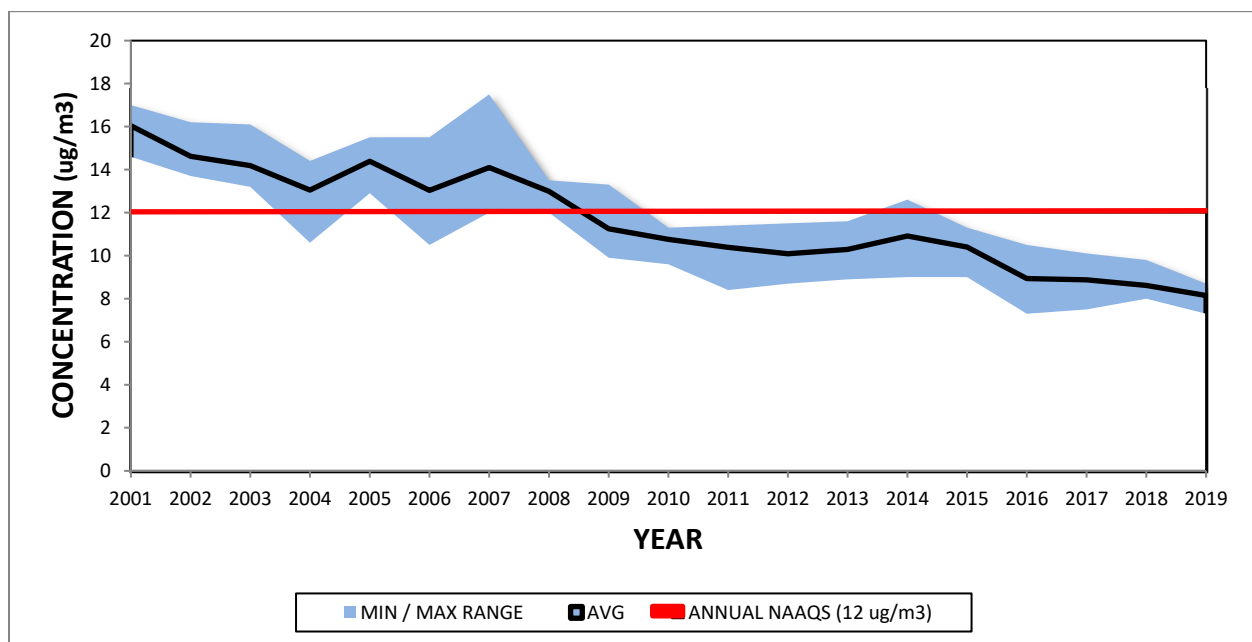
In 2019, there were six PM<sub>2.5</sub> monitoring sites in the network.

Figures 10.2 and 10.3 show the trends for the 24-hour concentration and the annual mean, respectively.

**Figure 10.2 – PM<sub>2.5</sub> Trends for 98<sup>th</sup> Percentile 24-Hour Concentration**



**Figure 10.3 – PM<sub>2.5</sub> Trends for Annual Mean Concentration**

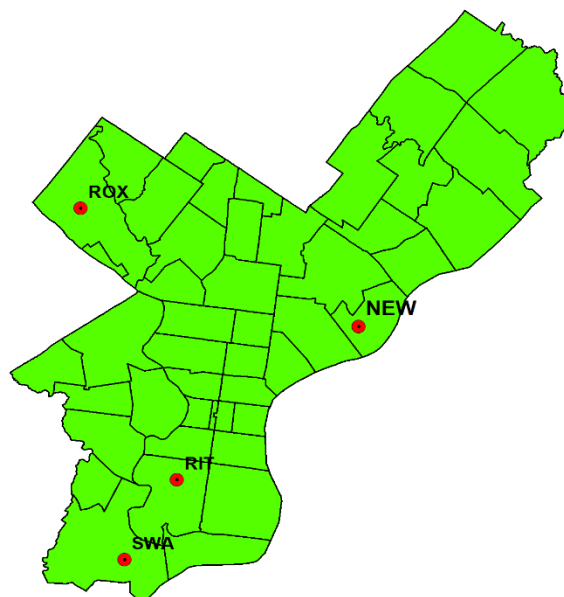




## Air Toxics

Air toxics, also referred to as toxic air pollutants or hazardous air pollutants (HAPs), are substances that cause adverse health effects or environmental damage. The Federal Clean Air Act Amendments (CAAA) of 1990 list 187 pollutants or chemical groups as HAPs. Examples of air toxics include heavy metals (such as beryllium), organic chemicals (such as formaldehyde), polycyclic organic matter (POM, which are formed primarily by combustion), benzene (which is found in gasoline), pesticides, fine mineral fibers, and asbestos. HAPs are emitted from stationary sources (large industrial facilities), nonpoint sources (dry cleaners, gas stations, and other small facilities), as well as mobile sources (trucks and buses).

Figure 11.1 – Air Toxics Monitoring Map



There is less information known about the health impact from the 187 HAPs than there are for criteria pollutants, and no national standards exist for them. However, a number of these pollutants are known or suspected to be carcinogenic, and there is no known “safe concentration.” The danger posed by toxics is often referred to in terms of risk. Risk is defined as the likelihood of a negative outcome from a certain level of a specific chemical, or the measure of a chance that health problems will occur. For example, many toxics cause cancer, while others cause respiratory problems, birth defects, neurological or immune response problems, and other health concerns. Toxics have varying degrees of danger, and some will cause harm with a very small amount of the substance while others require large amounts to have a negative effect. A cancer risk level of one in a million implies a likelihood that up to one person out of one million equally exposed people would contract cancer if exposed continuously (24 hours per day) to the specific concentration over 70 years (an assumed lifetime). This risk is calculated as additional to those cancer cases that would normally occur in an unexposed population of one million people.

AMS is helping to reduce HAPs in Philadelphia by enforcing Federal, State, and locally mandated programs that limit emissions from stationary and area sources. Many toxic emissions have been reduced by regulations designed to bring Philadelphia into compliance with the NAAQS for Ozone. In addition, Philadelphia enforces the National Emission Standards for Hazardous Air Pollutants (NESHAPs), a program designed to reduce emissions from existing major and area sources, as well as New Source Performance Standards (NSPS), which limit toxic emissions from new sources. In 2010, Air Management Regulation XIV – Control of Emissions from Dry Cleaning Facilities was promulgated. This regulation restricted the use of perchloroethylene (PERC) in dry

cleaning facilities that share a common wall with businesses or residences. Since 2010, ambient concentrations of PERC have been decreasing.

Since diesel emissions are a significant but unquantified contributing factor in determining health risks from toxic emissions, AMS continues working to promote voluntary emissions reductions from diesel vehicles and to bring clean diesel technology to the Philadelphia area. In addition, AMS is currently working with other City departments to enforce Mayor's Executive Order 1-07 which requires all public works and demolition contracts to use clean diesel technology. This program is expected to significantly reduce particulate matter, hydrocarbons, and carbon monoxide from diesel vehicles contracted by the City, resulting in substantial annual health benefits.

As part of EPA's National Air Toxics Assessment (NATA) activities, 180 air pollutants were assessed for either lifetime cancer risk or non-cancer hazard due to inhalation. NATA is EPA's ongoing comprehensive evaluation of air toxics in the U.S. These activities include: expansion of air toxics monitoring, improving and periodically updating emission inventories, improving national- and local-scale modeling, continued research on health effects and exposures to both ambient and indoor air, and improvement of assessment tools.

The goal of NATA is to identify air toxics which are of greatest potential concern, in terms of contribution to population risk. The results are used to establish strategies, priorities, and programs to reduce air toxics emissions. In 2018, EPA released the results of the NATA for 2014.

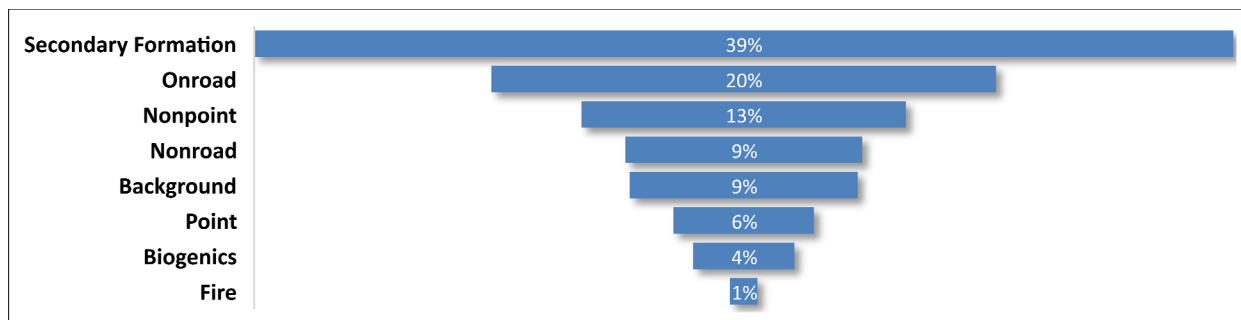
Nationwide, the results of the 2014 NATA can be summarized as follows:

- Nationwide, total emissions of air toxics are declining, and air quality monitoring data show that concentrations of many toxics in the air, such as benzene, also are trending downward.
- The 2014 NATA estimates that the nationwide average cancer risk from air toxics exposure is 30 in 1 million. About half of that risk comes from the formation of formaldehyde – produced when other pollutants chemically react in the air. This is known as secondary formation, and comes from emissions from industries, mobile sources, and natural sources. The other half of the nationwide cancer risk comes from pollution that is directly emitted to the air.

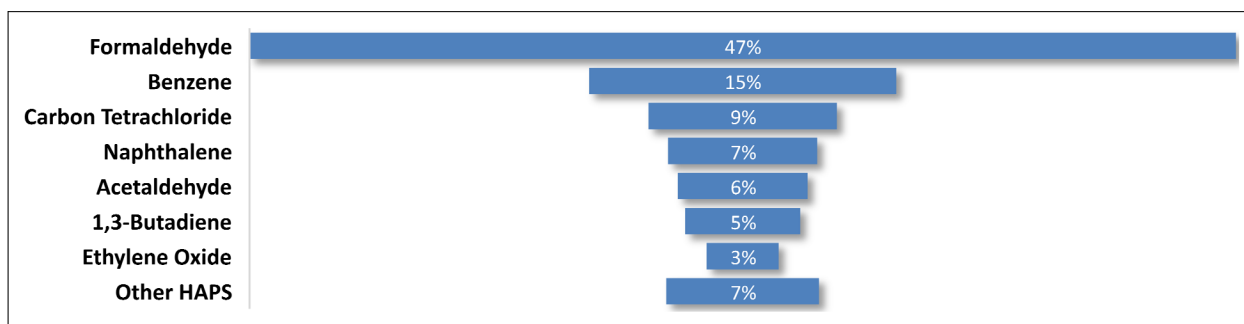
Specifically, the results of the 2014 NATA for Philadelphia County are summarized below:

- Total cancer risk was 36.8 in 1 million.
- Secondary formation accounted for 39 percent of cancer risk and on-road mobile sources accounted for 20 percent (see Figure 11.2).
- The pollutants contributing most to cancer risk were formaldehyde (47 percent), benzene (15 percent), and carbon tetrachloride (9 percent) (see Figure 11.3).

**Figure 11.2 – Source Category Contribution to 2014 NATA Cancer Risk**



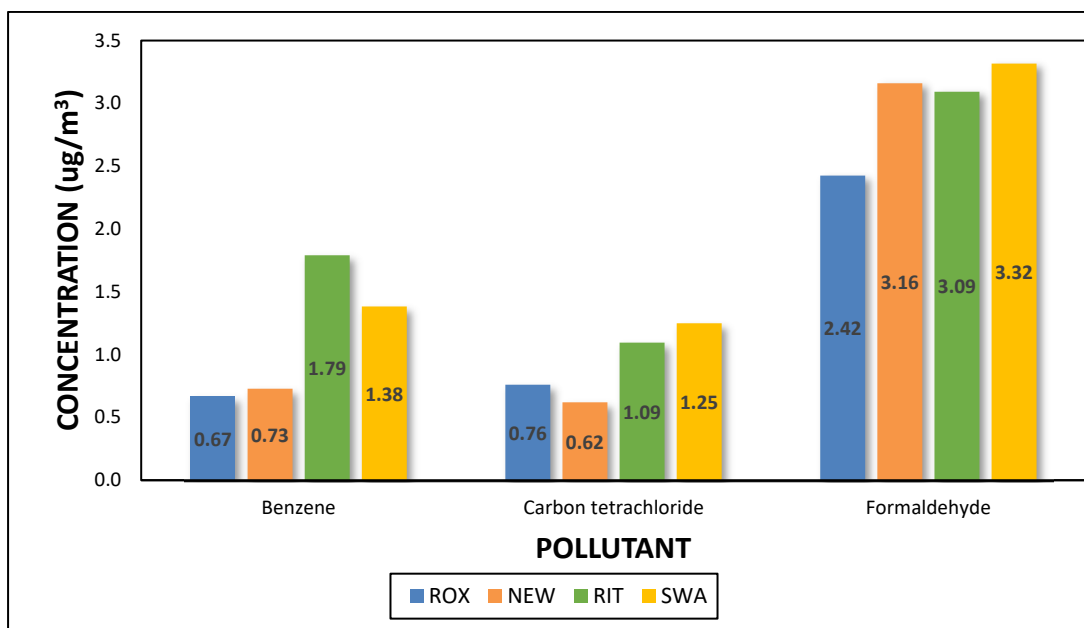
**Figure 11.3 – HAP Contributions to 2014 NATA Cancer Risk**



Addition information for the 2014 NATA can be found here: <https://www.epa.gov/national-air-toxics-assessment/2014-nata-assessment-results>.

In 2019, AMS monitored air toxics at 4 locations as seen in Figure 11.1. AMS monitors approximately 61 VOCs, 7 carbonyl compounds, and 59 hydrocarbons. Three different sampling methods are employed to obtain their concentrations in ambient air. VOCs are collected in 6-liter stainless steel canisters every six days and analyzed via Gas Chromatography/Mass Spectrometry. The carbonyls are also collected every six days on a cartridge and analyzed using a High Performance Liquid Chromatography. The hydrocarbons are collected and analyzed hourly every day via auto-Gas Chromatography. Figure 11.4 shows the 2019 mean concentrations for formaldehyde, benzene, and carbon tetrachloride which showed the highest contribution to cancer risk in the 2014 NATA.

Figure 11.4 – 2019 Mean Concentrations for Benzene, Carbon Tetrachloride, and Formaldehyde<sup>9</sup>



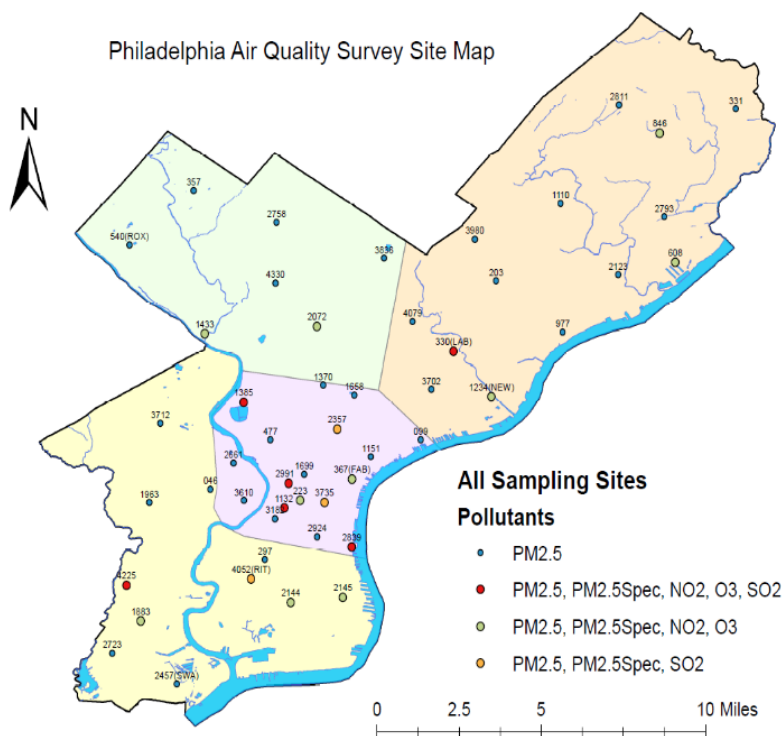
Data for air toxics can be found on the EPA websites <https://www.epa.gov/outdoor-air-quality-data/monitor-values-report-hazardous-air-pollutants> or [https://aqs.epa.gov/aqsweb/airdata/download\\_files.html](https://aqs.epa.gov/aqsweb/airdata/download_files.html).

<sup>9</sup> <https://www.epa.gov/outdoor-air-quality-data/monitor-values-report-hazardous-air-pollutants>. Downloaded 8/4/2020.

## Philadelphia Air Quality Survey

In 2019, AMS continued to collect monitoring data from the Philadelphia Air Quality Survey (PAQS). This project set up 50 street level, neighborhood-oriented air sampling sites throughout the City to sample the ambient air for PM<sub>2.5</sub>, NO<sub>2</sub>, SO<sub>2</sub>, and O<sub>3</sub>. The sites also contain meteorological sensors as well. PAQS captures the seasonal changes and neighborhood-to-neighborhood spatial variances in air quality. The data results may serve as a basis for future work, including: provide policy recommendations for reducing pollution from congested traffic, diesel vehicles and winter time fuel burning; analyze the relations between air quality and land use characteristics in different neighborhoods; and study public health impact of air pollution in the City. At the end of 2019, the project completed 26 two-week sampling sessions. Additional information regarding PAQS is provided in the latest version of the Air Monitoring Network Plan (<https://www.phila.gov/documents/air-management-reports-and-documents/>).

**Figure 12 – PAQS Survey Map**



## Appendix A: Websites

[www.airnow.gov](http://www.airnow.gov) – The AQI (Air Quality Index) tells you how clean the air is and whether it will affect your health. Through AIRNow, EPA, NOAA, NPS, state, and local agencies work together to report current and forecast conditions for ozone and particle pollution.

[www.airqualitypartnership.org](http://www.airqualitypartnership.org) – Ground level ozone and particle pollution forecasts.

[www.atsdr.cdc.gov](http://www.atsdr.cdc.gov) – Agency for Toxic Substances and Disease Registry: public health statements on specific toxics and the effects of exposure.

[www.cleanair.org](http://www.cleanair.org) – Clean Air Council.

[www.delawarevalley.enviroflash.info/about.cfm](http://www.delawarevalley.enviroflash.info/about.cfm) – Sign up for air quality forecasts.

[www.depweb.state.pa.us](http://www.depweb.state.pa.us) – Pennsylvania Department of Environmental Protection.

[www.epa.gov](http://www.epa.gov) – U.S. Environmental Protection Agency.

<https://www.epa.gov/outdoor-air-quality-data> – EPA's AQS (Air Quality System) database for air monitoring data.

[www.epa.gov/echo](http://www.epa.gov/echo) – EPA's ECHO (Enforcement & Compliance History Online) database for compliance inspections conducted by EPA or state/local government, violations, enforcement actions, and penalties assessments in response to environmental law violations.

[www.lungusa.org](http://www.lungusa.org) – American Lung Association website: Information on lung health, air pollution, and related matters.

<https://www.phila.gov/departments/departments-of-public-health/about-us/contact-us/> – Philadelphia Department of Public Health, Air Management Services.

[www.phila.gov/services/mental-physical-health/environmental-health-hazards/air-quality/](http://www.phila.gov/services/mental-physical-health/environmental-health-hazards/air-quality/) – Philadelphia's Air Quality Website, provides the most up-to-date information about the air quality in Philadelphia and lets you know what you should do to protect your health if the air quality is unhealthy.

## Appendix B: Air Quality Data Tables<sup>10</sup>

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<sup>10</sup> All data (including exceptional events data) in this appendix downloaded on 8/4/20 from EPA's AirData website (<https://www.epa.gov/outdoor-air-quality-data/monitor-values-report>).



**Table 2 – Ozone 4<sup>th</sup> Highest Daily Maximum 8-Hour Concentrations (ppm)**

Year	AQS SITE ID - POC												
	421010004-1	421010004-2	421010014-1	421010019-1	421010021-1	421010023-1	421010024-1	421010026-1	421010027-1	421010029-1	421010048-1	421010136-1	421011002-1
1980	0.032		0.113	0.037	0.032	0.062	0.128	0.025	0.026	0.028			
1981			0.100				0.108						
1982			0.111			0.087	0.103						
1983			0.120			0.102	0.121						
1984			0.106			0.093	0.108						
1985			0.111			0.083	0.116						
1986			0.103			0.080	0.105						
1987			0.104			0.085	0.118						
1988			0.130			0.088	0.128						
1989			0.085			0.097	0.077						
1990			0.097			0.097	0.101						
1991			0.106			0.095	0.112						
1992			0.087			0.076	0.091						
1993	0.086		0.092			0.090	0.097					0.103	
1994	0.080		0.095			0.070	0.092					0.085	
1995	0.091		0.096				0.113					0.092	
1996	0.087		0.093				0.092					0.072	
1997	0.067		0.096				0.101					0.074	
1998	0.077		0.095				0.093					0.088	
1999	0.073		0.081				0.060					0.098	
2000	0.067		0.086				0.089					0.082	
2001	0.074		0.097				0.097					0.086	
2002	0.082		0.098				0.110					0.094	
2003	0.069		0.084				0.086					0.074	
2004	0.057		0.077				0.091					0.073	
2005	0.066		0.083				0.094					0.068	
2006	0.066		0.076				0.085					0.081	
2007	0.073		0.081				0.095					0.082	
2008	0.062						0.087						
2009	0.059						0.072						
2010	0.077						0.088						
2011	0.070						0.089						0.088
2012	0.065						0.085						0.083
2013	0.047						0.068				0.036		0.071
2014	0.058						0.072				0.068		
2015	0.057						0.079				0.078		
2016	0.069						0.080				0.076		
2017	0.042						0.076				0.076		
2018		0.071					0.079				0.076		
2019		0.067					0.071				0.072		

**Table 3 – Carbon Monoxide 2nd Highest 8-Hour Average Concentrations (ppm)**

Year	AQS SITE ID - POC																	
	421010004-1	421010004-2	421010014-1	421010019-1	421010021-1	421010024-1	421010026-1	421010026-2	421010027-1	421010029-1	421010029-2	421010045-1	421010046-1	421010047-1	421010048-1	421010051-1	421010075-1	421010076-1
1980	7.1	4.6		4.9	6.8	5.5		5.3	5.8		5	8.2						
1981	9.9		3.3				7.4		9.3	9.4		9.5	7.6					
1982	9.9		3.9						11.9	7		7.9	7.6	7.3				
1983	6.6								10.6	6.1		6.1	5.6	5.6				
1984	9.5								10	6.3		7.8	6.6	6.2				
1985	6.8								7.6	5.1		6.3	4.9	5.8				
1986	6.3								7.9	5.3			5.3	6.3				
1987	7.8								10	7.3			5.3	5.4		8.7		
1988	5.9								6.3	4.9			3.9	4.3		6.9		
1989	11.6								6.4	6.9				6.9		9.5		
1990	6.8								6.4	4.7				5.4		6.1		
1991	5.6								7.1	4.4				4.3		5.1		
1992	6.4								6.8	5				4.1		5.4		
1993	6.6								4.6	5.6				5		5.4		
1994	8.3									5.1				5.4		5.9		
1995	4.5								4.8	4.1				3.9		5.6		
1996	5.6								5.4	4.5				3.6		5.6		
1997	5.3								5	3.3				3.1		3.6		
1998	4.6								4.9	3.2				3.3		3.8		
1999	4.9								4.9	4.4				3.3		4.2		
2000	3.3								4	3				3				
2001	4								4.7	2.9				2.5				
2002	2.9								2.9	2				1.9				
2003	2.4								3.9	2.2				2				
2004	3.1								3.4	2.5				2.7				
2005	2.1								3.4	0.8				2.1				
2006	2.3								3.2					1.4				
2007	2.6								3.2					5.5				
2008	1.5													0.7				
2009	2																	
2010	1.8																	
2011	1.7																	1.2
2012	1.5																	1.5
2013	1.7														1.1			1.3
2014	1.3														1.3		1.4	
2015	1.6														1.5		1.6	
2016	1.9														1.7		1.6	
2017	0.9														1.4		1.2	2.6
2018															1		1.3	1.1
2019															1.7		1.5	1.4

**Table 4 – Carbon Monoxide Highest 1-Hour Average Concentrations (ppm)**

Year	AQS SITE ID - POC																	
	421010004-1	421010004-2	421010014-1	421010019-1	421010021-1	421010024-1	421010026-1	421010026-2	421010027-1	421010029-1	421010029-2	421010045-1	421010046-1	421010047-1	421010048-1	421010051-1	421010075-1	421010076-1
1980	14	8		7	11	8		11	15		10	16						
1981	14		6				12		16	15		16	12					
1982	18		12						23	14		15	13	19				
1983	10								19	11		11	9	8				
1984	17								17	18		18	12	12				
1985	12								16	11		10	8	9				
1986	10								12	13			9	20				
1987	12								15	13			8	9		13		
1988	9								12	8			6	9		11		
1989	18								11	12				11		16		
1990	10								11	8				9		14		
1991	10								13	9				9		9		
1992	9								10	8				8		10		
1993	9								9	10				8		9		
1994	14									11				10		10		
1995	8								7	8				9		11		
1996	10								10	8				6		14		
1997	10								10	6.4				4.9		6.4		
1998	7.6								8.9	5.8				5.8		5.6		
1999	7.7								6.9	6				5.8		7		
2000	6.5								6.5	4.3				4.7				
2001	5.7								8	6.4				5				
2002	5.4								4.5	3.5				3.5				
2003	3.7								6	3.8				3.2				
2004	4.6								5.5	3.7				3.7				
2005	3.8								6.5	1.3				12.3				
2006	4.4								8.6					2.2				
2007	3.8								4.2					5.6				
2008	3.7													1.3				
2009	3.6																	
2010	3.1																	
2011	2.6																	2.2
2012	2.7																	2.1
2013	2.4														2.1			1.6
2014	2														1.8		2	
2015	2.1														2.5		1.9	
2016	3.6														2.6		2.2	
2017	1.4														2.3		1.9	4.3
2018															1.4		1.9	1.5
2019															2.3		2.4	1.8

**Table 5 – Nitrogen Dioxide Annual Average Concentrations (ppb)**

	AQS SITE ID - POC										
Year	421010004-3	421010022-1	421010023-2	421010024-1	421010026-2	421010027-1	421010029-2	421010047-1	421010048-1	421010075-1	421010076-1
1980	40.89			25.02	46.18	45.82	36.19				
1981	37.4	34.24		24.32	46.12		36.19				
1982	33.05	37.1		26.62			39.23	37.64			
1983	38.15	41.34					39.7	37.87			
1984	38.1	35.82					37.34	40.16			
1985	33.71	31.67					34.34	36.21			
1986	33.28		15.32				36.34	35.67			
1987	32.12						39.28	42.55			
1988	31.06						31.93	39.41			
1989	29.11						32.86	39.74			
1990	28.14						31.91	35.19			
1991	28.35						32.59	33.55			
1992	25.82						32.55	34.93			
1993	24.06						31.22	34.51			
1994	24.01						31.06	36.56			
1995	27.73						31.01	31.81			
1996	29.34						33.35	33.92			
1997	27.99						32.44	31.76			
1998	29.47						33.37	34			
1999	26.99						30.23	31.75			
2000	25.66						27.94	28.69			
2001	25.59						28.32	29.95			
2002	24.09						28.1	29.48			
2003	23.97						24.69	25.23			
2004	22.73						24.69	25.58			
2005	21.57						26.11	26.29			
2006	20.72							22.81			
2007	20.35							22.8			
2008	19.6							20.03			
2009	18.25							19.59			
2010	17.66							22.61			
2011	20.12							19.56			
2012	17.91							17.6			
2013	16.18							17.17			
2014	17.45							18.31		15.51	
2015	18.15							19.07		14.69	11.32
2016	15.73									13.55	11.28
2017	19.55									11.35	9.36
2018										10.69	10.76
2019									12.32	13.43	13.11

**Table 6 – Nitrogen Dioxide 98th Percentile Daily Maximum 1-Hour Concentrations (ppb)**

	AQS SITE ID - POC										
Year	421010004-3	421010022-1	421010023-2	421010024-1	421010026-2	421010027-1	421010029-2	421010047-1	421010048-1	421010075-1	421010076-1
1980	130			90	120	130	100				
1981	110	140		90	130		110				
1982	110	120		100			130	120			
1983	140	140					140	110			
1984	120	100					100	120			
1985	100	110					110	90			
1986	90		40				110	90			
1987	90						160	110			
1988	100						90	100			
1989	100						100	120			
1990	100						90	100			
1991	100						80	100			
1992	80						100	100			
1993	70						80	80			
1994	80						100	110			
1995	80						80	100			
1996	70						80	80			
1997	72						79	79			
1998	80						87	81			
1999	68						81	72			
2000	73						72	73			
2001	72						77	75			
2002	61						70	69			
2003	62						63	60			
2004	64						59	61			
2005	64						37	68			
2006	74							60			
2007	62							64			
2008	57							53			
2009	56							56			
2010	62							56			
2011	76							64			
2012	56							52			
2013	52							50			
2014	59							57		51	
2015	63							56		49	39
2016	58									49	43
2017	60									45	38
2018										37	43
2019									41	43	52

**Table 7 – Sulfur Dioxide 99th Percentile Daily Maximum 1-Hour Average Concentrations (ppb)**

	AQS SITE ID - POC																						
Year	421010004-4	421010004-5	421010014-1	421010019-1	421010020-1	421010021-1	421010022-1	421010022-2	421010023-2	421010023-3	421010024-1	421010026-2	421010026-3	421010027-1	421010029-2	421010029-3	421010047-1	421010048-1	421010048-2	421010055-1	421010136-1	421011002-1	
1980	160		120	150	120	90	300	180	160	140	110	170	110	150	170								
1981	140	120					270		100				140	130	170								
1982	120						130		100					120	160		150						
1983	90						130		80		80			80	150	100	100						
1984	120						180		90		100			130	140		120						
1985	100						100		70		70			100	90		120						
1986	80						110		80		60			100	90		100						
1987	90						150		80		80			90	90		80						
1988	100						130		110		90			110	130		130						
1989	80						100		80		60			80	100		120						
1990	90						110		90		80			100	110		140						
1991	70						100		90		60			70	100		90						
1992	80						100		60		60			80	110		80						
1993	80						110		60		70			60	90		80	230			80		
1994	80						90	80	80		60				100		90	330			90		
1995	60						80				50			50	60		60	180			40		
1996	60						100				50			60	70		90	330			50		
1997	54						87				56			47	65		54	377			52		
1998	53						53				48			48	55		52	43			50		
1999	58						73				37			51	54		84	53			60		
2000	44						56								60						45		
2001	48						60								67						67		
2002	49														58						53		
2003	52														63						53		
2004	47														56					39	51		
2005	51														55					42	56		
2006	40																			43	39		
2007	38																			35	50		
2008	33																			32			
2009	37																			32			
2010	18																			25			
2011	13																			15		8	
2012	7																			14		8	
2013	8																		15	9		5	
2014	11																		9	10			
2015																			9	10			
2016																			19	8			
2017																			10	10			
2018																			14	6			
2019																			17	6			

**Table 8 – Lead (Maximum) Rolling 3-Month Averages ( $\mu\text{g}/\text{m}^3$ )**

Year	AQS SITE ID - POC								
	421010004-5	421010014-1	421010047-1	421010048-1	421010055-1	421010063-1	421010076-1	421010136-1	421010449-1
2009									0.03
2010									0.03
2011									0.02
2012									0.02
2013				0.02					0.05
2014				0.02					0.04
2015				0.02					
2016				0.04					
2017									
2018									
2019									

**Table 9 – PM<sub>10</sub> Highest 24-Hour Average Concentrations (µg/m<sup>3</sup>)**

Year	AQS SITE ID - POC														
	421010003-1	421010004-1	421010004-2	421010037-1	421010038-1	421010047-1	421010048-1	421010048-2	421010049-1	421010136-1	421010149-1	421010449-1	421010449-2	421010649-1	421010649-2
1988	65	83		76	92	91									
1989		96		97	102	96									
1990		72		79	95	87									
1991		85	87	76	97	84									
1992		54		59	101	129					236				
1993		81		59	105	116			103		587	162			
1994		95		83	94	84			143	83	531	186			
1995		64		68		82			90	107	384	233			
1996		87		54		86			129	63	454	247			
1997		76		117		136			78	113	308	325		288	
1998		60		130		82			63	104	73	130		156	
1999		73		57		24			36	45	111	153		56	
2000		43		73						36	165	290		44	
2001		61		62						61	65	61	64	61	
2002		66		128						72	122	68	99	106	
2003		82		86						63		86		77	
2004		62		55						52		58		52	
2005		57		67						48		79		100	
2006		132								37		93		162	161
2007		44										37		61	62
2008		53										43		64	66
2009		49										41		80	78
2010		57										67		218	151
2011		46										59		64	90
2012		67										61			
2013		29					65					36			
2014							63					30			
2015							53								
2016							55								
2017							69								
2018							49								
2019								54							



**Table 10 – PM<sub>2.5</sub> 98th Percentile 24-Hour Concentrations (µg/m<sup>3</sup>)**

	AQS SITE ID - POC																							
Year	421010003-1	421010004-1	421010004-2	421010014-1	421010020-1	421010024-1	421010027-1	421010047-1	421010047-3	421010047-4	421010048-1	421010048-3	421010052-1	421010055-1	421010055-3	421010056-1	421010057-1	421010057-3	421010057-4	421010075-1	421010076-1	421010136-1	421011002-1	421011002-3
1999		39			30	33		33														34		
2000		41			32	38	57	39														39		
2001		40			36	37		40														46		
2002		41		30	40	38		39					51									43		
2003		40		39	39	39		42														36		
2004		34		15	29	33		32														30		
2005	29	36			33	36		39														31		
2006	29	38	36			35		48								#DIV/0!						38		
2007		35	36			34		40		38							33		28			32		
2008		35	33			31		38		33				35			33		33					
2009		26	34			26		37		29				29			28		28					
2010		28	25			25		32		27				29			28							
2011			24					32		28				31			31						25	16
2012			21					24		22				25			23						20	24
2013			35					26	22	34	36	35		35	25		19	33				24	24	
2014		20	25					28	26		26	28			31			32		21				
2015		25	27						24		26	26			30			27		27	30			
2016		17	24								19	22			22			22		22	21			
2017		22	21								22	20			20			20		22	20			
2018		19	21								17	20			21			18		23	19			
2019											25	23			22			18		22	20			

**Table 11 – PM<sub>2.5</sub> Annual Mean Concentrations (µg/m<sup>3</sup>)**

	AQS SITE ID - POC																							
	421010003-1	421010004-1	421010004-2	421010014-1	421010020-1	421010024-1	421010027-1	421010047-1	421010047-3	421010047-4	421010048-1	421010048-3	421010052-1	421010055-1	421010055-3	421010056-1	421010057-1	421010057-3	421010057-4	421010075-1	421010076-1	421010136-1	421011002-1	421011002-3
Year																								
1999		14.6			13.3	13		15.5														14.5		
2000		14.9			14.7	14.7	23.4	17.1														14.8		
2001		16.5			15.4	14.6		17														16.7		
2002		14.8		14.5	14.4	14.3		16.2					13.7									14.4		
2003		14.8		13.3	13.7	13.2		16.1														14		
2004		13.9		10.6	13.9	12.8		14.4														12.7		
2005	14.4	14.2			15.5	12.9		15.1														14.2		
2006	10.5	13.5	13.2			12.4		15.5														13.1		
2007		13.7	14.1			12.9		14.4		14.9							12		17.5			13.3		
2008		13	12.5			12		13.5		13.2				13.5			13.3		12.9					
2009		10.9	11.3			9.9		11.1		11.1				11.3			11.1		13.3					
2010		10.7	11			9.6		10.9		10.9				11.3			10.9							
2011			8.9					11.3		11.4				11.4			11.4						9.9	8.4
2012			9.7					10.2		10.1				10.3			10.1						8.7	11.5
2013			9.2					10	8.9	11	10.4	11		11	11.1		10.2	11.6				9.4	9.7	
2014		9	9.8					11.3	11.3		10.3	11.1			12.6			12.1		10.7				
2015		9.7	10.3						11.3		10.3	10.1			11.2			11		10.7	9			
2016		7.3	8								8.9	9.8			10.5			9.4		9.4	8.2			
2017		7.5	8.2								9.1	9			10			10.1		8.5	8.6			
2018		8.1	8								8	8.6			9.8			8		9.7	8.7			
2019											7.9	8.3			8.4			7.3		8.7	8.3			

**Table 12 – AQS Site ID information**

<b>AQS Site ID</b>	<b>Address</b>
421010003	Community Health Services; 500 S Broad Street
421010004	1501 E. Lycoming Ave.
421010014	Roxy Water Pump Station; Eva & Dearnley Streets
421010019	Fire Boat Station; Allegheny Ave & Del River
421010020	Ford Rd-Belmont Ave Water Treat Plant
421010021	Island Rd, East of Airport Circle PHL Int Airport
421010022	Defense Support Center; 20th & Oregon Ave
421010023	SE Sewage Plant; Front & Packer Streets
421010024	Grant & Ashton Roads; Phila NE Airport
421010026	Broad & Spruce Streets; Mobile Trailer
421010027	S W Corner Broad & Butler
421010029	20th & Race Streets
421010037	13th Street & Montgomery Avenue
421010038	4415 Almond (Near Orthodox Street)
421010045	1421 Arch Street
421010046	1206 Chestnut Street
421010047	500 South Broad Street-Parking Lot (CHS)
421010048	3000 Lewis St. (Near Bath St.), Philadelphia, Pa. 19137
421010049	Richmond St. & Wheatsheaf Lane
421010051	323 Race Street
421010052	1439 East Passyunk Avenue
421010055	24th & Ritner Streets
421010056	2851 Island Ave, Eastwick Free Library
421010057	240 Spring Garden Street
421010063	8200 Enterprise Ave
421010075	4901 Grant Avenue & James Street, Philadelphia, Pa. 19114
421010076	4100 Montgomery Drive
421010136	Amtrak, 5917 Elmwood Avenue
421010149	Castor & Carbon Streets on PGW Property
421010449	Castor & Delaware Avenues
421010649	Water Dept. Newpcp Lagoon Area (NEL)
421011002	5200 Pennypack Park, Philadelphia, Pa. 19136

## **Appendix C: History of the National Ambient Air Quality Standards**

**Table 13 – History of the National Ambient Air Quality Standards for Carbon Monoxide<sup>11</sup>**

Final Rule/Decision	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1971  36 FR 8186 Apr 30, 1971	Primary and Secondary	CO	1-Hour period	35 ppm	Maximum, not to be exceeded more than once in a year <sup>12</sup>
			8-hour period	9 ppm	Maximum, not to be exceeded more than once in a year
1985  50 FR 37484 Sept 13, 1985	Primary standards retained, without revision; secondary standards revoked.				
1994  <a href="#">59 FR 38906</a> <a href="#">Aug 1, 1994</a>	Primary standards retained, without revision.				
2011  <a href="#">76 FR 54294</a> <a href="#">Aug 31, 2011</a>	Primary standards retained, without revision.				

<sup>11</sup> <https://www.epa.gov/co-pollution/table-historical-carbon-monoxide-co-national-ambient-air-quality-standards-naaqs>

<sup>12</sup> Second highest, non-overlapping 8-hour average concentration.

**Table 14 – History of the NAAQS for Pb-TSP (Lead in total suspended particles)<sup>13</sup>**

<b>Final Rule/Decision</b>	<b>Primary/Secondary</b>	<b>Indicator</b>	<b>Averaging Time</b>	<b>Level</b>	<b>Form</b>
1978 <a href="#">43 FR 46246</a> Oct 5, 1978	Primary and Secondary	Pb-TSP	Calendar Quarter	1.5 µg/m <sup>3</sup>	Not to be exceeded
1991	Agency released multimedia " <a href="#">Strategy for Reducing Lead Exposures</a> "				
2008 <a href="#">73 FR 66964</a> Nov 12, 2008	Primary and Secondary	Pb-TSP	3-month period	0.15 µg/m <sup>3</sup>	Not to be exceeded
<a href="#">2016</a> <a href="#">81 FR 71906</a> Oct 18, 2016	Primary and secondary standards retained, without revision.				

<sup>13</sup> <https://www.epa.gov/lead-air-pollution/table-historical-lead-pb-national-ambient-air-quality-standards-naaqs>

**Table 15 – History of the NAAQS for Nitrogen Dioxide<sup>14</sup>**

<b>Final Rule/Decision</b>	<b>Primary/ Secondary</b>	<b>Indicator</b>	<b>Averaging Time</b>	<b>Level</b>	<b>Form</b>
1971  36 FR 8186 Apr 30, 1971	Primary and Secondary	NO <sub>2</sub>	Annual	53 ppb <sup>15</sup>	Annual arithmetic average
1985  50 FR 25532 Jun 19, 1985	Primary and secondary NO <sub>2</sub> standards retained, without revision.				
1996  61 FR 52852 Oct 8, 1996	Primary and secondary NO <sub>2</sub> standards retained, without revision.				
2010  75 FR 6474 Feb 9, 2010	Primary	NO <sub>2</sub>	1 hour	100 ppb	98th percentile, 1-hour daily maximum, averaged over 3 years <sup>16</sup>
		Primary annual NO <sub>2</sub> standard retained, without revision.			
2012  <a href="#">77 FR 20218</a> <a href="#">April 3, 2012</a>	Secondary	Existing secondary NO <sub>2</sub> standard (annual) retained, without revision.			

<sup>14</sup> <https://www.epa.gov/no2-pollution/table-historical-nitrogen-dioxide-national-ambient-air-quality-standards-naaqs>

<sup>15</sup> The official level of the annual NO<sub>2</sub> standard is 0.053 ppm, equal to 53 ppb, which is shown here for the purpose of clearer comparison to the 1-hour standard.

<sup>16</sup> The form of the 1-hour standard is the 3-year average of the 98th percentile of the yearly distribution of 1-hour daily maximum NO<sub>2</sub> concentrations.

**Table 16 – History of the NAAQS for Ozone<sup>17</sup>**

<b>Final Rule/Decision</b>	<b>Primary/Secondary</b>	<b>Indicator</b>	<b>Averaging Time</b>	<b>Level</b>	<b>Form</b>
1971 36 FR 8186 Apr 30, 1971	Primary and Secondary	Total photochemical oxidants	1 hour	0.08 ppm	Not to be exceeded more than one hour per year
1979 44 FR 8202 Feb 8, 1979	Primary and Secondary	O <sub>3</sub>	1 hour	0.12 ppm	Attainment is defined when the expected number of days per calendar year, with maximum hourly average concentration greater than 0.12 ppm, is equal to or less than 1
1993 58 FR 13008 Mar 9, 1993	EPA decided that revisions to the standards were not warranted at the time				
1997 62 FR 38856 Jul 18, 1997	Primary and Secondary	O <sub>3</sub>	8 hours	0.08 ppm	Annual fourth-highest daily maximum 8-hr concentration, averaged over 3 years
2008 73 FR 16483 Mar 27, 2008	Primary and Secondary	O <sub>3</sub>	8 hours	0.075 ppm	Annual fourth-highest daily maximum 8-hr concentration, averaged over 3 years
2015 <a href="#">80 FR 65292</a> <a href="#">Oct 26, 2015</a>	Primary and Secondary	O <sub>3</sub>	8 hours	0.070 ppm	Annual fourth-highest daily maximum 8 hour average concentration, averaged over 3 years

<sup>17</sup> <https://www.epa.gov/ozone-pollution/table-historical-ozone-national-ambient-air-quality-standards-naaqs>



**Table 17 – History of the NAAQS for Particulate Matter<sup>18</sup>**

<b>Final Rule/Decision</b>	<b>Primary/ Secondary</b>	<b>Indicator<sup>19</sup></b>	<b>Averaging Time</b>	<b>Level</b>	<b>Form</b>
1971 36 FR 8186 Apr 30, 1971	Primary	TSP	24 hour	260 µg/m <sup>3</sup>	Not to be exceeded more than once per year
1971 36 FR 8186 Apr 30, 1971	Primary	TSP	Annual	75 µg/m <sup>3</sup>	Annual geometric mean
1971 36 FR 8186 Apr 30, 1971	Secondary	TSP	24 hour	150 µg/m <sup>3</sup>	Not to be exceeded more than once per year
1971 36 FR 8186 Apr 30, 1971	Secondary	TSP	Annual	60 µg/m <sup>3</sup>	Annual geometric mean
1987 52 FR 24634 Jul 1, 1987	Primary and Secondary	PM <sub>10</sub>	24 hour	150 µg/m <sup>3</sup>	Not to be exceeded more than once per year on average over a 3-year period
1987 52 FR 24634 Jul 1, 1987	Primary and Secondary	PM <sub>10</sub>	Annual	50 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM <sub>2.5</sub>	24 hour	65 µg/m <sup>3</sup>	98th percentile, averaged over 3 years
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM <sub>2.5</sub>	Annual	15.0 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years <sup>3, 4</sup>
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM <sub>10</sub>	24 hour	150 µg/m <sup>3</sup>	Initially promulgated 99th percentile, averaged over 3 years; when 1997 standards for PM <sub>10</sub> were vacated, the form of 1987 standards remained in place (not to be exceeded more than once per year)

<sup>18</sup> <https://www.epa.gov/pm-pollution/table-historical-particulate-matter-pm-national-ambient-air-quality-standards-naaqs>

<sup>19</sup> TSP = Total Suspended Particles

					on average over a 3-year period) <sup>5</sup>
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM <sub>10</sub>	Annual	50 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years
2006 <a href="#">71 FR 61144</a> Oct 17, 2006	Primary and Secondary	PM <sub>2.5</sub>	24 hour	35 µg/m <sup>3</sup>	98th percentile, averaged over 3 years <sup>6</sup>
2006 <a href="#">71 FR 61144</a> Oct 17, 2006	Primary and Secondary	PM <sub>2.5</sub>	Annual	15.0 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years <sup>2,7</sup>
2006 <a href="#">71 FR 61144</a> Oct 17, 2006	Primary and Secondary	PM <sub>10</sub>	24 hour	150 µg/m <sup>3</sup>	Not to be exceeded more than once per year on average over a 3-year period
2012 <a href="#">78 FR 3085</a> Jan 15, 2013	Primary	PM <sub>2.5</sub>	Annual	12.0 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years <sup>2,7</sup>
2012 <a href="#">78 FR 3085</a> Jan 15, 2013	Secondary	PM <sub>2.5</sub>	Annual	15.0 µg/m <sup>3</sup>	Annual arithmetic mean, averaged over 3 years <sup>2,7</sup>
2012 <a href="#">78 FR 3085</a> Jan 15, 2013	Primary and Secondary	PM <sub>2.5</sub>	24 hour	35 µg/m <sup>3</sup>	98th percentile, averaged over 3 years <sup>6</sup>
2012 <a href="#">78 FR 3085</a> Jan 15, 2013	Primary and Secondary	PM <sub>10</sub>	24 hour <sup>8</sup>	150 µg/m <sup>3</sup>	Not to be exceeded more than once per year on average over a 3-year period