

Annual Ambient Air Assessment Report 2013

Arizona Department of Environmental Quality

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Table of Contents

Table of Contents	2
List of Figures	5
List of Tables	6
Section I – Introduction to ADEQ’s Ambient Air Monitoring Program	7
1.0 Purpose and Background	7
2.0 Standards and Guidelines	8
2.1 Air Quality Index	8
2.2 National Ambient Air Quality Standards	9
2.3 Monitoring Objectives	11
3.0 Quality Assurance	13
4.0 Monitoring Location Summary	15
Section II – Criteria Pollutants	18
1.0 Carbon Monoxide (CO)	18
1.1 Background	18
1.2 Monitoring Methods	19
1.3 Compliance/Summary of Design Values	19
1.4 Trends	21
2.0 Nitrogen Dioxide (NO ₂)	23
2.1 Background	23
2.2 Monitoring Methods	24
2.3 Compliance/Summary of Design Values	24
2.4 Trends	26
3.0 Ozone (O ₃)	28
3.1 Background	28
3.2 Monitoring Methods	29
3.3 Compliance/Summary of Design Values	29
3.4 Trends	31
4.0 Sulfur Dioxide (SO ₂)	33

4.1 Background	33
4.2 Monitoring Methods	34
4.3 Compliance/Summary of Design Values	34
4.4 Trends	36
5.0 Particulate Matter Smaller Than 10 Microns (PM ₁₀)	39
5.1 Background	39
5.2 Monitoring Methods	40
5.3 Compliance/Summary of Design Values	41
5.4 Trends	43
6.0 Particulate Matter Smaller Than 2.5 Microns (PM _{2.5})	47
6.1 Background	47
6.2 Monitoring Methods	48
6.3 Compliance/Summary of Design Values	48
6.4 Trends	51
7.0 Lead (Pb)	54
7.1 Background	54
7.2 Monitoring Methods	55
7.3 Compliance/Summary of Design Values	55
7.4 Trends	56
Section III – Non-Criteria Pollutants	58
1.0 Chemical Speciation Network	58
1.1 Background	58
1.2 Monitoring Methods	58
1.3 Graphs	59
1.4 Trends	62
2.0 National Air Toxics Trends Station (NATTS)	63
2.1 Background	63
2.2 Monitoring Methods	63
2.3 Tables	64
2.4 Trends	66
3.0 Photochemical Assessment Monitoring Stations (PAMS)	70
3.1 Background	70

3.2 Monitoring Methods	71
3.3 Trends	71
Section IV – Visibility	73
1.0 Urban Haze	73
1.1 Program Background	73
1.2 Monitoring Methods	74
1.3 Trends	74
2.0 IMPROVE	80
Appendix I – Abbreviations	81
Appendix II – References	83
Appendix III – 2013 Area Designations Map	84

List of Figures

Figure 1 – Chart of AQI Levels	8
Figure 2 – The locations of ADEQ’s monitoring sites	17
Figure 3 – Map of ADEQ’s CO sites	18
Figure 4 – CO One-Hour Average Trend	21
Figure 5 – CO Eight-Hour Average Trend	22
Figure 6 – Map of ADEQ’s NO ₂ sites.....	23
Figure 7 – NO ₂ Annual Mean Trend	26
Figure 8 – NO ₂ One-Hour Average Trend.....	27
Figure 9 – Map of ADEQ’s O ₃ sites	28
Figure 10 – O ₃ Eight-Hour Average Trend.....	32
Figure 11 – Map of ADEQ’s SO ₂ sites	33
Figure 12 – SO ₂ One-Hour Average Trend for Miami and Hayden Areas	37
Figure 13 – SO ₂ One-Hour Average Trend for JLG Supersite	38
Figure 14 – SO ₂ Annual Average Trend	38
Figure 15 – Map of ADEQ’s PM ₁₀ sites	39
Figure 16 – PM ₁₀ Annual Mean Average for the Southern Region	44
Figure 17 – PM ₁₀ Annual Mean Average for the Northern Region	45
Figure 18 – PM ₁₀ Annual Mean Average for the Central Region	46
Figure 19 – Map of ADEQ’s PM _{2.5} sites	47
Figure 20 – Nogales Post Office monitoring station.	48
Figure 21 – PM _{2.5} Annual Mean Trend.....	52
Figure 22 – PM _{2.5} 24-Hour Average Trend	53
Figure 23 – Map of ADEQ’s Pb sites	54
Figure 24 – Pb Three-Month Average Trend	57
Figure 25 – 2013 Annual Averages for Speciated PM _{2.5}	60
Figure 26 – 2013 Quarterly Averages for Speciated PM _{2.5}	61
Figure 27 – Speciated PM _{2.5} Annual Average Trend	62
Figure 28 – VOC Annual Mean Trend (0.0-1.4 ppb).....	66
Figure 29 – VOC Annual Mean Trend (0.00-0.25 ppb).....	67
Figure 30 – Aldehydes Annual Mean Trend.....	68
Figure 31 – Benzo (A) Pyrene Annual Mean Trend.....	69
Figure 32 – Naphthalene Annual Mean Trend.....	69
Figure 33 – TNMOC Annual Mean Trend.....	72
Figure 34 – Pleasant Valley monitoring station.	73
Figure 35 – Transmissometer Visual Range (All Hours) Trend.....	75
Figure 36 – Transmissometer Visual Range (Morning Hours) Trend.....	76
Figure 37 – Transmissometer Seasonal Average Trend.....	77
Figure 38 – Nephelometer Visual Range Trend	78
Figure 39 – Examples of Visual Condition in Phoenix.....	79

List of Tables

Table 1 – Current NAAQS.....	10
Table 2 – Site Index.....	15
Table 3 – History of the National Ambient Air Quality Standards for CO.....	20
Table 4 – CO One-Hour Compliance Summary.....	20
Table 5 – CO Eight-Hour Compliance Summary.....	20
Table 6 – History of the National Ambient Air Quality Standards for NO ₂	24
Table 7 – NO ₂ Annual Mean Compliance Summary.....	25
Table 8 – NO ₂ One-Hour Compliance Summary.....	25
Table 9 – History of the National Ambient Air Quality Standards for O ₃	30
Table 10 – O ₃ Compliance Summary.....	31
Table 11 – History of the National Ambient Air Quality Standards for SO ₂	34
Table 12 – SO ₂ One-Hour Compliance Summary.....	35
Table 13 – SO ₂ Three-Hour Compliance Summary.....	36
Table 14 – History of the National Ambient Air Quality Standards for PM ₁₀	41
Table 15 – PM ₁₀ Compliance Summary.....	42
Table 16 – History of the National Ambient Air Quality Standards for PM _{2.5}	49
Table 17 – PM _{2.5} Annual Average Compliance Summary.....	50
Table 18 – PM _{2.5} 24-Hour Average Compliance Summary.....	51
Table 19 – History of the National Ambient Air Quality Standards for Pb.....	56
Table 20 – Pb Compliance Summary.....	56
Table 21 – Air Toxics Data for JLG Supersite.....	65
Table 22 – Air Toxics Data for South Phoenix.....	65

Section I – Introduction to ADEQ’s Ambient Air Monitoring Program

1.0 Purpose and Background

The ADEQ Annual Ambient Air Assessment Report summarizes the air monitoring activities and the results of air quality monitoring conducted by the Arizona Department of Environmental Quality (ADEQ) for the calendar year 2013. This report shows monitoring locations, significance of monitoring, and monitoring methods. The report provides an overview of long-term monitoring trends and area/monitor compliance status based on air monitoring conducted by ADEQ in 2013. This report is an informational and technical document intended for use within ADEQ, other state and local agencies, other health organizations such as the American Lung Association.

Data from 44 sites located throughout Arizona are included in this report. Many of the sites have multiple instruments measuring a variety of gaseous, particulate, meteorological, and visibility parameters. The majority of the air quality measurements are for criteria pollutants (carbon monoxide (CO), nitrogen dioxide (NO₂), ozone (O₃), sulfur dioxide (SO₂), particulate matter (PM₁₀ and PM_{2.5}), and lead (Pb)) classified as State and Local Air Monitoring Stations (SLAMS) used for regulatory compliance. To show regulatory compliance, the U.S. Environmental Protection Agency (EPA) has established National Ambient Air Quality Standards (NAAQS). These standards are the concentrations at which each pollutant becomes harmful to human health and are unique to each pollutant.

In addition to sampling for criteria pollutants, ADEQ does special continuous monitoring for the optical characteristics of the atmosphere, manual sampling of O₃ forming compounds and other hazardous air pollutants, and speciated particulate matter sampling under the Photochemical Assessment Monitoring Stations (PAMS), Chemical Speciation Network (CSN), and National Air Toxics Trends Stations (NATTS). ADEQ also operates industrial networks to determine the effects of their emissions on local air quality and how well pollution control technologies are working. ADEQ also serves as an operator for the Interagency Monitoring of Protected Visual Environments (IMPROVE) which tracks visual conditions in and around national parks and monuments, as well as in some urban areas. Additionally, ADEQ operates a network of portable particulate monitors for public awareness to provide information regarding pollutant levels from wildfires and wood burning.

Air monitoring is commonly classified by networks based on individual pollutants or by a group of related pollutants. Monitoring networks for ambient air quality are established to sample pollution in a variety of representative settings, to assess health and welfare effects, and to assist in determining air pollution sources. The ambient monitoring networks cover both urban and rural areas of the state. They are composed of one or more monitoring sites whose data are compared to the NAAQS for compliance and statistically analyzed in various ways for trends analysis. ADEQ also tracks data recovery, quality control, and quality assurance parameters for the instruments operated at its various

sites. Most of the sites within each network also measure meteorological variables. ADEQ networks monitor a wide variety of pollutant and atmospheric characteristics including urban, industrial, rural, transport, and background surveillance.

2.0 Standards and Guidelines

The Federal Clean Air Act (CAA) of 1970 required EPA to assist states and local agencies in establishing ambient air quality monitoring networks to characterize human health exposure and public welfare effects of criteria pollutants. The way public welfare is measured is by analyzing ambient air conditions using a variety of instruments specifically designed to measure a certain pollutant. These instruments show pollution concentrations for a given time period and identify certain concentrations which can affect human health. Because different concentrations of pollutants affect human health at different levels, an Air Quality Index (AQI) is used for showing when a specific concentration can be bad for human health.

For data completeness, EPA requires 75 percent completeness over a set time period for values to be considered valid. This applies to the scales of hourly, daily, quarterly, and yearly, for which each averaging period must be 75 percent complete. For regulatory purposes, if all quarters for the year have completeness percent greater than 75 percent, then completeness criteria are met and data may be used for area designations.

2.1 Air Quality Index

The AQI is an index for reporting daily air quality to the general population. It tells you how clean or polluted the air is and what associated health effects might be a concern for you. The AQI focuses on health effects that may be experienced within a few hours or days after breathing polluted air. EPA calculates the AQI for the criteria air pollutants regulated by the CAA: ground-level O₃, PM₁₀, PM_{2.5}, CO, SO₂, and NO₂. For each of these pollutants, EPA has established national air quality standards to protect public health. When the AQI reaches 100, this indicates a concentration which exceeds the standards set forth by the EPA. Generally, ground-level O₃ and airborne particulate matter (PM) are the two pollutant types that pose the greatest threat to human health in this country. EPA's AQI website AIRNow is found at www.airnow.gov.

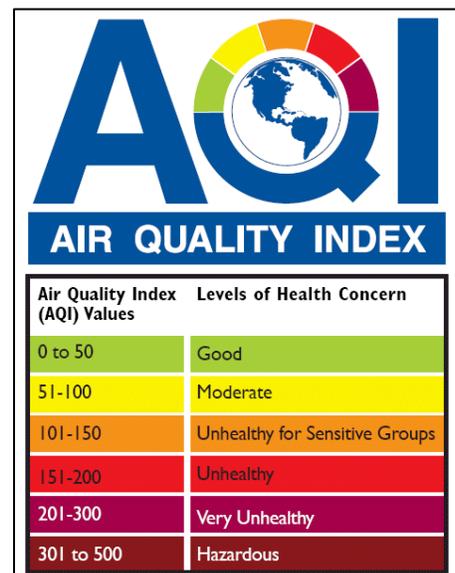


Figure 1 – Chart of AQI Levels

Each category in Figure 1 corresponds to a different level of health concern. The six levels of health concern and what they mean are:

- "Good" AQI is 0 - 50. Air quality is considered satisfactory, and air pollution poses little or no risk to human health.

- "Moderate" AQI is 51 - 100. Air quality is acceptable; however, for some pollutants there may be a moderate health concern for a very small number of people. For example, people who are unusually sensitive to O₃ may experience respiratory symptoms.
- "Unhealthy for Sensitive Groups" AQI is 101 - 150. Although general public is not likely to be affected at this AQI range, people with lung disease, older adults, and children are at a greater risk from exposure to O₃, whereas persons with heart and lung disease, older adults, and children are at greater risk from the presence of particles in the air.
- "Unhealthy" AQI is 151 - 200. Everyone may begin to experience some adverse health effects, and members of the sensitive groups may experience more serious effects.
- "Very Unhealthy" AQI is 201 - 300. This would trigger a health alert signifying that everyone may experience more serious health effects.
- "Hazardous" AQI greater than 300. This would trigger a health warning of emergency conditions. The entire population is more likely to be affected.

2.2 National Ambient Air Quality Standards

The NAAQS were first set by the EPA with the CAA in 1970 and are continuously evaluated and updated based on current scientific research on the effects of pollution exposure to the population. Focus is placed on those groups who are sensitive to air pollution. Table 1 shows the current NAAQS.

Table 1 – Current NAAQS
 (Source: USEPA TTN NAAQS)

Pollutant		Primary/ Secondary	Averaging Time	Level	Form
Carbon Monoxide		primary	8-hour	9 ppm	Not to be exceeded more than once per year
			1-hour	35 ppm	
Lead		primary and secondary	Rolling 3 month average	0.15 µg/m ³	Not to be exceeded
Nitrogen Dioxide		primary	1-hour	100 ppb	98 th percentile, averaged over 3 years
		primary and secondary	Annual	53 ppb	Annual Mean
Ozone		primary and secondary	8-hour	0.075 ppm	Annual fourth-highest daily maximum 8-hr concentration, averaged over 3 years
Particle Pollution	PM _{2.5}	primary	Annual	12 µg/m ³	annual mean, averaged over 3 years
		secondary	Annual	15 µg/m ³	annual mean, averaged over 3 years
		primary and secondary	24-hour	35 µg/m ³	98 th percentile, averaged over 3 years
	PM ₁₀	primary and secondary	24-hour	150 µg/m ³	Not to be exceeded more than once per year on average over 3 years
Sulfur Dioxide		primary	1-hour	75 ppb	99 th percentile of 1-hour daily maximum concentrations, averaged over 3 years
		secondary	3-hour	0.5 ppm	Not to be exceeded more than once per year

Compliance with the NAAQS is defined under the “Form” column in Table 1 and are calculated based on the averaging time stated. For many of the pollutants, there is a primary standard and a secondary standard. 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.

An area may be designated as nonattainment, attainment, or unclassifiable based on exceedances to the standard. Area designations occur after a revision to the NAAQS or at other times the EPA deems appropriate. At this time, the EPA will designate areas as previously stated.

- A nonattainment area is one in which either the primary or secondary standard has been violated, and both the local agency and the EPA have acknowledged the area as nonattainment. The EPA can designate that area as nonattainment with certain sanctions or penalties being placed in order to bring that area into attainment. When EPA designates an area nonattainment, a State Implementation Plan (SIP) is put in place. A SIP outlines the actions the state will take to improve air quality in the area. This can include instituting lower maximum pollutant allowances for industrial processes, paving of roads, replacing aging equipment, and other control strategies. If the controls outlined in the SIP do not achieve the standard or are inadequate, the EPA has the option to develop a Federal Implementation Plan (FIP) for the area.
- An attainment area is achieving the standard and no additional measures are taken.
- An unclassifiable area is a statistical area for which data are incomplete and do not support a designation of attainment or nonattainment. Data may be incomplete due to lack of monitoring or completeness criteria for the year.

It is important to understand the difference between an exceedance and a violation of a standard in order to define area designation. An exceedance occurs when the pollutant concentration reaches a level where the AQI is at or above 100 for a given time period. This is above the acceptable level that is defined in the NAAQS, but does not necessarily indicate that the NAAQS have been violated.

For example, the 24-hour $PM_{2.5}$ NAAQS level of exceedance is 35 micrograms per cubic meter ($\mu\text{g}/\text{m}^3$). If the concentration on a given day is above this level, an exceedance occurs, but a violation of standard does not necessarily occur. The form by which the $PM_{2.5}$ standard is calculated is the 98th percentile of samples averaged over three years. If the 24-hour average for a day goes above the 35 $\mu\text{g}/\text{m}^3$ level, but the 98th percentile averaged over three years is not above 35 $\mu\text{g}/\text{m}^3$, then there is an exceedance for that day, but there is no violation of the standard.

2.3 Monitoring Objectives

Ambient air monitoring has multiple purposes which have specific needs and requirements. There are three basic monitoring objectives, each containing multiple aspects and purposes: NAAQS comparison for regulatory comparison, research, and public information. There may be certain monitors which have more than one purpose due to crossover between different networks, for example a NAAQS comparison monitor may also be used for research purposes in some circumstances.

A. NAAQS Comparison

The majority of ADEQ's monitoring falls under the NAAQS comparison category. This monitoring is governed by the CAA and is explained under the Code of Federal Regulations (CFR) 40 part 58.

This monitoring is required based on rules spelled out in the CFR, including: total emissions in an area, population of an area, attainment/nonattainment areas, population weighted emissions, traffic counts, and pollutant point sources. ADEQ designates these monitors as SLAMS. These are long term monitors that operate for the primary purpose of comparison to the NAAQS and are used for regulatory decision making. They may also support compliance with air quality standards and emissions strategy development, and provide air pollution data to the general public in a timely manner. The SLAMS network includes monitors at stations classified as a National Core (NCore) monitoring network, PAMS, or Speciation, but does not include Special Purpose Monitors (SPM) and other monitors used for non-regulatory or industrial monitoring purposes. Once monitoring occurs, EPA designates areas as attainment or nonattainment based on the comparison of observed concentrations with the NAAQS.

B. Research

ADEQ operates a range of monitors used mainly for research purposes. These instruments may not be regulated by the EPA as part of the NAAQS, but represent various precursor and toxic pollutants. The programs that fall under research include PAMS, NATTS, CSN, IMPROVE, urban air toxics monitoring program (UATMP), and NCore. PAMS, NATTS, CSN, and IMPROVE will be discussed at length in this report (Sections III and IV). As there is cross over between NCore and the criteria pollutants, a brief description of NCore will be given here. Crossover also exists between UATMP and NATTS, which will also be described.

- **NCore** is a multi-pollutant network throughout the whole nation that integrates several advanced measurement systems for particles, pollutant gases, and meteorology. The stations are equipped with several measurement systems to monitor particulate matter (PM_{2.5}, PM₁₀, and PM_{10-2.5}), O₃, CO, SO₂, total reactive oxides of nitrogen (NO_y), and basic meteorological parameters (temperature, wind speed, wind direction, and relative humidity). The NCore stations should be perceived as developing a representative report card on air quality across the nation, capable of delineating differences among geographic and climatological regions. There are approximately 83 NCore stations nationwide. NCore meets a number of important data needs: improved flow and timely reporting of data to the public, including supporting air quality forecasting and information systems such as AIRNow; continued determination of NAAQS compliance; improved development of emissions control strategies; enhanced accountability for the effectiveness of emission control programs; and more complete information for scientific, public health, and ecosystem assessments.

Emphasis of NCore is placed on high sensitivity instruments with the capability to detect low levels of the precursor gases CO, SO₂, and NO_y. These gases play important roles in the formation of atmospheric O₃, air toxics, and PM, which are linked to human health issues. This interconnection among distinct air quality issues requires an integrated multiple pollutant air quality monitoring and management approach which NCore successfully incorporates. ADEQ

supports the NCore network by monitoring all required pollutants at its Phoenix JLG Supersite. Data will be shown and analyzed in Section II of this document.

- **UATMP** is a subset of the Air Toxics program, which includes NATTS, school air toxics monitoring initiative (SATMI), and community scale air toxics ambient monitoring (CSATAM) projects. All programs that are a part of Air Toxics utilize similar monitoring methods and are therefore directly comparable. All have similar monitoring objectives of characterizing toxics in an area of interest. UATMP specifically looks at toxics trends in urban areas. Data will be shown and analyzed in Section III of this document.

C. Public Information

Data generated by ADEQ using EPA approved monitoring techniques are reported to AIRNow on an hourly basis to show real-time conditions to the public. AIRNow data are used to report the AQI only, not to show regulatory status of an area.

ADEQ also operates networks that do not have EPA approved monitoring techniques, for the sole purpose of providing public health and visibility information to the general public. Although these networks do not use approved monitoring techniques, they give the public a general understanding of current air quality in their area.

For public health, a network of temporary PM_{2.5} monitors without an EPA method code are used to provide the public air quality conditions in areas that do not require monitoring under the CAA and in CFR 40. These areas are mainly located in the northeastern part of Arizona and are often impacted by summer wildfires. These monitors can be easily deployed around a wildfire to describe the impacts that it has on air pollution for the local population. There are also monitors located in areas where wood burning is the main source of heating in the winter. Wildfires and wood burning create small particulates that are harmful to human health. Information regarding these portable particulate monitors can be found at <http://phoenixvis.net/PPMmain.aspx>.

In the Phoenix metropolitan area, ADEQ operates a variety of instruments used to determine the visibility on a daily basis. Data and pictures for visibility will be shown and analyzed under Section IV.

3.0 Quality Assurance

ADEQ sustains a quality system as required by EPA to ensure high quality data are produced that meet the users' needs. The EPA primarily specifies the quality assurance (QA) requirements for operating SLAMS, SPM, CSN, NCore, NATTS, PAMS, and prevention of significant deterioration (PSD) air monitors in *40 CFR Part 58 Appendix A, the Quality Assurance Handbook for Air Pollution Measurement Systems: Volume II: Ambient Air Quality Monitoring Program*, technical assurance documents (TADs), and other supporting guidance documents. In response, ADEQ develops quality assurance plans for air monitoring

networks, which provide detailed information regarding the specifics of each air monitoring network and how data will be managed. Components of ADEQ's quality system include, but are not limited to:

- ADEQ being established as the primary quality assurance organization (PQAO) for the criteria and non-criteria pollutant air monitoring data collected and reported to EPA's air quality system (AQS).
- An agency-level Quality Management Plan (QMP), which is an "umbrella" document that details, in broad terms, the strategies used to carry out QA/QC in environmental data collection activities.
- Division-level quality assurance program plans (QAPPs) for each major, ongoing air monitoring network. Each QAPP describes:
 - purpose for operating the monitoring station or network
 - data quality objectives (DQOs) and/or measurement quality objectives (MQOs) along with data quality indicators (DQIs) that specify the amount of tolerable error in the data using statistical metrics
 - variety of regularly occurring quality control (QC) checks along with pass/fail criteria
 - types of QA assessments and reports needed from the network
 - data validation processes and data reporting requirements
- Unit-level standard operating procedures (SOPs) that document procedures to assure that work products are reliable, reproducible, and consistent in quality. SOPs also serve to clearly communicate any process customizations in-use, providing a means of attesting that work products are credible, legally defensible, and meet or exceed our customers' and/or stakeholders' needs or requirements.

ADEQ uses a multi-tiered approach to data validation to ensure consistent quality. It requires all data to move through different levels of QA by separate reviewers. Data have five different levels associated with each tier level.

- Raw – Original unchanged data recorded by the sampler or produced by laboratory analysis.
- Level 1 – Data are reviewed programmatically using software written to flag data. The data are flagged valid or invalid based on instrumentation parameters.
- Level 2 – Data are reviewed manually on a weekly basis by an initial data reviewer to flag any discrepancies found. This gives the data a preliminary validation decision and identifies outliers, anomalous data and instrumentation/laboratory problems.
- Level 3 – Data are reviewed manually on a quarterly basis by the final data reviewer by looking at the data spatially and temporally. QC measures are incorporated, environmental events are identified, and a final determination on the validity of data is made.

- Certified – Data are uploaded to AQS and are certified annually by ADEQ.

Periodically, EPA publishes reports for some of the criteria pollutant networks, and potentially non-criteria pollutant networks, that rate and/or rank monitoring organizations' performance over a three-year period. The QA Team, as well as other personnel in the data management and quality assurance (DM&QA) unit, reviews these reports to gauge how well ADEQ's networks are performing with those across the nation. If needed, corrective actions are taken to ensure data of the highest quality possible are collected.

4.0 Monitoring Location Summary

Table 2 contains all of ADEQ's sites associated including the site number from EPA's AQS database, the city the site is located, the main monitoring objectives for the site, the site coordinate location, and which parameters were recorded at the site in 2013.

Table 2 – Site Index

Site Name	Air Quality System ID	Location	Monitoring Objective(s)	Lat. Long. (Deg.)	Parameters Recorded
ADEQ Building	None	Phoenix, AZ	Visibility	33.4483, -112.088	Digital High Resolution Image
Ajo	04-019-0001	Ajo, AZ	NAAQS Comparison	32.3820, -112.858	PM ₁₀ , Wind
Alamo Lake	04-012-8000	Alamo Lake State Park, AZ	NAAQS Comparison	34.2439, -113.559	O ₃
Banner Mesa Medical Center	None	Mesa, AZ	Visibility	33.4335, -111.843	Digital High Resolution Image
Bullhead City	04-015-1003	Bullhead City, AZ	NAAQS Comparison	35.1539, -114.566	PM ₁₀
Chiricahua Entrance Station	04-003-8001	Wilcox, AZ	Visibility	32.0094, -109.389	IMPROVE
Douglas Red Cross	04-003-1005	Douglas, AZ	NAAQS Comparison	31.3492, -109.54	PM ₁₀ , PM _{2.5} , Temp/RH, Wind, IMPROVE
Dysart	04-013-4010	Surprise, AZ	Visibility	33.6370, -112.339	Bscat/PM _{2.5} , Temp/RH
Estrella	04-013-8005	Goodyear, AZ	Visibility	33.3833, -112.373	Bscat/PM _{2.5} , Temp/RH
Estrella Mountain Community College	None	Avondale, AZ	Visibility	33.4836, -112.350	Digital High Resolution Image
Flagstaff Middle School	04-005-1008	Flagstaff, AZ	NAAQS Comparison	35.2061, -111.653	O ₃ , PM _{2.5} , PM ₁₀
Globe Highway	04-007-1002	Winkelman, AZ	NAAQS Comparison	33.002, -110.765	Pb, Temp/RH, Wind
Grand Canyon NP Hance Camp	04-005-8102	Grand Canyon, AZ	Visibility	35.9731, -111.984	IMPROVE
Grand Canyon NP Indian Gardens	04-005-8101	Grand Canyon, AZ	Visibility	36.0776, -112.129	IMPROVE
Greer Water Treatment Plant	04-001-8001	Greer, AZ	Visibility	34.0583, -109.440	IMPROVE
Hayden Old Jail	04-007-1001	Hayden, AZ	NAAQS Comparison	33.0062, -110.786	SO ₂ , PM ₁₀ , Temp/RH, Wind
Ike's Backbone	04-025-8104	Strawberry, AZ	Visibility	34.3406, -111.683	IMPROVE

Site Name	Air Quality System ID	Location	Monitoring Objective(s)	Lat. Long. (Deg.)	Parameters Recorded
JLG Supersite	04-013-9997	Phoenix, AZ	NAAQS Comparison/ Research	33.5038, -112.096	CO, NO ₂ , NO _y , O ₃ , SO ₂ , Carbonyl, VOC, SVOC, Pb-PM ₁₀ , / PM ₁₀ metals speciation, PM ₁₀ , PM _{10-2.5} , PM _{2.5} , PM _{2.5} speciation, Temp/RH, Wind, IMPROVE
Meadview	04-015-9000	Meadview, AZ	Visibility	36.0193, -114.068	IMPROVE
Miami Golf Course	04-007-8000	Miami, AZ	NAAQS Comparison	33.4190, -110.83	Pb, PM ₁₀ , Temp/RH, Wind
Miami Jones Ranch	04-007-0011	Miami, AZ	NAAQS Comparison	33.3853, -110.867	SO ₂
Miami Ridgeline	04-007-0009	Miami, AZ	NAAQS Comparison	33.3992, -110.859	SO ₂
Miami Townsite	04-007-0012	Miami, AZ	NAAQS Comparison	33.3973, -110.874	SO ₂
Nogales Post Office	04-023-0004	Nogales, AZ	NAAQS Comparison	31.3372, -110.937	PM ₁₀ , PM _{2.5} , Temp/RH, Wind
North Mountain Summit	None	Phoenix, AZ	Visibility	33.5855, -112.072	Digital High Resolution Image
Organ Pipe National Monument	04-019-005	Ajo, AZ	Visibility	31.950, -112.80	IMPROVE
Paul Spur Chemical Lime Plant	04-003-0011	Paul Spur, AZ	NAAQS Comparison	31.366, -109.73	PM ₁₀ , Temp/RH, Wind
Payson Well Site	04-007-0008	Payson, AZ	NAAQS Comparison	34.230, -111.33	PM ₁₀ , PM _{2.5} , Temp/RH, Wind
Petrified Forest National Park	04-001-0012	Petrified Forest NP, AZ	Visibility	35.077, -109.77	IMPROVE
Phoenix Transmissometer Receiver	None	Phoenix, AZ	Visibility	33.490, -112.08	Bext, Temp/RH
Phoenix Transmissometer Transmitter	None	Phoenix, AZ	Visibility	33.525, -112.10	Bext
Pleasant Valley Ranger Station	04-007-8100	Young, AZ	Visibility	34.091, -110.94	IMPROVE
Prescott College AQD	04-025-8033	Prescott, AZ	NAAQS Comparison	34.547, -112.48	O ₃ , PM _{2.5}
Queen Valley	04-021-8001	Queen Valley, AZ	NAAQS Comparison	33.294, -111.29	O ₃ , NO _y , VOC, Temp/RH, Wind, IMPROVE
Rillito	04-019-0020	Rillito, AZ	NAAQS Comparison	32.414, -111.16	PM ₁₀ , Temp/RH, Wind
Saguaro National Park East	04-019-0021	Tucson, AZ	Visibility	32.174, -110.74	IMPROVE
Saguaro Nation Park West	04-019-9000	Tucson, AZ	Visibility	32.249, -111.22	IMPROVE
South Phoenix	04-013-4003	Phoenix, AZ	Research	33.403, -112.08	VOC
Sycamore Canyon	04-005-8103	Flagstaff, AZ	Visibility	35.141, -111.97	IMPROVE
Tonto National Monument	04-007-0010	Roosevelt, AZ	NAAQS Comparison	33.655, -111.11	O ₃ , IMPROVE
Vehicle Emissions Laboratory	04-013-9998	Phoenix, AZ	Research	33.455, -111.10	Bscat/PM _{2.5} , Delta Temp, Horizontal Solar Radiation, Ultraviolet Solar Radiation, Temp/RH, Wind
Yuma Supersite	04-027-8011	Yuma, AZ	NAAQS Comparison	32.690, -114.62	O ₃ , PM ₁₀ , PM _{2.5} , Temp/RH, Wind

ADEQ Monitoring Sites

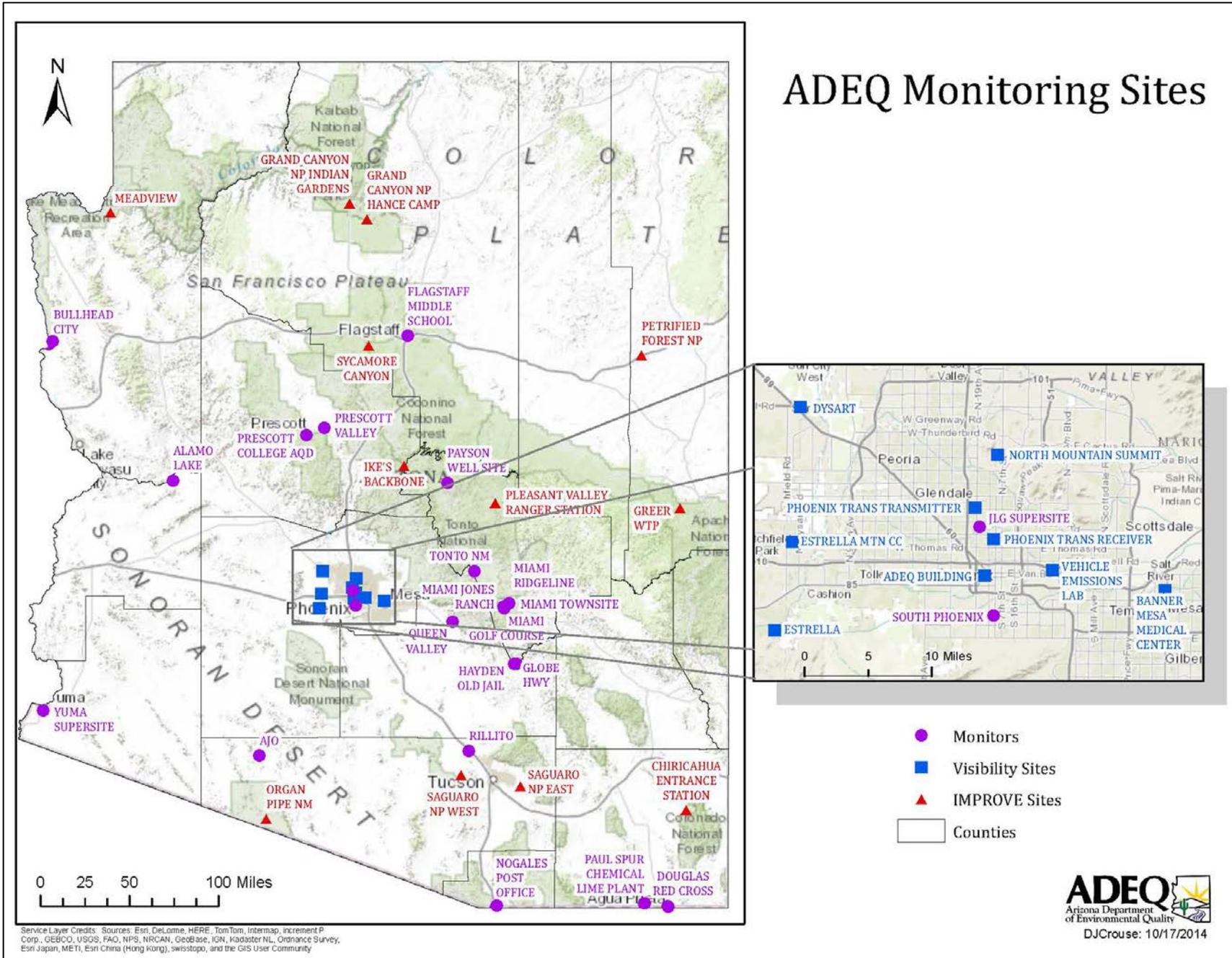


Figure 2 – The locations of ADEQ’s monitoring sites. Sites from other monitoring organizations are not included.
ADEQ Annual Ambient Air Assessment Report 2013 Page | 17

Section II – Criteria Pollutants

The six criteria pollutants as outlined in the CAA will be discussed at length in Section II. The health effect specific of each pollutant will be discussed, as well as the specific background of the pollutants in Arizona. Current monitoring techniques will also be explained. A discussion of 2013 data which shows a history of the pollutant standard, compliance to the current standard, as well as data completeness for the year. A length of record trend analysis provides a history of the monitors operating in Arizona in 2013. The trend analysis includes a qualitative summary of the trend and a quantitative increase/decrease of the pollutant over the years.

1.0 Carbon Monoxide (CO)

Carbon Monoxide is a colorless, odorless, tasteless gas that is produced in the incomplete combustion of fuels. It has a variety of adverse health effects that arise from its ability to chemically bind to blood hemoglobin. Carbon Monoxide successfully competes with oxygen for binding with hemoglobin and thereby impairs oxygen transport. This impaired transport leads to several central nervous system effects, such as headache, fatigue, and dizziness. Chronic CO exposure also contributes to or exacerbates arteriosclerotic heart disease. Chronic exposure to low levels of CO can lead to depression, confusion, and memory loss.

1.1 Background

According to the 2011 National Emissions Inventory (NEI), in Arizona, 63 percent of CO emissions come from fires, 35 percent from mobile sources including on-road motor vehicles, off-road vehicles, construction equipment, and lawn and garden equipment, with the remainder coming from point and area sources. This pollutant has low background levels, with the highest concentrations next to busy streets, and elevated neighborhood concentrations in locations with significant amounts of emissions transported from upwind areas. Concentrations peak from November to January because emissions are highest in cold weather. Automotive emissions of CO are greatest when engines operate in open loop, which occurs for longer periods of time in cold weather as the engine attempts to warm up and because the mixed layer of the atmosphere is most shallow in wintertime due to decreased solar heating. Hourly concentrations tend to be at their

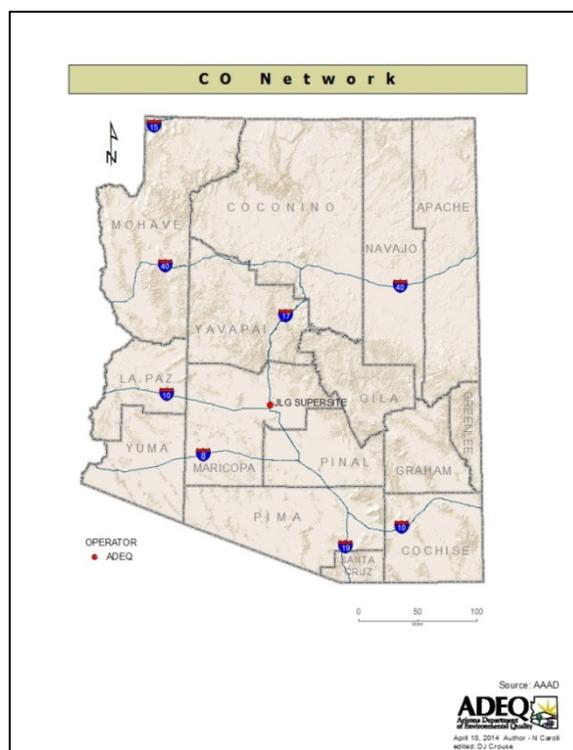


Figure 3 – Map of ADEQ’s CO sites

maximum during the morning rush hour and between 6 p.m. and midnight.

Emission controls have reduced overall CO emissions, and the standards have been achieved in the metropolitan Phoenix area since 1996, in stark contrast to the first half of the 1980s when more than 100 exceedances were recorded each year. Similar improvements have occurred in Tucson, where the last eight-hour exceedances were recorded in 1988. Equipping vehicles with catalytic converters and electronic ignition systems was the most effective control, but significant reductions can also be attributed to the vehicle emissions inspection program (beginning in 1976) and oxygenated fuels (beginning in 1989). On Jan. 3, 2005, EPA re-designated the Phoenix metropolitan area to attainment for the NAAQS for CO, and approved the attainment demonstration and maintenance plan showing maintenance of the CO standard through 2015.

Due to the successful nature of the emission controls placed on CO sources, ADEQ only operated one CO monitor in 2013 at its NCore station. Additional CO monitors are operated by other State and Local agencies as required, but will not be discussed in this report.

1.2 Monitoring Methods

Carbon Monoxide is monitored continuously with a nondispersive infrared (IR) instrument. The IR light passes through a gas filter correlation wheel that alternately uses a CO filled chamber and a chamber with no CO present. The light path travels through a sample cell following the correlation wheel. Carbon Monoxide absorbs a specific wavelength of IR light and the energy loss through the sample cell is compared with the zero reference provided by the gas filter correlation wheel to produce a signal that is proportional to concentration. Readings are averaged into hourly, daily, monthly, quarterly, and yearly averages for data analysis. Readings are retrieved by a data collection system and stored in a database.

1.3 Compliance/Summary of Design Values

On April 30, 1971, the EPA promulgated NAAQS for CO based on a criteria document published by the U.S. Department of Health, Education and Welfare in March 1970. Identical primary (health-based) and secondary (welfare-based) NAAQS for CO were set at 35 parts per million (ppm), one-hour average, and at 9 ppm, eight-hour average, neither to be exceeded more than once a year. After the most recent review of the CO NAAQS, on August 31, 2011, EPA proposed to retain the current primary standards. After review of the air quality criteria, EPA further concludes that no secondary standard should be set for CO at this time. Table 3 summarizes the history of the NAAQS for CO during the period 1971-2011. At present there are two primary standards for CO. The one-hour standard is 35 ppm and the eight-hour standard is 9 ppm (most critical for compliance).

Table 3 – History of the National Ambient Air Quality Standards for CO during the period 1971-2011
(Source: USEPA TTN NAAQS)

Final Rule	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
			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 59 FR 38906 Aug 1, 1994	Primary standards retained, without revision.				
2011 76 FR 54294 Aug 31, 2011	Primary standards retained, without revision.				

*Second highest, non-overlapping 8-hour average concentration of 9 ppm

Table 4 and Table 5 below show the 1st maximum and 2nd maximum values for both the current primary standards for the year 2012 and 2013. No exceedances of the one-hour or eight-hour standards were recorded in 2012 and 2013. The CO monitor at JLG Supersite is considered to be in compliance for the year 2013.

Table 4 – CO One-Hour Compliance Summary

2012 to 2013 One-Hour CO Compliance Summary (in ppm) (NAAQS primary one-hour standard 35 ppm)					
Site Name	2012		2013		Compliance Value
	1 st Max Value	2 nd Max Value	1 st Max Value	2 nd Max Value	
Maricopa County					
JLG Supersite	3.0	2.9	3.0	2.9	2.9
Number of Sites in Violation of the NAAQS					0

Table 5 – CO Eight-Hour Compliance Summary

2012 to 2013 Eight-Hour CO Compliance Summary (in ppm) (NAAQS primary eight-hour standard 9 ppm)					
Site Name	2012		2013		Compliance Value
	1 st Max Value	2 nd Max Value	1 st Max Value	2 nd Max Value	
Maricopa County					
JLG Supersite	1.8	1.6	2.0	1.9	1.9
Number of Sites in Violation of the NAAQS					0

1.4 Trends

Monitoring of CO throughout the state of Arizona contains the longest history of all the criteria pollutants. Most of this long-term monitoring was located in the highly urbanized areas of Phoenix and Tucson and several of these CO sites contain monitoring records dating back to the 1970s. For the purpose of this report, the examination of CO trends will include the ADEQ monitors only. ADEQ has monitored CO at JLG Supersite since 1999. Figure 4 and Figure 5 show a decreasing trend in Phoenix for primary one-hour and eight-hour CO respectively, both being under the NAAQS since monitoring started at the location. Most of the improvements can be attributed to emission control programs as stated in 1.1 of this section. ADEQ started trace-level monitoring of CO in 2010 as part of the NCore program.

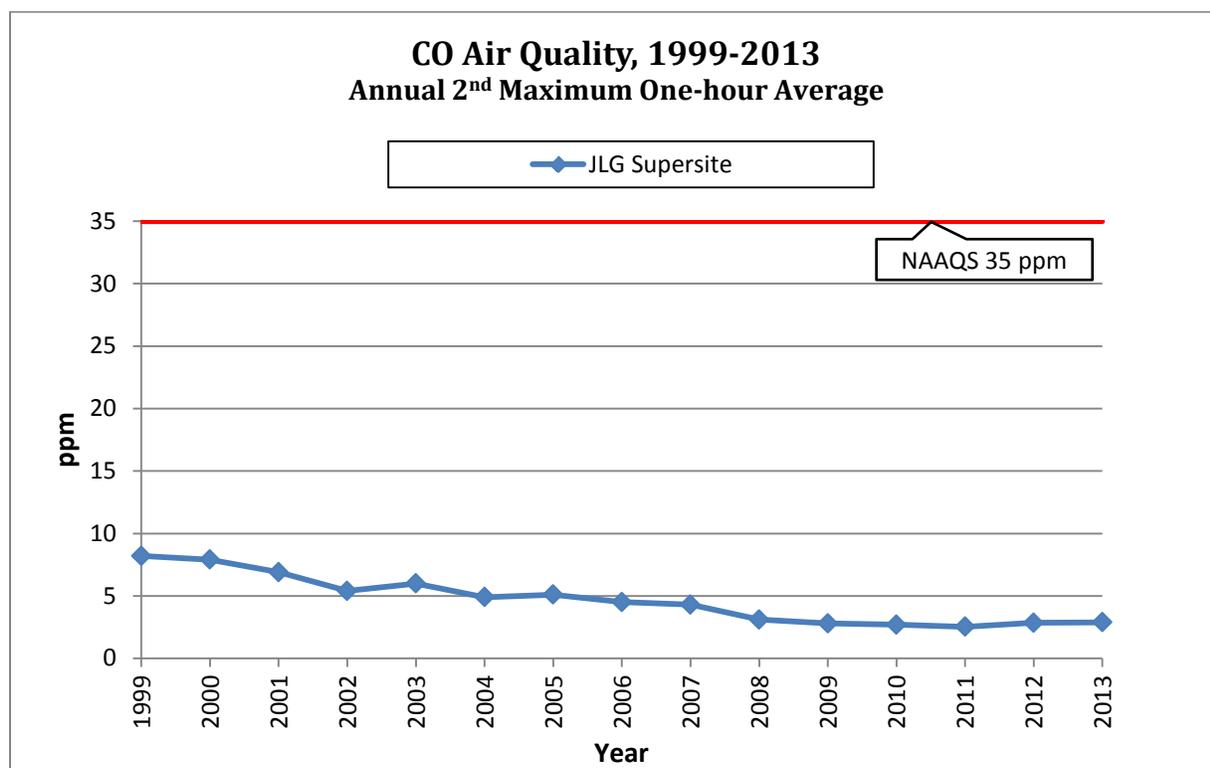


Figure 4 – CO One-Hour Average Trend

1999-2013: 77% decrease for JLG Supersite

Note: Some years might not satisfy completeness criteria.

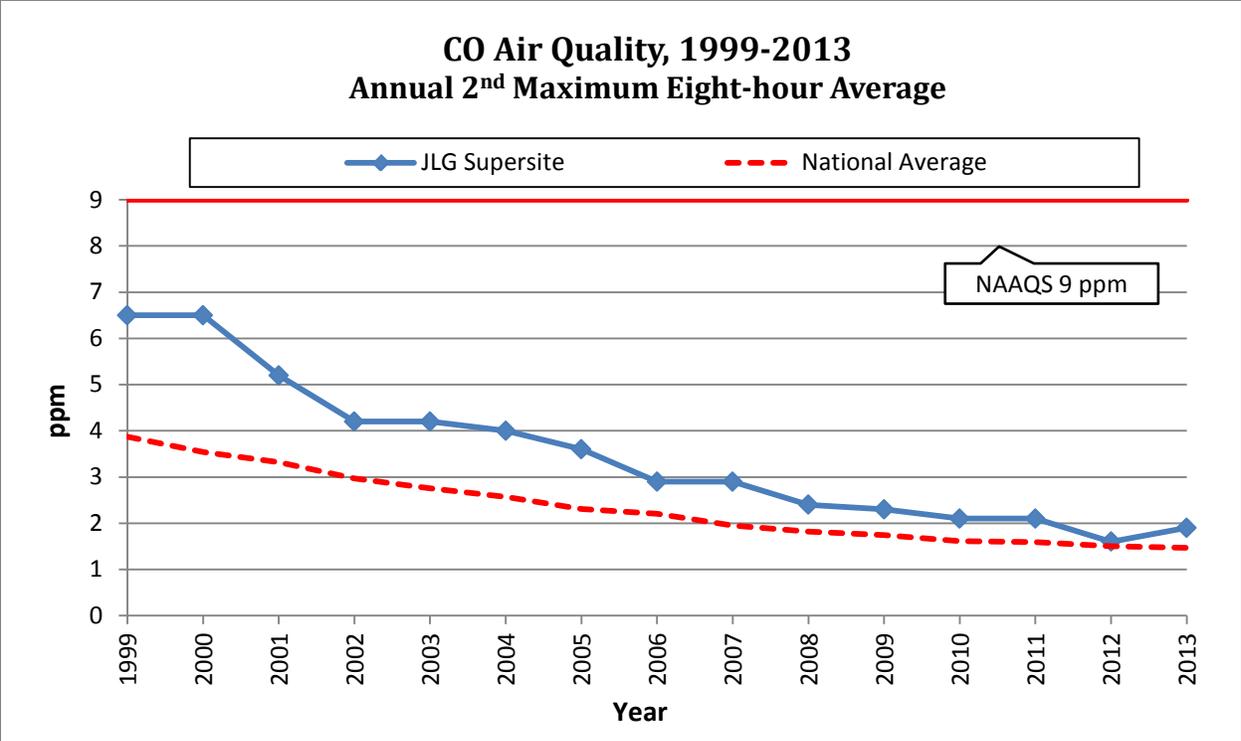


Figure 5 – CO Eight-Hour Average Trend
 1999-2013: 82% decrease for JLG Supersite
 69% decrease in the National average
Note: Some years might not satisfy completeness criteria.

2.0 Nitrogen Dioxide (NO₂)

NO₂ is a reddish-brown gas that is formed by the oxidation of nitric oxide (NO) - a byproduct of combustion. Adverse health effects associated with NO₂ include risk of respiratory illness in children and vary depending on the level of NO₂ and exposure time. Short exposure to low levels may result in changes to airway responsiveness and decreased lung function in individuals with pre-existing conditions. Irreversible changes may occur to lungs due to long-term exposure to higher levels. This pollutant also of concern in its reduction of visibility (it causes five percent of the visibility reduction in Phoenix) and its contributory role in the photochemical formation of ground level O₃ and acid rain.

2.1 Background

Combustion emissions of nitrogen oxides (NO_x) are 95 percent NO and five percent NO₂. Since NO rapidly oxidizes to NO₂, NO₂ concentrations often serve as the indicator for the larger group of nitrogen oxides. In the 2011 NEI, Arizona NO emissions were led by the transportation sector with 66 percent of the emissions from mobile sources such as cars and trucks; 17 percent came from fuel combustion processes such as utility power plants; and 15 percent from other sources, including fires, biogenic emissions from soil, stationary combustion sources and other industrial processes. NO and NO₂ concentrations are highest near major roadways. NO concentrations decrease rapidly with distance from the roadway, whereas NO₂ concentrations are more evenly distributed because of their formation through oxidation and their subsequent transport. Concentrations of NO₂ are highest in the late afternoon and early evening of winter, when rush

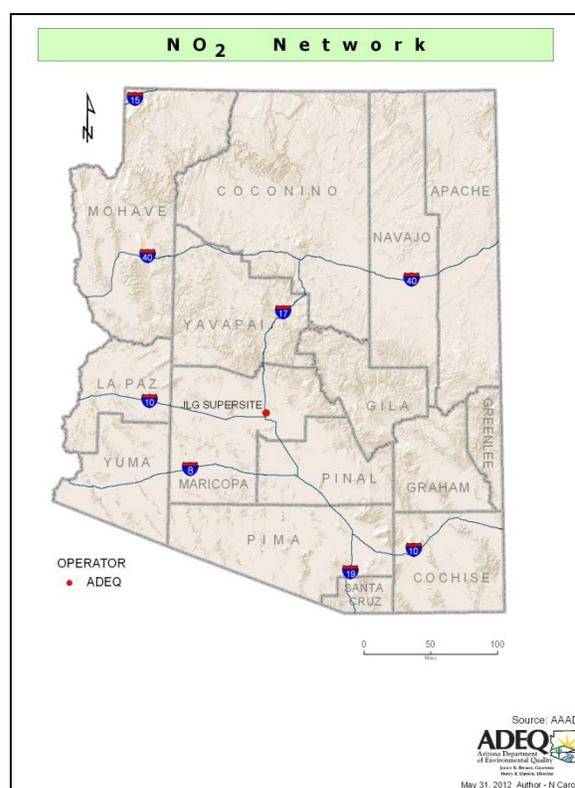


Figure 6 – Map of ADEQ’s NO₂ sites

hour emissions of NO are converted to NO₂ under relatively stable atmospheric conditions. Because NO reacts rapidly with O₃, nocturnal O₃ concentrations in cities are often reduced to near zero levels, while concentrations at background sites remain higher.

NO emissions have been reduced over time using several different techniques. NO emissions from motor vehicles have been reduced through retardation of spark timing, lowering the compression ratio, exhaust gas recirculation systems, and three-way catalysts. Also, the vehicle inspection program’s NO_x test for light-duty gasoline vehicles age 1981 and newer (in Phoenix only) has helped reduce emissions. Reformulated gasolines also decrease NO emissions: Federal Phase II gasoline, by 1.5 percent for

vehicular and 0.5 percent for off-road equipment; California Phase 2 gasoline, by 6.4 percent for vehicular and 7.7 percent for off-road equipment.

One NO₂ monitor was operated by ADEQ in 2013 and is located in a local neighborhood which represents a typical Phoenix area community. This monitor is part of the PAMS at JLG Supersite. This is atypical for the normal siting of NO₂ monitors which normally are located near high traffic roadways. Additionally, two total NO_y monitors were operated by ADEQ. One was operated as part of the NCore station at JLG supersite, and the other at Queen Valley as part of the PAMS network. However, NO_y is not a criteria pollutant and is not further evaluated in this section.

2.2 Monitoring Methods

NO₂ is monitored continuously with chemiluminescence instruments which operate on the principle that when two chemicals combine, a unique wavelength of light is emitted. This wavelength of light is detected using a sensor, and the intensity of that light is a direct correlation to the concentration of the target chemical species (NO₂). An NO₂ analyzer is based on the chemiluminescence of an excited NO₂ molecule which determines NO and NO_x (the sum of NO₂ and NO) concentrations. Readings from all instruments are averaged into hourly, daily, monthly, quarterly, and yearly averages for data analysis. Readings are retrieved by a data collection system and stored in a database.

2.3 Compliance/Summary of Design Values

The NAAQS for NO₂ was reviewed in 2010 and the original 1971 primary NAAQS of 53 parts per billion (ppb) for the annual mean was retained. However, a new primary one-hour NO₂ standard of 100 ppb was added. The annual standard is attained when the annual arithmetic mean concentration in a calendar year is less than or equal to 53 ppb. The one-hour standard is attained when the three-year average of the 98th percentile of the yearly distribution of one-hour daily maximum NO₂ concentrations is below 100 ppb. A history of the NAAQS for NO₂ is provided in Table 6.

Table 6 – History of the National Ambient Air Quality Standards for NO₂ during the period 1971-2010
(Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1971 36 FR 8186 Apr 30, 1971	Primary and Secondary	NO ₂	Annual	53 ppb	Annual arithmetic average
1985 50 FR 25532 Jun 19, 1985	Primary and secondary NO ₂ standards retained, without revision.				

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1996 61 FR 52852 Oct 8, 1996	Primary and secondary NO ₂ standards retained, without revision.				
2010 75 FR 6474 Feb 9, 2010	Primary	NO ₂	1-hour	100 ppb	98 th percentile, averaged over 3 years
		Primary annual NO ₂ standard retained, without revision.			

The 2013 NO₂ annual mean was approximately a third of the 53 ppb standard at JLG Supersite and is in compliance with the NAAQS. Refer to Table 7 for the 2013 annual mean.

Table 7 – NO₂ Annual Mean Compliance Summary

2013 NO₂ Annual Mean (in ppb) (NAAQS Annual Mean 53 ppb)	
Site Name	2013 Annual Mean
Maricopa County	
JLG Supersite	16.98
Number of Sites in Violation of the NAAQS	0

The NO₂ three-year average of the one-hour averages at the 98th percentile was slightly over half the 100 ppb standard at JLG Supersite and is in compliance with the NAAQS. Refer to Table 8 for the 2013 three-year average.

Table 8 – NO₂ One-Hour Compliance Summary

2011 to 2013 One-Hour Average NO₂ Compliance (in ppb) (NAAQS One-Hour Average 100 ppb)				
Site Name	98 th Percentile Samples			Three- Year Average
	2011	2012	2013	
Maricopa County				
JLG Supersite	54.0	58.0	57.0	56
Number of Sites in Violation of the NAAQS				0

2.4 Trends

Monitoring for NO₂ by ADEQ in Phoenix at JLG Supersite began in 1993. However, data are not readily available prior to 1999. As a result, the assessment of trends in NO₂ uses a period of fifteen years from 1999 to 2013. Figures 7 and 8 illustrate the temporal variability of JLG Supersite over the 1999 to 2013 period in the form of annual means and one-hour averages at the 98th percentile. The NO₂ trend can be described as decreasing over this fifteen-year period.

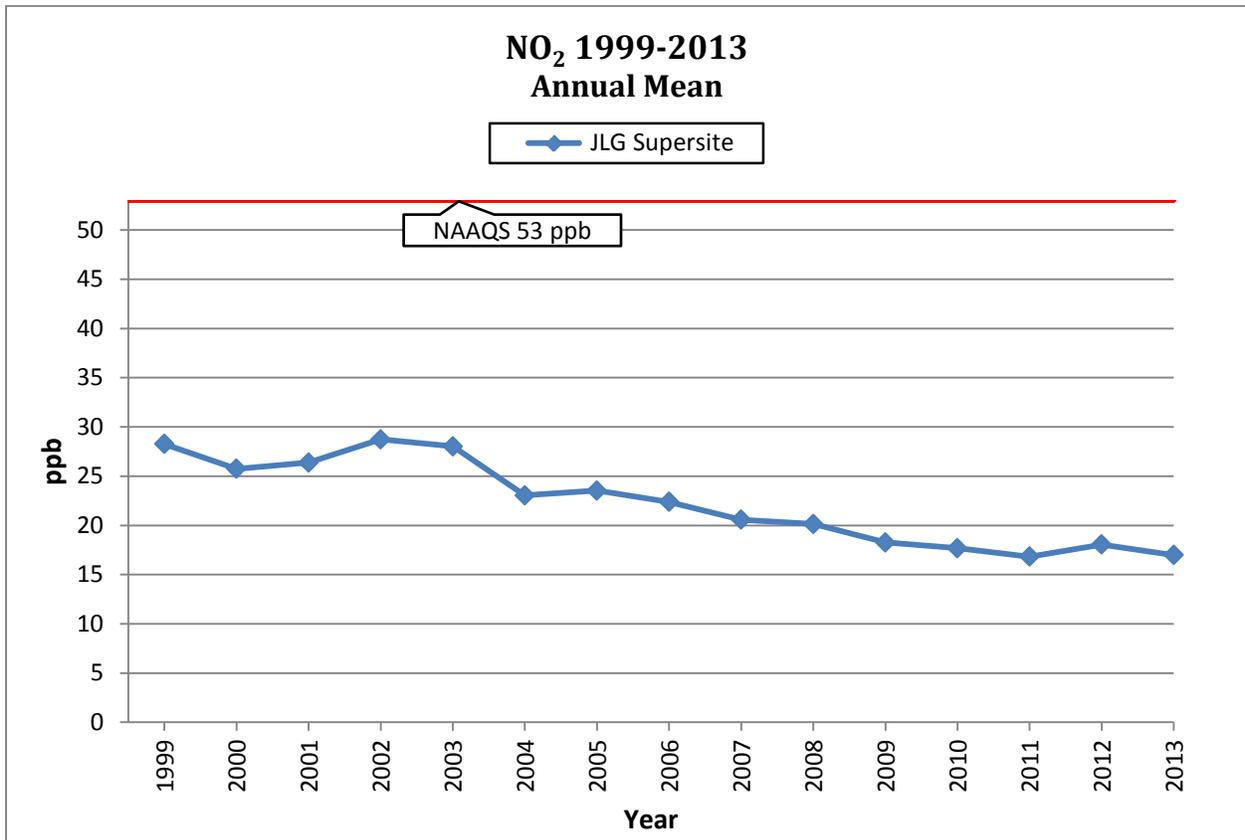


Figure 7 – NO₂ Annual Mean Trend
1999-2013: 46.2% decrease for JLG Supersite
Note: Some years might not satisfy completeness criteria.

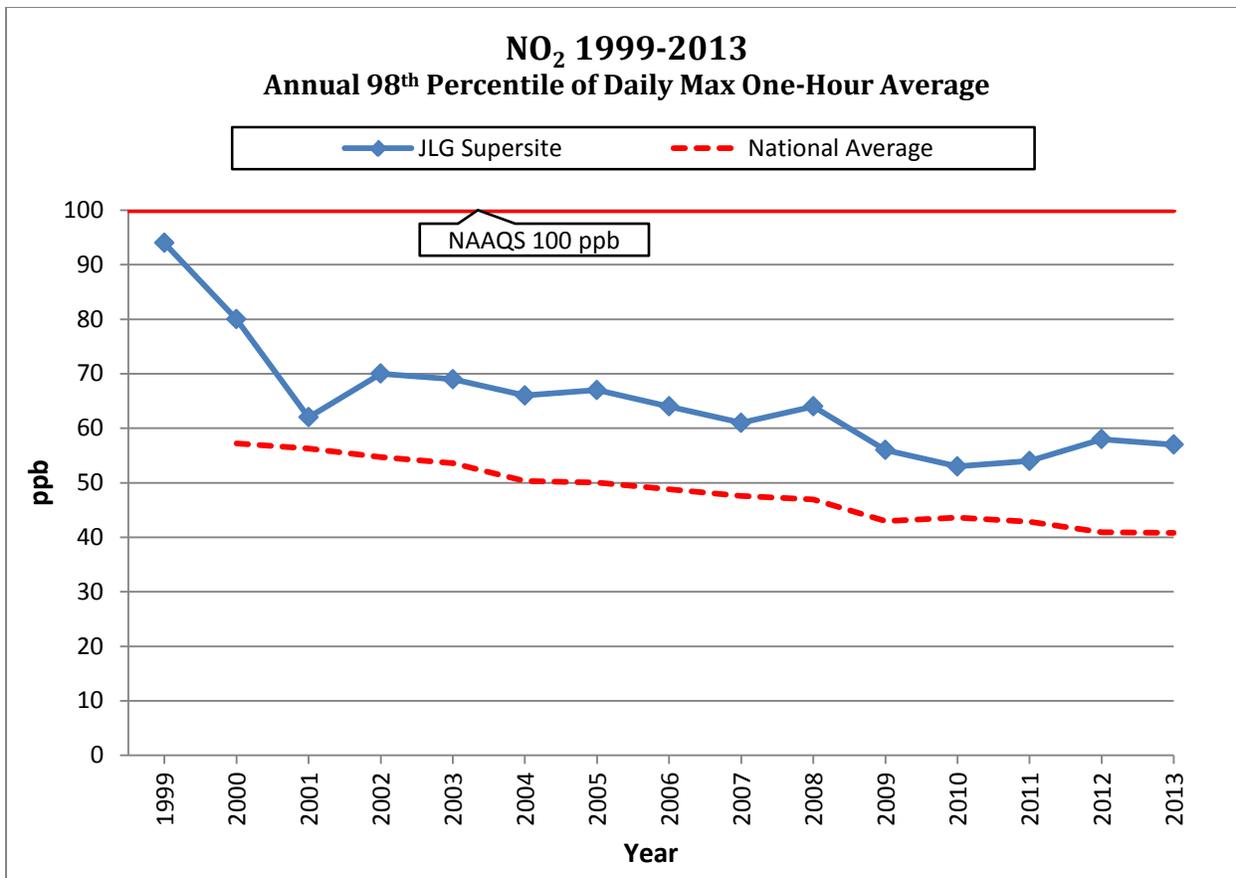


Figure 8 – NO₂ One-Hour Average Trend
 1999-2013: 36.4% decrease for JLG Supersite
 2000-2013: 29.0% decrease in the National Average
Note: Some years might not satisfy completeness criteria.

3.0 Ozone (O₃)

Ozone is a colorless, slightly odorous gas that is both a natural component of the upper atmosphere and a key air contaminant in the lower atmosphere. In the stratosphere, O₃ blocks harmful ultraviolet radiation. In the lower atmosphere, its photochemical formation by the reaction of volatile organic compounds (VOC), hydrocarbons (HC), and NO_x, leads to concentrations harmful to people, animals, plants, and materials (plastics, tires, o-rings). In both animals and humans O₃ causes significant physiological and pathological changes at concentrations present in many urban environments. Short-term (one to two hours) exposures to concentrations in the range of 100 ppb to 400 ppb induce changes in lung function, including increased respiratory rates, increased pulmonary resistance, decreased volume of air, and changes in lung mechanics. Symptomatic responses in exercising adults include throat dryness, chest tightness, substernal pain, coughing, wheezing, pain on deep inspiration, shortness of breath, and headache. These symptoms also have been observed at lower concentrations for longer exposures. Evidence suggests that O₃ exposure makes the respiratory airways more susceptible to other bronchoconstrictive challenges and interferes with or inhibits the immune system. Ozone at ambient concentrations also injures the stomata of plants, which are the cells that regulate plant respiration, resulting in flecks on the upper leaf surfaces of dichotomous plants and the death of the tips of coniferous needles. Plant scientists consider O₃ to be the most important of all of the phytotoxic air pollutants, causing over 90 percent of all plant injury from air pollution on a global basis.

3.1 Background

High O₃ concentrations are a summer phenomenon caused when sunlight, emissions from plant life, and evaporating fuel emissions peak. VOC emissions in greater Phoenix come from cars and trucks, off-road vehicles and equipment such as lawn mowers, small stationary sources, biogenic emissions from grass, shrubs, and trees, and point sources. Nitrogen oxides come from cars and trucks, off-road vehicles such as construction equipment and trains, electric power plants, small stationary sources, and biogenic emissions from soil. Ozone has relatively high background levels, with the daily maximum in remote areas being about one-half to three-quarters of the daily maximum in the urban areas. In an urban area, the highest O₃ concentrations tend to occur on the downwind edge, although high concentrations do occur less frequently in the central city. Urban O₃

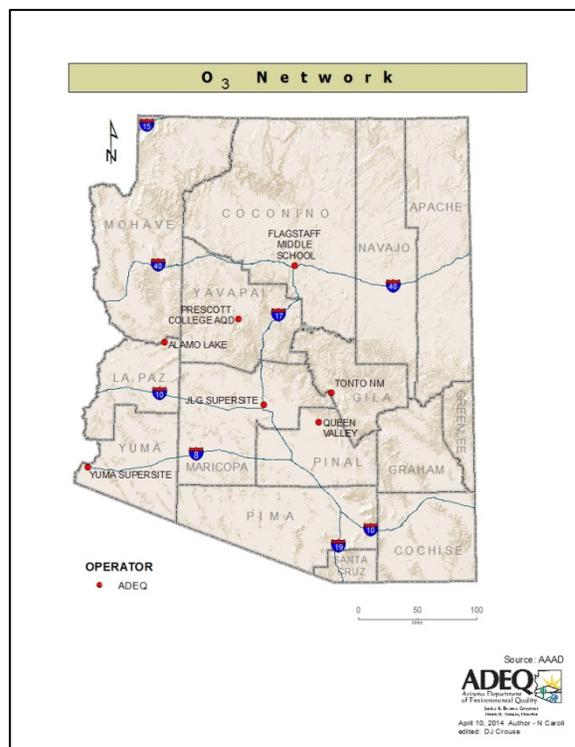


Figure 9 – Map of ADEQ's O₃ sites

concentrations are low to near zero at night and rise rapidly through the morning and peak in the afternoon.

Controls to reduce the precursors of O₃, VOC, and NO_x have been successfully implemented for years. Nitrogen oxides and VOC from vehicular exhaust have been reduced through engine modifications and three-way catalytic converters. Evaporative HC from vehicles have been reduced through better engineered fuel tanks and auxiliary plumbing combined with carbon absorption canisters. Additional reductions of vehicular VOC have come through ADEQ's vehicle emissions inspection program, which tests all gasoline fueled vehicles for HC (Phoenix and Tucson), through vapor capturing equipment for gasoline tankers, vapor recovery systems at retail gas stations (Phoenix area only), and cleaner burning gasoline (Phoenix area only). Stationary sources of HC have been reduced through a variety of better control equipment required by stricter regulations. Despite these efforts, the continued population growth in Arizona combined with the high natural background O₃, may make achieving the eight-hour standard difficult.

In 2013, ADEQ operated a network of seven O₃ monitors to support a variety of monitoring objectives; chief among them is for NAAQS compliance. Other monitoring objectives include support for the NCore station, PAMS, and to show O₃ transport coming from across Arizona's border.

3.2 Monitoring Methods

Continuous monitoring for O₃ is done with an ultraviolet absorption instrument. A specific ultraviolet wavelength of light which O₃ absorbs is passed through a sample cell. A drop in light intensity is detected by a sensor and that drop is a direct correlation to the concentration of O₃ in the sample cell. This results in accurate readings of O₃ concentrations continuously. Readings are averaged into hourly, daily, monthly, quarterly, and yearly averages for data analysis. Readings are retrieved by a data collection system and stored in a database.

3.3 Compliance/Summary of Design Values

On May 21, 2012, EPA published two rules; (1) the final implementation rule of the 2008 NAAQS for O₃: Nonattainment Area Classifications Approach, Attainment Deadlines, and Revocation of the 1997 O₃ Standards for Transportation Conformity Purposes and (2) Air Quality Designations for the 2008 O₃ NAAQS. The implementation rule establishes classifications and associated attainment deadlines and revoked the 1997 O₃ standards for transportation conformity purposes. The designation rule finalized the nonattainment area boundaries for areas that do not meet the 0.075 ppm standard. The standard is met when the 4th highest rolling eight-hour average for the year, averaged over three years is less than the 0.075 ppm standard. Thus an exceedance above the 0.075 ppm standard for a given year may not cause a violation of standard, if the three-year average is still below 0.075 ppm. The designations are based on air quality monitoring data and the history of the NAAQS for O₃ is provided in Table 9 below.

Table 9 – History of the National Ambient Air Quality Standards for O₃ during the period 1971-2008
(Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
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 ₃	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 ₃	8-hour	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 ₃	8-hour	0.075 ppm	Annual fourth-highest daily maximum 8-hr concentration, averaged over 3 years

Area designations occur after the new rules are promulgated and the last designations occurred in 2012 for the 2008 rule. Parts of Maricopa and Pinal counties are the only areas in Arizona designated nonattainment for the 2008 rule. All the other areas are classified as attainment for the standard. Yuma is still classified as attainment for the 2008 rule since additional area designations have not occurred since 2012. The three-year average for Yuma is above the standard for the time period of 2011-2013; however, area designations made in 2012 were based on 2008-2010 data and showed Yuma in attainment. See Attachment III for a map of area designations in Arizona.

The data in Table 10 are from the sites in operation in 2011 to 2013 and have been evaluated based on the 2008 O₃ standard (0.075 ppm). Three sites did not meet this standard: JLG Supersite, Queen Valley, and Yuma Supersite. The other four sites were in compliance with the 0.075 ppm O₃ standard.

Table 10 – O₃ Compliance Summary

2011 to 2013 Eight-Hour O₃ Compliance (in ppm) <i>(NAAQS eight-hour 0.075 ppm)</i>				
Site Name	Fourth-Highest Value			Three-Year Average
	2011	2012	2013	
Coconino County				
Flagstaff Middle School ¹	0.068	0.072	0.069	0.069
Number of Sites in Violation of the NAAQS				0
Gila County				
Tonto NM ¹	0.076	0.078	0.072	0.075
Number of Sites in Violation of the NAAQS				0
La Paz County				
Alamo Lake ¹	0.072	0.075	0.071	0.072
Number of Sites in Violation of the NAAQS				0
Maricopa County				
JLG Supersite	0.078	0.076	0.079	0.077
Number of Sites in Violation of the NAAQS				1
Pinal County				
Queen Valley ¹	0.078	0.078	0.073	0.076
Number of Sites in Violation of the NAAQS				1
Yavapai County				
Prescott College AQD ¹	0.070	0.072	0.065	0.069
Number of Sites in Violation of the NAAQS				0
Yuma County				
Yuma Supersite ¹	0.076	0.080	0.073	0.076
Number of Sites in Violation of the NAAQS				1

¹ Seasonal monitor, operational during April 1st to November 1st.

Bold denotes exceedance of the 2008 NAAQS of 0.075 ppm for the time period of 2011-2013.

3.4 Trends

Monitoring for O₃ has occurred for many years in Arizona, mainly in the urbanized centers of Phoenix and Tucson. For the purposes of this report, the examination of O₃ trends will include the monitors being run by ADEQ only for the years 2005-2013. The sites included in the trends graphs are Flagstaff Middle School, Tonto National Park, Alamo Lake, JLG Phoenix Supersite, Queen Valley, Prescott College, and Yuma Supersite. In general, the trend shown in Figure 10 can be described as slightly decreasing from 2005 to 2013. In 2009 all sites reported a substantial drop in USG days (Unhealthy for Sensitive Groups). According to the National Oceanic and Atmospheric Administration (www.noaa.gov) observations indicated that O₃ was on the low side of historical observations all year long. In 2008, the Yuma instrument was moved from the Yuma Game & Fish site to Yuma Supersite. The annual values of both sites met the completeness criteria and were averaged for the purpose of the trends graph.

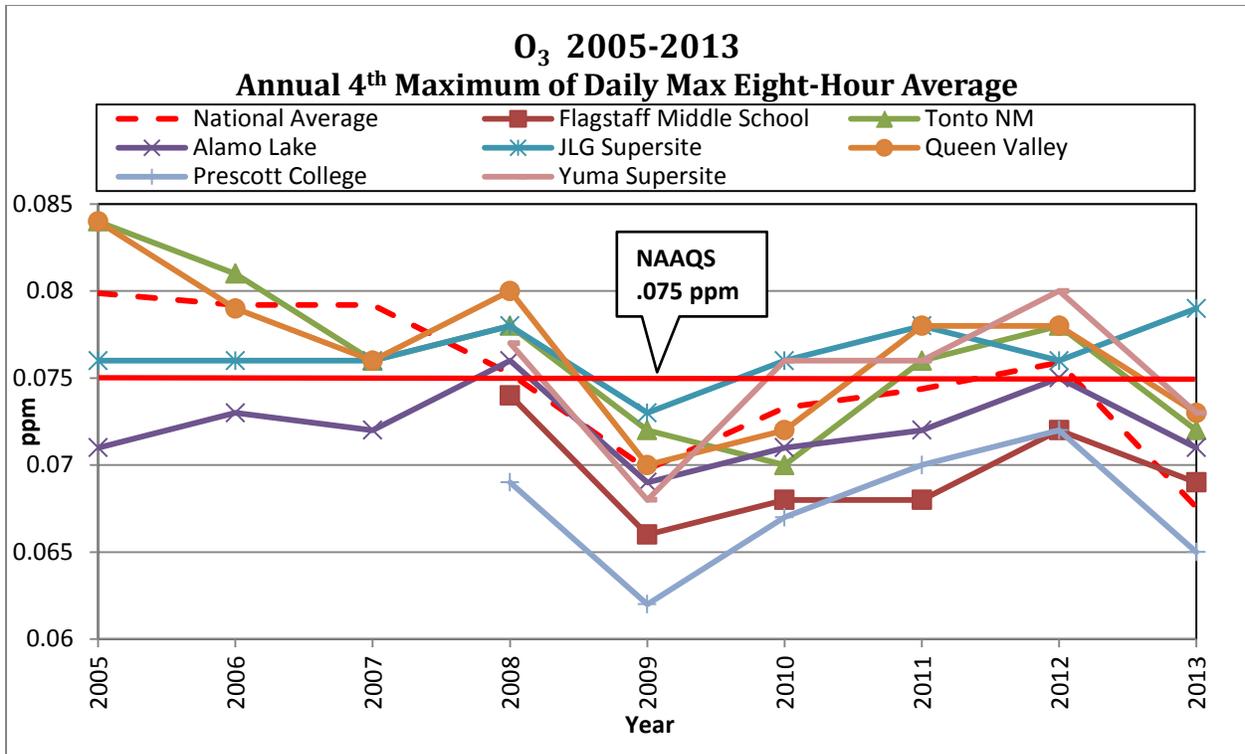


Figure 10 – O₃ Eight-Hour Average Trend

2005-2013: 4.6% decrease for the average of Tonto NM, Alamo Lake, JLG Supersite, and Queen Valley

2005-2013: 7.5% decrease in the National Average

Note: Some years might not satisfy completeness criteria. All O₃ sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least seven consecutive years of data and the same number of consecutive years.

4.0 Sulfur Dioxide (SO₂)

Sulfur Dioxide is a colorless gas with a pungent, irritating odor at elevated concentrations. It mainly alters the mechanical function of the upper airway, including increasing the nasal flow resistance and decreasing the nasal mucus flow rate. Current scientific evidence links short-term exposures to SO₂, ranging from five minutes to 24 hours, with an array of adverse respiratory effects, including bronchoconstriction and increased asthma symptoms. These effects are particularly significant for asthmatics at elevated ventilation rates. Studies have shown a connection between short-term exposure and increased visits to emergency departments and hospital admissions for respiratory illnesses, particularly in at-risk populations including children, the elderly, and asthmatics.

4.1 Background

In Arizona, the principal source of SO₂ emissions has been the smelting of sulfide copper ore. Most fuels contain trace quantities of sulfur and their combustion releases both gaseous SO₂ and particulate sulfate. In the 2011 NEI, Arizona showed 45 percent of SO₂ emissions came from industrial point sources including smelting, 38 percent from fuel combustion, 15 percent from fires, and 2 percent from mobile sources such as off-road vehicles and on-road motor vehicles. Sulfur Dioxide is removed from the atmosphere through dry deposition on plants and is converted to sulfuric acid and eventually to sulfate. Sulfur Dioxide has extremely low background levels, with elevated concentrations found downwind of large point sources. Concentrations in urban areas are low and are homogeneously distributed, with annual averages varying from 0.0010 ppm to 0.0020 ppm, well within the annual standard of 0.03 ppm.

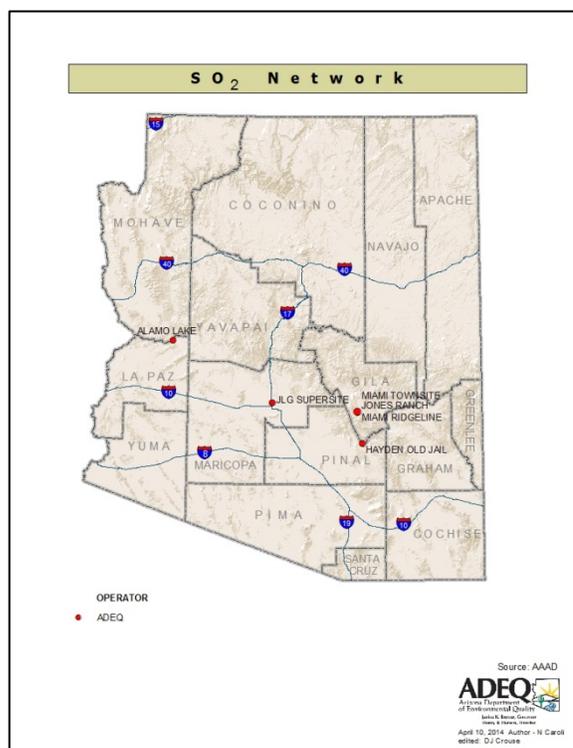


Figure 11 – Map of ADEQ's SO₂ sites

Major controls were installed in Arizona's copper smelters in the 1980s, which reduced SO₂ emissions substantially. Vehicular emissions of SO₂ and sulfate have been reduced through lowering the sulfur content in diesel fuel and gasoline.

Controls have reduced SO₂ emissions throughout the state in recent years, but there are still two significant point sources which are affecting nearby air quality. Copper smelting operations have caused the areas in Miami, AZ and Hayden, AZ to be designated by the EPA as nonattainment areas for the 2010 standard. Compliance with the new NAAQS standard is achieved throughout the whole state besides these two areas.

ADEQ operated four source oriented monitors in 2013. Three are located in and around the Miami, AZ area and one is located in Hayden, AZ. One additional monitor was operated at JLG supersite as part of the NCore station.

4.2 Monitoring Methods

Continuous monitoring for SO₂ is done with pulsed fluorescence instruments. Sulfur Dioxide absorbs a specific wavelength of ultraviolet light. Absorption of light at this wavelength results in the fluorescent emission of light at a different wavelength. This second wavelength is directly proportional to the SO₂ concentration in the sample. The wavelengths of light are isolated using bandpass filters. Readings are averaged into hourly, daily, monthly, quarterly, and yearly averages for data analysis and can be retrieved by a data collection system and stored in a database.

4.3 Compliance/Summary of Design Values

On April 30, 1971, the EPA promulgated primary and secondary NAAQS for sulfur oxides, measured as SO₂ under section 109 of the Act (36 FR 8186). After periodic reviews of additional scientific information, EPA announced first in 1986 and then in 1996, its decision not to revise the NAAQS for SO₂. In 2010, EPA replaced both the 24-hour and annual standards with a new short-term standard based on the three-year average of the 99th percentile of the yearly distribution of one-hour daily maximum SO₂ concentrations. EPA set the level of this new standard at 75 ppb. The one-hour SO₂ standard added in 2010 is a primary standard. The revision of the SO₂ NAAQS in 2010 did not address the secondary standard, which remains a three-hour standard with a level of 0.5 ppm. Table 11 summarizes the history of the NAAQS for oxides of sulfur during the period 1971-2010.

Table 11 – History of the National Ambient Air Quality Standards for SO₂ during the period 1971-2010
(Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1971 36 FR 8186 Apr 30, 1971	Primary	SO ₂	24-Hour	0.14 ppm	Not to be exceeded more than once per year
			Annual	0.03 ppm	Annual arithmetic average
	Secondary		3-Hour	0.5 ppm	Not to be exceeded more than once per year
			Annual	0.02 ppm	Annual arithmetic average
1973 38 FR 25678 Sept 14, 1973	Secondary	Secondary 3-hour SO ₂ standard retained, without revision; secondary annual SO ₂ standard revoked.			
1996 61 FR 25566 May 22, 1996	Primary	Existing primary SO ₂ standards retained, without revision.			

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
2010 75 FR 35520 Jun 22, 2010	Primary	SO ₂	1-hour	75 ppb	99 th percentile, averaged over 3 years
Primary annual and 24-hour SO ₂ standards revoked.					

Table 12 summarizes the status for the annual 99th percentile of one-hour daily maximum for SO₂ for the years 2011 to 2013. ADEQ collocated SO₂ monitors at Miami Jones Ranch and Miami Townsite locations of Freeport McMoRan Inc. (FMMI) Miami stations in 2013 hence data are not available for the years 2011 and 2012. Hayden Old Jail and Miami Ridgeline are sites in violation of the 2011 NAAQS of 75 ppb.

Table 12 – SO₂ One-Hour Compliance Summary

2011 to 2013 Annual 99th % of One-Hour daily max. SO₂ Compliance (in ppb) (NAAQS primary one-hour 75 ppb)				
Site Name	2011	2012	2013	Three- Year Average
Gila County				
Miami Jones Ranch	N/A	N/A	148*	N/A
Miami Townsite	N/A	N/A	117*	N/A
Hayden Old Jail	189	353	256	266
Miami Ridgeline	96	102	117	105
Number of sites in violation of NAAQS				2
Maricopa County				
JLG Supersite	5*	6	6*	N/A
Number of sites in violation of NAAQS				N/A

*Indicates the data do not satisfy EPA's summary criteria.

N/A-Data are not available

Bold denotes exceedances and sites in violation of the 2011 NAAQS of 75 ppb.

Table 13 summarizes the status for the three-hour secondary standard for SO₂ for the years 2011 to 2013. All sites are well below the secondary standard of 0.5 ppm. ADEQ collocated SO₂ monitors at Miami Jones Ranch and Miami Townsite location of FMMI Miami stations in 2013 hence data are not available for the years 2011 and 2012.

Table 13 – SO₂ Three-Hour Compliance Summary

2011 to 2013 Three-Hour SO₂ Compliance (in ppm) (NAAQS secondary three-hour 0.5 ppm)						
Site Name	2011		2012		2013	
	1st max value	2nd max value	1st max value	2nd max value	1st max value	2nd max value
Gila County						
Miami Jones Ranch	N/A	N/A	N/A	N/A	0.1266*	0.1193*
Miami Townsite	N/A	N/A	N/A	N/A	0.0766	0.0396
Hayden Old Jail	0.1746	0.1163	0.2656	0.2163	0.1656	0.158
Miami Ridgeline	0.0943	0.0843	0.119	0.045	0.0903	0.0806
Number of sites in violation of NAAQS						0
Maricopa County						
JLG Supersite	0.0039*	0.0038*	0.0062	0.005	0.0061*	0.0043*
Number of sites in violation of NAAQS						0

*Indicates the data do not satisfy EPA's summary criteria.

N/A-Data are not available

4.4 Trends

In Arizona SO₂ monitoring began in the late 1960s and early 1970s. These early sites were predominantly located near facilities where smelting of sulfide copper ore occurred. Monitoring SO₂ at several of these smelting facilities no longer occurs due to the termination of smelting operations. However, a long SO₂ monitoring record does exist for the Hayden and Miami areas due to continued smelting operations in these areas. Only monitors operated by ADEQ are used for the purpose of assessing SO₂ trends in this report. For the period of 1975-2013, three sites were used to assess trends in SO₂ as shown in Figure 12. These sites are Hayden Old Jail, Miami Ridgeline, and JLG Supersite. Figure 12 does not reflect the trends from Miami Jones Ranch and Miami Townsite as ADEQ started monitoring at these locations in 2013. Beginning 2011, trace-level SO₂ monitoring began in the Phoenix area as part of EPA's NCore program. The sensitivity of this monitor is far greater than the monitors used earlier, and low concentrations of SO₂ in the urban environments can be assessed with greater confidence, therefore a separate trend graph will be shown (Figure 13). The annual average trend for Hayden Old Jail is shown in Figure 14, which indicates that total output of the sources has not changed over the past 15 years. However, the increase in the daily max one-hour average trend at Hayden Old Jail from 1999-2013 (Figure 12) indicates that the one-hour averages have increased. This means that the highs are higher, and the lows are lower.

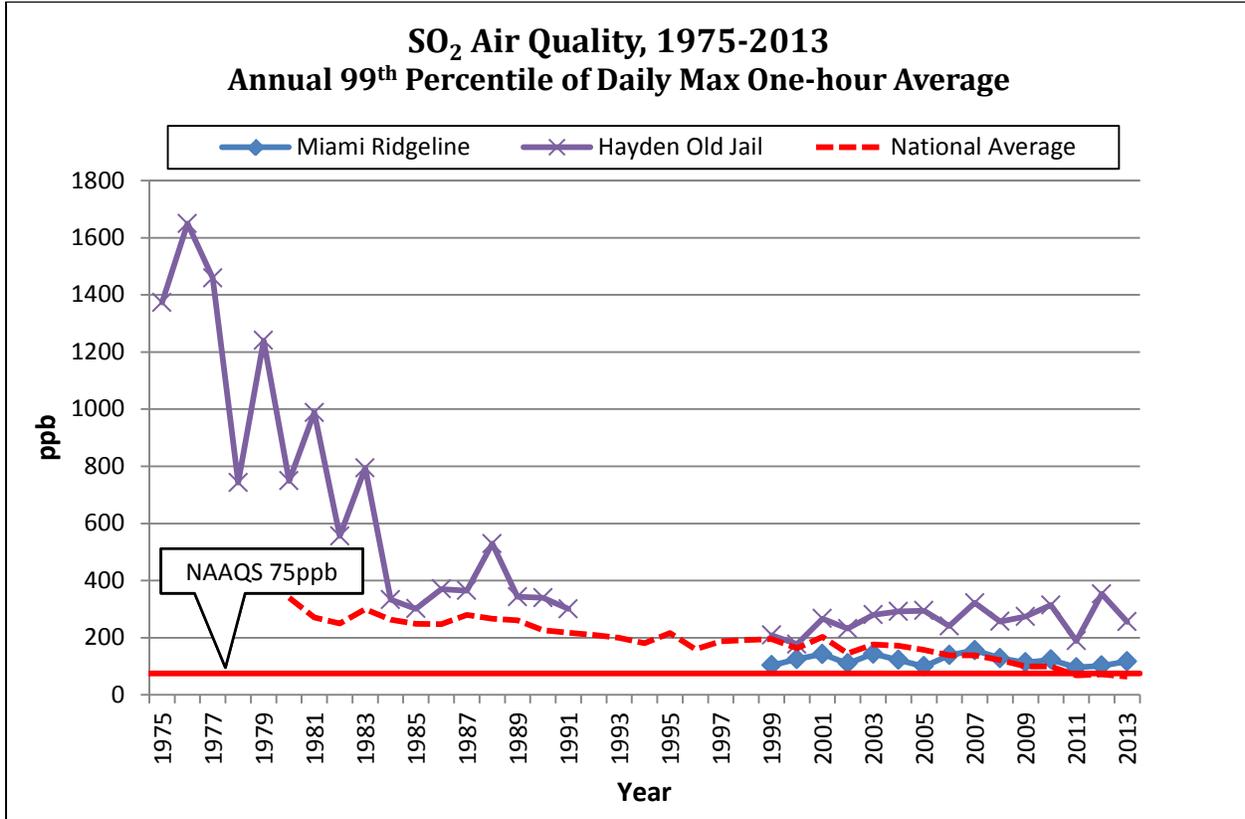


Figure 12 – SO₂ One-Hour Average Trend for Miami and Hayden Areas

1975-1991: 92.38% decrease for Hayden Old Jail

1980-1991: 24.56% decrease in the National average

1999-2013: 10.34% decrease for Miami Ridgeline

1999-2013: 29.68% increase for Hayden Old Jail

1999-2013: 67.92% decrease in the National average

Note: Some years might not satisfy completeness criteria.

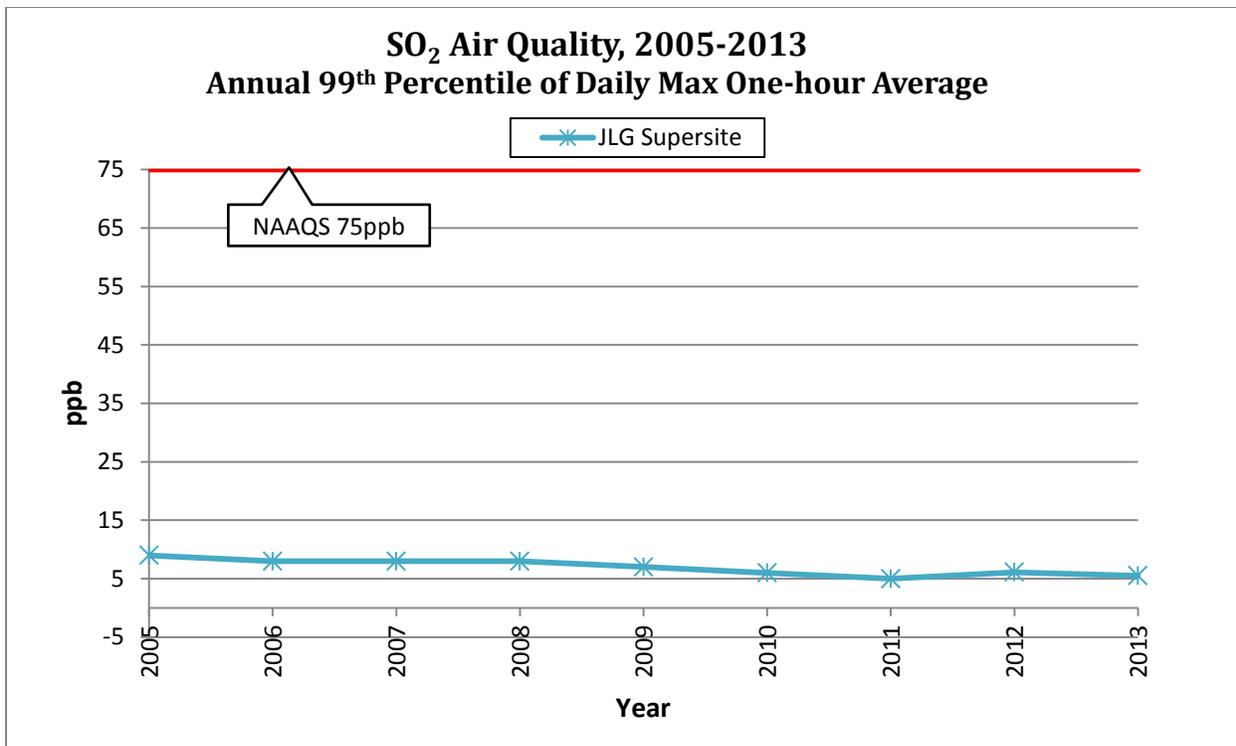


Figure 13 – SO₂ One-Hour Average Trend for JLG Supersite
2005-2013: 3.987% decrease for JLG Supersite
Note: Some years might not satisfy completeness criteria.

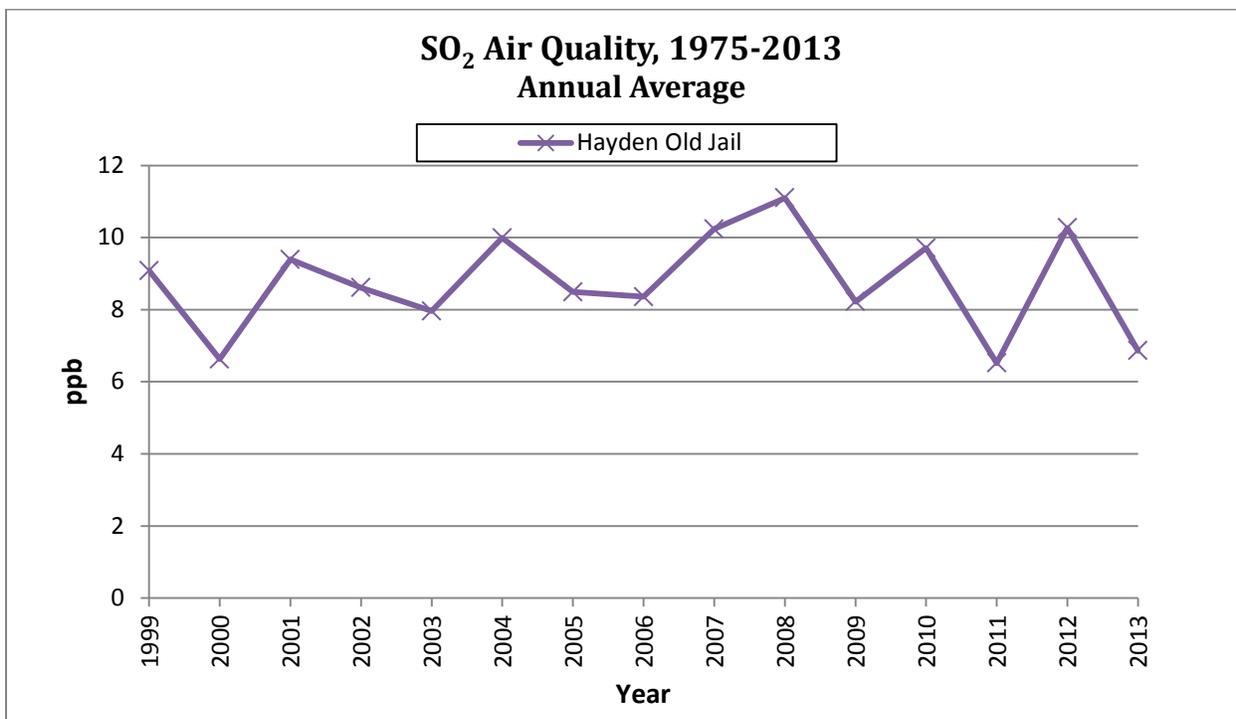


Figure 14 – SO₂ Annual Average Trend
1999-2013: 0.67% negligible increase for Hayden Old Jail
Note: Some years might not satisfy completeness criteria.

5.0 Particulate Matter Smaller Than 10 Microns (PM₁₀)

Particulate matter is a collective term describing very small solid or liquid particles that vary considerably in size, geometry, chemical composition, and physical properties. PM₁₀ describes particles that are less than 10 microns in diameter. Produced by natural processes (pollen and wind erosion) and by human activity (soot, fly ash, and dust from paved and unpaved roads), particulates contribute to visibility reduction, pose a threat to public health, and cause economic damage through soil disturbances. The size, shape, and chemical composition of particulates determine their health effects. Particles from 2.5 to 10 microns are inhaled and deposited in the upper parts of the respiratory system. Epidemiological studies have shown causal relationships between particulates and excess mortality, aggravation of bronchitis, and small reversible changes in pulmonary function in children.

5.1 Background

PM₁₀ emissions in Arizona are mostly geological in origin and are dominated by dust from three activities: the constant grinding (re-entraining) of dust from paved roads, driving on unpaved roads, and earth moving associated with construction. On days with winds in excess of 15 miles per hour, wind erosion of soil may contribute to this loading. Soil dust from these and other sources, such as high winds, contribute to about 52 percent of the PM₁₀ in Arizona. Additionally agricultural sources make up about 22 percent, fires make up about 13 percent, and the remaining 12 percent includes fuel combustion, mobile sources, and other industrial processes.

PM₁₀ concentrations are not spatially distributed evenly because each monitoring site is strongly influenced by the degree of localized emissions of particulates. PM₁₀ maximum concentrations can occur in any season, provided nearby sources of coarse particulates are

present or when strong and gusty winds suspend soil disturbed by human activities. Hourly concentrations of particulates tend to peak during those hours of the worst distribution, which is from sunset to midmorning.

Controls to reduce particulates in Arizona have been in place for decades, beginning in the 1960s with a Pima County ordinance that required watering to reduce dust from construction. Maricopa County's umbrella dust abatement rule, Rule 310, has been revised many times through the years and now regulates construction dust, trackout dust (accumulation on tires of vehicles) from construction sites,



Figure 15 – Map of ADEQ's PM₁₀ sites

and dust from unpaved parking and vacant lots. Efforts to reduce dust re-suspended from paved roads have concentrated on eliminating trackout from construction sites, curbing and stabilizing road shoulders, and investigating more efficient street sweepers. In Maricopa County, the Governor's Agricultural Best Management Practices Committee developed a rule containing best management practices for agricultural activities (AgBMP) to reduce particulate emissions from tilling and harvesting activities of cropland and non-cropland. In a recent PM₁₀ SIP, the Maricopa Association of Governments (MAG) committed to implement 77 new measures including enhanced enforcement of the county's dust rules, implementation of AgBMP, diesel engine replacement and retirement programs, and requirements for cleaner burning fireplaces.

Controls on PM₁₀ have been successful in limiting anthropogenic emissions in recent years. Due to natural sources of PM₁₀ in Arizona, much of the state is in nonattainment for PM₁₀ for the 1987 standard. PM₁₀ is one of the major pollutants affecting the health of the people of Arizona and as such is a major part of ADEQ's ambient air monitoring. In 2013, ADEQ operated a network of 15 monitors throughout the state.

5.2 Monitoring Methods

ADEQ is utilizing several methods for measuring PM₁₀. The two general types are filter based and continuous instruments.

Particulates can be monitored using a weighing and filter based method. This is done by pulling ambient air through a filter for 24 hours every sixth day (or as designated for the monitor per the CFRs), weighing the filter before and after sampling and measuring the volume of air sampled. Pollutant concentrations are measured by the weight of pollutant within a standard volume of air, for example micrograms per cubic meter ($\mu\text{g}/\text{m}^3$) or milligrams per cubic meter (mg/m^3). Weight and volume of air sampled are calculated to give a daily concentration. Weight is determined from an automated weighing system in ADEQ's air filter lab. The 24-hour monitoring instruments are fitted with different aerodynamic devices (inlets) to segregate particle size fractions. The particles can be segregated into two size ranges (less than 10 microns and less than 2.5 microns). This method provides one 24-hour concentration.

Particulates can also be monitored continuously with a tapered element oscillating microbalance (TEOM) instrument which utilizes the same inlets as the filter instruments to segregate particle sizes. PM particles are deposited onto an oscillating filter, which changes its frequency of oscillation based on the weight of particles deposited. This change in frequency is a direct correlation to the concentration of PM in ambient air based on the volume of air sampled in a given time period. This change is recorded by the instrument every hour to get an hourly PM concentration.

Another type of instrument called a beta attenuation mass monitor (BAM), utilizes the same inlet as filter instruments to segregate particles. This method of detection utilizes a radioactive beta source to measure the particles through a paper tape. The rate that a beta source decomposes is known and does not change; therefore a sensor can detect this rate of radiation. Any changes to that rate can be recorded and this change in readings is a direct correlation to the concentration of PM. Particles block

or interfere with the beta radiation by absorbing or causing the beta radiation to change. Readings on blank part of the paper tape is taken before sampling and with an airflow rate of 16.7 liters per minute particles are deposited on the paper tape for about one hour. A reading is taken on the tape where the particles are deposited to determine the concentration of PM. The volume of air sampled is also used to determine the concentration.

Readings from continuous types of instruments (TEOM and BAM) are averaged into hourly, daily, monthly, quarterly, and yearly averages for data analysis. Readings are retrieved by a data collection system and stored in a database. Filter data are stored by air filter lab weighing equipment and uploaded to the database.

5.3 Compliance/Summary of Design Values

In 2012, EPA revised the suite of standards for PM. EPA retained the 24-hour PM₁₀ standard of 150 µg/m³ which has been in place since 1987 and revoked the annual PM₁₀ standard. The 24-hour PM₁₀ standard is met when the 24-hour average (rounded to the nearest 10 µg/m³) does not exceed 150 µg/m³ more than once per year on average over a three-year period. A history of the NAAQS for PM₁₀ is provided in Table 14 below.

Table 14 – History of the National Ambient Air Quality Standards for PM₁₀ during the period 1971-2012
(Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1971 36 FR 8186 Apr 30, 1971	Primary	TSP	24-hour	260 µg/m ³	Not to be exceeded more than once per year
			Annual	75 µg/m ³	Annual Average
1987 52 FR 24634 Jul 1, 1987	Primary and Secondary	PM ₁₀	24-hour	150 µg/m ³	Not to be exceeded more than once per year on average over a 3-year period
			Annual	50 µg/m ³	Annual arithmetic mean, averaged over 3 years
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM ₁₀	24-hour	150 µg/m ³	Initially promulgated 99th percentile, averaged over 3 years; when 1997 standards for PM10 were vacated, the form of 1987 standards remained in place (not to be exceeded more than once per year on average over a 3-year period)
			Annual	50 µg/m ³	Annual arithmetic mean, averaged over 3 years

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
2006 71 FR 61144 Oct 17, 2006	Primary and Secondary	PM ₁₀	24-hour	150 µg/m ³	Not to be exceeded more than once per year on average over a 3-year period
2012	Primary and Secondary	PM ₁₀	24-hour	150 µg/m ³	Not to be exceeded more than once per year on average over a 3-year period

Table 15 presents the 2011 to 2013 expected exceedance rates for the PM₁₀ sites in Arizona, along with the annual maximum 24-hour concentration. The 24-hour primary PM₁₀ NAAQS of 150 µg/m³ has been exceeded in several counties in the 2011 to 2013 time period. Parts of the following counties are currently designated nonattainment with the 1987 PM₁₀ NAAQS of 150 µg/m³: Gila, Pima and Yuma. See Appendix III for the nonattainment area map.

Table 15 – PM₁₀ Compliance Summary

2011 to 2013 Maximum 24-Hour Average PM₁₀ Compliance (in µg/m³)							
Bold denotes value above the standard.							
(NAAQS 24-hour Average 150 µg/m³)							
Site Name	2011		2012		2013		Three-Year Avg Exp. Rate of Exc.
	Max 24-Hr Avg	Exp. Exceed.	Max 24-Hr Avg	Exp. Exceed.	Max 24-Hr Avg	Exp. Exceed	
Cochise County							
Douglas Red Cross ¹	138	0	71	0	251	2.0	0.7
Paul Spur Chemical Lime Plant ¹	85	0	194[#]	1.0	165[#]	1.0	0.7
Number of Sites in Violation of the NAAQS							0
Coconino County							
Flagstaff Middle School ²	61	0	38	0	30	0	0
Number of Sites in Violation of the NAAQS							0
Gila County							
Hayden Old Jail ³	210	2.0	250	1.0	407	1.0	1.4
Miami Golf Course ¹	N/A	N/A	52 [#]	0	129	0	0
Payson Well Site ²	39	0	44	0	58	0	0
Number of Sites in Violation of the NAAQS							1
Maricopa County							
JLG Supersite ³	150	2.0	120	0	262	0	0.7
Number of Sites in Violation of the NAAQS							0
Mohave County							
Bullhead City ¹	132	0	185[#]	1.0	208	1.0	0.7
Number of Sites in Violation of the NAAQS							0
Pima County							
Ajo ³	213	4.1	138	0	299	1.0	1.7
Rillito ³	242	5.0	239	2.0	421	4.1	3.7
Number of Sites in Violation of the NAAQS							2

2011 to 2013 Maximum 24-Hour Average PM₁₀ Compliance (in µg/m³)							
Bold denotes value above the standard.							
(NAAQS 24-hour Average 150 µg/m³)							
Site Name	2011		2012		2013		Three-Year Avg Exp. Rate of Exc.
	Max 24-Hr Avg	Exp. Exceed.	Max 24-Hr Avg	Exp. Exceed.	Max 24-Hr Avg	Exp. Exceed	
Santa Cruz County							
Nogales Post Office ¹	126	0	102	0	89	0	0
Nogales Post Office ³	161	2.0	169[#]	1.0	272[#]	2.0	1.7
Number of Sites in Violation of the NAAQS							1
Yavapai County							
Prescott Valley ² Site closed 12/31/2013	90	0	39	0	51	0	0
Number of Sites in Violation of the NAAQS							0
Yuma County							
Yuma Supersite ²	225	2.0	274	10.0	640	5.0	5.7
Number of Sites in Violation of the NAAQS							1

¹Switched from filter-based sampler to continuous monitor in 2012.

²Samples collected every sixth day - 61 sample days in non-leap years.

³ Samples collected every day - 365 sample days in non-leap years

[#]Indicates the data do not satisfy EPA's summary criteria, usually meaning less than 75 percent valid data recovery available in one or more calendar quarters.

Exp.-Expected

N/A-Data are not available

5.4 Trends

The analysis of trends in PM₁₀ concentrations were divided into three different regions of Arizona: Southern Region, Northern Region, and Central Region. The division into regions helps to group sites that have similar PM₁₀ sources and characteristics.

PM₁₀ Southern Region

Sites evaluated in the southern region of Arizona include Ajo, Douglas Red Cross, Nogales Post Office, Paul Spur Chemical Lime Plant, Rillito, and Yuma. The site identified as “Yuma area” included records from Yuma Courthouse for the years 2008 and 2009 and from Yuma Supersite for the years 2010 to 2013. Figure 16 illustrates the temporal variability of PM₁₀ in the southern region over the 1987 to 2013 period in the form of annual means. The PM₁₀ trend in this region can be described as a significant decrease over the 1987 to 1997 period and a negligible decrease over the 1998-2013 period.

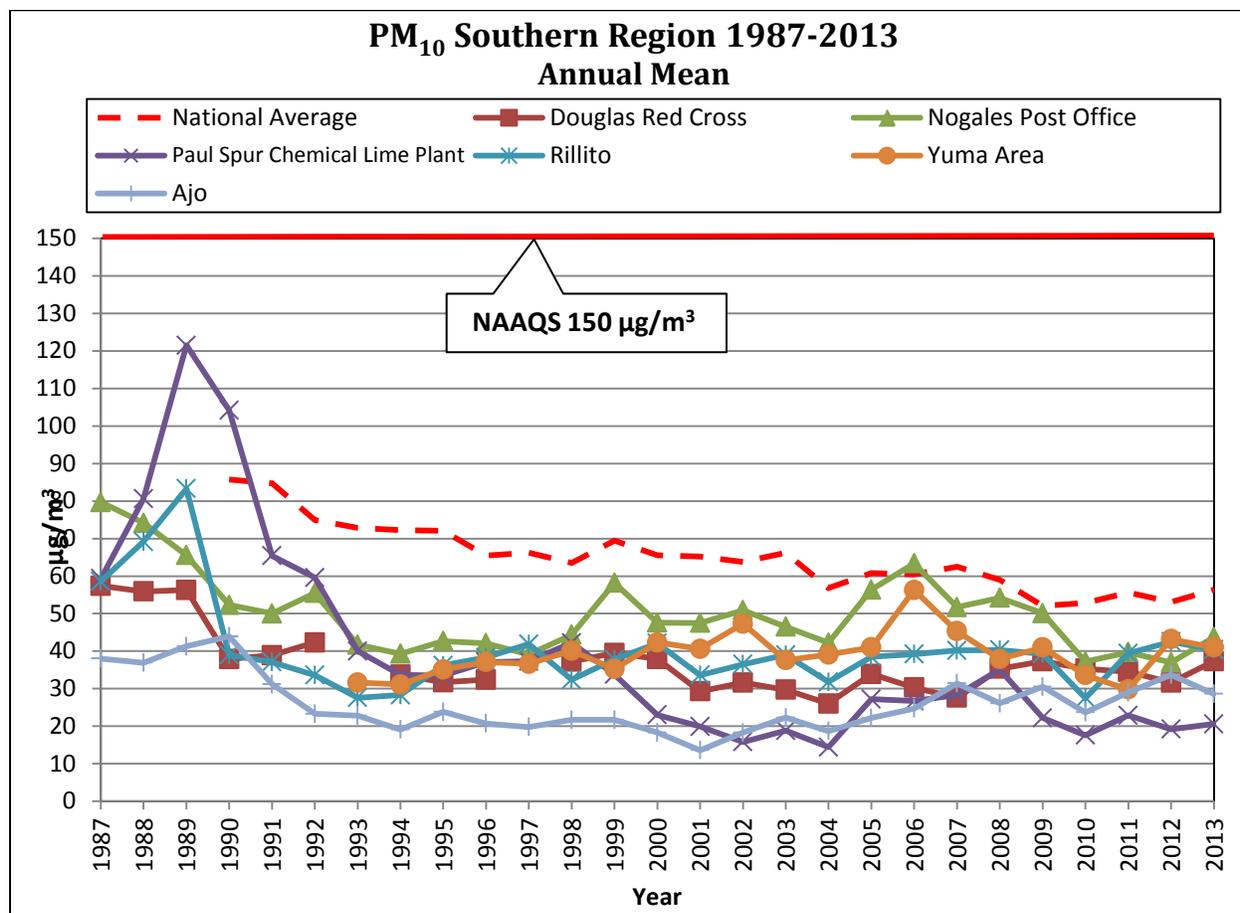


Figure 16 – PM₁₀ Annual Mean Average for the Southern Region

1987-1997: 60.9% decrease for the Southern Region Average

1998-2013: 1.25% negligible decrease for the Southern Region Average

2000-2013: 34% decrease in the National Average

Note: Some years might not satisfy completeness criteria. All PM₁₀ Southern Region sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least 7 consecutive years of data and the same number of consecutive years.

PM₁₀ Northern Region

Sites evaluated in the northern region of Arizona include Bullhead City, Flagstaff Middle School, Payson Well Site, and Prescott Valley. Figure 17 illustrates the temporal variability of PM₁₀ in the northern region over the 1992 to 2013 period in the form of annual means. The PM₁₀ trend in this region can be described as decreasing over the 2003 to 2013 period.

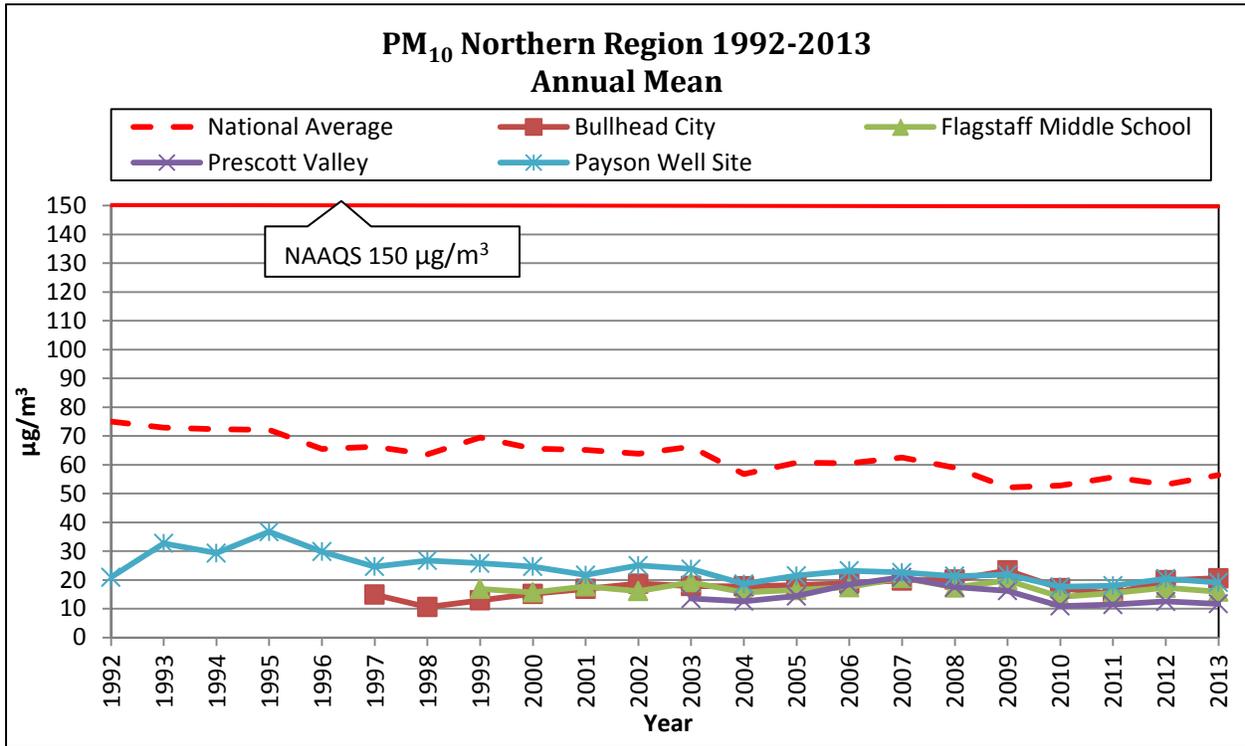


Figure 17 – PM₁₀ Annual Mean Average for the Northern Region
 2003-2013: 11.18% decrease for the Northern Region Average
 2003-2013: 16.6% decrease in the National Average

Note: Some years might not satisfy completeness criteria. All PM₁₀ Northern Region sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least 7 consecutive years of data and the same number of consecutive years.

PM₁₀ Central Region

Sites evaluated in the central region of Arizona include Hayden Old Jail, JLG Supersite, and Miami Golf Course. Figure 18 illustrates the temporal variability of PM₁₀ in the central region over the 1990 to 2013 period in the form of annual means. The PM₁₀ trend in this region can be described as fairly constant, with a slight increase, over the 1999 to 2013 period.

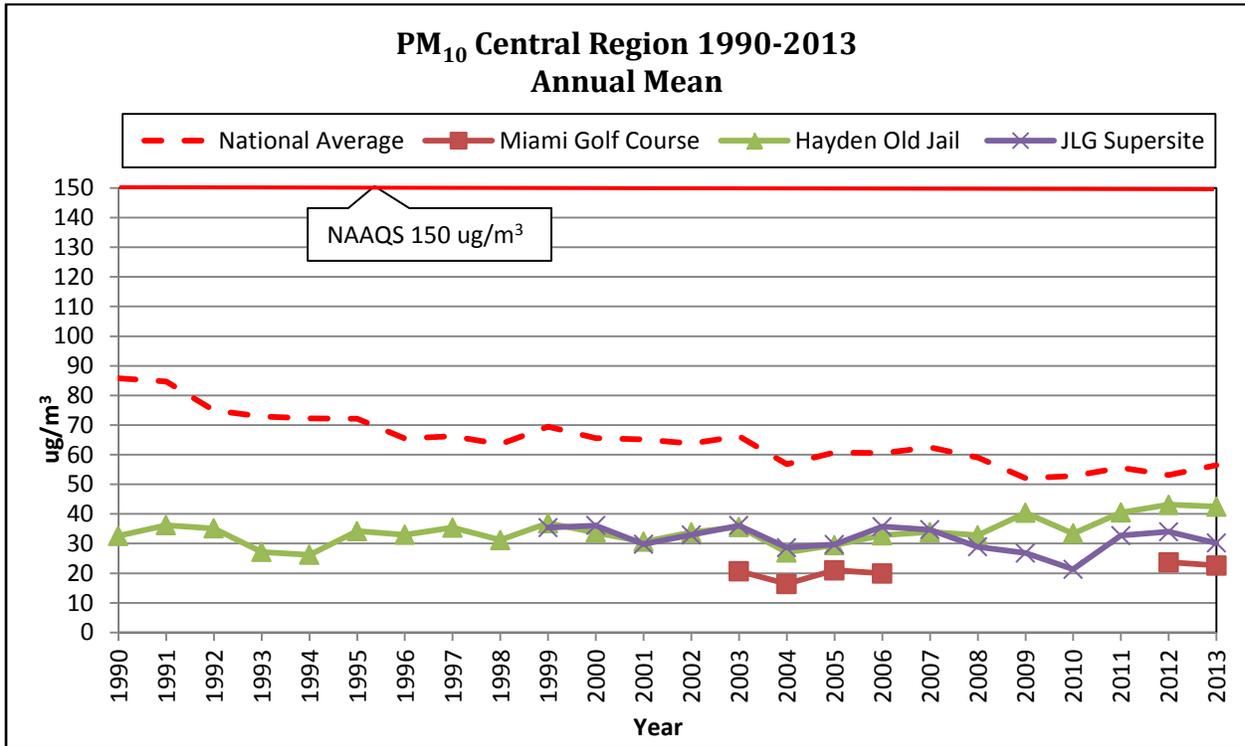


Figure 18 – PM₁₀ Annual Mean Average for the Central Region
 1999-2013: 5.86% increase for the Central Region Average
 2000-2013: 30% decrease in the National Average

Note: Some years might not satisfy completeness criteria. All PM₁₀ Central Region sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least 7 consecutive years of data and the same number of consecutive years.

6.0 Particulate Matter Smaller Than 2.5 Microns (PM_{2.5})

Particulate matter smaller than 2.5 microns (PM_{2.5}) describes particles that are less than 2.5 microns in diameter. These fine particulates are formed by the condensation of vapors or by their subsequent growth through coagulation or agglomeration. Others are emitted directly from sources, either by combustion or from mechanical grinding of soils. Fine particulates are classified as: primary - produced within and emitted from a source with little subsequent change; or secondary - formed in the atmosphere from gaseous emissions. For example, secondary particulate nitrates and sulfates form in the atmosphere from the oxidation of gaseous SO₂ and NO₂. In contrast, most atmospheric carbon is primary, having been emitted directly from combustion sources, although some of the organic carbon in aerosols is secondary, having been formed by the complex photochemistry of gaseous VOCs.

For particulate matter, the primary route of entry into the body is inhalation. Particles smaller than 2.5 microns are respired and enter the pulmonary tissues where they are deposited. Particles in the size range of 0.1 to 2.5 microns are most efficiently deposited in the alveoli, where their effective toxicity is greater than larger particles because of the higher relative content of toxic heavy metals, sulfates, and nitrates. Health studies have shown a significant association between exposure to fine particles and premature death from heart or lung disease. Fine particles can aggravate heart and lung diseases and have been linked to effects such as: cardiovascular symptoms; cardiac arrhythmias; heart attacks; respiratory symptoms; asthma attacks; and bronchitis. Individuals that may be particularly sensitive to fine particle exposure include people with heart or lung disease, older adults, and children.

6.1 Background

With a more diverse chemical composition, fine particulate emissions are distributed among a larger number of sources. According to the 2011 NEI, 72 percent of all PM_{2.5} emissions come from wildfires and wood burning, 10 percent from dust, 6 percent from industrial sources, 5 percent from mobile sources including diesel trucks and off highway vehicles, 3 percent from fuel combustion, 2 percent from agriculture, and 2 percent from miscellaneous industrial sources in Arizona.

Concentrations of PM_{2.5} tend to be at their highest in the central portions of urban areas, diminishing to background levels at the urban fringe. Background concentrations of PM_{2.5} are about 5 µg/m³, in contrast to the urban maximum of 12 to 15 µg/m³. Concentrations of fine particulates tend to be higher in the late fall and winter, when atmospheric dispersion is

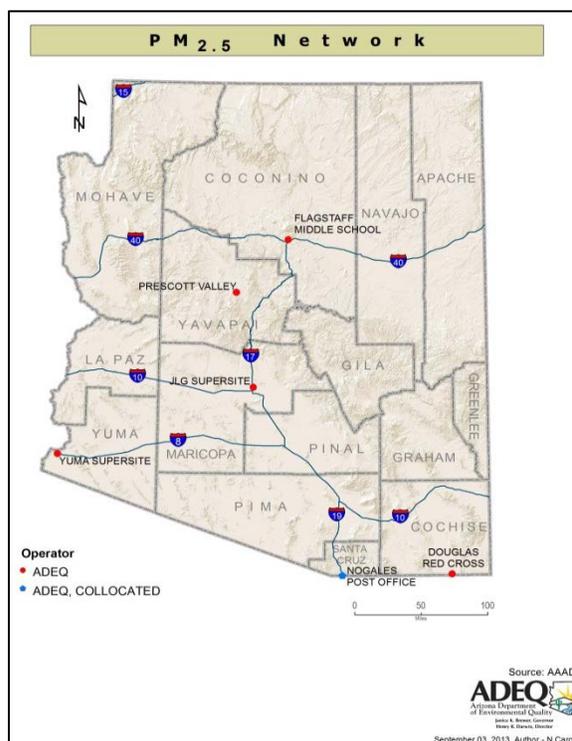


Figure 19 – Map of ADEC's PM_{2.5} sites

at a seasonal low, which traps the particulates and allows for a build-up in the ambient air.

Controls to reduce particulates in Arizona have been in place for decades, beginning in the 1960s with a Pima County ordinance that required watering to reduce dust from construction. Secondary fine particulates have been reduced by vehicle emission controls, which reduce precursor gases. For example, reducing gaseous HC emissions has led to reductions in ambient concentrations of secondary organic carbon.

Due to the successful nature of the controls on $PM_{2.5}$, most of Arizona is in attainment for both of the 2012 primary $PM_{2.5}$ NAAQS. Since fires and blowing dust make up the majority of $PM_{2.5}$ in Arizona, levels can be elevated above the 24-hour standard and are beyond controls on anthropogenic sources.

6.2 Monitoring Methods

Monitors measuring $PM_{2.5}$ operate using the same methods as PM_{10} instruments. The instruments are also the same, using BAMs, and filter-based samplers for monitoring. The difference being that the instruments are fitted with different aerodynamic devices to segregate particle size fractions.



Figure 20 – Nogales Post Office monitoring station.

6.3 Compliance/Summary of Design Values

Effective December 14, 2012 the EPA changed the annual primary $PM_{2.5}$ NAAQS from $15.0 \mu\text{g}/\text{m}^3$ (set in 1997) to $12.0 \mu\text{g}/\text{m}^3$, thereby strengthening this annual health standard. The annual $PM_{2.5}$ standard is met when the three-year average (rounded to the nearest $0.1 \mu\text{g}/\text{m}^3$) of annual means is less than or equal to $12.0 \mu\text{g}/\text{m}^3$. This three-year average is determined by calculating the quarterly averages for each year to determine the calendar year average and then averaging the three years together. A history of the NAAQS for $PM_{2.5}$ is provided in Table 16 below.

Table 16 – History of the National Ambient Air Quality Standards for PM_{2.5} during the period 1997-2012
 (Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1997 62 FR 38652 Jul 18, 1997	Primary and Secondary	PM _{2.5}	24-hour	65 µg/m ³	98th percentile, averaged over 3 years
			Annual	15.0 µg/m ³	Annual arithmetic mean, averaged over 3 years
2006 71 FR 61144 Oct 17, 2006	Primary and Secondary	PM _{2.5}	24-hour	35 µg/m ³	98th percentile, averaged over 3 years
			Annual	15.0 µg/m ³	Annual arithmetic mean, averaged over 3 years
2012	Primary	PM _{2.5}	Annual	12.0 µg/m ³	Annual arithmetic mean, averaged over 3 years
	Secondary		Annual	15.0 µg/m ³	Annual arithmetic mean, averaged over 3 years
	Primary and Secondary		24-hour	35 µg/m ³	98th percentile, averaged over 3 years

The annual primary PM_{2.5} NAAQS of 12.0 µg/m³ was met in 2013 by all six sites operated by ADEQ. See Table 17 for a summary of the annual three-year averages at these sites. In order to calculate the three-year average, there must be a designation of a primary monitor at each site. In some cases, this resulted in the combination of data from filter monitors and data from continuous monitors to complete this three-year average. The primary monitors at Douglas Red Cross, Nogales Post Office, and Yuma Supersite were all filter monitors until continuous monitors were installed in 2013.

Table 17 – PM_{2.5} Annual Average Compliance Summary

2011 to 2013 Annual Average PM_{2.5} Compliance (in µg/m³)				
Federal Reference Monitors and Federal Equivalent Monitors				
Bold denotes a value above the standard. (NAAQS Annual Average 12 µg/m ³)				
Site Name	2011	2012	2013	Three-Year Average
Cochise County				
Douglas Red Cross ¹	6.5	6.7	6.9	6.7
Number of Sites in Violation of the NAAQS				0
Coconino County				
Flagstaff Middle School ²	5.2	5.4	5.4	5.3
Number of Sites in Violation of the NAAQS				0
Maricopa County				
JLG Supersite ³	9.7	8.2	7.6	8.5
Number of Sites in Violation of the NAAQS				0
Santa Cruz County				
Nogales Post Office ¹	9.7	9.6	8.9 [#]	9.4
Number of Sites in Violation of the NAAQS				0
Yavapai County				
Prescott Valley ² (closed 12/31/2013)	3.8	4.4	4.2	4.2
Number of Sites in Violation of the NAAQS				0
Yuma County				
Yuma Supersite ¹	7.6	8.5	7.4	7.8
Number of Sites in Violation of the NAAQS				0

¹Switched from filter-based sampler to continuous monitor in 2013.

²Samples collected every sixth day - 61 sample days in non-leap years.

³Samples collected every day - 365 sample days in non-leap years.

[#]Indicates the data do not satisfy EPA's summary criteria, usually meaning less than 75 percent valid data recovery available in one or more calendar quarters.

As shown in Table 16, the 24-hour primary PM_{2.5} NAAQS of 35 µg/m³ was retained in December 2012. EPA originally issued this 24-hour standard in 2006 when they changed it from 65 µg/m³ to 35 µg/m³. The 24-hour standard is met when the three-year average (rounded to the nearest 1 µg/m³) of the yearly 98th percentile value is less than or equal to 35 µg/m³.

In 2013, all six sites operated by ADEQ were in compliance with the 24-hour primary PM_{2.5} NAAQS of 35 µg/m³. See Table 18 for a summary of the 24-hour three-year averages at these sites. Design calculations for this standard were also based on the designated primary monitor, as they were for the annual standard discussed above.

Table 18 – PM_{2.5} 24-Hour Average Compliance Summary

2011 to 2013 24-Hour Average PM_{2.5} Compliance (in µg/m³)				
Federal Reference Monitors and Federal Equivalent Monitors				
Bold denotes a value above the standard.				
(NAAQS 24-hour Average 35 µg/m³)				
Site Name	98th Percentile Samples			Three-Year Average
	2011	2012	2013	
Cochise County				
Douglas Red Cross ¹	13.0	12.1	12.2	12
Number of Sites in Violation of the NAAQS				0
Coconino County				
Flagstaff Middle School ²	14.0	11.9	10.2	12
Number of Sites in Violation of the NAAQS				0
Maricopa County				
JLG Supersite ³	26.4	19.0	21.6	22
Number of Sites in Violation of the NAAQS				0
Santa Cruz County				
Nogales Post Office ¹	27.2	25.9	27.2	27
Number of Sites in Violation of the NAAQS				0
Yavapai County				
Prescott Valley ² (closed 12/31/2013)	10.2	10.8	8.4	10
Number of Sites in Violation of the NAAQS				0
Yuma County				
Yuma Supersite ¹	15.6	15.8	17.0	16
Number of Sites in Violation of the NAAQS				0

¹Switched from filter-based sampler to continuous monitor in 2013.

²Samples collected every sixth day - 61 sample days in non-leap years.

³ Samples collected every day - 365 sample days in non-leap years

[#]Indicates the data do not satisfy EPA's summary criteria, usually meaning less than 75 percent valid data recovery available in one or more calendar quarters.

6.4 Trends

For this trends assessment, data from Douglas Red Cross, Flagstaff Middle School, JLG Supersite, and Nogales Post Office were evaluated for a period of fifteen years from 1999 to 2013. A shorter length of records for Prescott Valley and Yuma area sites were evaluated for a period of six years from 2008 to 2013. The site identified as "Yuma area" included records from Yuma Courthouse for the years 2008 and 2009 and from Yuma Supersite for the years 2010 to 2013.

Figures 21 and 22 illustrate the temporal variability of PM_{2.5} over the 1999 to 2013 period in the form of annual means and the 24-hour averages at the 98th percentile. The PM_{2.5} trend can be described as decreasing over the 1999 to 2013 period. While the overall trend for the six sites is decreasing, it is important to note that JLG Supersite and Nogales Post Office do not have the same magnitude of PM_{2.5} concentration reductions. This pollutant will continue to be of concern at these sites and will be further analyzed with additional years of data.

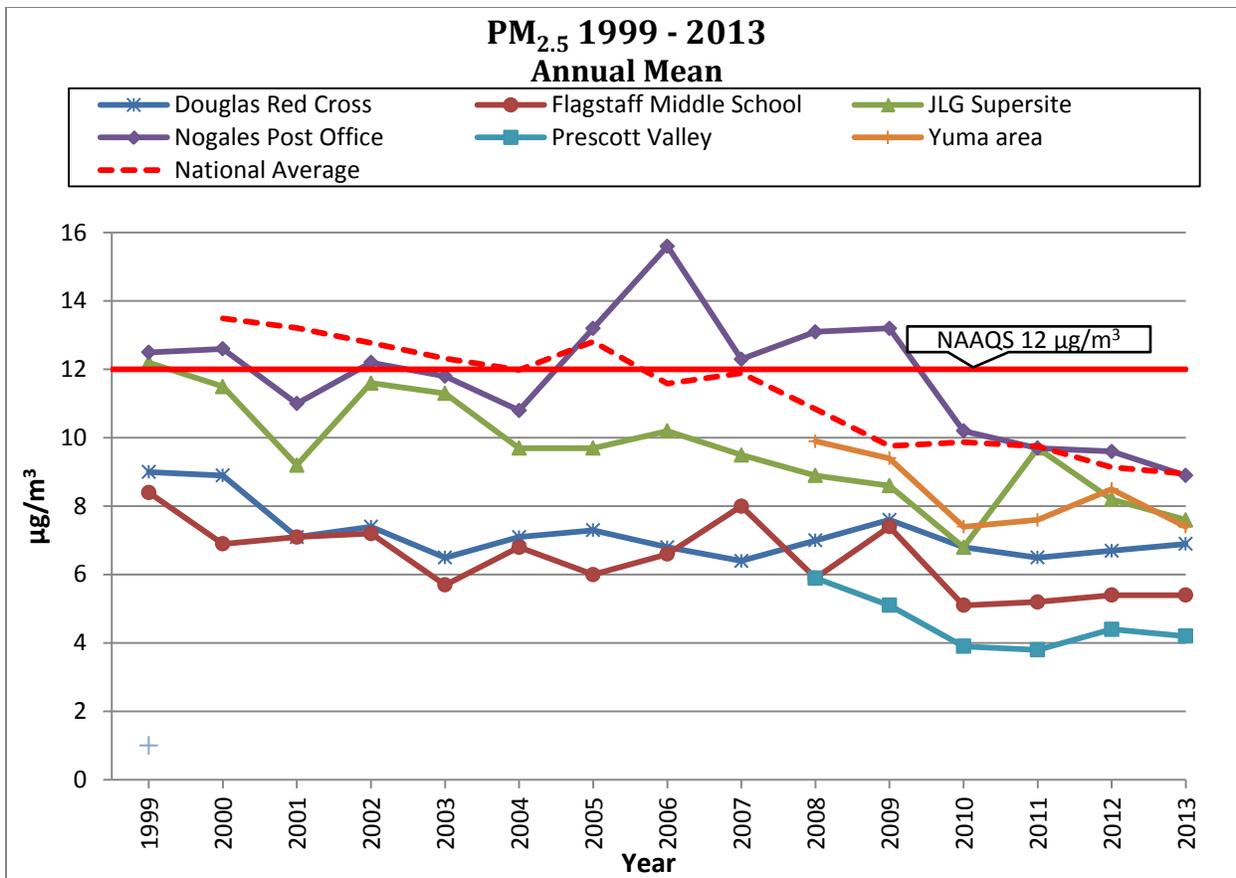


Figure 21 – PM_{2.5} Annual Mean Trend

1999 - 2013: 26.48% decrease for Douglas Red Cross, Flagstaff Middle School, JLG Supersite, and Nogales Post Office

2000 - 2013: 34% decrease in the National Average

Note: Some years might not satisfy completeness criteria. All PM_{2.5} sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least 7 consecutive years of data and the same number of consecutive years.

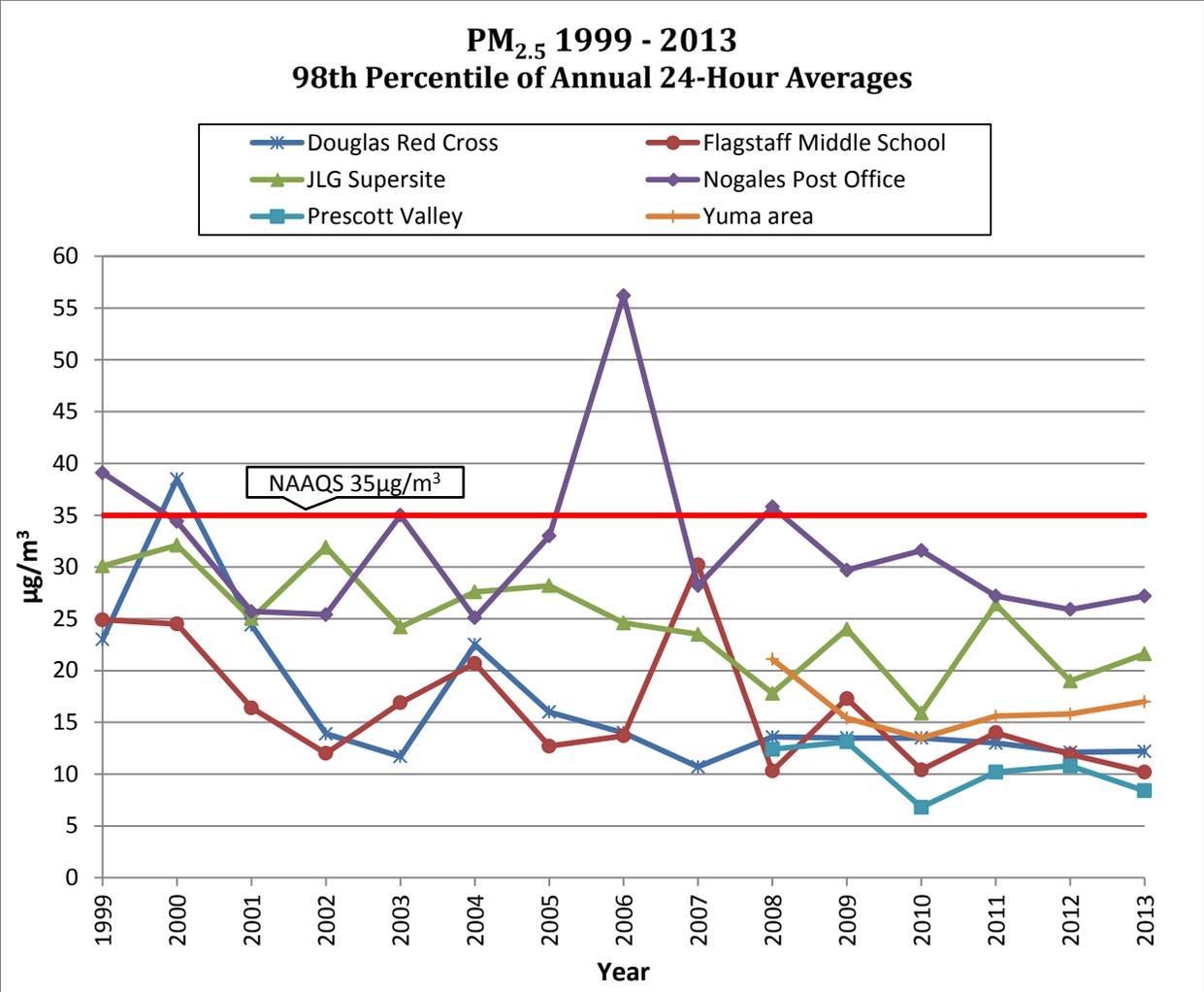


Figure 22 – PM_{2.5} 24-Hour Average Trend
 1999 - 2013: 39.12% decrease for Douglas Red Cross, Flagstaff Middle School, JLG Supersite, and Nogales Post Office
Note: Some years might not satisfy completeness criteria. All PM_{2.5} sites were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all sites must have at least 7 consecutive years of data and the same number of consecutive years.

7.0 Lead (Pb)

Lead is a metal found naturally in the earth and a key element in the composition of many rocks and soils. It is a very dense metal that is used in a variety of industrial processes in many forms. It can be used as anodes in car batteries or as ballast in boats or in scuba diving. Today, the major sources of Pb emissions are ore and metals processing and piston-engine aircraft operating on leaded aviation gasoline. The highest ambient air concentrations of Pb in Arizona are usually found near Pb smelters.

In addition to exposure to Pb in air, other major exposure pathways include ingestion of Pb in drinking water and Pb-contaminated food as well as incidental ingestion of Pb-contaminated soil and dust. Lead-based paint remains a major exposure pathway in older homes. Once taken into the body in any form, Pb distributes throughout the body in the blood and is accumulated in the bones. Depending on the level of exposure, Pb can adversely affect the nervous system, kidney function, immune system, reproductive and developmental systems and the cardiovascular system. Lead exposure also affects the oxygen carrying capacity of the blood. The Pb effects most commonly encountered in current populations are neurological effects in children and cardiovascular effects in adults. Infants and young children are especially sensitive to even low levels of Pb, which may contribute to behavioral problems, learning deficits, and lowered IQ.

7.1 Background

Lead emissions in Arizona are made up of two main sources: mobile sources including piston-engine aircraft and other older engines which use leaded gasoline, and copper and metals smelting. According to the 2011 NEI, mobile sources contribute about 64 percent of Pb emissions in Arizona, smelting and other industrial processes contribute about 33 percent, and all other sources including fuel combustion contribute about three percent.

Lead has historically been used in paints, as an additive to fuel, in electronics, and in various other industrial applications. U.S. Regulations have eliminated the use of Pb in almost all of these applications due to its toxicity.

The large percentage of mobile sources is spread out over many different small sources in the state. There are around 100 small airports that still have some airplanes that require leaded-fuel, but since this is spread out over the whole state, no single airport

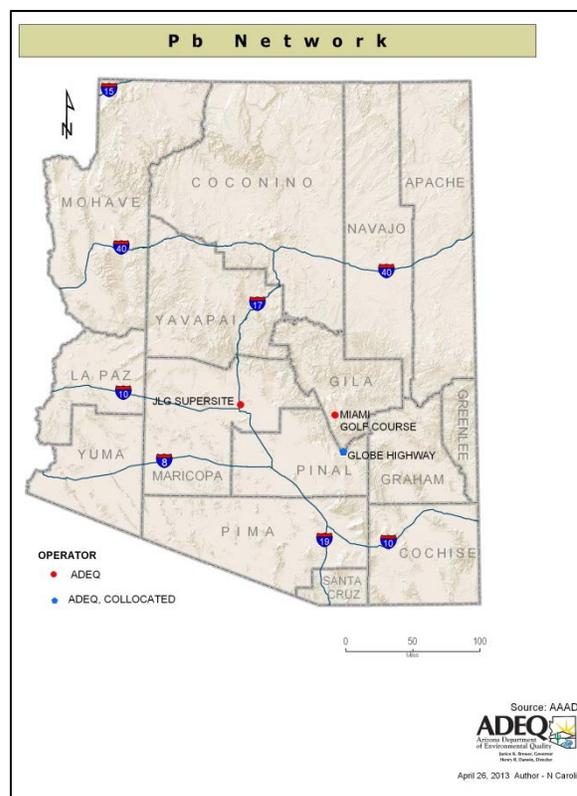


Figure 23 -- Map of ADEQ's Pb sites

contributes to a high concentration of Pb at a time. The only other large sources of Pb are due to metals smelting. Since Arizona has large copper smelting operations, Pb continues to be a concern in the areas surrounding these sources.

7.2 Monitoring Methods

Lead is monitored using two different sampling techniques. The first uses the same manual filter instruments that are used for PM₁₀ sampling. The same PM₁₀ inlet is used, and this method for using the PM₁₀ sampler is approved by the EPA as Pb-PM₁₀. The difference between sampling for Pb and sampling for PM₁₀ is that the filters are not pre-weighed or post weighed; the filters are analyzed for Pb content using an EPA approved method involving the digestion or dissolving of other contaminants and the isolation of Pb on the filter. Once the Pb is isolated, it can be weighed using a precise scale. Based on the total flow of the sample and the weight, a concentration is calculated.

The other method of sampling uses a total suspended particle (TSP) sampler that does not have an inlet attached. All particles in the air are captured by the sampler on a large filter using a high flow rate. Once the 24-hour sampling period is over, the filter is cut, digested, and Pb is isolated in the same way as with the Pb-PM₁₀ sample filter.

Readings from all the types of instruments are averaged into daily, monthly, quarterly, and yearly averages for data analysis. Readings are retrieved on the samplers manually and by weighing equipment then stored in a database.

7.3 Compliance/Summary of Design Values

In 2008, the Pb standard was revised from the 1978 rule. This lowered the three-month rolling average from 1.5 µg/m³ to 0.15 µg/m³. The rule also set forth a monitoring requirement based on an emissions threshold for each source. This threshold states that sources require monitoring if they are above 0.5 tons per year based on the most recent emissions inventory. Lead concentrations in Arizona generally do not exceed this standard since leaded fuels were banned in the U.S. starting in 1996. Due to the 2008 rule change, ADEQ operated three source oriented monitors in 2013 around two copper smelters in Miami and Hayden. ADEQ also operated one Pb monitor at JLG Supersite as part of NCore. A history of the NAAQS for Pb is provided in Table 19 below:

Table 19 – History of the National Ambient Air Quality Standards for Pb during the period 1978-2008
(Source: USEPA TTN NAAQS)

Final Rule	Primary/ Secondary	Indicator	Averaging Time	Level	Form
1978 43 FR 46246 Oct 5, 1978	Primary and Secondary	Pb-TSP	Calendar Quarter	1.5 µg/m ³	Not to be exceeded
<i>Feb 21, 1991 – Agency released multimedia “Strategy for Reducing Lead Exposures”</i>					
2008 73 FR 66964 Nov 12, 2008	Primary and Secondary	Pb-TSP	3-month period	0.15 µg/m ³	Not to be exceeded

Table 20 summarizes the maximum three-month rolling average Pb concentrations for the year from 2011 to 2013. The PM₁₀ sampler at JLG Supersite was approved by the EPA as a Pb-TSP equivalent method in July of 2012, so the maximum three-month rolling average data are unavailable for 2011 and 2012.

Table 20 – Pb Compliance Summary

2011 to 2013 Maximum 3 - Month Average Pb-TSP Compliance (in µg/m³)				
Bold denotes value above the standard. (NAAQS 3-Month Rolling Average 0.15 µg/m ³)				
Site Name	Max 3-Month Rolling 24-Hr Avg			Three-Year Design Value
	2011	2012	2013	
Gila County				
Globe Highway	0.27	0.20	0.12	0.27
Miami Golf Course	0.06	0.04	0.03	0.06
Number of Sites in Violation of the NAAQS				1
Maricopa County				
JLG Supersite	N/A	N/A	0.0058	N/A
Number of Sites in Violation of the NAAQS				N/A

N/A-Data are not available

7.4 Trends

ADEQ began monitoring for Pb in 2011. During the 2011-2013 time period there was a decrease in the Maximum Rolling Three-Month Average at Miami and Hayden (Figure 24). This decrease is far above the national decrease of 15% during the same time period. The relatively sharp decrease can be attributed to tighter emission controls enacted at the source level.

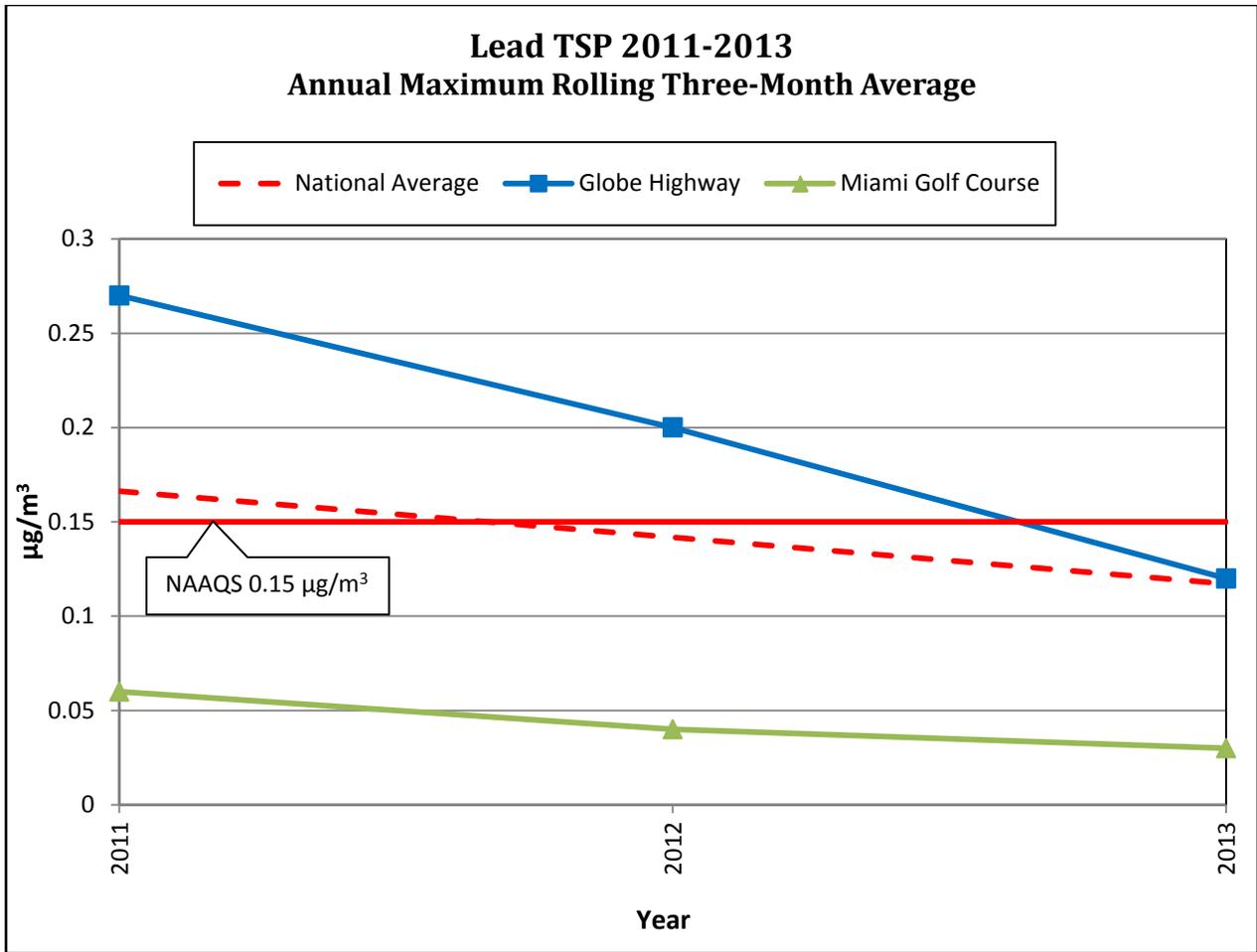


Figure 24 – Pb Three-Month Average Trend

2011 to 2013: 64% decrease in the Maximum Rolling Three-Month Average at Miami and Hayden
 15% decrease in the National Maximum Rolling Three-Month Average

Note: Some years might not satisfy completeness criteria.

Section III – Non-Criteria Pollutants

This section covers the non-criteria pollutant networks monitored by ADEQ. Non-criteria pollutants are all pollutants not classified under the CAA as criteria pollutants. These include, but are not limited to, hazardous air pollutants (HAPs) sometimes referred to as toxics, heavy metal pollutants, and many other contaminants and carcinogens. The following sections will discuss the CSN, the NATTS, and the PAMS programs. Each subsection will provide a background for each program and a synopsis of current monitoring techniques. There will be a discussion of 2013 data and trend analysis on certain pollutants for the length of record for each program.

1.0 Chemical Speciation Network

PM_{2.5}, as described previously, are particles with a diameter less than 2.5 microns and are associated with respiratory and other health-related problems. These negative effects have been correlated to the total mass concentration in ambient air, but whether these negative effects are more correlated to total mass concentration or to the concentration of specific chemical species is not fully understood. To better understand and manage PM_{2.5}, EPA commissioned a companion network in 1997 called the CSN, in addition to the NAAQS PM_{2.5} network in 1997, to ascertain the chemical composition of fine particulates. Chemical speciation includes the identification and quantification of individual chemical elements, compounds, or classes of compounds that make up PM_{2.5} aerosols. The PM_{2.5} CSN targets analytes or species that are of particular interest to health officials, epidemiological researchers and other interested parties.

1.1 Background

ADEQ supports the CSN network by serving as the operator for a CSN site in Arizona at JLG Supersite.

The program objectives for the CSN are:

- to characterize annual and seasonal spatial characterization of aerosols
- air quality trends analysis
- tracking progress of control programs
- to compare the chemical speciation data set to the data collected from the IMPROVE network
- development of emission control strategies

The data generated from this network are not used for regulatory decisions concerning the PM_{2.5} NAAQS, but may be used to supplement the PM_{2.5} program.

1.2 Monitoring Methods

CSN instruments operate using the same principles as PM filter-based samplers. A filter is sampled on a specific date for a period of 24-hours, after which the sample is collected and sent to an EPA contracted

lab for speciation analysis. EPA sets the schedule so that every CSN site in the U.S. is sampling on the same day. Currently, two different types of speciation samplers produce three 24-hour PM_{2.5} filter samples every three days at JLG Supersite. Samples are collected on three different filter types. Polytetrafluoroethylene (PTFE, commonly known as Teflon) filters are used to determine trace elements/crustal constituents that include metals, metalloids, and non-metals. Samples collected on nylon filters are used to determine the cations (sodium, potassium, and ammonium, and the anions: nitrate and sulfate). Samples collected on quartz filters are used to determine elemental and organic carbons.

1.3 Graphs

PM_{2.5} CSN characterization summaries for JLG Supersite are shown in Figures 25 and 26. The JLG Supersite CSN site far exceeded the 75% data completeness requirement in 2013. Major elements show seasonal shifts, but organic carbon emerges as the dominant element throughout the year. Sulfates are more abundant during the spring and summer months, while nitrates are greater during the cooler months. Unidentified constituents are classified as "Other" and tend to increase in cooler months as well. This category exists because the analytical tests do not yield results for all possible species of PM_{2.5} aerosol. Analyzing for all species in PM_{2.5} would be impractical; however, the analytes selected are carefully chosen and often serve as indicators of other species. As more information concerning the chemical composition of fine particulates becomes available to researchers and regulators, the selected analytes may change as they did in early 2009.

Speciated PM_{2.5} at JLG Supersite 2013
Annual Average Concentration (µg/m³) Percentage

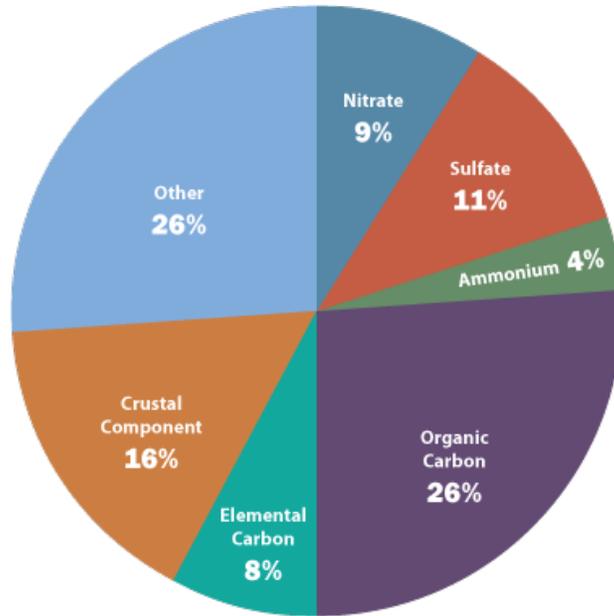


Figure 25 – 2013 Annual Averages for Speciated PM_{2.5} major elements expressed as percentages of the total PM_{2.5} concentration (µg/m³) at JLG Supersite

Speciated PM_{2.5} Quarterly Averages at JLG Supersite 2013
Annual Average Concentration (µg/m³) Percentage

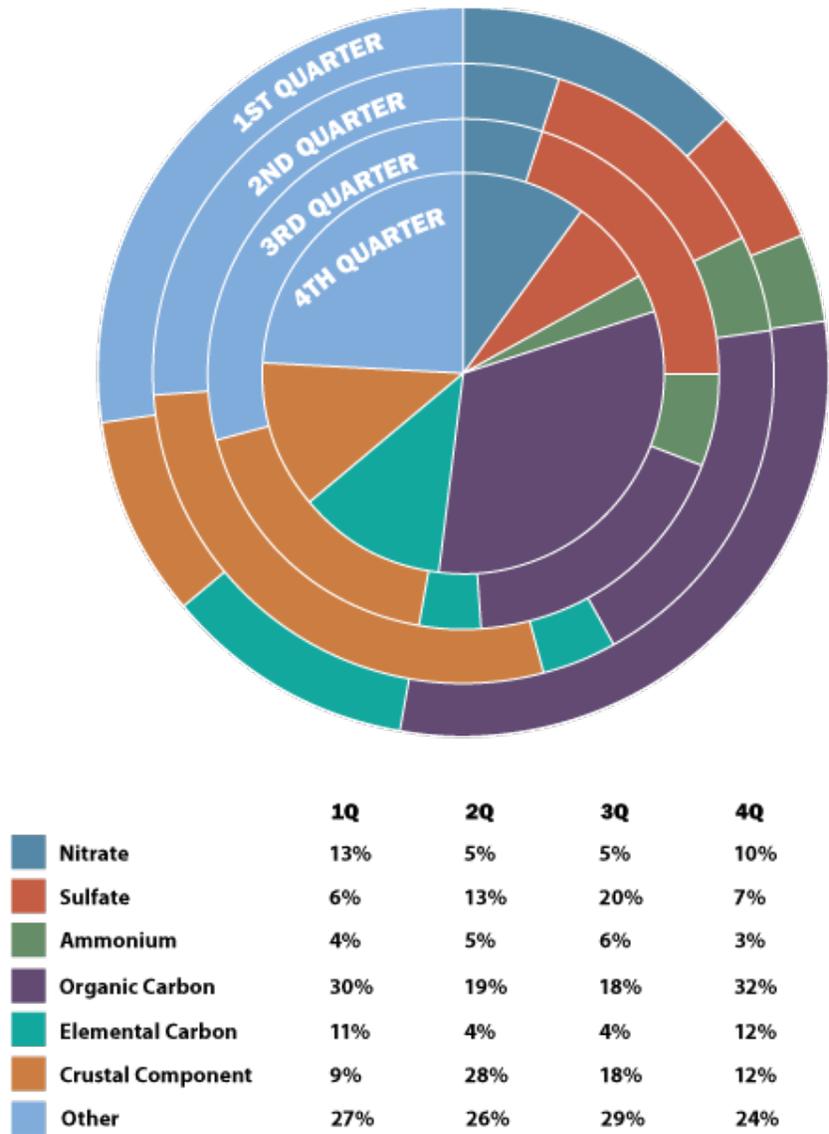


Figure 26 – 2013 Quarterly Averages for Speciated PM_{2.5} major elements as percentages of the total PM_{2.5} concentration (µg/m³) at JLG Supersite in 2013

1.4 Trends

In order to improve data comparability, in 2009 the EPA switched the sampling and analytical methods for organic carbon and elemental carbon nationwide. The organic carbon, elemental carbon, and “Other” concentrations reported from 2010 to 2013 are slightly more accurate and more comparable to national data than those reported prior to 2010. The percentages of each species compared to the total PM_{2.5} concentration, had a negligible change from 2000 to 2013 at JLG Supersite (Figure 27). This indicates that the annual average composition of PM_{2.5} at JLG Supersite has shown no significant changes during the time period.

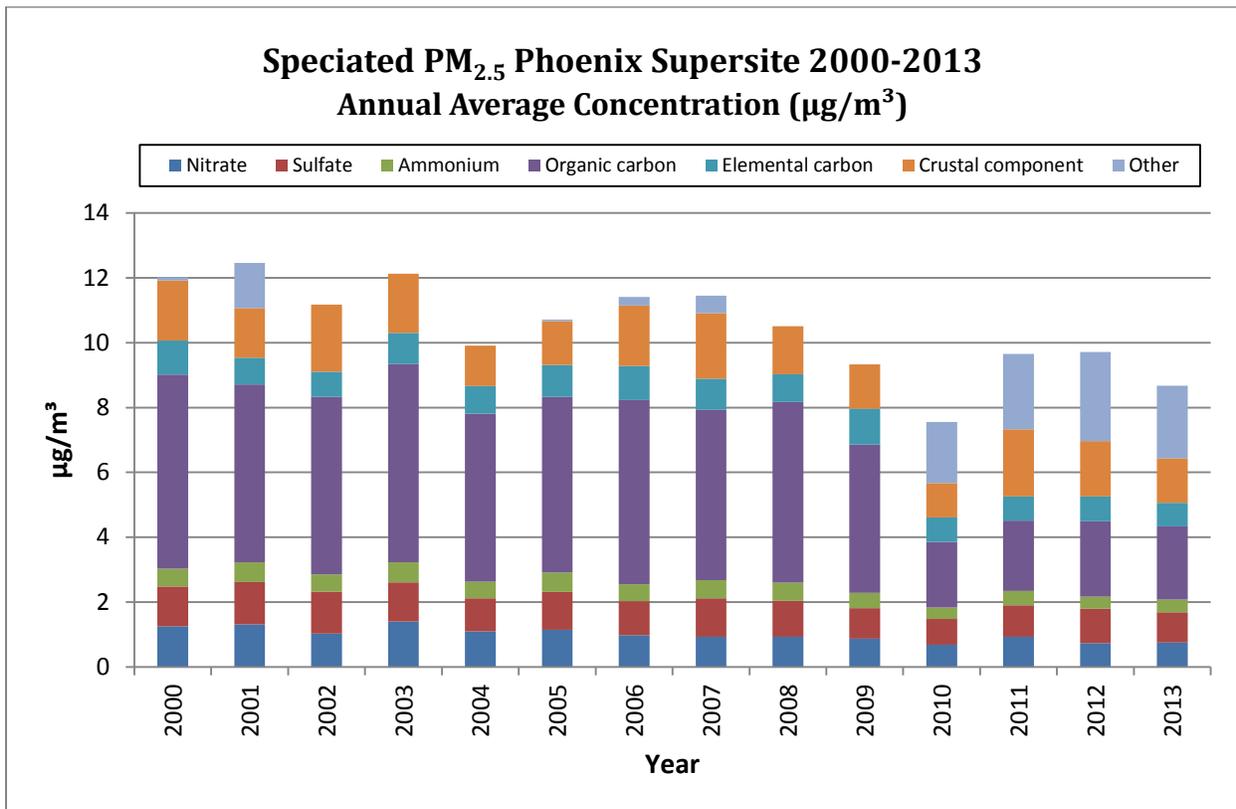


Figure 27 – Speciated PM_{2.5} Annual Average Trend
1999-2013: Less than 1% variation in average Annual PM_{2.5} composition at JLG Supersite

2.0 National Air Toxics Trends Station (NATTS)

Air toxics, also known as HAPs, are compounds or elements known or suspected to cause cancer or other serious health effects, such as reproductive, neurological, birth defects, or adverse environmental effects. There are currently 187 HAPs regulated under the CAA. Examples of HAPs include benzene, which is found in gasoline; perchlorethylene, which is emitted from some dry cleaning facilities; and methylene chloride, which is used as a solvent and paint stripper by a number of industries. Examples of other known air toxics include chloroform, formaldehyde, and metals such as cadmium, chromium, and Pb compounds.

2.1 Background

ADEQ monitors many HAPs including VOC, carbonyls, and PM₁₀ metals at JLG Supersite as part of the NATTS program. NATTS is a national network of 27 monitors that was developed to fulfill the need for long-term, high-quality toxics data to estimate national toxics averages. The principle purpose of the NATTS program is tracking toxics trends to assess progress towards emission and risk reduction. The goal is to reduce air toxics nationally by 15 percent over rolling three-year periods. ADEQ entered the NATTS program in 2003 but had been monitoring some of the toxics in prior years. Funding for the monitoring of hexavalent chromium was removed from the NATTS grants budget due its removal from the NATTS list of required pollutants. This decreased the minimum pollutant monitoring from 19 to 18 pollutants in July, 2013. With the minimum pollutants monitored being met, ADEQ monitors a total of 104 pollutants.

The UATMP monitoring program began in 1987 to characterize the magnitude and composition of urban air pollution through extensive monitoring in various cities across the U.S. Urban air pollution consists of many components from a wide range of industrial, motor vehicle, and biogenic and natural sources, some of which are toxics. ADEQ monitored VOCs at the South Phoenix site as part of the EPA's UATMP from 2001-2003 and again from 2007-2013. As stated earlier in this report, UATMP data are directly comparable to NATTS data as many of the program objectives overlap.

2.2 Monitoring Methods

To monitor VOCs, ambient air is captured in special evacuated canisters at a constant flow rate over a 24-hour period every sixth day in accordance with the EPA monitoring schedule. The canisters are collected and sent to a laboratory for analysis by a standard procedure (TO-15) appropriate for VOCs, and the results are reported to ADEQ for review before submittal to EPA.

For carbonyls, ambient air is drawn through small cartridges at a specific flow rate for 24-hours every sixth day in accordance with the EPA monitoring schedule. The cartridges are sent to a laboratory for analysis using a standard procedure (TO-11A) appropriate for carbonyls analysis. The results are reviewed by ADEQ before submittal to EPA.

PM₁₀ metals are collected by drawing ambient air through a filter at a specific flow rate for 24-hours, as done for PM measurement discussed in Section II of this report, every sixth day in accordance with the EPA monitoring schedule. The filters are sent to a laboratory and processed using a standard procedure (TO-3.5) that is appropriate for metals analysis. The results are reviewed by ADEQ before submittal to EPA.

2.3 Tables

EPA has not issued monitoring compliance criteria for air toxics (as it has for O₃, for example). Instead it addresses control issues through rules covering emissions from industrial sources such as chemical plants as well as smaller sources, such as dry cleaners and chromium electroplating facilities. As can be seen in Tables 21 and 22, air toxics concentrations at JLG Supersite and South Phoenix are comparable to national averages with some values being higher, and some lower, but none with extreme differences. For the pollutants that are in common between the two sites (i.e., VOCs), the summary data are very similar, which is an indication that the sites do a reasonable job of representing ambient air and are not significantly affected by nearby sources. These data also show that in comparison to the 2011 National Averages, manganese may be of local concern as the 2013 annual average exceeds the 2011 national average by more than a factor of two. Air Toxics is a relatively new field within ambient air monitoring, and these data will continue to be collected to provide more information about urban air pollution.

Table 21 – Air Toxics Data for JLG Supersite

2013 Air Toxics data for JLG Supersite						
Pollutant	Quarterly Average					National (2011)
	Q1	Q2	Q3	Q4	Annual	
VOC (ppb)						
1,3-Butadiene	0.13	0.04	0.03	0.18	0.10	0.04
Acrolein	0.61	0.41	0.49	0.51	0.50	0.49
Benzene	0.44	0.17	0.17	0.52	0.33	0.31
Carbon Tetrachloride	0.10	0.10	0.10	0.09	0.10	0.10
Chloroform	0.05	0.07	0.07	0.09	0.07	0.08
Tetrachloroethylene	0.05	0.02	0.02	0.06	0.04	0.02
Trichloroethylene	0.00	0.00	0.00	0.00	0.00	0.01
Aldehydes (ppb)						
Acetaldehyde	1.49	1.39	1.24	1.98	1.54	1.11
Formaldehyde	2.88	3.13	3.17	3.43	3.16	2.33
PAH(ng/m³)						
Benzo[A]Pyrene (Tsp) Stp	0.14	0.02	0.01	0.24	0.11	0.08
Naphthalene (Tsp) Stp	116	52	42	157	93	81.7
PM₁₀ Metals (ng/m³)						
Arsenic	0.57	0.31	0.51	0.55	0.49	0.59
Beryllium	0.02	0.04	0.04	0.03	0.03	0.01
Cadmium	0.17	0.06	0.09	0.15	0.12	0.16
Chromium	3.81	3.77	1.01	1.76	2.58	2.68
Lead	3.67	2.37	2.45	4.71	3.32	3.82
Manganese	15.25	17.71	21.25	17.78	17.99	8.81
Nickel	1.43	1.39	1.44	1.70	1.50	1.27

Table 22 – Air Toxics Data for South Phoenix

Air Toxics data for South Phoenix site						
Pollutant	Quarterly Average VOC (ppb)					National (2011)
	Q1	Q2	Q3	Q4	Annual	
1,3-Butadiene	0.11	0.05	0.05	0.18	0.10	0.04
Acrolein	0.45	0.30	0.36	0.44	0.39	0.49
Benzene	0.38	0.21	0.21	0.51	0.33	0.31
Carbon Tetrachloride	0.09	0.10	0.10	0.09	0.10	0.10
Chloroform	0.02	0.05	0.04	0.05	0.04	0.08
Tetrachloroethylene	0.02	0.01	0.02	0.04	0.02	0.02
Trichloroethylene	0.00	0.00	0.00	0.00	0.00	0.01

2.4 Trends

The VOC trends include data from the JLG Supersite and South Phoenix sites. Seven chemical species were used to calculate the VOC trends data shown in Figures 28 and 29. These species were chosen due to their high frequency of detections in Arizona and nationwide. South Phoenix VOC concentrations are unavailable from 2005-2006 because ADEQ did not monitor for the UATMP during that time period. ADEQ began monitoring for Acrolein at JLG Supersite and South Phoenix in 2007, so Acrolein was not used in the 2001 to 2013 average concentration calculations for JLG Supersite. At JLG Supersite and South Phoenix, there was an overall decrease in the average annual concentrations of the seven species selected for trends data from 2001 to 2013.

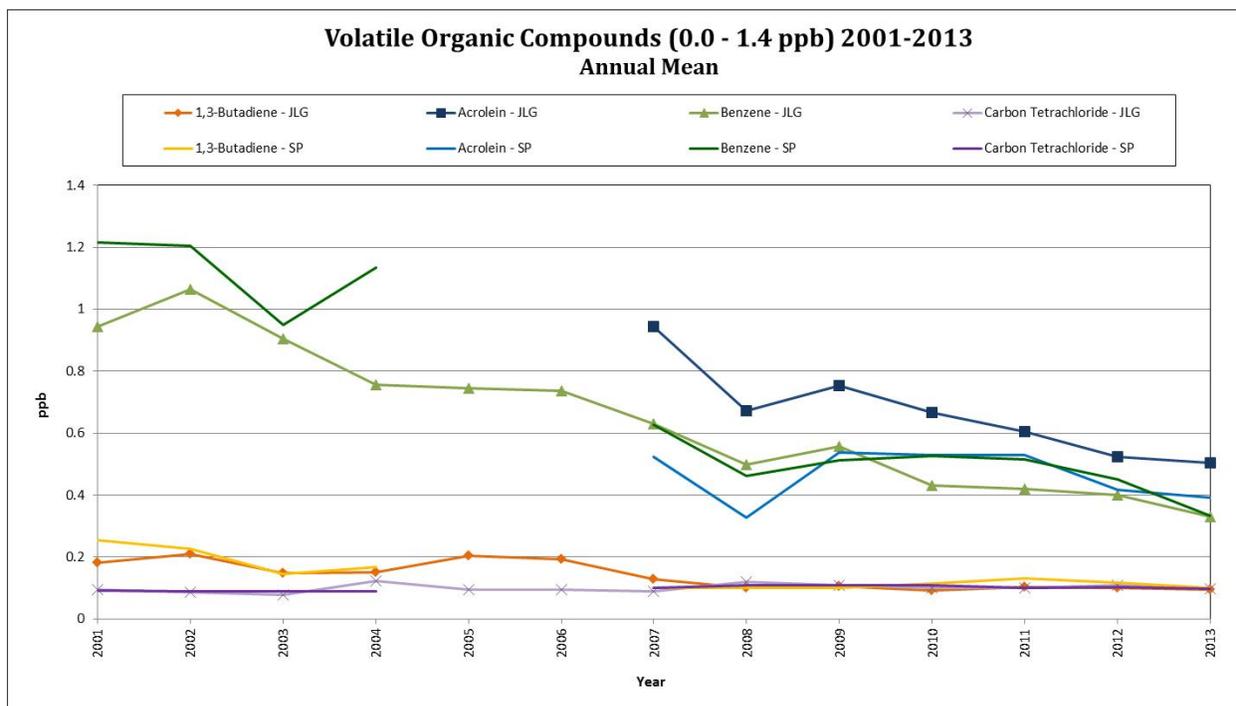


Figure 28 – VOC Annual Mean Trend (0.0-1.4 ppb)

2001-2013: 63.0% decrease at JLG Supersite

2007-2013: 48.0% decrease in Acrolein at JLG Supersite

2007-2013: 23.5% decrease at South Phoenix

Note: Some years might not satisfy completeness criteria. Seven species were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all species must have at least seven consecutive years of data and the same number of consecutive years.

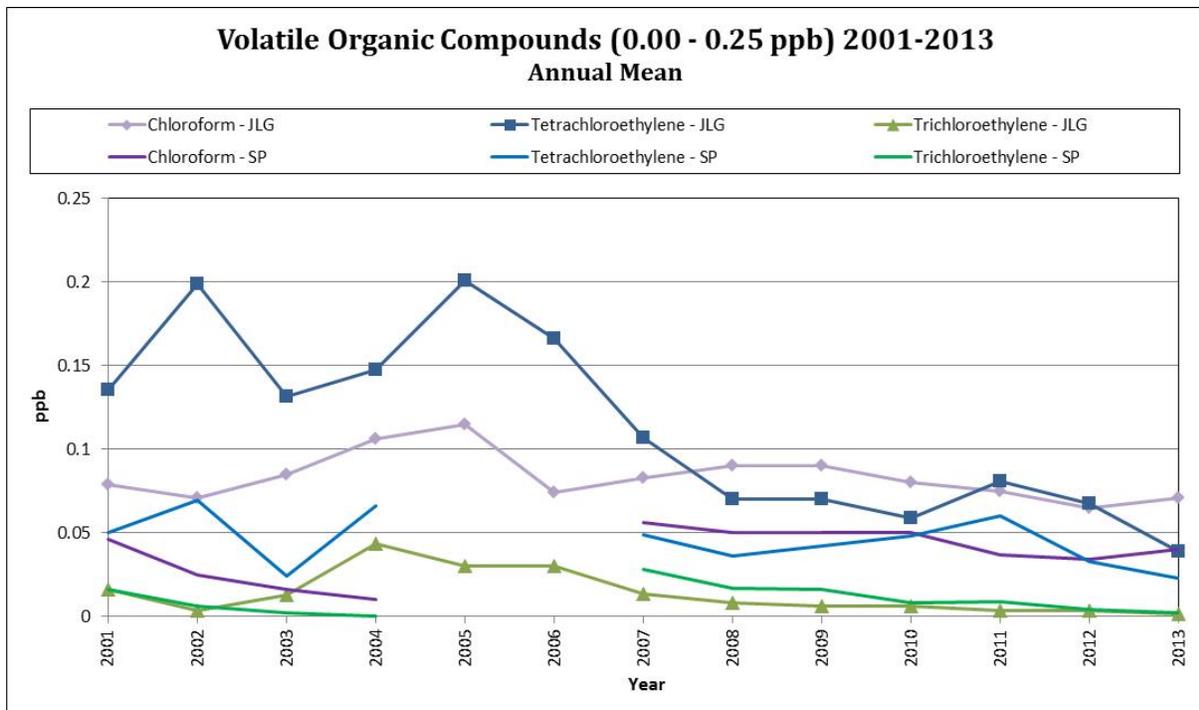


Figure 29 – VOC Annual Mean Trend (0.00-0.25 ppb)

2001-2013: 63.0% decrease at JLG Supersite

2007-2013: 23.5% decrease at South Phoenix

Note: Some years might not satisfy completeness criteria. Seven species were averaged to calculate a best-fit straight line, which was used to determine the average percent change. In order to reduce bias in averaging, all species must have at least seven consecutive years of data and the same number of consecutive years.

The aldehydes and polycyclic aromatic hydrocarbon (PAH) trends data were taken from JLG Supersite. There was a decrease in the average annual concentrations of both aldehydes (acetaldehyde and formaldehyde) from 2003 to 2013 (Figure 30) and PAH benzo(A)pyrene from 2007 to 2013 (Figure 31). There was an increase in the average annual concentration of the PAH naphthalene from 2007 to 2013 (Figure 32).

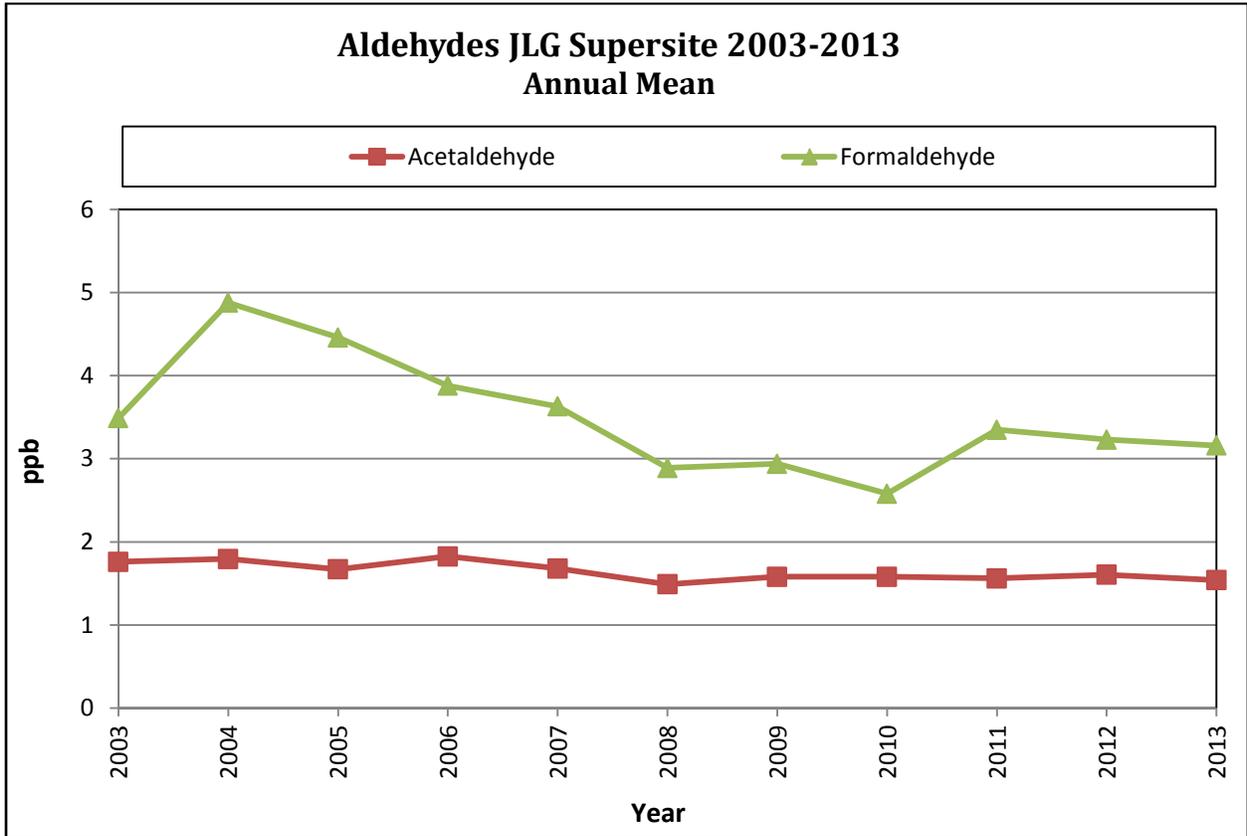


Figure 30 – Aldehydes Annual Mean Trend
 2003-2013: 28.9% decrease at JLG Supersite
Note: Some years might not satisfy completeness criteria.

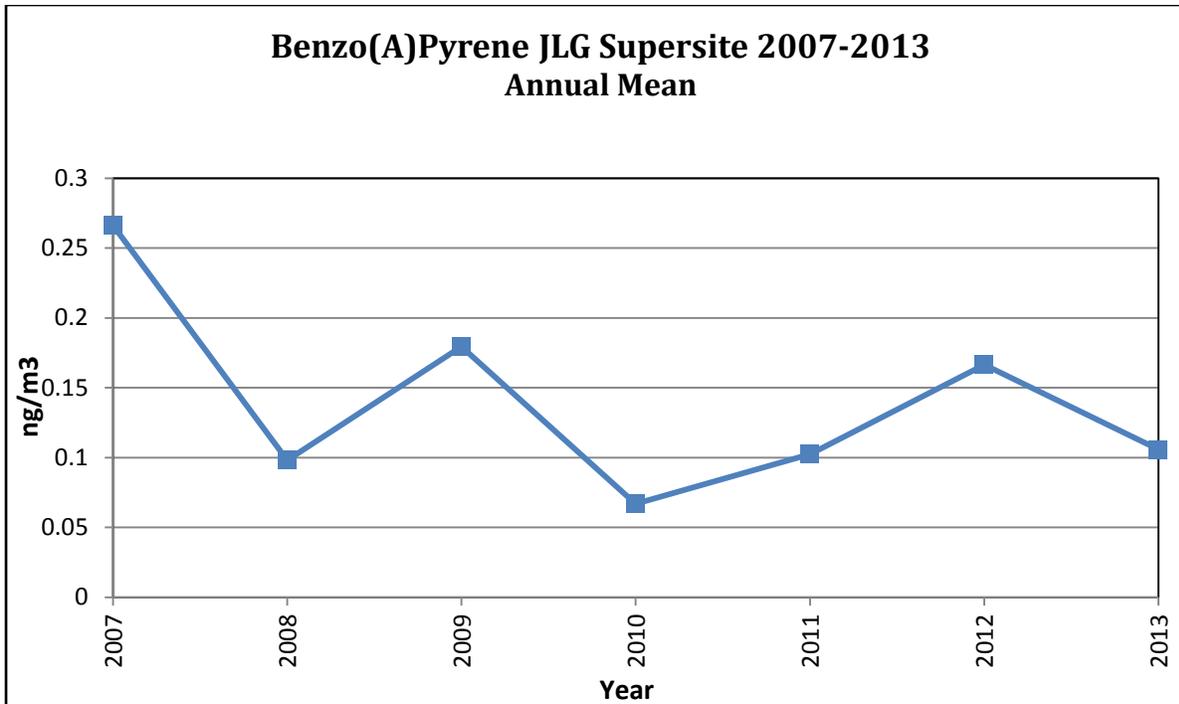


Figure 31 – Benzo (A) Pyrene Annual Mean Trend
 2007-2013: 52.6% decrease at JLG Supersite
Note: Some years might not satisfy completeness criteria.

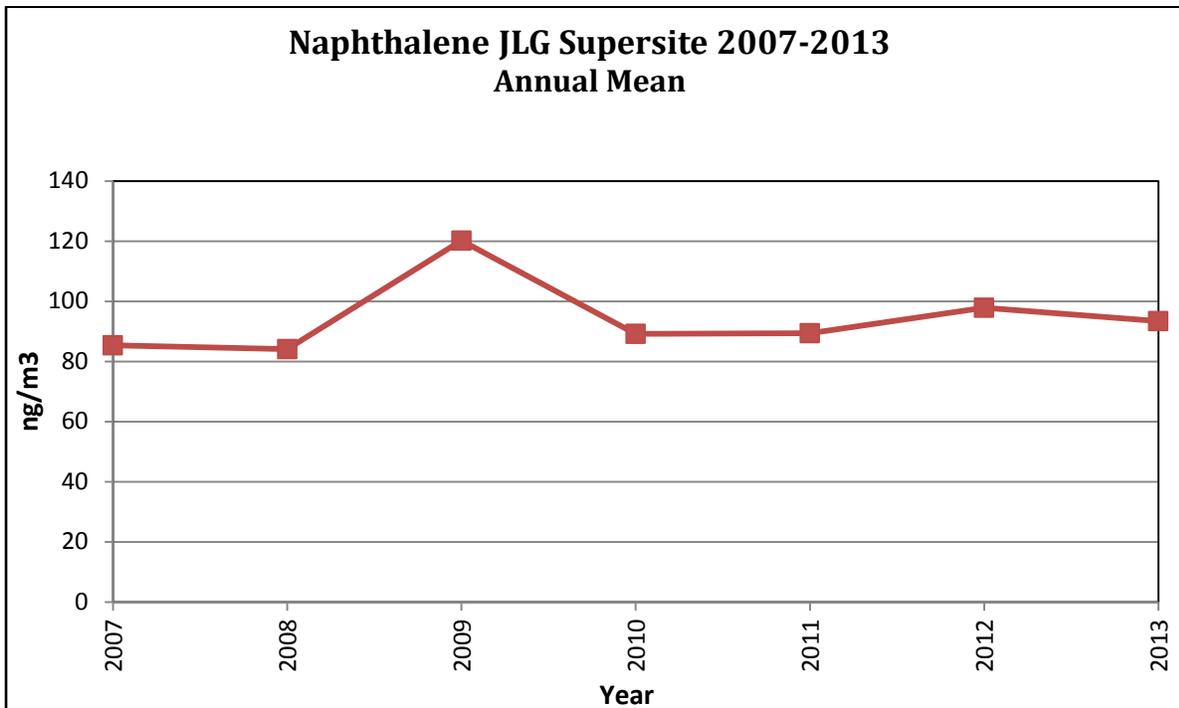


Figure 32 – Naphthalene Annual Mean Trend
 2007-2013: 5.7% increase at JLG Supersite
Note: Some years might not satisfy completeness criteria.

3.0 Photochemical Assessment Monitoring Stations (PAMS)

The PAMS are a collection of monitoring stations that are used to obtain comprehensive and representative data on O₃ air pollution. The main purpose of PAMS is to provide a more comprehensive database for O₃ and its precursors due to nonattainment of the NAAQS in many areas around the country. The larger database of O₃ and its precursors is used to assist state and local agencies in evaluating, tracking, and refining controls for attaining the O₃ NAAQS. In order to fully describe an O₃ area, a PAMS network may include four different types of sites. Type 1 is an upwind background site, Type 2 is a maximum O₃ precursor concentration site, Type 3 is a maximum O₃ concentration site, and Type 4 is an extreme downwind site. The different types of PAMS sites are used to show a complete picture of O₃ formation. The 40 CFR Part 58 requires states to establish PAMS as part of their monitoring networks in O₃ nonattainment areas classified as serious, severe, or extreme. Pollutants measured include O₃, NO_x, CO, NO_y, and VOCs. Health effects from these pollutants are described in their individual sections of this report and range from short term to long term effects.

3.1 Background

High O₃ concentrations are caused when sunlight and precursor pollutants react in the lower atmosphere. This is typically a summer phenomenon occurring from March-October in Arizona. Highest concentrations occur during the June-August months, due to higher temperatures and more direct sunlight. The PAMS monitoring season is during this high concentration time from June-August, which allows for many more samples to be taken in order to characterize peak levels of O₃ and O₃ precursor pollutants. The sources for O₃ precursor pollutants include combustion vehicles, biogenics from plants, industrial processes, and electric power plants. Ozone conditions and typical concentrations in Arizona are described in more detail in the O₃ section of this report.

ADEQ maintains the PAMS network in the metropolitan statistical area (MSA) of Phoenix-Mesa-Scottsdale. This MSA was designated as marginal nonattainment for the 2008 O₃ standards as stated in Section II of this report. ADEQ operates a Type 2 station and a Type 3 station, to support the PAMS network. The Type 2 site is JLG Supersite. The Pollutants monitored are VOCs, Carbonyls, NO_x, CO, O₃, and surface meteorology. JLG Supersite is used to monitor the magnitude and type of precursor emissions in the area and is representative of the MSA. It is located near the boundary and downwind of the central business district of downtown Phoenix and downwind of many industrial sources of precursor emissions in the Phoenix MSA. The Type 3 site is Queen Valley. The pollutants monitored are VOCs, NO_y, O₃, and surface meteorology. The Queen Valley location is designed to capture maximum O₃ concentrations. Typical Type 3 sites are located 10-30 miles from the fringe of the urban area. Queen Valley is located 30 miles from downtown Mesa.

3.2 Monitoring Methods

Methods for monitoring pollutants that are part of PAMS are the same as with other networks. PAMS O₃ is monitored using the same instruments and ultraviolet absorption method as described in Section II of this report.

VOCs and Carbonyls use the same instruments and methods as the air toxics network. In addition to a 24-hour sample, three three-hour samples are collected for analysis during the morning and early afternoon in order to better characterize precursor pollutant concentrations during the day. Peak levels generally occur during the morning rush hour; therefore, samples of both VOCs and Carbonyls are collected in three three-hour blocks from 5:00 am to 8:00 am, 8:00 am to 11:00 am, and 11:00 am to 2:00 pm during PAMS season.

Nitrogen oxides and NO_y monitoring use the same chemiluminescence instruments and methods as the NO₂ network. Reactive nitrogen oxides differ from NO_x due to the sampling height. Since O₃ is a scavenger of some oxides of nitrogen the sample inlet is located above ground level O₃ (~10 m). The reactive forms of nitrogen that are normally scavenged by O₃ are measured at this elevated height to give a better representative value of the total nitrogen precursor pollutants.

PAMS CO is monitored using the same nondispersive infrared instruments and methods as described in Section II of this report.

All readings from continuous gas analyzers are averaged into hourly, daily, quarterly, and yearly averages for data analysis. Readings are retrieved by a data collection system and stored in a database. Readings from sample monitors (VOCs and Carbonyls) are sent to ADEQ by the analysis lab and are also averaged for data analysis.

3.3 Trends

Total Non-Methane Organic Compounds (TNMOC) are defined in the PAMS TAD as the unspiciated total concentration of VOC (C2 through C12) in ambient air as determined by “summation of peaks” from GC/FID analysis, expressed in parts per billion carbon (ppbC). Parts per billion carbon is the concentration (in ppb) of the compounds multiplied by the number of carbon molecules in the compound. This unit is useful as it gives the concentration of the individual carbon molecules available to react to form O₃. TNMOC are a precursor to O₃ and emission sources include fossil fuel burning, landfills, and solvents. The annual means used for trend analysis were calculated from 24-hour samples taken during the June 1st through August 31st PAMS sampling season of each year. There was a decrease in the average TNMOC concentrations at JLG Supersite and Queen Valley from 2007 to 2013 (Figure 33).

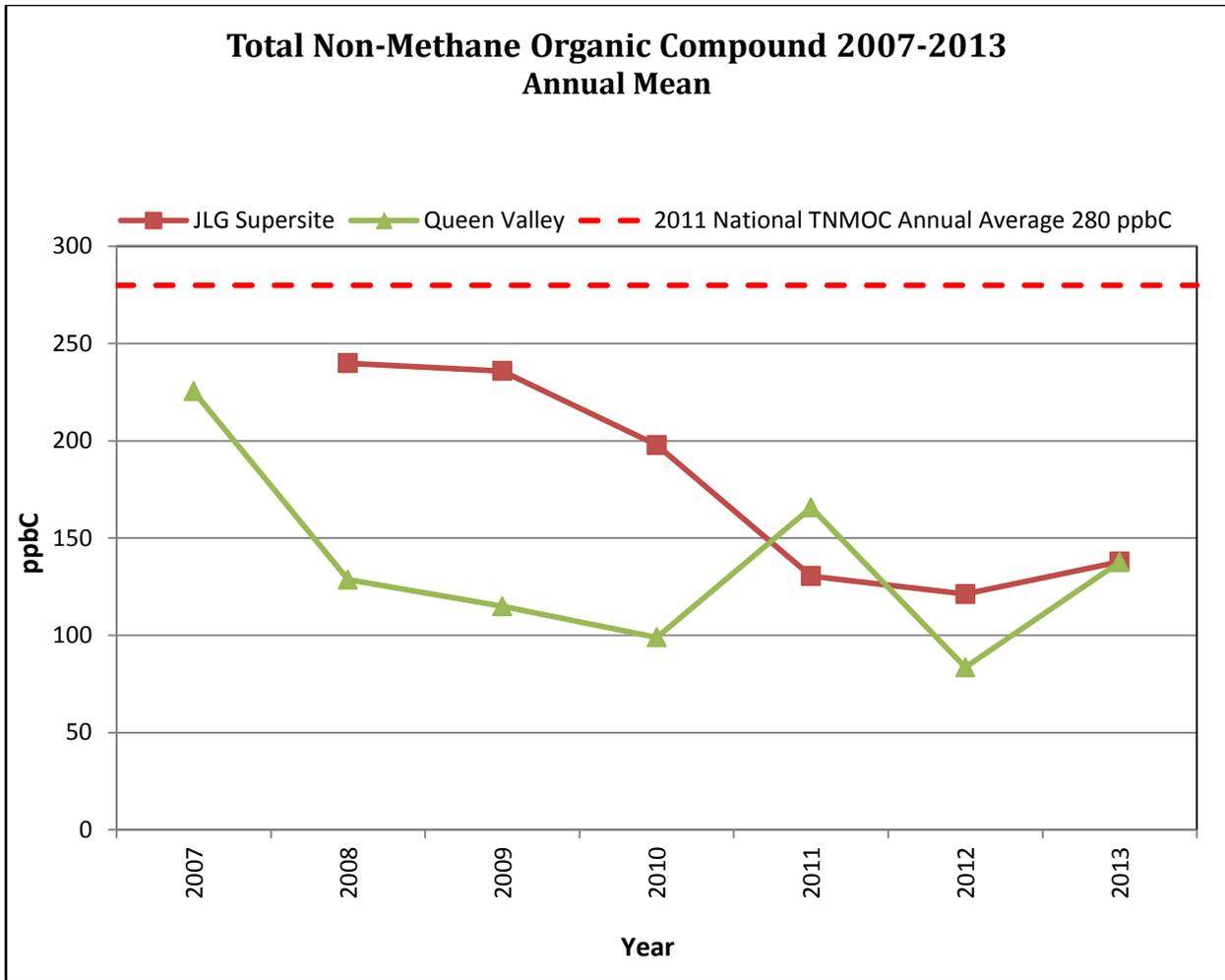


Figure 33 – TNMOC Annual Mean Trend

2007-2013: 50.4% decrease at JLG Supersite and Queen Valley

Note: Some years may not satisfy completeness criteria.

Section IV – Visibility

Visibility has historically been defined as the maximum distance that a human eye can view a contrasting object. This is a technical definition that also can include other quantitative measurements. Defining visibility does not always lend itself to the quantitative measures that many instruments can produce. Visibility can also be based on the judgments of an observer viewing a beautiful vista. It is the combination of both the technical and the judgmental definition that most closely describes visibility.

Visibility and haze are two terms that are closely related, as haze is the form of air pollution that degrades visibility. Haze is caused when sunlight encounters tiny particles in the air, which reduces the color and clarity of what is seen. Since

1988, the U.S. EPA, States, and Federal land management agencies have conducted monitoring of air pollution and visibility impairment at a number of national parks and wilderness areas across the U.S. In 1999, the EPA announced a major effort to improve air quality in national parks and wilderness areas. The Regional Haze Rule (RHR) calls for the improvement of visibility in 156 Class I national parks and wilderness areas. The RHR outlines the requirements for states to follow in order to address haze impairing pollutants. The purpose of the rule is to improve visibility at the 156 national parks and wilderness areas throughout the U.S. (formally known as mandatory Federal Class I areas). States are required to demonstrate reasonable progress towards the national visibility goal established in 1977 by the CAA (Section 169A and 169B): "The prevention of any future, and the remedying of any existing impairment of visibility in mandatory Class I Federal areas which impairment results from man-made air pollution." ADEQ submitted its Regional Haze SIP in 2011 which satisfies the planning portion of the RHR with an emphasis on the human-caused sources of pollutants.



Figure 34 – Pleasant Valley monitoring station

1.0 Urban Haze

Haze not only affects natural vistas, but can also affect urban skylines. Reducing urban haze mirrors the objectives of the RHR in improving visibility in the urban environment by reducing emissions that directly contribute to haze.

1.1 Program Background

ADEQ operates a network of urban visibility instruments that are designed to characterize different optical phenomena in the Phoenix metropolitan area. This network measures the amount of visibility impairing haze using different optical measurements which show the amount of light scattered from

one point to another. ADEQ operates several instruments as part of its Urban Haze program including one transmissometer, two nephelometers, and five high resolution digital cameras throughout the Phoenix metropolitan area. The transmissometer is located in downtown Phoenix, the nephelometers are the in the west valley, and the cameras are located in various locations throughout the entire Phoenix metropolitan area.

1.2 Monitoring Methods

The transmissometer is used to assess visibility impairment by measuring the amount of light lost over a known distance. The instrument consists of a light source (transmitter) and light detector (receiver) which are generally located on an elevated surface, such as a tall building, and are separated by a distance of 1-3 miles. The Phoenix transmissometer transmitter is located on the roof of Phoenix Baptist Hospital and the receiver is 2.96 miles away on the roof of the Holiday Inn on the corner of Osborn and 3rd Avenue in downtown Phoenix. The transmitter emits a uniform light beam of constant intensity that is carefully aimed at the receiver. The amount of light transmitted and received is precisely measured. The receiver includes a telescope that gathers the transmitter's light and a computer that compares the measured light intensity with the known transmitter light intensity to calculate the transmission of the intervening atmosphere. The measured transmission can be related to the light lost along the path due to scattering and absorption.

The nephelometer is used to assess visibility impairment by estimating the particle scattering coefficient at a point location. The nephelometer provides a direct measurement of the light scattered by aerosols and gases in a sampled air volume. It employs a light source and a detector set to the side of the source. Light is scattered by particles over a large range of scattering angles, in a defined band of visible wavelengths. The detector picks up the specific wavelengths of scattered light to the side. These wavelengths are particular to a certain type of particle, PM_{2.5}. Because the total light scattered out of a path is the same as the reduction of light along a path due to scattering, the integrating nephelometer gives a direct estimate of the particle scattering coefficient and hence impaired visibility.

Photographic documentation is an important aspect of evaluating visibility. Photography is an effective way to document events and trends on a media that is easily interpreted. ADEQ uses high-resolution digital images from locations around Phoenix to document visibility conditions. The digital images are readily available for viewing on a computer at www.phoenixvis.net and can be conveniently distributed via the Internet, easily stored, managed, and duplicated without degradation. Each site consists of a high-resolution digital camera housed in a weatherproof, environmental enclosure, and a supporting image capture computer. Digital images are captured every 15 minutes, stored on the system's internal hard drive, and uploaded to the Web site.

1.3 Trends

Visibility data from these monitors can be expressed by several different measurement units: deciview, inverse megameters, and visual range. An inverse megameter (Mm⁻¹) (units used by ADEQ) is a

representation of the ratio between how much light is not received by a sensor compared to the amount of light that leaves a source. Higher numbers mean worse visibility. As a Mm^{-1} is an uncommon unit of measure, the visual range will be reported.

Optical measurements of urban visibility have been made continuously in Phoenix since 1994. Light extinction, the degree to which light is reduced by its interaction with particles and gases in the atmosphere, is measured continuously with transmissometers. The units of measurement are Mm^{-1} : the higher the light extinction value in Mm^{-1} , the more visibility is reduced. In Figure 35, these light extinction data have been plotted as a yearly average and converted to the more practical units of visual range in miles.

In Phoenix, when taking into consideration all hours of the day, transmissometer data indicate a steady trend toward clearer air for the mean, cleanest 20%, and the dirtiest 20% categories over the past 18 years (Figure 35). The data trends have shown that the morning hours visual range (Figure 36) have increased less when compared with the all hours visual range. This indicates that the overall increasing trend of visual range has been influenced by the other parts of the day more than the morning hours. Visual range peaked during the year 2010, which was an above average year for rainfall.

The trend in downtown Phoenix has an increase which shows an increase in visual range of approximately 15.7 miles over the past 18 years. Haze and visibility in the Phoenix downtown area have improved steadily.

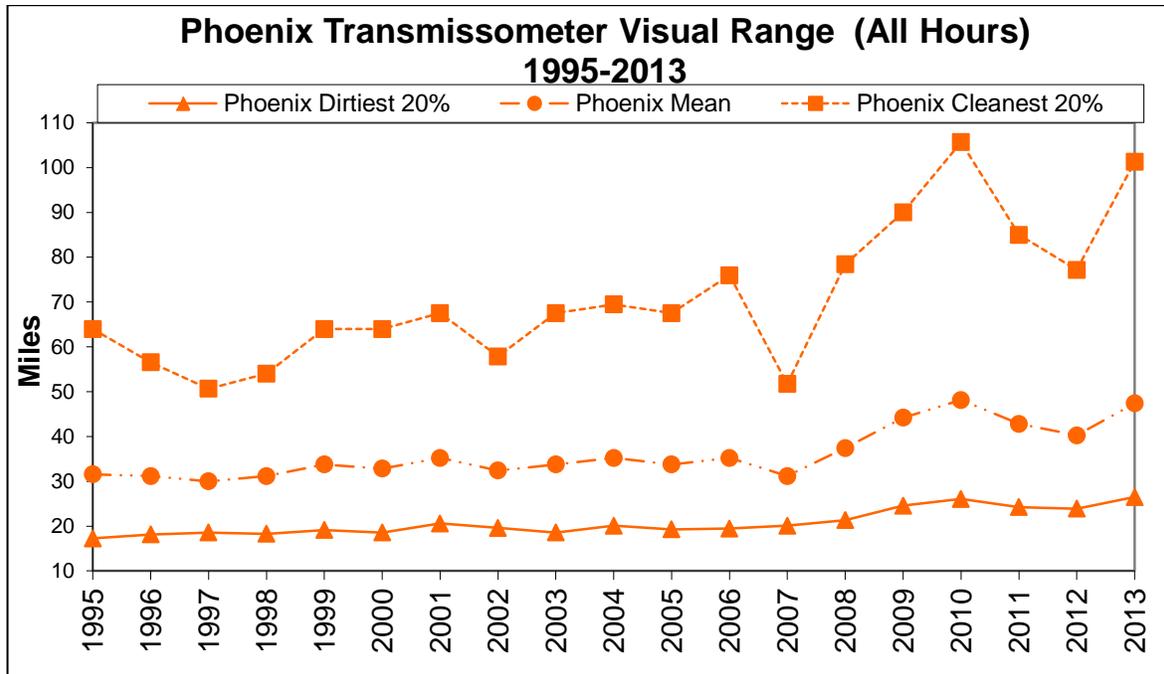


Figure 35 – Transmissometer Visual Range (All Hours) Trend
 1995-2013: 56% increase in the Mean visual range or 15.7 miles
 83% increase in the 20% Cleanest times
 52% increase in the 20% Dirtiest times

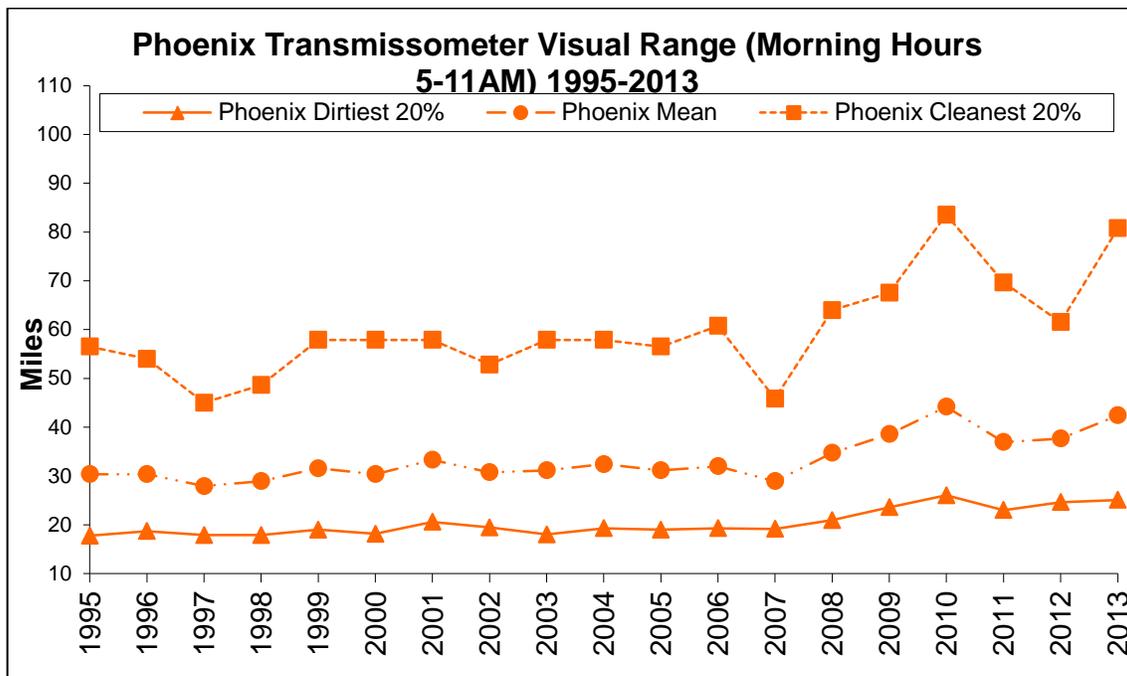


Figure 36 – Transmissometer Visual Range (Morning Hours) Trend

1995-2013: 45% increase in the Mean visual range trend in the morning hours or 12.1 miles

50% increase in the 20% Cleanest times

47% increase in the 20% Dirtiest times

Transmissometer seasonal variability show which seasons have higher visual range (Figure 37). The seasons shown are Winter (Jan-Mar), Spring (Apr-June), Summer (July-Sept), and Fall (Oct-Dec). Data for seasonal variability is currently available for the years 2006-2012. The visual range is greater during the spring and summer months than during the fall and winter months. Visual range peaked during the year 2010 in the summer months at around 75 miles. A contributing factor for this was that 2010 was an above normal year for precipitation. Rainfall cleans the air, improving visual range. The overall visual range trend is increasing over the past seven years.

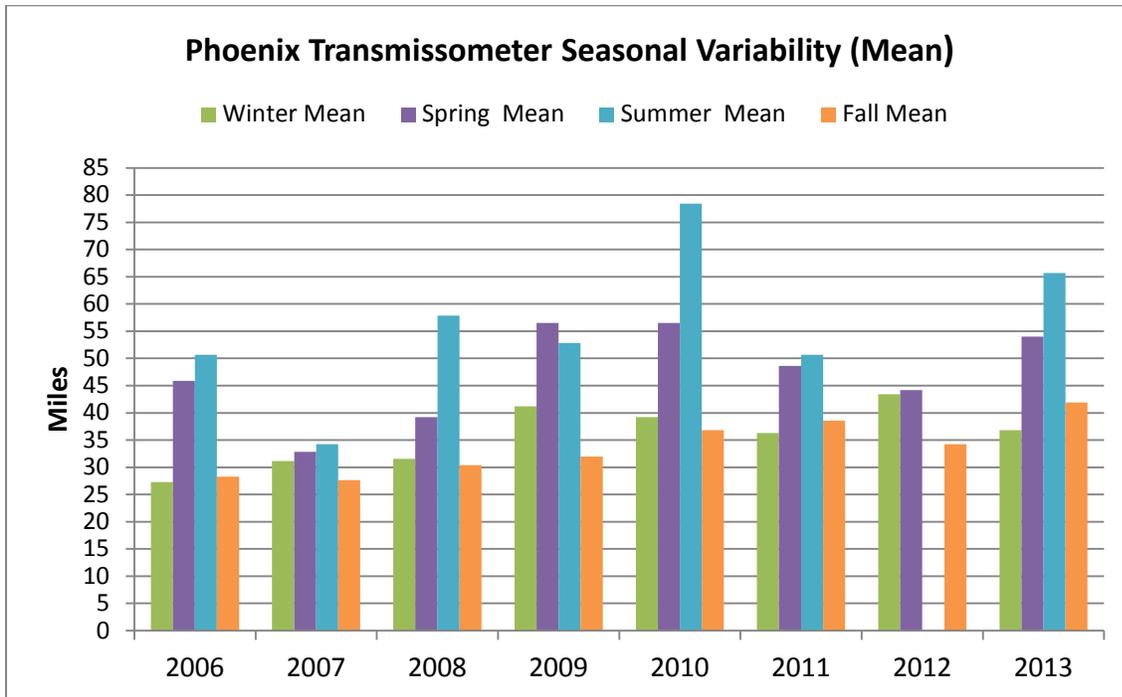


Figure 37 – Transmissometer Seasonal Average Trend

2006-2013: 14% increase in the Transmissometer Mean visual range trend in the spring months

58% increase in the summer months

60% increase in the fall months

47% increase in the winter months

Note: Data did not meet completeness criteria for the summer of 2012; therefore, it was excluded from analysis.

Nephelometer visual range also shows an increasing trend over the years 2006-2012 (Figure 38). Spatial variation shows sites increase in visual range at different rates, with the greatest improvement at the Vehicle Emissions Laboratory site, close to downtown Phoenix. While there steady improvement at Estrella Mountain Park and Dysart, the biggest increases have been in the downtown area. Possible causes for this can be that decreases in vehicle pollution have increased visual range. Whereas in the areas where vehicle density is lower (Estrella and Dysart), the increases come from other pollutant controls which are less substantial.

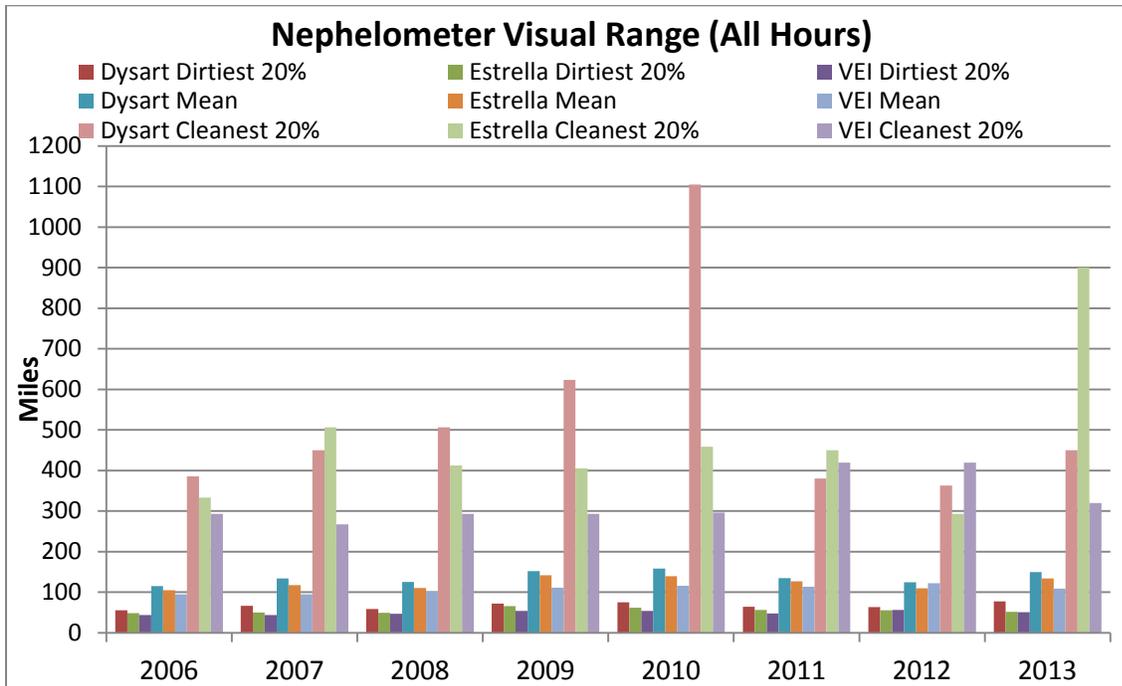


Figure 38 – Nephelometer Visual Range Trend
 2006-2013: 25% increase in the Nephelometer Mean visual range trend or 23 miles for VEI
 16% increase for Estrella
 15% increase for Dysart

Examples of visibility conditions are shown in Figure 39. As visibility decreases, the clarity and resolution of the mountains and downtown area decreases and a milky appearance covers the vista. On poor visibility days, the mountain ridgeline and most of the downtown area are no longer visible.



Figure 39 – Examples of Visual Condition in Phoenix. Excellent to Very Poor visibility days taken from the high resolution digital picture. All are images of South Mountain taken from the camera located at North Mountain. Center-Excellent, Top left-Good, Top right-Fair, Bottom left-Poor, Bottom right-Very Poor.

2.0 IMPROVE

The IMPROVE program is the main supporter of the RHR across the nation. It uses monitors specific to the objectives to increase visibility and decrease haze in the national parks and wilderness areas. The program objectives of IMPROVE are:

- to establish current visibility and aerosol condition in mandatory Class I areas;
- to identify chemical species and emission sources responsible for existing man-made visibility impairment;
- to document long-term trends for assessing progress towards the national visibility goal; and,
- with the enactment of the Regional Haze Rule, to provide regional haze monitoring representing all visibility-protected federal class I areas where practical.

The IMPROVE monitors are closely related to those in the CSN in that they use different types of sample filters to measure the amount of speciated pollutants in the air. Ambient air is drawn into the sampler, where the pollutants are deposited onto the sample filters. There are four different sample filters that are run on the same day, each collecting different types of pollutants. Different aerosols and particulates contribute to haze more than others: therefore, the monitors are designed to capture these specific pollutants.

The IMPROVE network in Arizona has 16 samplers in 12 Class I areas. This program is governed by a steering committee composed of representatives from federal and regional/state organizations. This network was established in 1985 to aid the creation of federal and state implementation plans. ADEQ supports the IMPROVE network by performing filter sample changes at JLG Supersite, Queen Valley, and Douglas Red Cross sites and by conducting performance audits on protocol samplers in Arizona. ADEQ is one of the many government agencies involved in the IMPROVE network. Users of these data include, but are not limited to, the EPA, visibility researchers, and land management agencies.

More information of the IMPROVE program and data can be accessed at the following website: <http://vista.cira.colostate.edu/improve/>.

Appendix I – Abbreviations

ADEQ	Arizona Department of Environmental Quality
AgBMP	Best Management Practices for Agricultural Activities
AQI	Air Quality Index
AQS	Air Quality System (EPA database)
BAM	Beta Attenuation Monitor
Bext	Total Light Extinction
Bscat	Light Scattering
CAA	Clean Air Act
CFR	Code of Federal Regulations
CO	Carbon Monoxide
CSATAM	Community Scale Toxics Ambient Monitoring
CSN	Chemical Speciation Network
DM&QA	Data Management & Quality Assurance Unit
DQO	Data Quality Objective
EPA	Environmental Protection Agency
FIP	Federal Implementation Plan
FMMI	Freeport McMoRan Copper and Gold Inc.
HAP	Hazardous Air Pollutant
HC	Hydrocarbons
IMPROVE	<u>I</u> nteragency <u>M</u> onitoring of <u>P</u> ROtected <u>V</u> isual <u>E</u> nvironments
IR	Infrared
MAG	Maricopa Association of Governments
MQO	Measurement Quality Objective
MSA	Metropolitan Statistical Area
mM ⁻¹	Inverse Megameter
mg/m ³	Milligrams per Cubic Meter
µg/m ³	Micrograms per Cubic Meter
NAAQS	National Ambient Air Quality Standard
NATTS	National Air Toxics Trends Station
NCore	National Core multipollutant monitoring stations
NEI	National Emissions Inventory
NM	National Monument
NO	Nitric Oxide
NO ₂	Nitrogen Dioxide
NO _x	Nitrogen Oxides
NO _y	Total Reactive Oxides of Nitrogen
O ₃	Ozone
PAH	Polycyclic Aromatic Hydrocarbon
PAMS	Photochemical Assessment Monitoring Station
Pb	Lead
PM	Particulate Matter
PM ₁₀	Particulate Matter ≤ 10 microns
PM _{10-2.5}	Coarse Particulate Matter between 2.5 to 10 micrometers aerodynamic diameter, may also be denoted as PM _{coarse}

PM _{2.5}	Particulate Matter ≤ 2.5 microns
ppb	Parts Per Billion
ppm	Parts Per Million
PQAO	Primary Quality Assurance Organization
PSD	Prevention of Significant Deterioration
PTFE	Polytetrafluoroethylene
QA	Quality Assurance
QAPP	Quality Assurance Program Plan
QC	Quality Control
QMP	Quality Management Plan
RH	Relative Humidity
RHR	Regional Haze Rule
SATMI	School Air Toxics Monitoring Initiative
SIP	State Implementation Plan
SLAMS	State and Local Air Monitoring Stations
SO ₂	Sulfur Dioxide
SOP	Standard Operating Procedure
SPM	Special Purpose Monitor
SVOC	Semi-Volatile Organic Compound
TAD	Technical Assistance Document
TEOM	Tapered Element Oscillating Microbalance
TNMOC	Total Non-Methane Organic Compound
TSP	Total Suspended Particle
UATMP	Urban Air Toxics Monitoring Program
USG	Unhealthy for Sensitive Groups
VOC	Volatile Organic Compound

Appendix II – References

<http://www.airnow.gov/> - AQI Forecast

<http://www.ecy.wa.gov/programs/air/sips/pollutants/naaqs.htm> - National and State Ambient Air Quality Standards, Department of Ecology, State of Washington

<http://www.epa.gov/air/criteria.html> - National Ambient Air Quality Standards (NAAQS)

<http://www.epa.gov/airquality/carbonmonoxide/health.html> - Carbon Monoxide Health Effects

<http://www.epa.gov/airquality/ozonepollution/health.html> - Ozone (O₃) Health Effects

<http://www.epa.gov/airtrends/index.html> - National Trends.

http://www.epa.gov/ttn/naaqs/standards/co/s_co_history.html - Carbon Monoxide (CO) Standards - Table of Historical CO NAAQS

http://www.epa.gov/ttn/naaqs/standards/nox/s_nox_history.html - Nitrogen Dioxide (NO₂) Standards - Table of Historical NO₂ NAAQS

http://www.epa.gov/ttn/naaqs/standards/ozone/s_o3_history.html - Ozone (O₃) Standards - Table of Historical O₃ NAAQS

http://www.epa.gov/ttn/naaqs/standards/pb/s_pb_history.html - Lead (Pb) Standards - Table of Historical Pb NAAQS

http://www.epa.gov/ttn/naaqs/standards/pm/s_pm_history.html - Particulate Matter (PM) Standards - Table of Historical PM NAAQS

<http://www.epa.gov/ttn/naaqs/standards/so2/data/so2final.pdf> - National Ambient Air Quality Standards for Sulfur Oxides (Sulfur Dioxide)—Final Decision

http://www.epa.gov/ttn/naaqs/standards/so2/s_so2_history.html - Sulfur Dioxide (SO₂) Primary Standards - Table of Historical SO₂ NAAQS

<http://www.phoenixvis.net/PPMmain.aspx> - Non-Regulatory Portable Particulate Monitors

<https://aqs.epa.gov/aqs/> - EPA's Air Quality System

www.epa.gov/air/criteria.html - National Ambient Air Quality Standards

Appendix III – 2013 Area Designations Map

