

ANALYSIS OF INDOOR AND OUTDOOR PARTICULATE MATTER AT
VARIOUS RESIDENCES IN THE EL PASO REGION

JOSEPH C. PIÑON

Department of Civil Engineering

APPROVED:

Wen-Whai Li, Ph.D., Chair

Luis Trueba, Ph.D.

Hector Olvera, Ph.D.

Benjamin C. Flores, Ph.D.
Acting Dean of the Graduate School

Copyright ©

by

JOSEPH PIÑON

2011

PREVIEW

Dedication

Without the support system I received from my family, none of this would have been made possible. Thank you all for believing in me and standing by my side when I needed you all the most. This thesis is dedicated to my loving parents who always had a warm welcome place for me; to my brothers, for being the best a brother could ask for; to Nadia, for being patient with me and for always being there for me when I needed you; to my grandparents, for being great pillars of support; and to my Aunt Lupe and Uncle Richard for their unbridled support. Thank you all!

PREVIEW

ANALYSIS OF INDOOR AND OUTDOOR PARTICULATE MATTER AT VARIOUS
RESIDENCES IN THE EL PASO REGION

by

JOSEPH C. PIÑON, B.S.C.E

THESIS

Presented to the Faculty of the Graduate School of
The University of Texas at El Paso
in Partial Fulfillment
of the Requirements
for the Degree of

MASTER OF SCIENCE IN ENVIRONMENTAL ENGINEERING

Department of Civil Engineering
THE UNIVERSITY OF TEXAS AT EL PASO

August 2011

UMI Number: 1498310

All rights reserved

INFORMATION TO ALL USERS

The quality of this reproduction is dependent on the quality of the copy submitted.

In the unlikely event that the author did not send a complete manuscript and there are missing pages, these will be noted. Also, if material had to be removed, a note will indicate the deletion.



UMI 1498310

Copyright 2011 by ProQuest LLC.

All rights reserved. This edition of the work is protected against unauthorized copying under Title 17, United States Code.



ProQuest LLC.
789 East Eisenhower Parkway
P.O. Box 1346
Ann Arbor, MI 48106 - 1346

Acknowledgements

This research was made possible by the dedication and guidance provided by Dr. Wen-Whai Li of the Department of Civil Engineering at the University of Texas at El Paso (UTEP). I would also like to acknowledge Dr. Hector Olvera, who was always willing to give a helping hand and guidance whenever it was needed.

In addition, the help of the undergraduate and graduate members of the UTEP Air Quality Research Laboratory was paramount in ensuring everything was completed as per the full potential of the project.

Partial student support was received by the Center for Environmental Resource Management (CERM) and the Southwest Consortium for Environmental Research and Policy (SCERP). The UTEP Civil Engineering Department had also provided partial support throughout the duration of my graduate studies.

Abstract

This thesis is a presentation of particulate matter data collected from February 2007 to June 2008. Personal environment monitors (PEM), which measured 7-day $PM_{2.5}$ concentrations, had been deployed at 14 homes during the 2007 year and deployed in 16 different homes during the 2008 sampling year. Within each sampling year, sampling occurred for two one-week (7-day) periods during each season. In addition to PEM sampling, a tapered element oscillating microbalance (TEOM) continuous sampler had been deployed in both indoors and outdoors at one pre-selected household per week. Gravimetric, chemical, and black carbon analysis had been performed on the PEM $PM_{2.5}$ filters while time-series plots of the TEOM PM_{10} and $PM_{2.5}$ data were constructed to extract diurnal patterns from the data. Results from the TEOM data affirm the purported existence of a strong diurnal pattern in particulate matter observations in the El Paso region.

Strong increases in particulate matter were seen, on average, between the hours of 06:00 and 10:00 (morning) and once again in the evening from 16:00 to 20:00. Of the TEOM data, the mean outdoor and indoor $PM_{2.5}$ concentrations were found to be $23.9 \mu\text{g}/\text{m}^3$ and $17.8 \mu\text{g}/\text{m}^3$, while the mean outdoor and indoor PM_{10} concentrations were found to be $27.3 \mu\text{g}/\text{m}^3$ and $18.0 \mu\text{g}/\text{m}^3$, respectively. Mean outdoor and indoor TEOM $PM_{2.5}$ concentrations were found to be, on average, greater during the summer seasons. The mean TEOM outdoor PM_{10} was also found to be greater during the summer seasons over the winter seasons, while the mean indoor PM_{10} TEOM concentration was found to be only slightly greater during the winter seasons over the summer seasons. Indoor/Outdoor (I/O) ratios for PM_{10} for all TEOM sampling locations during the entire study period were found to be greater, on average, than (I/O) ratios for $PM_{2.5}$ with values of 1.15 and 1.05, respectively.

A mean $PM_{2.5}$ concentration of $19.8 \mu\text{g}/\text{m}^3$ was seen in the PEM indoor samples collected at all locations during the entire study period, while the mean winter concentration $22.9 \mu\text{g}/\text{m}^3$ was shown to

be much higher than the mean summer concentration $16.8 \mu\text{g}/\text{m}^3$. Results of an EDXRF analysis found that geologic elements (Al, Si, Ca, Fe, Ti, and K) were found to compose a majority of the total $\text{PM}_{2.5}$ mass at 17.4%, while toxic trace elements (Cu, Cr, As, Cd, and Pb) were found to compose only 0.19% of the total $\text{PM}_{2.5}$ mass. All analyzed elements (Σ Elements) were found to compose, on average, 24% of the total $\text{PM}_{2.5}$ mass collected using the PEM sampler. The mean black carbon (BC) concentration on the PEM filters for the entire study period was found to be $0.30 \mu\text{g}/\text{m}^3$; BC concentrations were also found to be higher during the winter seasons, $0.38 \mu\text{g}/\text{m}^3$, than the summer seasons, $0.21 \mu\text{g}/\text{m}^3$. Black carbon was also found to compose, on average, 1.8% of the total $\text{PM}_{2.5}$ mass.

Principle components analysis (PCA) was conducted on the EDXRF data and presented four primary components within the elemental data. The first component included Al, Ba, Ca, Fe, K, Mg, Mn, Pb, Sr, Ti, and V, while the second component included Cr, Ga, Mo, and Ni. The third component consists of As, Rh, and Te, while the final component was composed of what is believed to be salt (Na and Cl). The first component, composed of mostly geologic material, composed the greatest majority of elements found in the total PM mass, while the other components were mostly hypothesized to be related to industrial release in the area.

Table of Contents

Acknowledgements.....	v
Abstract.....	vi
1. Introduction.....	1
1.1. Introduction.....	1
1.2. Research Objective	7
2. Particulate Matter and Related Health Effects.....	9
2.1. Particulate Matter.....	9
2.2. Sources and Characteristics of Particulate Matter	12
2.3. Local Particulate Matter Studies.....	15
2.4. Health Effects	19
3. The Monitoring Program	21
3.1. Site Description	21
3.2. Sampling Period.....	30
3.3. Personal Environmental Monitors (PEM)	33
3.4. TEOM.....	34
3.5. Sample Collection and Gravimetric Analysis.....	35
3.5.1. Filters	36
3.6. Energy Dispersive X-Ray Fluorescence	36
3.7. Reflectance Analysis	37
3.8. TCEQ CAMS Data	38
3.9. EPA Toxic Release Inventory (TRI) Program.....	38
4. Results & Discussion.....	39
4.1. PEM Summary Statistics	39
4.2. Elemental Data Comparison	49
4.3. Principle Components Analysis.....	52
4.3.1. PCA Method	54
4.3.2. PCA Results and Discussion	61
4.3.3. PCA Components Summary Statistics	63
4.4. TEOM Diurnal Variation PM ₁₀ and PM _{2.5} Data	64
4.4.1. Year 2007.....	65

4.4.2. Year 2008.....	76
4.5. TEOM PM ₁₀ and PM _{2.5} Indoor and Outdoor Summary Statistics	91
4.6. TEOM PM ₁₀ and PM _{2.5} Indoor /Outdoor Ratio Summary Statistics ...	93
5. Conclusions.....	95
References.....	104
Appendix A.....	108
Appendix B.....	206
Appendix C.....	218
Appendix D.....	226
Appendix E.....	229
Vita	235

List of Tables

Table 1.1: Meteorological Data Averages	2
Table 2.1: National Ambient Air Quality Standards	11
Table 3.1: PEM Sample Sites and Sample Periods for 2007	31
Table 3.2: PEM Sample Sites and Sample Periods for 2008	32
Table 3.3: TEOM Sampling Periods and Locations	33
Table 4.1: Summary Statistics by Season and Year for Indoor PEM $PM_{2.5}$, Σ Elements, and BC	39
Table 4.2: Percent (%) Composition by Year, Season, and Study Period	41
Table 4.3: Summary Statistics by Location for $PM_{2.5}$	45
Table 4.4: Mean Concentrations for Each Element for the Entire Period	47
Table 4.5: Mean Elemental Concentrations by Year and Season	48
Table 4.6: Study Elemental Means vs. UTEP Study (2001) & Paschold (2003)	51
Table 4.7: Critical Values for a Correlation Coefficient at $\alpha=.01$ for a Two-Tailed Test	53
Table 4.8: KMO and Bartlett's Test	55
Table 4.9: Total Variance Explained	56
Table 4.10: Rotated Component Matrix ^a	58
Table 4.11: Total Variance Explained	59
Table 4.12: PCA Results	61
Table 4.13: PCA Component Percent Composition by Year, Season, and Study Period	64
Table 4.14: TEOM $PM_{2.5}$ & PM_{10} Summary Statistics	91
Table 4.15: TEOM Indoor /Outdoor Ratio Summary Statistics	93
Table 4.16: TEOM Mean Concentrations by Location	94

List of Figures

Figure 1.1: National PM _{2.5} Air Quality, 2000-2009	5
Figure 1.2: National PM ₁₀ Air Quality, 1990-2009	5
Figure 1.3: El Paso PM ₁₀ Air Quality; 1990-2009	6
Figure 1.4: El Paso PM _{2.5} Air Quality; 1990-2009	7
Figure 3.1: 2009 Toxic Release Inventory (TRI) Sites in the El Paso Region	23
Figure 3.2: PEM Sampling Locations for Summer 2007	24
Figure 3.3: PEM Sampling Locations for Winter 2007	25
Figure 3.4: PEM Sampling Locations for Summer 2008	26
Figure 3.5: PEM Sampling Locations for Winter 2008	27
Figure 3.6: TEOM Sampling Sites for Year 2007	28
Figure 3.7: TEOM Sampling Sites for Year 2008	29
Figure 4.1: Boxplot of PEM Samples for Entire Study Period (Units µg/m ³)	43
Figure 4.2: PCA Cattell's Scree Plot	57
Figure 4.3: Angie Bombach PM ₁₀	65
Figure 4.4: Angie Bombach PM _{2.5}	66
Figure 4.5: Angie Bombach PM ₁₀ Indoor vs. Outdoor	66
Figure 4.6: Angie Bombach PM _{2.5} Indoor vs. Outdoor	67
Figure 4.7: Juliandra PM ₁₀	69
Figure 4.8: Juliandra PM _{2.5}	69
Figure 4.9: Juliandra Indoor vs. Outdoor PM ₁₀	69
Figure 4.10: Juliandra Indoor vs. Outdoor PM _{2.5}	70
Figure 4.11: Juliandra PM ₁₀	71
Figure 4.12: Juliandra PM _{2.5}	71
Figure 4.13: Juliandra Indoor vs. Outdoor PM ₁₀	72
Figure 4.14: Juliandra Indoor vs. Outdoor PM _{2.5}	72
Figure 4.15: Angie Bombach PM ₁₀	74
Figure 4.16: Angie Bombach PM _{2.5}	74
Figure 4.17: Angie Bombach Indoor vs. Outdoor PM ₁₀	75
Figure 4.18: Angie Bombach Indoor vs. Outdoor PM _{2.5}	75
Figure 4.19: Sundial PM ₁₀	77

Figure 4.20: Sundial PM _{2.5}	77
Figure 4.21: Sundial Indoor vs. Outdoor PM ₁₀	78
Figure 4.22: Sundial Indoor vs. Outdoor PM _{2.5}	78
Figure 4.23: University PM ₁₀	80
Figure 4.24: University PM _{2.5}	80
Figure 4.25: University PM ₁₀	80
Figure 4.26: University PM _{2.5}	81
Figure 4.27: University Indoor vs. Outdoor PM ₁₀	81
Figure 4.28: University Indoor vs. Outdoor PM _{2.5}	82
Figure 4.29: University Outdoor vs. CAMS 12 PM ₁₀	82
Figure 4.30: University Outdoor vs. CAMS 12 PM _{2.5}	83
Figure 4.31: University PM ₁₀	84
Figure 4.32: University PM _{2.5}	85
Figure 4.33: University Outdoor vs. CAMS 12 PM ₁₀	85
Figure 4.34: University Indoor vs. Outdoor PM ₁₀	85
Figure 4.35: Starr PM ₁₀	87
Figure 4.36: Starr PM _{2.5}	87
Figure 4.37: Starr PM _{2.5} (Reduced Y-Axis)	88
Figure 4.38: Starr Indoor vs. Outdoor PM ₁₀	88
Figure 4.39: Starr Indoor vs. Outdoor PM _{2.5}	89

1. Introduction

1.1. Introduction

El Paso County is located at the westernmost edge of Texas, adjoining the state of New Mexico and the Mexican state of Chihuahua. El Paso, TX is a predominately semi-arid area with an average (30 years) precipitation of 8.65 inches and average high and low temperatures of 76.8 °F and 50.6 °F, respectively. El Paso sees an average of 202 days with clear skies and 49 days with precipitation more than 0.01 inches. Mean wind directions in the area are from the southwest (229 degrees) direction. Table 1.1 depicts the meteorology variation found in El Paso according to each respective season.

El Paso averages 14.5 significant dust events per year; consequences of these episodes range from simple irritations and increased particulate matter concentrations to serious disruptive events. These events aggravate respiratory health problems and can even turn deadly, resulting in fatal collisions as a result of near-zero visibility on city roadways and highways in the surrounding desert (Novlan et al., 2006). El Paso is generally a flat desert area with a large range of mountains, known as the Franklin Mountains, which rise to over 3280 ft. (1000m) above the surrounding area and are a north-south oriented mountain chain that is approximately 23.1 km (14.4 miles) long and 5.0 km (3.1 miles) wide. The Franklins create a divide between the western one-third of the city of El Paso and the central and eastern two-thirds of the city.

Table 1.1: Meteorological Data Averages

Period	Average Temperature ² (°C)	Average Precipitation ² (cm) ²	Average Wind Speed ² (km/hr)	Prevailing Wind Direction ² (degrees)
Annual	17.3	22.4	14.2	360
Winter (Dec.-Feb.)	7.2	1.2	13.6	333
Spring (March-May)	17.4	0.6	17.3	267
Summer (June-August)	27.2	3.2	13.6	183
Fall (Sept.-Nov.)	17.6	2.5	12.4	243

1: Source: National Climactic Data Center, Ashville, NC

2: Data based on 30 year averages

The El Paso region, which consists of El Paso, Texas, Sunland Park, New Mexico and Ciudad Juarez, Chihuahua, Mexico, has a population exceeding two million and is one of the largest metropolitan areas along the border. The region generally has poor visibility, especially in the winters which are susceptible to inversions, and respiratory problems are common place. The regional pollution problem is not only generated by inhabitants in the United States but also by those in surrounding areas such as Ciudad Juarez. In order to ensure that both countries work together to solve the regional pollution problem, the Border Environment Cooperation Commission (BECC) was created in 1993 under a side-agreement to the North American Free Trade Agreement (NAFTA). BECC aims at effectively enforce bi-national policies that aims to improve the air quality in the region. In concert with efforts by BECC, the United States Environmental Protection Agency (EPA) and the Texas Commission on Environmental Quality (TCEQ), the regional air quality in the El Paso region has progressively improved since 1990.

The use of fossil fuels or wood for heating and cooking in the area, the semi-arid weather, adverse meteorological conditions, and complex terrestrial features of the El Paso region are all contributors to the air pollution situation in the city. Poverty and industrialization in the Ciudad Juarez part of the El Paso/Juarez region are contributors to PM concentrations in the region (Espino et al., 2005). Higher levels of PM₁₀ and PM_{2.5} are witnessed in El Paso during the winter months when temperature inversions, along with calm conditions, trap PM in the ambient air.

The US EPA's regulatory PM₁₀ standard was frequently violated in the border region, including El Paso, with far fewer reported violations of the PM_{2.5} standard (EPA, 2011a). As a result of the frequent PM₁₀ violations, El Paso has been classified as a moderate "non-attainment" area with regard to PM₁₀ as of 1991 (EPA, 2011a, 2011b). It has been shown that episodic (short-lived, high concentration) events contribute many of the violations of the PM₁₀ National Ambient Air Quality Standards (NAAQS). PM₁₀ NAAQS are based on a maximum 24-h averaged concentration of 150 µg/m³ as well as an annual-averaged cap of 50.0 µg/m³. The aforementioned averaging obscures short-term peak doses and may misrepresent epidemiological dangers of PM (Staniswalis et al., 2005).

There are many local electronics, transportation, textile, machinery and refining industries in the El Paso area that contribute to the local PM pollution problem. The surrounding desert landscape is also another major source for the area's pollution problem and it has long been noted that there is a high natural ambient mineral material loading in the region (Dattner, 1994; Einfeld & Church, 1995). Unregulated kilns, for the manufacturing of bricks in particular, are reported to burn toxic waste fuels such as tires or used motor oil and are considered major contributors to PM₁₀ pollution in the region (Lauer et al., 2009). The preparation of food with wood, propane, or natural gas coupled with frequent major meteorological events; along with the arid weather, further complicate the local PM pollution problem. Analysis of PM_{2.5} samples collected using indoor personal environment monitors may help to

provide an insight as to the primary components of pollutants found in indoor $PM_{2.5}$ which may be contributed to the releases of the aforementioned activities in El Paso.

There are several commonly agreed on factors which can be attributed to the pollution problem in El Paso. El Paso is mostly surrounded by desert land which produces large amounts of dust during episodes of high winds. There are also a large number of unpaved roads in the area surrounding El Paso as well as in areas in Mexico located along the border with El Paso which can also be attributed to the creation of fugitive dust in the area. Many of the same areas along the border in Mexico burn scrap wood and other refuse material for home heating and cooking in the much colder winter months of the year. The burning of such material may release metals into the atmosphere and may exacerbate PM concentrations in the region (Espino et al., 2005). Commercial activities such as scrap metal foundries, refining, smelting, various agricultural activities and brick kilns also contribute to the local pollution problem.

In the United States, PM concentrations have decreased on average since interest for certain size of particles invoked national monitoring programs. Monitoring for $PM_{2.5}$ began nationwide as of the year 2000 and according to Figure 1.1, the $PM_{2.5}$ levels in the United States has decreased by 27%. Based on Figure 1.2, PM_{10} levels in the United States have decreased 38% since nationwide monitoring began in 1990. The decrease in PM in the United States is a direct result of the efforts of the EPA, state, tribal and local agencies which use the monitoring data to ensure that PM in the air is at respectable levels that protect public health and the environment.

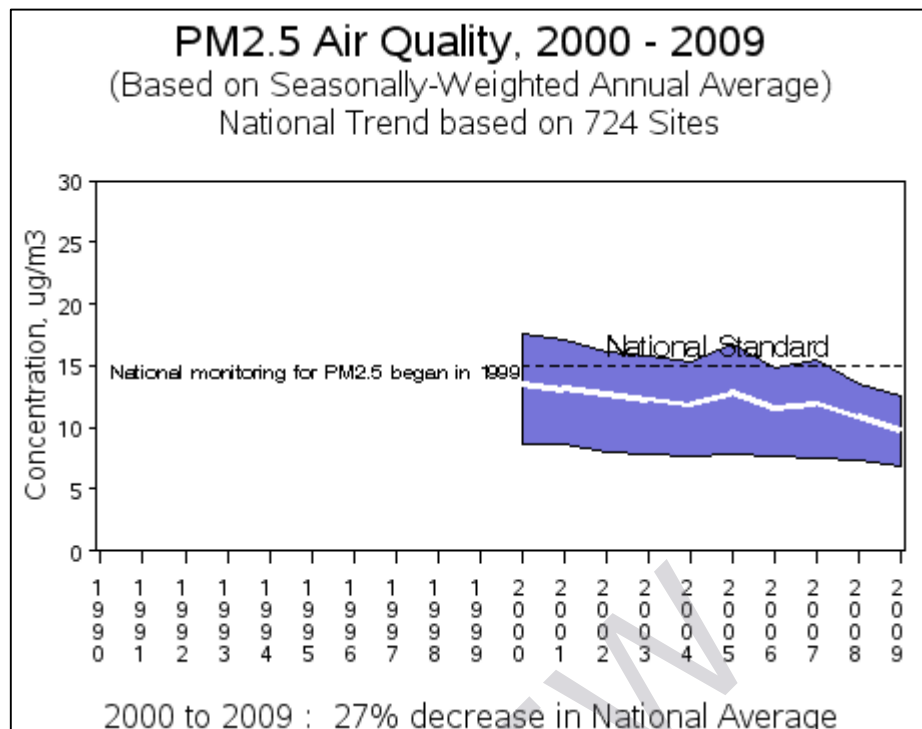


Figure 1.1: National PM_{2.5} Air Quality, 2000-2009

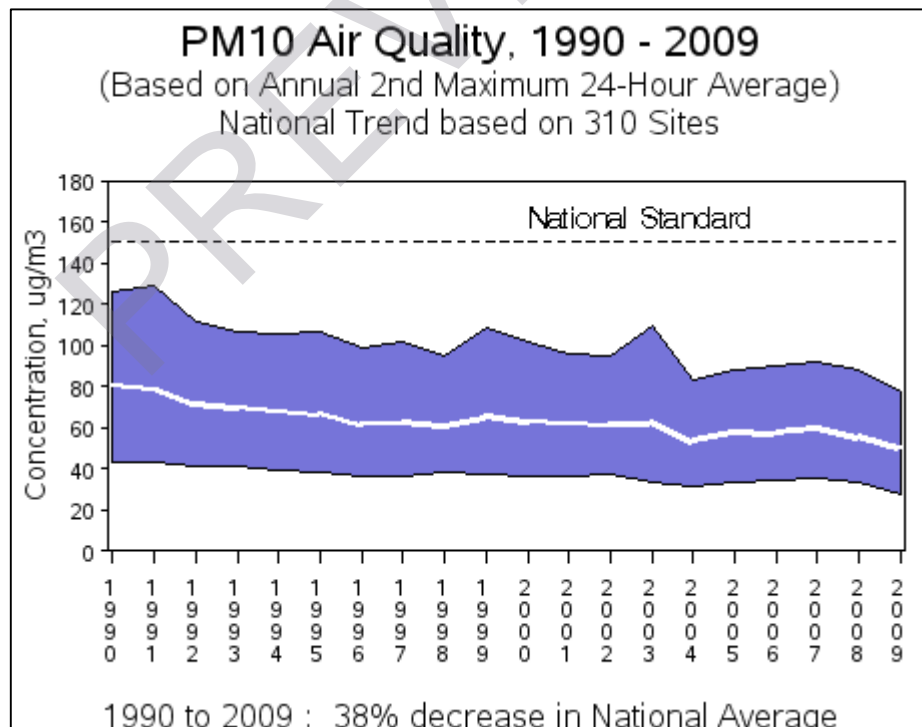


Figure 1.2: National PM₁₀ Air Quality, 1990-2009

In the El Paso region, PM levels have decreased as well with regard to PM₁₀ and PM_{2.5} since 2003. Figure 1.3 is a plot of the annual 2nd maximum 24-hour average PM₁₀ concentrations for El Paso over the span of about 20 years. Figure 1.4 is a plot of the weighted annual mean concentrations for PM_{2.5} for El Paso since 1999, which is when the air quality monitoring program began for PM_{2.5}. The weighted annual mean is the weighted arithmetic mean of 24-hour values for the year and compensates for scheduled sampling that did not occur.

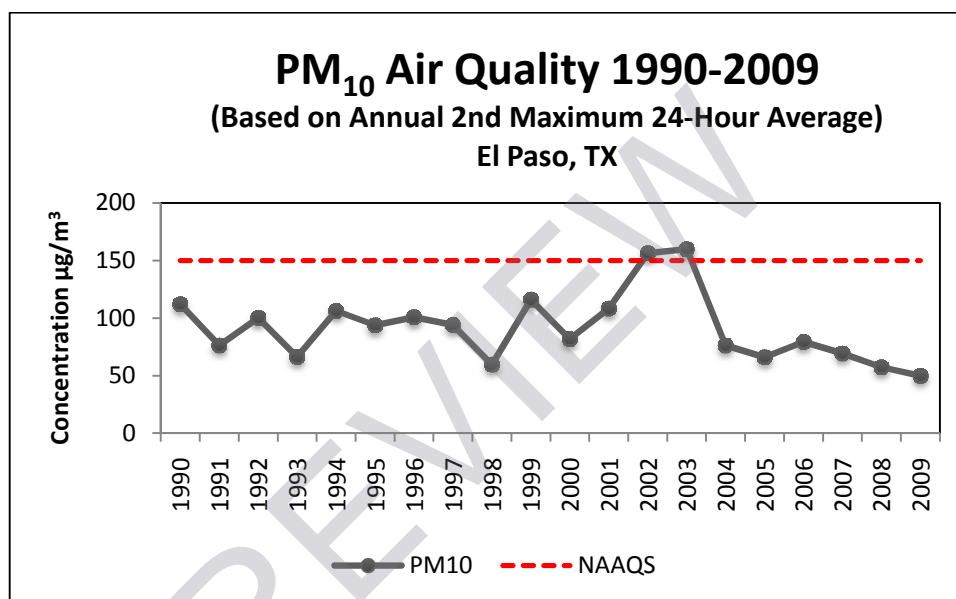


Figure 1.3: El Paso PM₁₀ Air Quality; 1990-2009

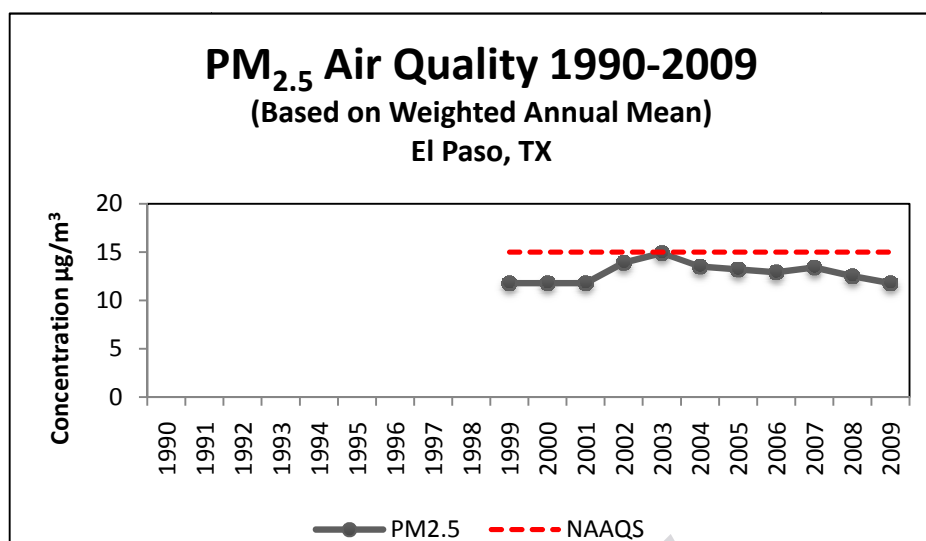


Figure 1.4: El Paso PM_{2.5} Air Quality; 1990-2009

It should be noted that the variation in weighted annual mean PM_{2.5} concentration are not as pronounced as the variation in annual 2nd maximum 24-hour average PM₁₀ concentrations. This may be due to either increases or decreases in severe meteorological conditions which predominantly affect PM₁₀ concentrations.

Based on commercial and industrial activities in the area, coupled with chronic PM₁₀ “non-attainment” status, El Paso can only benefit from any research studies which aim to characterize indoor and outdoor PM in the region. Basic understating of any diurnal variations that exist in PM in El Paso may help investigators to understand when and why PM may be troublesome. Also, any insight into the elemental constitution of PM in the region may also help investigators pinpoint possible sources of PM which may possibly contribute to the local PM pollution problem.

1.2. Research Objective

This study was designed to monitor indoor and outdoor air quality by randomly selecting 12 houses from a cohort constructed for the use in an epidemiological study in the area which would be representative of different strata. It is part of a National Institute of Health (NIH) sponsored project to study the impacts of PM pollution on asthmatic children in El Paso. The scope of the NIH project is an attempt to characterize the time-resolved indoor and outdoor (I/O) PM, nitrogen dioxide (NO₂), and

ozone concentrations in spatially selected Latino households throughout the El Paso region while this project aimed to define the diurnal pattern of I/O PM mass concentrations in two major fractions: coarse PM (PM_{10} , PM with aerodynamic diameters of less than 10 μm) and fine PM ($PM_{2.5}$, PM with aerodynamic diameters of less than 2.5 μm). The chemical composition of PM was of interest in addition to the mass characteristics. Energy dispersive x-ray fluorescence (EDXRF), a preferred method for the analysis of trace elements in PM collected on filters, was used to characterize the chemical composition of $PM_{2.5}$. In addition, the black carbon (BC) content of $PM_{2.5}$ was investigated as well.

PREVIEW

2. Particulate Matter and Related Health Effects

2.1. Particulate Matter

Particulate matter is a complex mixture of extremely small particles and liquid droplets that vary in size, shape, surface area, solubility, chemical composition and origin. PM is composed of a number of components, including acids (such as nitrates and sulfates), organic chemicals, metals, and soil or dust particles. The size of particles is directly linked to their potential for causing health problems. Once particles smaller than 10 microns are inhaled, these particles affect the heart and lungs and cause serious health effects. Through inertial impaction, coarse PM is generally removed in the upper respiratory tract, while fine PM has the ability to penetrate much further into the lower respiratory tract (Kennedy, 2007).

The Environmental Protection Agency (EPA) groups PM into two categories:

- **"Inhalable coarse particles,"** such as those found near roadways and dusty industries, have aerodynamic diameters larger than 2.5 μm and smaller than 10 μm in diameter. Usually comprised of suspended or re-suspended dust from industrial processes, agriculture, construction, road traffic, plant pollen and other natural sources.
- **"Fine particles,"** such as those found in smoke and haze, have an aerodynamic diameter less than or equal to 2.5 μm . Fine particles are responsible for reduced visibility in certain parts of the United States. These particles can be directly emitted from combustion sources such as automobiles, wood burning, or they can form when gases emitted from power plants, cement plants, or steel mills react in the air. This mode of particles also consists of transformation products such as sulfates and nitrates which are formed of primary sulfur and nitrogen oxide emissions along with secondary organic aerosols from volatile organic compounds (VOC).

Fine-mode and coarse-mode particles differ not only in size and morphology (e.g., smooth droplets vs. rough solid particles) but also in formation mechanisms and sources; along with differences in chemical, physical, and biological properties. They also differ in terms of dosimetry (deposition in the respiratory system), toxicity, and health effects as observed by epidemiologic studies. PM transport generally occurs through convection, diffusion and, for particles larger than $1\mu\text{m}$, diffusion (Kennedy, 2007)

Fine particles are a major cause of reduced visibility in parts of the United States. The EPA regulates inhalable particles (fine and coarse) while particles larger than $10\mu\text{m}$ (sand and large dust) are not regulated by EPA. The Clean Air Act requires the EPA to set National Ambient Air Quality Standards (NAAQS) for six criteria pollutants, of which PM is one of these. The remaining criteria pollutants are ozone, carbon monoxide, nitrogen dioxides (NO_2), sulfur dioxide and lead. The purpose of regulation for the aforementioned criteria pollutants is that epidemiologic studies have attributed daily exposure to one or more of the criteria pollutants with an increase in lower respiratory tract effects, exacerbation of asthma and increased daily mortality (Pope, 2000). The Clean Air Act established two types of national air quality standards for PM pollution. Primary standards set limits to protect public health, including the health of "sensitive" populations such as asthmatics, children, and the elderly. Secondary standards set limits to protect public welfare, including protection against visibility impairment, damage to animals, crops, vegetation, and buildings. The air quality standards for PM in 2006 were revised by the EPA to tighten the 24-hour fine PM standard from the level of $65.0\mu\text{g}/\text{m}^3$ to $35.0\mu\text{g}/\text{m}^3$, and retain the current annual fine PM standard at $15\mu\text{g}/\text{m}^3$. The EPA decided to retain the existing 24-hour PM_{10} standard of $150\mu\text{g}/\text{m}^3$ and revoked the annual PM_{10} standard, because available evidence does not suggest a link between long-term exposure to PM_{10} and health problems.