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Analysis of the Fine Fraction of Particulate Matter in Fugitive Dust

Final Report

**For
Western Governors' Association
Western Regional Air Partnership (WRAP)**

MRI Project No. 110397

October 12, 2005

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**For
Western Governors' Association
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October 12, 2005

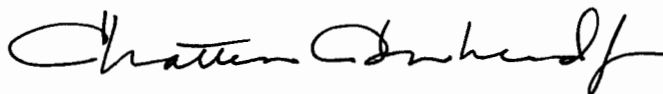
Preface

Midwest Research Institute (MRI) is pleased to submit this project report to the Western Governors' Association, Western Regional Air Partnership, to describe an *Analysis of the Fine Fraction of Particulate Matter in Fugitive Dust*. This report was prepared by Dr. Chatten Cowherd, who has served as Program Manager for the project. He was assisted by John Donaldson, who was responsible for processing of data from the wind tunnel tests of sampler performance. Dr. Greg Muleski served as test advisor for this project. Mary Ann Grelinger served as quality assurance coordinator.

This report contains revisions in response to peer review comments on the draft version dated August 17, 2005.

If you have any technical questions on this report, please call Dr. Cowherd at 816 753 7600, ext. 1586.

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October 12, 2005

Executive Summary

The WRAP Dust Emissions Joint Forum (DEJF) is engaged in gathering and improving data pertaining to the $PM_{2.5}$ and PM_{10} components of fugitive dust emissions. Most of the $PM_{2.5}$ emission factors in EPA's AP-42 guidance for fugitive dust sources were determined by using high-volume samplers, each fitted with a cyclone precollector and cascade impactor. Typically, AP-42 recommends that $PM_{2.5}$ emission factors be calculated by using PM_{10} emission factor equations along with $PM_{2.5}/PM_{10}$ ratios that have been published by EPA in AP-42 based largely on data from the high-volume cyclone/cascade impactor system.

Beginning with the introduction of the cyclone/impactor method, it was realized particle bounce from the cascade impactor stages to the backup filter may have resulted in inflated $PM_{2.5}$ concentrations, even though steps were taken to minimize particle bounce. This led to an EPA-funded field study in the late 1990s to gather comparative particle sizing data in dust plumes downwind of paved and unpaved roads around the country. The test results indicated that Federal Reference Method (FRM) dichotomous samplers produced consistently lower $PM_{2.5}/PM_{10}$ ratios than with the cyclone/impactor system. As a result, the decision was made that the true ratios would best be represented by an averaging of the cyclone/impactor data with the dichotomous sampler data.

Based on the results of the EPA-funded field program, modifications were made to the appropriate sections of AP-42 for dust emissions from paved and unpaved roads. The $PM_{2.5}/PM_{10}$ ratio for paved roads was reduced from 0.46 to 0.25 (realizing that much of the fine particle component was associated with vehicle exhaust), and the ratio for unpaved roads was reduced from 0.26 to 0.15. Subsequent to these modifications to the $PM_{2.5}/PM_{10}$ ratios in AP-42, additional evidence has been compiled in support of further reduction to the ratios. For example, although AP-42-based $PM_{2.5}$ emission estimates seem to show about 20 percent of the PM_{10} fugitive dust mass is in the fine fraction, ambient air monitoring data suggest that it may be on the order of 10 percent.

This led DEJF to fund the subject controlled study to determine if the $PM_{2.5}$ to PM_{10} ratio measured by the cyclone/impactor system has a measurement bias as compared to FRM monitors, and if so, what the fine fraction ratio should be for fugitive dust sources. For this purpose, an exposure chamber with a recirculating feed was used in conjunction with a fluidization system for generating dust plumes from a variety of western soils and road surface materials. The R&P Model 2000 Partisol sampler was selected as the ground-truthing FRM sampler for PM_{10} and $PM_{2.5}$. This study was performed in two phases.

In the first testing phase of the project, $PM_{2.5}$ measurements using the high-volume cascade impactors were compared to simultaneous measurements obtained using EPA reference-method samplers for $PM_{2.5}$. As stated above, these tests were conducted in a flow-through wind tunnel and exposure chamber, where concentration level and uniformity were controlled. The test levels of PM_{10} concentration extended to the range of $5,000 \mu\text{g}/\text{m}^3$, which is representative of uncontrolled dust plume core concentrations.

With the same test setup, a second phase of testing was performed with reference method samplers, for the purpose of measuring PM_{2.5} to PM₁₀ ratios in fugitive dust plumes from different geologic sources in the West. This testing provided needed information on the magnitude and variability of this ratio, especially for source materials that are recognized as problematic with regard to application of mitigative dust control measures. Once again, test PM₁₀ concentrations extended to the range of 5,000 µg/m³.

Based on the 100 wind tunnel tests that were performed in this study, the findings support the following conclusions:

1. PM_{2.5} concentrations measured by the high-volume cyclone/impactor system used to develop AP-42 emission factors for fugitive dust sources have a positive bias by a factor of 2, as compared to the PM_{2.5} concentration measurements from reference-method samplers. (The geometric mean bias is 2.01 and the arithmetic mean bias is 2.15.)
2. The PM_{2.5} bias associated with the cyclone/impactor system as measured under controlled laboratory conditions with dust concentrations held at nearly steady values, closely replicates the bias observed in the prior EPA-funded field study at distributed geographic locations across the country.
3. The PM_{2.5}/PM₁₀ ratios measured in the current study for a variety of western soils show a decrease in magnitude with increasing PM₁₀ concentration. Soils with a nominally spherical shape are observed to have somewhat lower ratios (at given PM₁₀ concentrations) than soils with angular shape. A very similar dependence of PM_{2.5}/PM₁₀ ratio on PM₁₀ concentration was also observed in the prior field study that used dichotomous samplers as FRM devices.
4. The test data from the current study support a PM_{2.5}/PM₁₀ ratio of 0.1 for typical fugitive dust sources. This ratio takes into account the fact that during AP-42 source tests most PM₁₀ sample mass from uncontrolled dust sources is collected at plume core PM₁₀ concentrations exceeding 5,000 µg/m³.
5. The PM_{2.5}/PM₁₀ ratio of 0.1 is also supported by numerous other studies including the prior field study that used dichotomous samplers as reference devices. It is possible that a ratio as low as 0.05 (as was found in the prior field tests of unpaved roads) might be appropriate, but this would require extrapolation of the current test data to higher PM₁₀ concentrations.

The results of this project are needed to ensure development of the most accurate PM_{2.5} and PM₁₀ fugitive dust emissions inventories that are possible for regional haze regulatory purposes. In particular, a reduction in the quantity of dust apportioned to the fine versus coarse size modes will significantly affect models for visibility and long-range transport.

The results will also be helpful in developing accurate emission inventories for PM nonattainment, maintenance, and action plan areas in the WRAP region. Finally, if appropriate, results may be used to seek modifications to the EPA's AP-42 emission

factors to ensure widespread availability of the most recent and accurate scientific information.

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Section 1.

Introduction

The WRAP Dust Emissions Joint Forum (DEJF) is engaged in gathering and improving data pertaining to the $PM_{2.5}$ and PM_{10} components of fugitive dust emissions. Most of the $PM_{2.5}$ emission factors in EPA's AP-42 guidance for fugitive dust sources [1] were determined by using high-volume samplers fitted with a cyclone precollector and cascade impactor. Under the same conditions, the $PM_{2.5}$ mass measured using these cyclone/cascade impactors should be nearly identical to the mass collected using an ambient $PM_{2.5}$ sampler that meets EPA ambient air monitoring requirements.

The MRI cyclone/impactor system (see Figure 1) has been recognized as the standard method to characterize fugitive dust emissions for EPA and to develop predictive emission factor equations for AP-42. A sample of particles is drawn at a flow rate of $34 \text{ m}^3/\text{hr}$ (20 acfm) into a directional probe tip with an inlet velocity of 2.2 m/s (5 mph) and then through a cyclonic preseparator to remove particles larger than $15 \mu\text{m}$ in diameter. The PM_{10} in the exit airflow from the cyclone is channeled through multiple impactor stages before being collected on a back-up filter, from which time-integrated PM_{10} and $PM_{2.5}$ concentrations can be calculated.

Historically, data from the MRI high-volume cyclone/impactor system have provided the basis for $PM_{2.5}/PM_{10}$ ratios that have been published by EPA in AP-42 for various categories of fugitive dust sources. The advantage of the high volume cyclone/impactor systems relates to its directional sampling capability and higher analytical sensitivity for particle size measurement. However, particle bounce from the cascade impactor stages to the backup filter may have resulted in higher $PM_{2.5}$ measurements than the actual $PM_{2.5}$ concentrations, even though steps were taken by MRI to minimize particle bounce. Those steps included halving the flow rate, greasing the substrates prior to use, and limiting the sampling time so that substrates are not overloaded with collected particle mass.

Some comparative data collected in dust plumes downwind of paved and unpaved roads in the late 1990s indicate that $PM_{2.5}/PM_{10}$ ratios determined with other particle sizing systems produced consistently lower ratios than with the cyclone/impactor system. This comparative particle sizing work was performed by MRI under contract to EPA [2]. As a result, modifications to reduce the ratios were made by EPA to the appropriate sections of AP-42.

Even with the adjusted particle size ratios imbedded in the current version of AP-42, there is still evidence of a potential bias due to residual measurement error associated with particle bounce in the high-volume cascade impactor system. This possible measurement error may help to explain why the AP-42-based $PM_{2.5}$ emission estimates seem to show about 20 percent of the fugitive dust mass is in the fine fraction, while ambient air monitoring data suggest that it may be on the order of 10 percent [3].

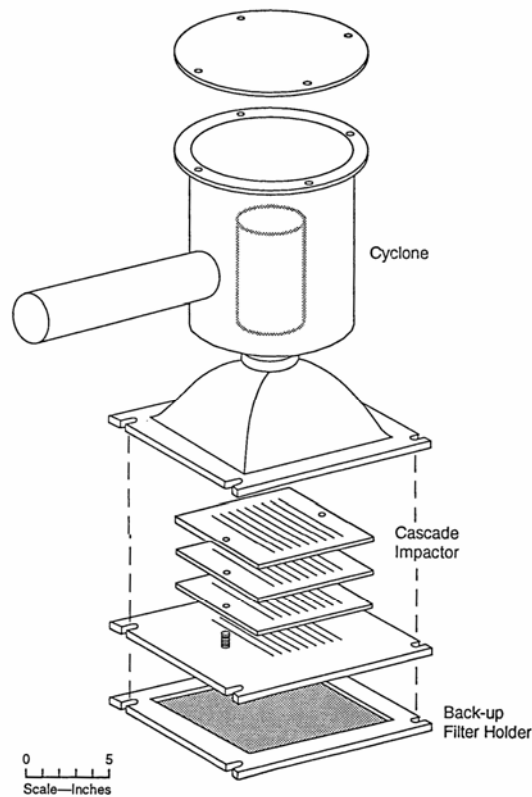


Figure 1. Cyclone/Impactor

In the first testing phase of this project, $PM_{2.5}$ measurements using the high-volume cascade impactors were compared to simultaneous measurements obtained using EPA reference-method samplers for $PM_{2.5}$. These tests were conducted in a flow-through wind tunnel and exposure chamber, where concentration level and uniformity were controlled. The results of the tests provide the basis for quantifying more effectively any sampling bias associated with the cascade impactor system.

With the same test setup, a second phase of testing was performed with reference method samplers, for the purpose of measuring $PM_{2.5}$ to PM_{10} ratios for fugitive dust from different geologic sources in the West. This testing provided needed information on the magnitude and variability of this ratio, especially for source materials that are recognized as problematic with regard to application of mitigative dust control measures.

The results of this project are needed to ensure the most accurate $PM_{2.5}$ and PM_{10} fugitive dust emissions inventories that are possible for regional haze regulatory purposes, given the available resources and the significant contribution of fugitive dust to visibility impairment. In particular, the results of this project may affect the quantity of dust apportioned to the fine versus coarse size modes, which have significantly different effects on visibility and long-range transport potentials. The results will also be helpful

in developing accurate emission inventories for PM nonattainment, maintenance, and action plan areas in the WRAP region. Finally, if appropriate, results may be used to seek modifications to the EPA's AP-42 emission factors to ensure widespread availability of the most recent and accurate scientific information.

Section 2.

Test Facility and Air Samplers

2.1 Aerosol Test Facility

The testing was conducted in the Aerosol Test Facility (ATF) in Building 2 at MRI's Deramus Field Station in Grandview, Missouri. This system for generation and control of dust concentrations in an exposure chamber used equipment similar to that used in the recent performance evaluation of the MetOne GT-641 aerosol monitor [4] for use at Owens Lake, except that the MRI ATF is a recirculating flow system rather than a once-through system. The recirculation feature of the ATF promotes the establishment of well-mixed and steady target aerosol concentration levels in exposure section where the PM_{10} and $PM_{2.5}$ monitors are positioned. The ATF also limits the consumption of test dust so that much smaller amounts of starting soil or road dust samples are sufficient for extensive testing.

The MRI test facility consists of a push-through flow system (3 ft by 3 ft cross section) with an exposure chamber at the downstream end of the flow tunnel, as shown in Figure 2. The flow system can be operated at low air speeds (as low as 0.5 m/s), using an electronic motor speed controller. An 18-in diameter air return loop connects the outlet of the exposure chamber with the inlet to the blower, as shown in Figure 3. In this study, the tunnel was operated at a speed of 1 m/s (2 mph). The wind speed was highly uniform in the core of the flow (into which the inlets to the air samplers were placed). When a Kestral vane anemometer was used to measure the wind speed at the center points of nine equal areas of the tunnel cross-section, all measured point values were within about 10 percent of the average value.

A fluidized bed injects dust through a vertical copper tube (1-in internal diameter) into the return line to the blower inlet that feeds the flow tunnel (Figure 4). The fluidized bed aerosolizes fine dust from samples of loose, dry soil from a cylindrical glass container (Puff cartridge) that is 10.2 cm tall and 5.5 cm internal diameter. A glass fiber filter is sandwiched between screens at the bottom of the chamber, and a conical section is positioned on top of the screens, directing the dust material to a circular area in the center of the screen. The fluidization rate is controlled by the upward airflow through the bed. The injection point to the tunnel flow is just downstream of the center of an 8-in orifice in the 18-in return line. The turbulent effluent from the orifice aids in mixing the injected dust into the tunnel airflow.

The PM_{10} dust concentration in the exposure chamber is continuously tracked with a TSI Model 8520 DustTrak monitor that draws air from the centerline position in the exposure section. The DustTrak sampling line was standard equipment supplied by TSI. Based on a 9-point equal-area profile transecting the core of the flow, the PM_{10} concentration at each point is within 10 percent of the average value. A second DustTrak monitor (with its own sampling line) tracks the $PM_{2.5}$ concentration at the centerline.



Figure 2. MRI Flow Tunnel/Test Chamber



Figure 3. Return Line to Wind Tunnel Blower



Figure 4. Fluidization System for Injecting Fine Fraction Dust Into the Flow System

2.2 Air Samplers

Table 1 lists the air sampling equipment that was used in the two testing phases. The EPA reference method samplers [5] consisted of Rupprecht & Patashnick Model 2000 Partisols for $PM_{2.5}$ and PM_{10} measurements. The approximately 3 ft by 3 ft working cross section of the exposure chamber (Figure 5) and the two sampling stations within the chamber provided adequate space for the inlets of two cyclone/impactors, two Partisol samplers, and two DustTRAKs, depending on the requirements of the test being performed.

The MRI cyclone/impactor system, shown earlier in Figure 1, consists of a Sierra Model 230 cyclone with a directional intake and a Sierro Model 230CP three-stage slotted-type cascade impactor. When operated at 20 acfm, the cyclone has an aerodynamic cut point of 15 microns, and the cascade impactor stages have cut points of 10.2, 4.2, and 2.1 microns, respectively [2]. The slotted glass fiber substrates were coated with a thin film of grease before tare weighing (using the same procedure as in the past), to mitigate against particle bounce problems. For the same reason, care was taken not to overload the substrates with collected dust. Pretest estimation of the dust loading on the impactor stages must take into account that the PM_{10} sample is distributed over three separate impactor stages plus the back-up filter when the cyclone is operated at 20 acfm. Typically, the cyclone/impactor system has been utilized to generate $PM_{2.5}/PM_{10}$ ratios that can be used in combination with PM_{10} plume profiles to generate $PM_{2.5}$ emission factors.

Table 1. Air Samplers Used for Testing

Phase	No. in use	Sampler	Manufacturer/model	Flow rate	Comments
1	2	Cyclone preseparators	Sierra Model 230 CP	20 acfm	Third stage has D ₅₀ cut of 2.1 µmA, which MRI has used surrogate for PM _{2.5} .
	2	Multistage impactor	Sierra Model 230	20 acfm	First three stages used.
	2	Partisol (FRM)	R&P Model 2000	16.7 alpm	Device uses WINS impactor to provide PM _{2.5} cut point.
	2	DustTrak	TSI Model 8520	5 alpm	Used to continuously track concentration level and uniformity within exposure chamber
2	2	Partisol (FRM)	R&P Model 2000	16.7 alpm	Reference sampler uses WINS impactor to provide PM _{2.5} cut point.
	2	Partisol (FRM)	R&P Model 2000	16.7 alpm	Reference sampler uses dichot inlet, which has D ₅₀ cut point of 10 µmA. Sampler is fitted with R&P part to bypass WINS impactor.
	2	DustTRAK	TSI Model 8520	5 alpm	Used to continuously track concentration level and uniformity within exposure chamber.



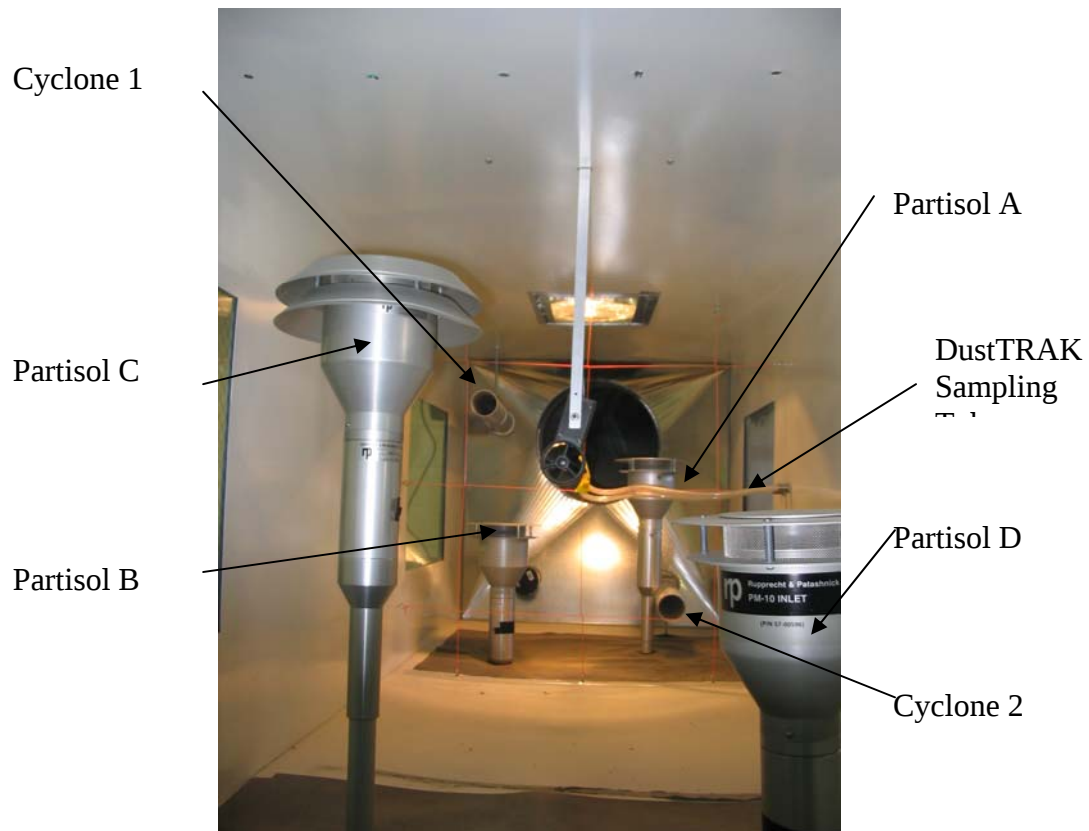
**Figure 5. Test Chamber Interior With Flow Straightener
(Viewed from downstream end of exposure section)**

EPA reference-method R&P Partisol analyzers were used to measure the concentrations of PM_{10} and $PM_{2.5}$ in the dust exposure chamber. The chamber was large enough to accommodate four Partisol inlets and two cyclone/impactor inlets (see Figures 6 and 7). By locating sampler inlets in opposing quadrants at the two sampling stations, any interference effects were negligible. The low flow in the exposure section of the wind tunnel minimized the propagation of turbulent wakes created by the sampling inlets. In Figure 6, the large opening at the rear of the tunnel is the inlet to the 18-in diameter return line. The bodies of the air samplers were placed underneath the exposure chamber (Figure 8).

It should be noted that because the sampling systems (except for the DustTrak monitors) involved collection of PM on filters (and impactor substrates), the measured concentrations represented averages over each test period. With regard to the filter medium, a fibrous rather than a membrane-type filter was used for better retention of dust particles. Whatman EPM 2000 glass fiber filter and impactor substrate media were used throughout the testing.

Prior to testing, required filter and impactor substrate media were prepared for air sampling. A temperature- and humidity-controlled gravimetrics laboratory was used for obtaining tare and final weights of filters and greased impactor substrates. The

temperature and humidity were maintained within the limits recommended by the EPA for filter weighing in association with ambient PM monitoring [6].



**Figure 6. Partisol, Cyclone/Impactor, and DustTrak Inlets in Exposure Section
(as viewed from upstream)**



**Figure 7. View of Partisol, Cyclone, and DustTrak Inlets
Through Observation Window**



Figure 8. Partisol Sampler Bodies Below Exposure Section of Wind Tunnel

2.3 Test Matrix

Table 2 shows the matrix of tests that were specified in the scope of work for this project. At least 10 percent of tests utilized “field” blanks for all particle collection media, as is customary to meet quality control requirements.

Table 2. Preliminary Test Matrix

Phase	Source materials	Concentration levels	Replication	Total No. of tests	Sampling media used
1	3 (Arizona road dust—coarse and fine, Owens Lake surface material)	3 (Low, moderate, high)	3 (triplicates)	27	8x10 filters 4 x 5 substrates 47-mm filters plus > 10% field blanks for all media
2	5 (Representative soils or road surface materials)	3 (low, moderate, high)	3 (triplicates)	45	47-mm filters plus > 10% field blanks

2.3.1 Phase I—AP-42 PM_{2.5} Emission Factor Evaluation

As indicated in Table 2, three dust source materials were tested under Phase I. Owens Dry Lake surface soil was used to provide one dust source material. The other two dust source materials were reference standards referred to as Arizona road dust, coarse and fine fractions. Arizona road dust is ground from Arizona sand using ball mills, elutriation, and blending to achieve reproducible size distributions for a variety of applications including performance testing of automotive air cleaners.

In order to entrain a steady stream of fine particles from the Arizona road dust samples, it was necessary to add a small amount of sand to the dust generation chamber. Because of the fine texture of Arizona road dust (both size fractions consisting entirely of particles smaller than 75 microns), it is impractical to aerosolize these test dusts in any consistent fashion without adding sand to the fluidized bed. The powder tends to clump because of strong interparticle binding forces. If this powder were exposed to the atmosphere, its release as fine particles would require vigorous mechanical contact (e.g., rolling tires) or sandblasting by saltating particles in the size range of 100 microns.

Fixed PM₁₀ concentration levels in the range of 1,000, 2,500, and 5,000 micrograms per cubic meter (each with its naturally occurring PM_{2.5} level) were tested. These PM₁₀ concentration levels were selected as representative of dust plume concentrations under which major particle mass contributions to plume samples occur in emission factor development.

The PM₁₀ concentration level during each 20- to 120-min test was maintained as closely as possible to a predetermined target value. This was accomplished by adjusting the airflow in the dust aerosolization system. Tests at each concentration level were performed in triplicate.

Because filter-based reference-method PM_{2.5} monitors with relatively low sampling rates were used in the study, care was taken so that the test periods were sufficient so that minimum quantifiable mass is collected on the PM_{2.5} (47-mm) filters. However, certain test soils did not have sufficient dust emission potential to achieve the higher concentrations using the dust aerosolization system.

2.3.2 Phase II—PM_{2.5} to PM₁₀ Ratios for Different Soil Samples

In Phase II, reference-method Partisols measured both PM₁₀ and PM_{2.5} concentrations. Once again, fixed PM₁₀ concentrations (each with its naturally occurring PM_{2.5} level) were tested.

In accordance with the test protocol and Quality Assurance Project Plan (QAPP), MRI measured the ratio of PM_{2.5} to PM₁₀ for fugitive dust from different geologic soil types. A total of seven source materials were tested, which was two more than originally specified in the scope of work. Test results included the calculation of the average PM_{2.5}

concentration and the collocated PM₁₀ concentration. It was intended that any variation in PM_{2.5}/PM₁₀ ratio be evaluated as a function of the test soil properties (for example, position in soil texture triangle).

As stated under Section 3, MRI worked with the DEJF in selecting the individual soil types to be tested. These soils were provided by WRAP members. The types and locations of test soils for Phase II are listed in Table 3.

Table 3. Test Soils for Phase II

Code	State	Location	Type of material
AK	Alaska	MAT-SU Knik River Bed	Sediments
AZal	Arizona	Phoenix Area	Alluvial Channel
AZag	Arizona	Phoenix Area	Agricultural Soil
NMr	New Mexico	Las Cruces Landfill	Road Dust
NMs	New Mexico	Radium Springs	Grazing Soil
SS	California	Salton Sea	Shoreline Soils
WY	Wyoming	Thunder Basin Mine	Barrow Pit for Access Road Surface Material

Section 3.

Pretest Activities

3.1 Test Protocol and Quality Assurance Project Plan

Prior to the testing, MRI submitted to the DEJF a combined test protocol/QAPP [7] according to EPA standards. The test protocol included design of the dust generation system and exposure chamber, specification of monitoring equipment, procedures for sampling, testing, and data analysis, and other procedures or design information essential to the completion of the project. Because the results of this project may be considered in revisions to EPA-published emission factors, adherence to EPA quality assurance documentation procedures was an important part of this project. The test protocol and quality assurance plans complied with EPA standards and will specify all procedures to be followed in collecting, recovering, transferring, and analyzing samples.

These plans were prepared in compliance with EPA document QA/R-5 (***EPA Requirements for QA Project Plans***) as well as the guidance document QA/G-5 (***Guidance for Quality Assurance Project Plans***). To aid the EPA in review of the plans, the documents were structured to mimic the required groups and elements of QA/G-5.

The outline for the combined plan is given as Table 4. Note that the labels in parentheses (such as “A3”) after each section refer to the group/element labels in QA/G-5 and are included to facilitate review.

Table 4. Annotated Outline for Combined Test Protocol/QAPP

Preface

Distribution of QAPP (A3)—*includes list of persons who have received the QAPP, signature approval page, and revision history of the document*

Figures—*includes Test Facility Schematic, Sampler Locations, Amendment Record for QAPP, Corrective Action Report Form*

Tables—*includes Data Quality Objectives, Test Design, Test Schedule, Critical and Non-Critical Measurements, and Quality Control Procedures for Sampling Media, Quality Control Procedures for Sampling Equipment, Quality Assurance for Miscellaneous Equipment*

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3.2 Test Dust Sample Collection and Handling

Soil and road surface material samples for the subject testing were collected from a variety of locations. Some existing samples were available from prior studies, such as those involving reference Arizona road dust and surface materials from Owens Lake. However, the remaining samples came from other sources. MRI worked with the DEJF to determine the most appropriate types of samples to be used in this study. Samples that were not available from other studies were collected specifically for this study. Again, MRI worked with the DEJF to determine the best locations and surfaces from which to collect such samples.

It was recommended that test materials represent troublesome western soils that were observed to be high dust emitters. It was pointed out that soils with high dustiness potential tend to be subject to frequent mechanical disturbances by agricultural operations, construction operations, or other operations involving vehicle travel across exposed areas. Therefore, only samples of unconsolidated (uncrusted and uncompacted) soils or road aggregate materials are likely to have high dustiness potential.

Two 5-gal containers of each dusty soil or aggregate material were requested to sustain a series of tests to determine $PM_{2.5}/PM_{10}$ ratios under a variety of test conditions. It was stated that only loose, dry (less than 1% moisture) soils should be collected. For soils, the depth of sampling should not exceed 2 in (5 cm). All particles greater than 4-mesh (0.47 cm) are considered nonerodible, and so it was instructed that they be removed from the sample by dry sieving prior to shipment of the sample to MRI.

Members of the WRAP (e.g., state, local, and tribal air quality professionals) collected and shipped the samples to MRI. A soil screening device, soil collection procedures, and associated data forms were provided by MRI as specified in the test protocol and QAPP. An example procedure is supplied in Figure 9.

The properties of the test soils provided for this study are summarized in Table 5. This includes the moisture content and the dry silt content (fraction passing a 200-mesh screen upon dry sieving) using the procedure specified in AP-42 [1]. The dry silt content has been used as a surrogate for dustiness in the AP-42 emission factor equations for fugitive dust.

A standard agricultural soil analysis [8] was performed on well-mixed subsamples of each test material. When plotted on the soil texture triangle, most of the samples fell into a relatively small region, as shown in Figure 10. This texture region is characterized by high wet silt content and moderate sand content.

Two 5-gal containers of a dusty soil or aggregate material are needed for each test material to sustain a series of tests to determine $PM_{2.5}/PM_{10}$ ratios under a variety of test conditions. Only loose, dry (less than 1% moisture) soils should be collected. For soils, the depth of sampling should not exceed 2 in (5 cm). All particles greater than 4-mesh (0.47 cm) are considered nonerodible and should be removed from the sample by dry sieving prior to shipment of the sample to MRI. MRI will provide a special screen for this purpose.

Collection of Surface Soil Samples

The following steps outline the procedure to collect a two 5-gal soil samples of pulverized soil from an open area:

1. Define and document the area of interest:
 - a. Size of sampled field
 - b. Lat/long or UTM coordinates of approximate field centroid
 - c. Land use and recent disturbance history
 - d. Recent and current weather
 - e. Observed field surface texture/appearance
 - f. USDA surface soil classification
2. Collect surface soil sample
 - a. Collect two 5-gal containers of soil by compositing approximately equal amounts from a minimum of 10 locations in the same agricultural field
 - i. Use a straight-edge shovel to collect each incremental sample
 - ii. Sample to a loose soil depth not exceeding 2 in (5 cm)
 - iii. Empty each incremental sample from the shovel into the screen that covers a 5-gal plastic bucket
 - iv. Measure the approximate volume of coarse material that is screened from the total composite sample
 - b. Seal the bucket lid using tape and label the bucket with a permanent marker
 - i. Field name
 - ii. Date/time of collection
 - iii. Sample number corresponding to the data sheet
 - iv. Name of person collecting the sample
 - c. Document the sample collection on a data sheet
 - i. Location of sampled areas (e.g., GPS coordinates, sketch of field locations)
 - ii. Approximate area (sq ft) from which each incremental sample is taken

Figure 9. Sampling Procedure for Collecting Test Soil or Road Surface Material

Collection of Surface Samples of Unpaved Road Aggregate

The following steps outline the procedure to collect two 5-gal samples of surface aggregate from an unpaved road.

3. Define and document the area of interest:
 - a. Length and width of sampled road segment
 - b. Lat/long or UTM coordinates of approximate road segment centroid
 - c. Road use and recent maintenance history
 - d. Recent and current weather
 - e. Observed road surface texture/appearance
 - f. Road aggregate type and origin
4. Collect surface soil sample
 - a. Collect 5 gal of loose surface material by compositing equal amounts of uncrusted soil from a minimum of 10 edge-to-edge steps across in the traveled area of the road
 - i. Use a whisk broom and dust pan
 - ii. Sample to a 1 cm depth or to the hardpan
 - iii. Empty each incremental sample from the dust pan into the screen that covers a 5-gal plastic bucket
 - b. Seal the bucket lid using tape and label the bucket with a permanent marker
 - i. Road name
 - ii. Date/time of collection
 - iii. Sample number corresponding to the data sheet
 - iv. Name of person collecting the sample
 - c. Document the sample collection on a data sheet
 - i. Location of sampled areas (e.g., GPS coordinates, sketch of field locations)
 - ii. Approximate area (sq ft) from which each incremental sample is taken

Figure 9. Sampling Procedure for Collecting Test Soil or Road Surface Material (Concluded)

Table 5. Properties of Test Soils

Code	State	Location	Type of material	Moisture content (%)	Dry silt content (%)	Dry Silt rank
TF	Arizona	—	Standard Test Dust—Fine	—	—	—
TC	Arizona	—	Standard Test Dust—Coarse	0.60	87.6	1
AK	Alaska	MAT-SU Knik River Bed	Sediments	0.80	8.69	6
AZal	Arizona	Phoenix Area	Alluvial Channel	0.33	17.3	3
AZag	Arizona	Phoenix Area	Agricultural Soil	1.06	21.6	2
NMr	New Mexico	Las Cruces Landfill	Road Dust	1.27	12.2	4
NMs	New Mexico	Radium Springs	Grazing Soil	0.47	10.9	5
OW	California	Owens Dry Lake	Lakebed Soil	0.27	3.14	9
SS	California	Salton Sea	Shoreline Soils	5.46*	3.63	8
WY	Wyoming	Thunder Basin Mine	Barrow Pit for Access Road Surface Material	2.47*	6.83	7

* Required drying prior to testing.

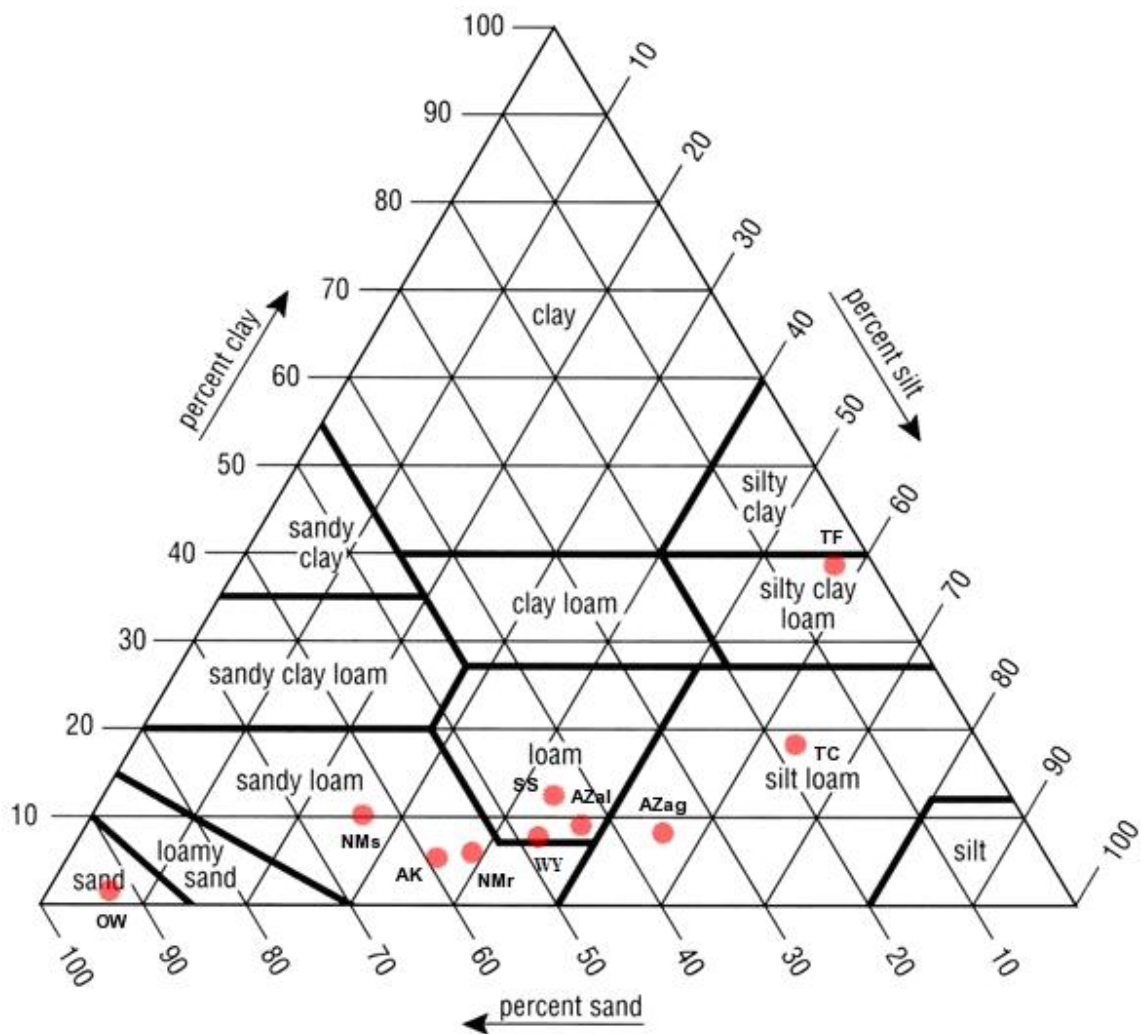


Figure 10. Agricultural Texture of Test Soils

Section 4.

Test Results

The list of tests that were performed is shown in Tables 6 and 7 for Phase I and Phase II tests, respectively. The Phase I tests were performed in March and April of 2005. The Phase II tests were performed in June through August of 2005. Although the scope of work specified five test materials for Phase II, seven materials were tested (with the last two at one target concentration level). Although it was desired to test the last two materials in the PM_{10} concentration range of $5,000 \mu\text{g}/\text{m}^3$, the perimeter soil from the Salton Sea was found to have a low dustiness index, so that only a concentration in the range of $1,000 \mu\text{g}/\text{m}^3$ could be achieved.

A total of 100 individual tests were performed, including 17 blank runs (for quality assurance purposes). The raw and intermediate test data are summarized in the tables presented in Appendix B of this report. Static blank filters were collected by loading filters into samplers with inlets positioned in a still air environment. Dynamic blank filters were collected by loading filters into samplers with inlets positioned in a moving air environment. In neither case was air drawn through the samplers.

4.1 Phase I

As noted above, the testing in Phase I was directed to determining whether a bias existed in the $PM_{2.5}$ concentrations historically measured by the high-volume cyclone/impactor system in fugitive dust plumes. Data from the cyclone/impactor system were balanced against dichotomous sampler results [2] in developing particle size multipliers for EPA's fugitive dust emission factor equations currently published in AP-42.

In Phase I, the flow through the dust generation chamber was varied as needed to maintain the target PM_{10} concentration in the exposure section of the wind tunnel, as measured by the DustTrak monitor with an inlet line extended to the centerline position. An auger was used periodically to add test soil to the chamber, once the supply was depleted to less than half of the original amount added at the beginning of a test.

Prior to each test series, the wind tunnel was purged with clean air that was exhausted outside the building, to remove any loose accumulation of dust on the interior surfaces of the flow system. In addition, all sampler flow rates were checked, as well as the impactors in the $PM_{2.5}$ Partisol samplers and the $PM_{2.5}$ DustTrak monitors.

The results of the Phase I testing are summarized in Figure 11, which shows that the $PM_{2.5}$ concentrations measured by the cyclone/impactor system are consistently biased by a factor of about 2 relative the $PM_{2.5}$ concentrations measured by the Partisol samplers. (The geometric mean bias is 2.01, and the arithmetic mean bias is 2.15.) This bias is also reflected in the $PM_{2.5}/PM_{10}$ ratios measured by the cyclone/impactor system in relation to the reference samplers. The one exception occurs for the Arizona course test dust at the high concentration, for which the ratio is nearly a factor of 4.

Table 6. List of Tests Performed in Phase I

Date	Run	Test dust	Duration of test (min)	Actual PM ₁₀ concentration (µg/m ³) ^a
3/2/2005	1	AZ Fine Dust	60	2220
3/24/2005	2	AZ Fine Dust	60	1060
3/29/2005	3	AZ Fine Dust	60	1080
3/29/2005	4,5	AZ Fine Dust	120	1190, 1120
3/29/2005	6	Dynamic Blank Run	–	–
3/30/2005	7	AZ Fine Dust	120	1020
3/30/2005	8,9,10	AZ Fine Dust	15	5001, 5600, 4990
3/30/2005	11	Static Blank Run	–	–
4/1/2005	12,13,14	AZ Coarse Dust	15	4580, 3860, 4850
4/1/2005	15	Static Blank Run	–	–
4/6/2005	16,17,18	AZ Coarse Dust	60	2210, 2170, 1950
4/6/2005	19	Dynamic Blank Run	–	–
4/7/2005	20,21,22	AZ Coarse Dust	120	575, 582, 348
4/7/2005	23	Static Blank Run	–	–
4/21/2005	24	Owens Dry Lake Bed	20	3460
4/21/2005	25	Owens Dry Lake Bed	22	2840
4/21/2005	26	Owens Dry Lake Bed	20	3720
4/21/2005	27,28,29	Owens Dry Lake Bed	60	2350
4/21/2005	30	Static Blank Run	–	–
4/22/2005	31,32,33	Owens Dry Lake Bed	120	492, 559, 464
4/22/2005	34	Dynamic Blank Run	–	–

^a PM₁₀ concentrations measured by DustTRAK monitor through run WR-15 and by Partisol samplers for subsequent tests.

Table 7. List of Tests Performed in Phase II

Date	Run	Test dust	Duration of test (min)	Actual PM ₁₀ concentration (µg/m ³)
6/9/2005	35	AZ Ag Field	60	927
6/10/2005	36,37	AZ Ag Field	60	874, 889
6/10/2005	38,39,40	AZ Ag Field	40	2676, 3246, 2473
6/10/2005	41,42,43	AZ Ag Field	20	5197, 4576, 4852
6/10/2005	44	Dynamic Blank Run	—	—
6/14/2005	45	Knik River Sediment	40	783
6/14/2005	46	Knik River Sediment	60	340
6/14/2005	47	Knik River Sediment	120	487
6/14/2005	48,49	Knik River Sediment	20	3658, 4800
6/14/2005	50	Dynamic Blank Run	—	—
6/16/2005	51,52	Knik River Sediment	40	3119, 3254
6/16/2005	53,54	Knik River Sediment	80	1211, 1316
6/16/2005	55	Dynamic Blank Run	—	—
6/17/2005	56,57,58	Las Cruces Landfill Road	20	5807, 5736, 5737
6/17/2005	59,60	Las Cruces Landfill Road	40	2784, 2719
6/17/2005	61	Dynamic Blank Run	—	—
6/22/2005	62	Las Cruces Landfill Road	40	2662
6/22/2005	63,64,65	Las Cruces Landfill Road	80	1050, 1000, 1011
6/22/2005	66	Dynamic Blank Run	—	—
6/23/2005	67	Thunder Basin Barrow Pit	40	2600
6/23/2005	68,69	Thunder Basin Barrow Pit	40	2473, 2180
6/23/2005	70,71	Thunder Basin Barrow Pit	60	1270, 682
6/23/2005	72	Dynamic Blank Run	—	—
6/24/2005	73	Thunder Basin Barrow Pit	60	1113
6/24/2005	74,75,76	Thunder Basin Barrow Pit	120	473, 616, 509
6/24/2005	77	Dynamic Blank Run	—	—
6/30/2005	78,79,80	AZ Alluvial Channel	20	6141, 6755, 6711
6/30/2005	81,82,83	AZ Alluvial Channel	40	3343, 3201, 2907
6/30/2005	84	AZ Alluvial Channel	80	1278
6/30/2005	85	Dynamic Blank Run	—	—
7/1/2005	86,87	AZ Alluvial Channel	80	1088, 1112
7/1/2005	88	Dynamic Blank Run	—	—
8/1/2005	89	Radium Springs	40	2605
8/1/2005	90,91	Radium Springs	40	2346, 2914
8/1/2005	92	Dynamic Blank Run	—	—
8/1/2005	93,94,95	Salton Sea	80	805, 595, 548
8/2/2005	96	Dynamic Blank Run	—	—
8/2/2005	97,98,99	AZ Ag Field	20	5475, 7087, 5603
8/2/2005	100	Dynamic Blank Run	—	—

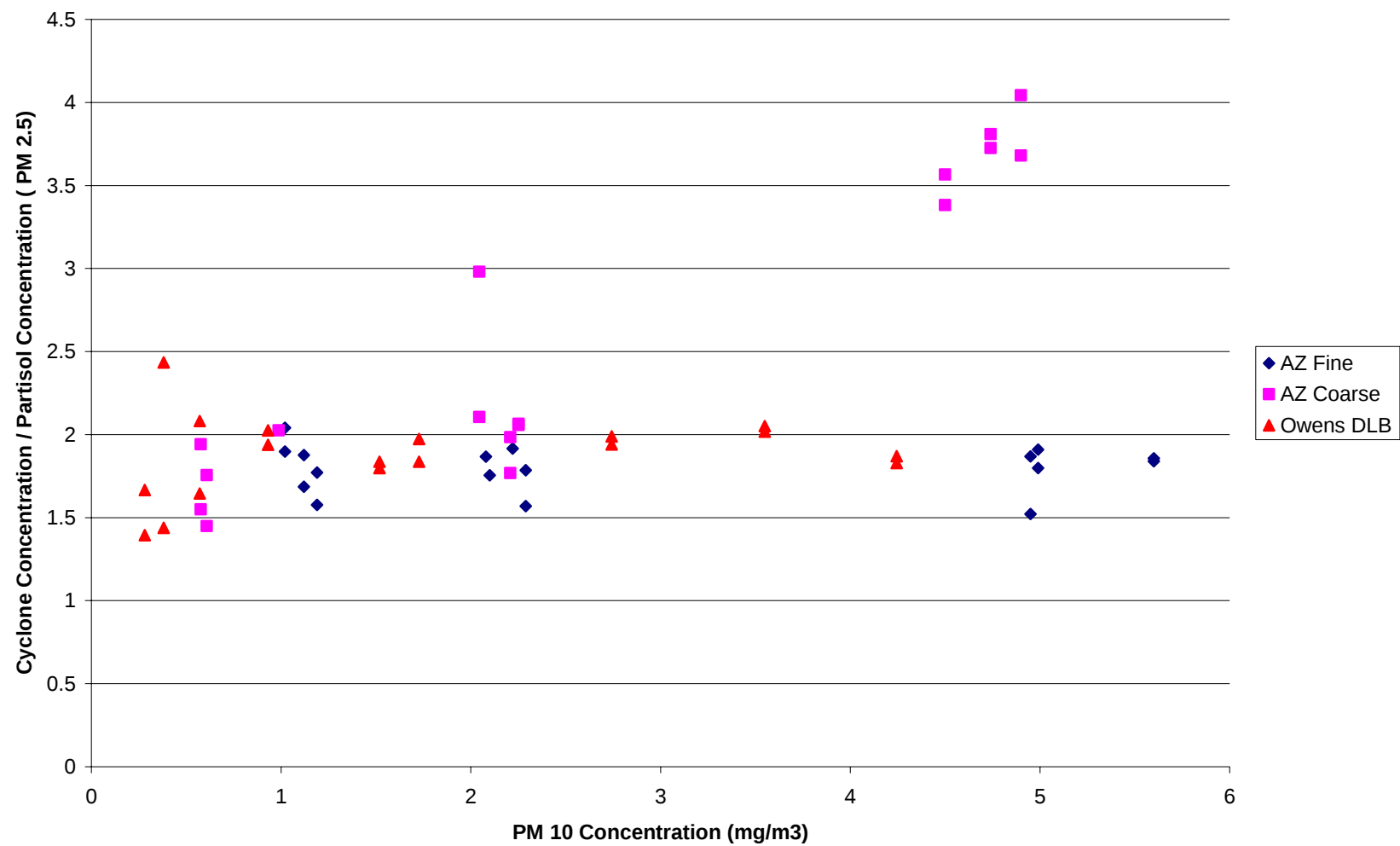


Figure 11. Correlation Between Cyclone/Impactor and Partisol PM_{2.5} Concentrations

This bias was also observed in an EPA-funded field testing program [2] performed on dust emissions from paved and unpaved roads at three geographic locations across the country (Raleigh, North Carolina; Grandview, Missouri; Reno, Nevada). In that study, reference-method dichotomous samplers were used as the standard, with either Teflon (DT) or quartz fiber (DQ) filters. The bias in the $PM_{2.5}$ measurements with the cyclone/impactor system was reflected in the calculated $PM_{2.5}/PM_{10}$ ratios as compared with the ratios calculated from the collocated dichotomous samplers.

As shown in Figure 12 for EPA-funded tests of unpaved roads, the $PM_{2.5}/PM_{10}$ ratios measured by the cyclone/impactors clustered around a value of 0.25, whereas the dichotomous samplers gave much lower values that decreased with increasing PM_{10} concentrations. Based on the test results from the EPA program, a new default value of 0.15 for the $PM_{2.5}/PM_{10}$ ratio was used to replace the previous value of 0.26 for dust emissions from unpaved roads. A default value of 0.15 was also recommended [2] for most of the other fugitive dust source categories, the exceptions being paved roads (0.25) and agricultural crops (0.20).

It is important to note that the PM_{10} concentrations in Figure 12 are averages during the testing rather than peak values encountered with each plume passage. Because vehicles passed the test site at the rate of about one per minute (with a plume passage time of less than 6 seconds), the peak values of PM_{10} concentration were at least 10 times the average values.

In the first test series of Phase I (through Test WR-15), a DustTrak monitor was used to continuously measure the PM_{10} concentration in the exposure section of the wind tunnel. Later, when a Partisol sampler was operated concurrently to measure the PM_{10} concentration, a running comparison was made between the DustTrak PM_{10} concentration and the Partisol PM_{10} concentrations, as shown in Figure 13. Except for PM_{10} concentrations below $1,000 \mu\text{g}/\text{m}^3$, all ratios fell within one relative standard deviation of 30 percent about the mean of 1.06.

4.2 Phase II

The purpose of the Phase II testing was to determine whether $PM_{2.5}/PM_{10}$ ratios measured with reference method samplers depend on (a) the test material [soil or road surface aggregate] that generates the dust, or (b) the dust concentration to which the samplers are exposed.

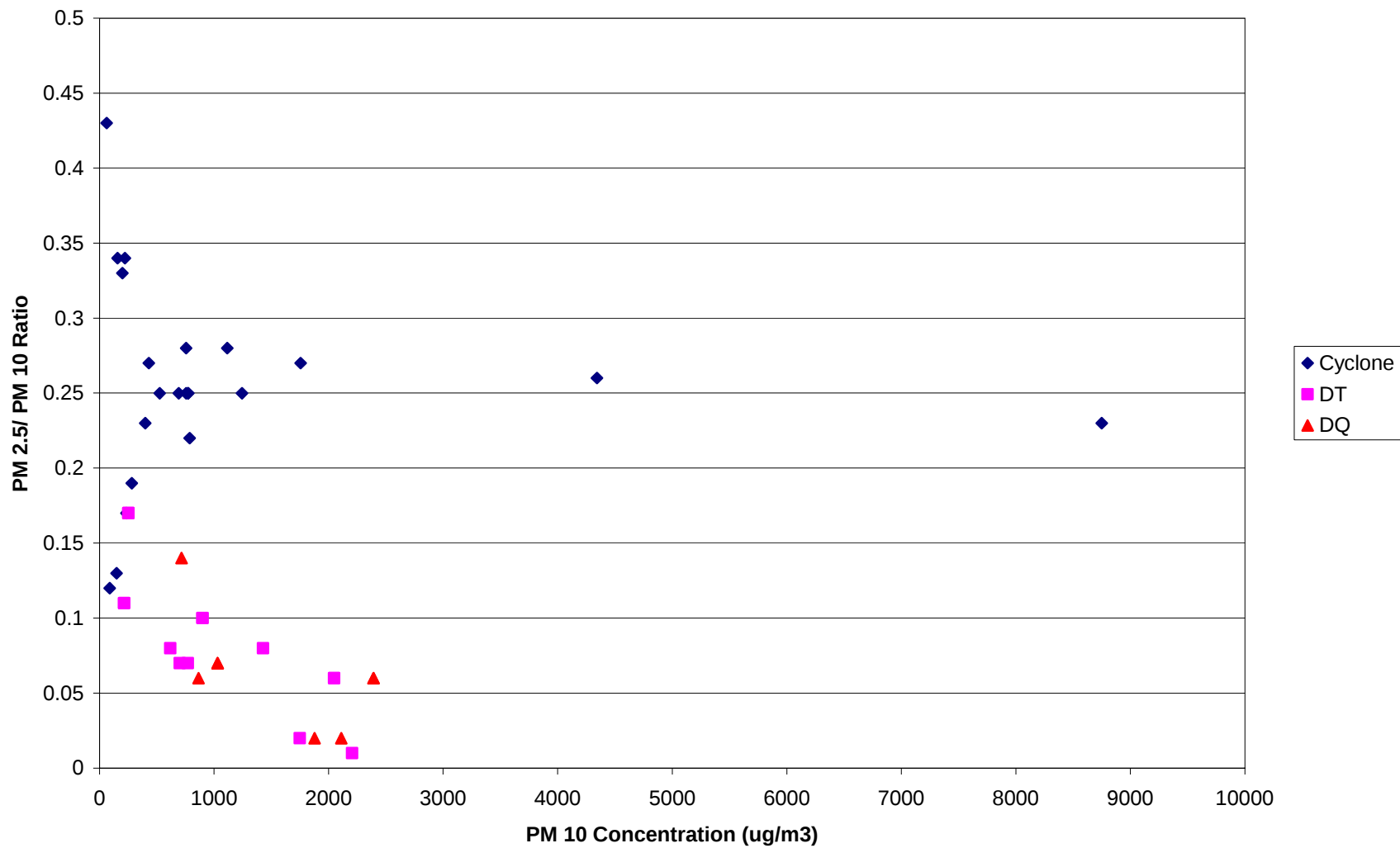


Figure 12. PM_{2.5}/PM₁₀ Ratio vs. PM₁₀ Concentration from Prior Field Study [2]

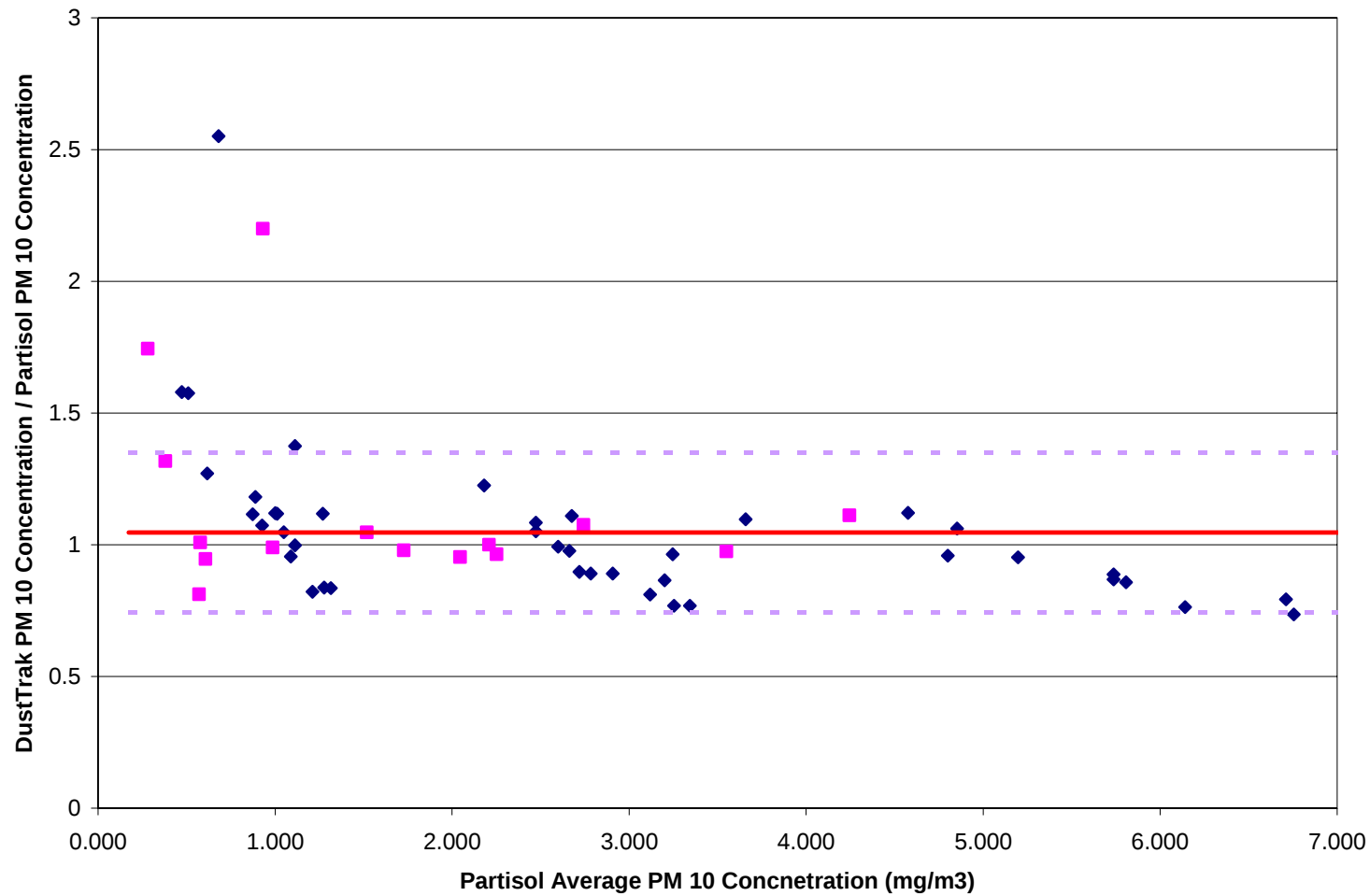


Figure 13. Ratio of DustTrak to Partisol PM₁₀ Concentration Versus Average PM₁₀ Concentration

In this phase, known amounts of test material were added to the dust chamber at the beginning of the test, so the auger (used in Phase I to introduce additional material) was eliminated. Whenever the airflow through the dust suspension chamber was activated, it was held at a constant value of 19 lpm, so that the energy input to the dust generation process did not vary. Each test soil was manually dry sieved through a 20-mesh screen, to eliminate observed erratic behavior in the dust generation process. For example, when the Knik River Sediment was tested without presieving, highly variable results were obtained (Runs WR-45 through WR-47).

Prior to each test series, the wind tunnel was purged with clean air to remove any loose accumulation of dust on the interior surfaces of the flow system. In addition, all sampler flow rates were checked, as well as the impactors in the PM_{2.5} Partisol samplers and the PM_{2.5} DustTrak monitors. The DustTrak impactors were regreased before proceeding.

The results of the Phase II testing are summarized in Figure 14. Although some data separation of different test materials is evident, there is an obvious tendency of the measured PM_{2.5}/PM₁₀ ratio to decrease with increasing PM₁₀ concentration. One observation of note is that the results for the more spherical test soils (river sediment and alluvial channel) are separated from the results for the other materials, which are more angular in nature.

One possible explanation for the decrease in PM_{2.5}/PM₁₀ ratio with increasing PM₁₀ concentration is the increased agglomeration of the coarse and fine modes of PM₁₀. This enhanced agglomeration would occur near the point of dust release from an open source. As the plume disperses, the PM₁₀ concentration decreases because of enhanced deposition of the coarse mode and the PM_{2.5}/PM₁₀ ratio increases.

In specifying the appropriate PM_{2.5}/PM₁₀ ratio to associate with emission factor determination for fugitive dust sources, an important factor is the PM₁₀ concentration range under which most of the plume mass is collected on sampler substrates. For uncontrolled emissions from unpaved roads, which accounts for the majority of PM₁₀ emissions from mechanically generated dust across the country, most of the plume mass is obtained at concentrations exceeding 5,000 µg/m³.

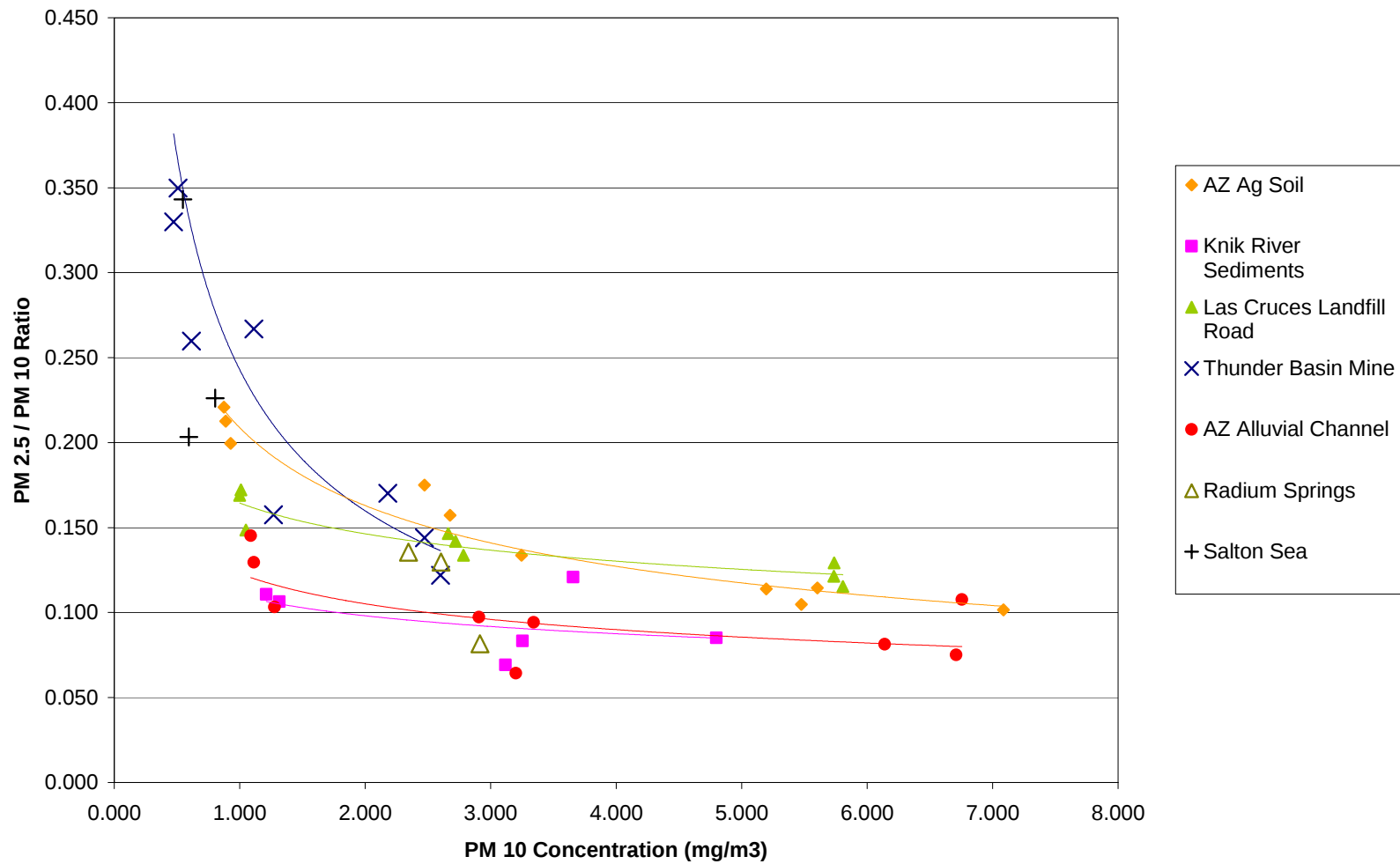


Figure 14. PM_{2.5}/PM₁₀ Ratio Versus PM₁₀ Concentration

An example of peak PM_{10} concentration data from uncontrolled road dust plumes is shown in Figure 15, which was obtained alongside an unpaved road at Ft. Riley, Kansas, in July 2005 [9]. Note that the alternate 6-sec peaks in this 20-pass test corresponded to 15-mph passes as opposed to the 30-mph passes. The latter passes correspond to the default vehicle speed used in AP-42 for the unpaved road emission factor equation. It should be noted that the peak 1-sec concentrations exceed the peak 6-sec concentrations by at least a factor of 3.

It is evident from Figure 14, that at PM_{10} concentrations above $5,000 \mu\text{g}/\text{m}^3$, at which most of the plume mass is collected for emission factor determination, the representative ratio of $PM_{2.5}/PM_{10}$ is 0.1. This conclusion is supported by the results of the earlier field testing program [2], which are shown above in Figure 12. The fact that the ratios from the reference-method samplers in the field study were as low as 0.05 is explained in part by the fact that the effective PM_{10} concentrations under which the samples were collected were in the range of $10,000 \mu\text{g}/\text{m}^3$ and above. In fact, Reference 2 states that dust plume core concentrations adjacent to unpaved roads often exceed $20,000 \mu\text{g}/\text{m}^3$. A $PM_{2.5}/PM_{10}$ of 0.1 is also consistent with observed data using FRM samplers during high-wind dust events on Owens Dry Lake [10].

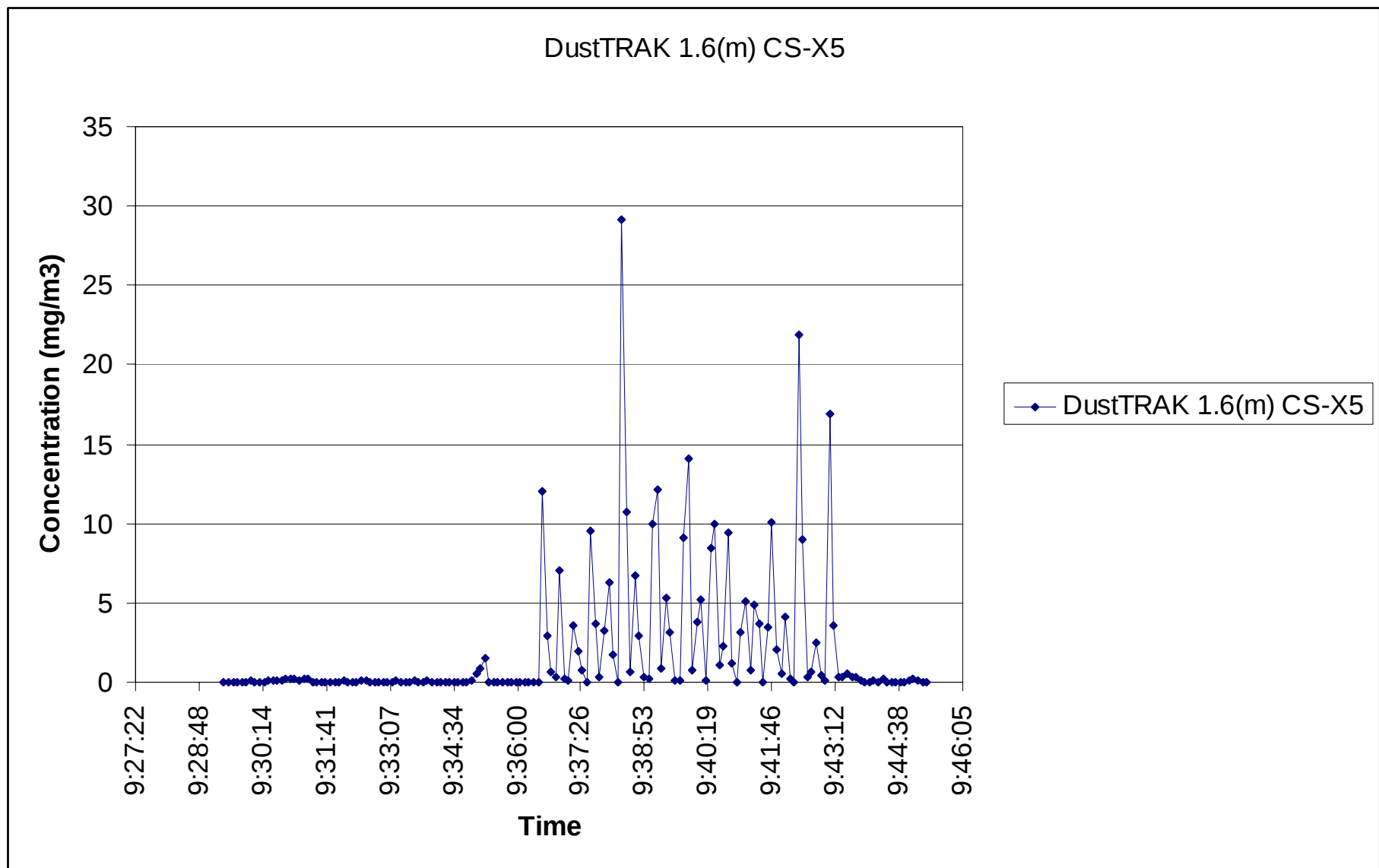


Figure 15. Example Peak (6-sec) Plume PM₁₀ Concentrations at Roadside Plume Profiling Location

Section 5.

Conclusions

Based on the 100 wind tunnel tests that were performed in this study, the findings support the following conclusions:

- $PM_{2.5}$ concentrations measured by the high-volume cyclone/impactor system used to develop AP-42 emission factors for fugitive dust sources have a positive bias in the range of a factor of 2, as compared to the $PM_{2.5}$ concentration measurements from reference-method samplers. (The geometric mean bias is 2.01, and the arithmetic mean bias is 2.15.)
- The $PM_{2.5}$ bias associated with the cyclone/impactor system as measured under controlled laboratory conditions with dust concentrations held at nearly steady values, closely replicates the bias observed in a prior study of dust emissions from unpaved roads. That study examined particle size ratios under field conditions at distributed geographic locations across the country.
- The $PM_{2.5}/PM_{10}$ ratios measured in this study for a variety of western soils show a decrease in magnitude with increasing PM_{10} concentration. Soils with a nominally spherical shape are observed to have somewhat lower ratios (at given PM_{10} concentrations) than soils with angular shape. A very similar dependence of $PM_{2.5}/PM_{10}$ ratio on PM_{10} concentration was also observed in the prior field study that used dichotomous samplers as FRM devices.
- The test data from the current study support a $PM_{2.5}/PM_{10}$ ratio of 0.1 for typical fugitive dust sources. This ratio takes into account the fact that most PM_{10} sample mass from uncontrolled dust sources is collected at PM_{10} concentrations exceeding $5,000 \mu\text{g}/\text{m}^3$.
- The $PM_{2.5}/PM_{10}$ ratio of 0.1 is also supported by numerous other studies including the prior field study that used dichotomous samplers as reference devices. It is possible that a ratio as low as 0.05 (as was found in the prior field tests of unpaved roads) might be appropriate, but this would require extrapolation of the current test data to higher PM_{10} concentrations.

Section 6.

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10. Ono, Duane. 2005. "Ambient PM_{2.5}/PM₁₀ ratios for Dust Events from the Keeler Dunes." Great Baron UAPCD, Bishop, CA. **[Describes FRM test results for high-wind events on Owens Dry Lake.]**

Appendix A

Summaries of Test Data

Table A-1. Test Data From Phase I

		Concentration (mg/m ³)										
Run	Test dust	DustTRAKS		Partisols		Average Partisol	Cyclone/Impactors	Average Cyclone /Impactor	PM-2.5/PM-10 Ratios			
		PM-10	PM-2.5	PM-10	PM-2.5	PM-2.5	PM-2.5	PM-2.5	DustTRAKS	Partisols	Cyclone/impactors	Average cyclone/impactor
WR-1	AZ Fine	2.15			0.6	0.595	1.15	1.125		0.28	0.5	0.51
WR-2	AZ Fine	2.29	1.06		0.589	0.645	1.1	1.075	0.46	0.28	0.52	0.48
WR-3	AZ Fine	2.1	1.05		0.701	0.627	1.1	1.110	0.50	0.30	0.48	0.58
WR-4	AZ Fine	1.19	0.581		[2.295]	0.627	1.12	1.110	0.50	0.30	0.58	0.58
WR-5	AZ Fine	1.12	0.385		0.627	0.318	1.1	0.530	0.49	0.27	0.57	0.51
WR-6	BLANK				0.333	0.296	0.525	0.524	0.34	0.26	0.5	0.45
WR-7	AZ Fine	1.02	0.323		0.302	0.296	0.535	0.524	0.34	0.26	0.51	0.45
WR-8	AZ Fine	4.95	1.46		0.332	0.296	0.56	0.524	0.34	0.26	0.46	0.44
WR-9	AZ Fine	5.6	1.77		0.26	0.296	0.488	0.524	0.34	0.26	0.44	0.45
WR-10	AZ Fine	4.99	1.44			0.241	0.506	0.475	0.32	0.24	0.42	0.42
WR-11	BLANK				0.234	0.241	0.444	0.475	0.32	0.24	0.42	0.42
WR-12	AZ Coarse	4.74	0.662		1.613	1.426	2.455	2.385	0.29	0.29	0.46	0.47
WR-13	AZ Coarse	4.5	0.615		1.238	1.426	2.314	2.385	0.29	0.29	0.46	0.47
WR-14	AZ Coarse	4.9	0.769		1.693	1.613	3.144	2.982	0.32	0.29	0.49	0.48
WR-15	BLANK				1.533	1.290	2.82	2.982	0.32	0.29	0.47	0.48
WR-16	AZ Coarse	2.21	0.395		1.338	1.290	2.408	2.391	0.29	0.26	0.45	0.46
WR-17	AZ Coarse	2.17	0.225		1.242	1.290	2.373	2.391	0.29	0.26	0.46	0.46
WR-18	AZ Coarse	1.95	0.182			0.457	1.919	1.872	0.14	0.10	0.33	0.33
WR-19	BLANK				0.479	0.457	1.825	1.872	0.14	0.10	0.33	0.33
WR-20	AZ Coarse	0.575	0.169		0.376	0.566	2.108	1.905	0.14	0.13	0.37	0.36
WR-21	AZ Coarse	0.582	0.16		0.591	0.566	1.701	1.905	0.14	0.13	0.34	0.34
WR-22	AZ Coarse	0.977	0.234		0.503	0.566	2.072	2.037	0.16	0.09	0.34	0.35
WR-23	BLANK				0.603	0.417	2.002	2.037	0.16	0.09	0.35	0.35
WR-24	Owens DLB	3.46	0.945		0.563	0.417	2.072	2.037	0.16	0.09	0.34	0.35
WR-25	Owens DLB	2.95	0.941		0.495	0.417	2.002	2.037	0.16	0.09	0.35	0.35
WR-26	Owens DLB	4.72	1.54		0.192	0.417	2.002	2.037	0.16	0.09	0.35	0.35
WR-27	Owens DLB	2.05	0.845			0.378	0.764	0.710	0.18	0.17	0.31	0.30
WR-28	Owens DLB	1.59	0.565		0.385	0.378	0.656	0.710	0.18	0.17	0.31	0.30
WR-29	Owens DLB	1.69	0.687		0.371	0.378	0.64	0.710	0.18	0.17	0.29	0.24
					0.31	0.286	0.537	0.589	0.10	0.13	0.25	0.24
					0.261	0.213	0.556	0.520	0.09	0.10	0.24	0.24
					0.162	0.213	0.483	0.520	0.09	0.10	0.24	0.24
						0.120	0.195	0.191	0.29	0.20	0.32	0.31
						0.122	0.187	0.210	0.27	0.21	0.29	0.32
						0.156	0.217	0.210	0.27	0.21	0.31	0.32
						0.156	0.202	0.303	0.24	0.16	0.33	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
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						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
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						0.156	0.29	0.303	0.24	0.16	0.28	0.29
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						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
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						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29
						0.156	0.29	0.303	0.24	0.16	0.28	0.29
						0.156	0.316	0.303	0.24	0.16	0.29	0.29

Table A-1. Test Data From Phase I (Continued)

		Concentration (mg/m ³)										
Run	Test dust	DustTRAKS		Partisols		Average Partisol	Cyclone/Impactors	Average Cyclone/Impactor	PM-2.5/PM-10 Ratios			Average cyclone/impactor
		PM-10	PM-2.5	PM-10	PM-2.5	PM-2.5	PM-2.5	PM-2.5	DustTRAKS	Partisols	Cyclone/impactors	
WR-30	BLANK				0.374			0.738			0.46	
WR-31	Owens DLB	0.492	0.419	[0.282]	0.147	0.135	0.205	0.205	0.85	0.27	0.59	0.61
WR-32	Owens DLB	0.502	0.319	0.381	0.123	0.111	0.205	0.198	0.64	0.29	0.62	0.45
WR-33	Owens DLB	0.464	0.219	0.571	0.146	0.131	0.21	0.185			0.45	
WR-34	BLANK				0.076		0.252	0.242	0.47	0.23	0.44	0.38
					0.121		0.232				0.38	
					0.141							

[] Denotes outlier not used in calculations

Table A-2. Test Data From Phase II

Run	Test dust	DustTRAKS		Partisol concentration				Average of Partisols		PM-2.5/PM-10 ratios	
		PM-10	PM-2.5	A (PM-2.5)	B (PM-2.5)	C (PM-10)	D (PM-10)	PM-10	PM-2.5	DustTRAKS	Partisols
WR-35	AZ Ag Field	0.995	0.241	0.19	0.18	0.927	0.926	0.927	0.185	0.242	0.200
WR-36	AZ Ag Field	0.975	0.247	0.189	0.198	0.883	0.866	0.875	0.194	0.253	0.221
WR-37	AZ Ag Field	1.050	0.262	0.189	0.211	1.723	0.899	1.311	0.200	0.250	0.153
WR-38	AZ Ag Field	2.970	0.720	0.433	0.408	2.752	2.6	2.676	0.421	0.242	0.157
WR-39	AZ Ag Field	3.130	0.631	0.439	0.43	3.501	2.991	3.246	0.435	0.202	0.134
WR-40	AZ Ag Field	2.600	0.738	0.433	0.433	2.514	2.431	2.473	0.433	0.284	0.175
WR-41	AZ Ag Field	4.950	1.030	0.579	0.605	5.548	4.847	5.198	0.592	0.208	0.114
WR-42	AZ Ag Field	5.130	1.580	0.974	0.968	4.694	4.458	4.576	0.971	0.308	0.212
WR-43	AZ Ag Field	5.150	1.650	1.054	1.126	5.124	4.581	4.853	1.090	0.320	0.225
WR-44	BLANK										
WR-48	Knik River Sediment	4.010	1.460	0.433	0.451	3.746	3.571	3.659	0.442	0.364	0.121
WR-49	Knik River Sediment	4.600	1.160	0.415	0.403	5.016	4.585	4.801	0.409	0.252	0.085
WR-50	BLANK										
WR-51	Knik River Sediment	2.530	0.385	0.228	0.205	3.336	2.901	3.119	0.217	0.152	0.069
WR-52	Knik River Sediment	2.500	0.544	0.28	0.263	3.344	3.165	3.255	0.272	0.218	0.083
WR-53	Knik River Sediment	0.995	0.280	0.135	0.134	1.289	1.134	1.212	0.135	0.281	0.111
WR-54	Knik River Sediment	1.100	0.314	0.138	0.141	1.344	1.289	1.317	0.140	0.285	0.106
WR-55	BLANK										
WR-56	Las Cruces Landfill Road	4.980	1.060	0.676	0.662	5.893	5.722	5.808	0.669	0.213	0.115
WR-57	Las Cruces Landfill Road	4.980	1.140	0.689	0.704	5.873	5.599	5.736	0.697	0.229	0.121
WR-58	Las Cruces Landfill Road	5.090	1.260	0.75	0.734	5.802	5.671	5.737	0.742	0.248	0.129
WR-59	Las Cruces Landfill Road	2.480	0.654	0.369	0.376	2.834	2.735	2.785	0.373	0.264	0.134
WR-60	Las Cruces Landfill Road	2.440	0.680	0.384	0.388	2.718	2.72	2.719	0.386	0.279	0.142
WR-61	BLANK										

Table A-2. Test Data From Phase II (Continued)

Run	Test dust	DustTRAKS		Partisol concentration				Average of Partisols		PM-2.5/PM-10 ratios	
		PM-10	PM-2.5	A (PM-2.5)	B (PM-2.5)	C (PM-10)	D (PM-10)	PM-10	PM-2.5	DustTRAKS	Partisols
WR-62	Las Cruces Landfill Road	2.6	0.62	0.388	0.392	2.755	2.569	2.662	0.390	0.238	0.147
WR-63	Las Cruces Landfill Road	1.1	0.4	0.144	0.169	1.103	0.996	1.050	0.157	0.364	0.149
WR-64	Las Cruces Landfill Road	1.12	0.361	0.169	0.169	1.007	0.993	1.000	0.169	0.322	0.169
WR-65	Las Cruces Landfill Road	1.13	0.377	0.169	0.178	1.048	0.973	1.011	0.174	0.334	0.172
WR-66	BLANK										
WR-67	Thunder Basin Mine	2.58	1.03	0.319	0.314	2.702	2.497	2.600	0.317	0.399	0.122
WR-68	Thunder Basin Mine	2.68	1.23	0.375	0.337	2.534	2.413	2.474	0.356	0.459	0.144
WR-69	Thunder Basin Mine	2.67	1.36	0.378	0.364	2.218	2.141	2.180	0.371	0.509	0.170
WR-70	Thunder Basin Mine	1.42	0.695	0.197	0.203	1.296	1.245	1.271	0.200	0.489	0.157
WR-71	Thunder Basin Mine	1.74	1.54	0.369	0.334	0.701	0.663	0.682	0.352	0.885	0.515
WR-72	BLANK										
WR-73	Thunder Basin Mine	1.53	0.996	0.303	0.29	1.196	1.03	1.113	0.297	0.651	0.266
WR-74	Thunder Basin Mine	0.747	0.552	0.16	0.152	0.475	0.471	0.473	0.156	0.739	0.330
WR-75	Thunder Basin Mine	0.783	0.51	0.164	0.157	0.63	0.602	0.616	0.161	0.651	0.261
WR-76	Thunder Basin Mine	0.802	0.63	0.18	0.177	0.513	0.505	0.509	0.179	0.786	0.351
WR-77	Blank										
WR-78	AZ Alluvial Channel	4.69	0.893	0.344	0.656	6.335	5.946	6.141	0.500	0.190	0.081
WR-79	AZ Alluvial Channel	4.97	0.989	0.498	0.955	6.966	6.543	6.755	0.727	0.199	0.108
WR-80	AZ Alluvial Channel	5.32	0.975	0.476	0.532	6.884	6.537	6.711	0.504	0.183	0.075
WR-81	AZ Alluvial Channel	2.57	0.54	0.337	0.292	3.431	3.254	3.343	0.315	0.210	0.094
WR-82	AZ Alluvial Channel	2.77	0.69	0.284	0.127	3.246	3.155	3.201	0.206	0.249	0.064
WR-83	AZ Alluvial Channel	2.59	0.879	0.298	0.268	2.976	2.839	2.908	0.283	0.339	0.097

Table A-2. Test Data From Phase II (Continued)

Run	Test dust	DustTRAKS		Partisol concentration				Average of Partisols		PM-2.5/PM-10 ratios	
		PM-10	PM-2.5	A (PM-2.5)	B (PM-2.5)	C (PM-10)	D (PM-10)	PM-10	PM-2.5	DustTRAKS	Partisols
WR-84	AZ Alluvial Channel	1.07	0.365	0.0139	0.0125	1.291	1.265	1.278	0.013	0.341	0.010
WR-85	Blank										
WR-86	AZ Alluvial Channel	1.04	0.399	0.162	0.153	1.111	1.065	1.088	0.158	0.384	0.145
WR-87	AZ Alluvial Channel	1.11	0.42	0.144	0.145	1.141	1.083	1.112	0.145	0.378	0.130
WR-88	Blank										
WR-89	Radium Springs	2.75	1.04	0.338	[-0.122]	2.728	2.483	2.606	0.338	0.378	0.130
WR-90	Radium Springs	2.75	1.06	0.321	0.315	2.514	2.179	2.347	0.318	0.385	0.136
WR-91	Radium Springs	3.05	0.883	[-0.096]	0.238	2.874	2.982	2.928	0.238	0.290	0.081
WR-92	Blank										
WR-93	Salton Sea	1.57	0.823	0.163	0.202	0.846	0.765	0.806	0.183	0.524	0.227
WR-94	Salton Sea	1.12	0.605	0.119	0.123	0.628	0.561	0.595	0.121	0.540	0.204
WR-95	Salton Sea	1.17	0.697	0.195	0.18	0.581	0.515	0.548	0.188	0.596	0.342
WR-96	Blank										
WR-97	AZ Ag Field	4.83	0.96	0.521	0.627	5.725	5.224	5.475	0.574	0.199	0.105
WR-98	AZ Ag Field	5.42	1.05	0.694	0.746	7.264	6.91	7.087	0.720	0.194	0.102
WR-99	AZ Ag Field	5.1	1.08	0.61	0.673	5.846	5.36	5.603	0.642	0.212	0.114
WR-100	Blank										

Table A-3. Field Blank Corrections

Phase 1	Filter type	Blank correction	Comments
	8 X 10 in	-0.143	An overall blank correction was applied to each filter type in Phase 1
	4 X 5 in ^a	0.040	
	47 mm	-0.053	
Phase 2	Runs	Blank correction	
	WR-35 to 43	0.044	Only 47-mm filters were used in Phase 2 of testing
	WR-45 to 50	0.044	
	WR-51 to 54	0.008	
	WR-56 to 60	-0.002	
	WR-62 to 65	-0.004	
	WR- 67 to 71	0.013	
	WR-73 to 76	-0.031	
	WR-78 to 85	0.002	
	WR-86 to 87	0.007	
	WR-89 to 92	0.044	
	WR-93 to 95	0.035	
	WR 97 to 99	0.033	

^a Impactor substrate.

Appendix B

QA/QC Activities

B.1 QA/QC Procedure

The Quality Assurance Project Plan (QAPP) prepared for this test program is a separate document that describes all the QA/QC activities for the project. An outline of that document is attached to the end of this section.

B.2 QA/QC Activities

As part of the QA program for this study, audits of sampling and analysis procedures were performed. The purpose of the audits was to demonstrate that measurements are made within acceptable control conditions for particulate source sampling and to assess the source testing data for precision and accuracy. Examples of items audited included gravimetric analysis, flow rate calibration, data processing, and concentration calculation. The mandatory use of specially designed reporting forms for sampling and analysis data obtained in the field and laboratory aided in the auditing procedure.

Requirements for high-volume (hi-vol) sampler flow rates rely on the use of secondary and primary flow standards. The Roots meter is the primary volumetric standard and the BGI orifice is the secondary standard for calibration of hi-vol sampler flow rates. The Roots meter is calibrated and traceable to a NIST standard by the manufacturer. The BGI orifice is calibrated against the primary standard on an annual basis. Before going to the field, the BGI orifice is first checked to assure that it has not been damaged. In the wind tunnel laboratory, the orifice is used to calibrate the flow rate of each hi-vol sampler. Table B-1 specifies the frequency of calibration and other QA checks regarding air samplers.

A second pretest activity is the preparation of the hi-vol filters for use in the testing. In this preparation, the filters are weighed under stable temperature and humidity conditions. After they are weighed and have passed audit weighing, the filters are packaged for shipment to the field. Table B-2 outlines the general requirements for conditioning and weighing sampling media. Note the audit weighing is performed by a second, independent analyst.

Table B-1. Quality Assurance Procedures for Sampling Equipment

Activity	QC check/requirement
Flow Rate Calibration	Prior to start of test series,
• High-volume samplers	Use BGI transfer standard (calibrated orifice) to set flow rate of each sampler
• Partisol samplers	Use Gillian transfer standard (bubble meter) to check flow rate of each sampler
• TSI DustTRAK monitors	Use TSI transfer standard (rotameter) to set flow rate of each sampler
Operation	
• Timing	Start and stop all downwind samplers during time span not exceeding 1 min.

Table B-2. Quality Assurance Procedures for Sampling Media

Activity	QA check/requirement
Preparation	Inspect and imprint glass fiber media with identification numbers.
Conditioning	Equilibrate media for 24 h in clean controlled room with relative humidity of 40% (variation of less than $\pm 5\%$ RH) and with temperature of 23°C (variation of less than $\pm 1^\circ\text{C}$).
Weighing	Weigh hi-vol back-up filters to nearest 0.05 mg. Weigh Partisol filters to nearest 0.001 mg. Weigh cascade impactor substrates to nearest 0.05 mg.
Auditing of weights	Independently verify final weights of all filters and substrates. Reweigh entire batch if weights of any hi-vol filters deviate by more than ± 2.0 mg. For tare weights, conduct a 100% audit by a second analyst. Reweigh any high-volume filter whose weight deviates by more than ± 1.0 mg. Follow same procedures for impactor substrates used for sizing tests. Audit limits for impactor substrates are ± 1.0 and ± 0.5 mg for final and tare weights, respectively.
Correction for handling effects	Weigh and handle at least one blank for each 1 to 10 filters of each type used to test.
Calibration of balance	Balance to be calibrated once per year by certified manufacturer's representative. Check prior to each use with laboratory Class S weights.

As indicated in Table B-2, a minimum of 10% field blanks were collected for QC purposes. This involves handling at least one blank filter for every 10 exposed filters in an identical manner to determine systematic weight changes due to handling steps alone. These changes are used to mathematically correct the net weight gain due to handling. A field blank filter is loaded into a sampler and then immediately recovered without any air being passed through the media. Blanks have been successfully used in many MRI programs to account for systematic weight changes due to handling.

After the particulate matter samples and blank filters are collected and returned from the field, the collection media are placed in the gravimetric laboratory and allowed to come to equilibrium. Each filter is weighed, allowed to return to equilibrium for an additional 24 h, and then all of the exposed filters are reweighed by a second analyst. If a filter fails the audit criterion, the entire lot will be allowed to condition in the gravimetric laboratory an additional 24 h and then reweighed. The tare and first weight criteria for filters (Table B-2) are based on an internal MRI study conducted in the early 1980s to evaluate the stability of several hundred 8- x 10-in glass fiber filters used in exposure profiling studies.

B.3 QA/QC Checks for Data Reduction and Validation

Whenever practical, all data collected in the study were entered directly onto standard data forms. All data were recorded on standard data forms using permanent black ink and signed/dated by sampling personnel. Data forms were inspected for completeness and accuracy by the appropriate field supervisor at the end of each test. At that time, data forms were grouped by test number and bound into 3-ring binders.

The data analysis procedures that were used for this project are procedures that have been through several layers of validation in substantiating the performance of the method. It should be noted that blank-corrected sample mass is considered quantifiable (and usable for concentration calculation) only if it equals or exceeds three times the standard deviation of the average net weight change of the field blanks.

An independent auditor performed a check of the calculations in the computer data reduction programs. The Field Team Leader or his/her designee conducted an on-site spot check to ensure that data were being recorded accurately. After the field test, an independent auditor checked data input to assure accurate transfer of the raw data.

For this project, all records were evaluated for the adherence to all procedures and requirements. The items that were reviewed include:

- Gravimetric audit weighing for the assessment of the particulate data,
- Calibration and calibration criterion checks,
- The results of all blanks, and
- The validation of data process systems or procedures.

Selected data were reconstructed, including tracing the calibration back to the primary standards. Any software (spreadsheets) used to determine numerical values was checked by hand calculating all intermediate and final results for one run by referring to original sources of data (i.e., field filter logs, filter weight logs, run sheets, sampler look-up tables).

B.4 Sample Identification and Traceability

To maintain sample integrity, the following procedures were used:

- Each filter was issued a unique identification number. MRI SOP EET-610 describes the numbering system that is employed to identify filter type, project, and other information.
- The sample number was recorded in a sample logbook along with the date the sample was obtained. The sample number was coded to indicate the sample location and test series.

- Other pertinent information that was recorded included short descriptions of sample type or location, storage location, condition of sample, any special instructions, and signatures of personnel who received the sample for analysis.
- In order to conduct traceability, all sample transfers were recorded in a notebook or on forms. The following information was recorded: the assigned sample codes, date of transfer, location of storage site, and the name of the person initiating and accepting the transfer.

All documented work was reviewed by the project leader for completeness. The field technical coordinator was responsible for assuring that all samples are accounted for and that proper traceability/tracking procedures were followed.