SAT Initiative: Elm Street Elementary School (Wauseon, OH)

This document describes the analysis of air monitoring and other data collected under EPA's initiative to assess potentially elevated air toxics levels at some of our nation's schools. The document has been prepared for technical audiences (e.g., risk assessors, meteorologists) and their management. It is intended to describe the technical analysis of data collected for this school in clear, but generally technical, terms. A summary of this analysis is presented on the page focused on this school on EPA's website (www.epa.gov/schoolair).

I. Executive Summary

- Air monitoring has been conducted at Elm Street Elementary School as part of the EPA initiative to monitor specific air toxics in the outdoor air around priority schools in 22 states and 2 tribal areas.
- This school was selected for monitoring based on information indicating the potential for elevated ambient concentrations of 1,6-hexamethylene diisocyanate (1,6-HDI); 4,4- methylenediphenyl diisocyanate (MDI); and 2,4-toluene diisocyanate (2,4-TDI) in air outside the school. The school was ranked in the top 25 on the USA Today list due to 2005 Toxics Release Inventory (TRI) estimates of diisocyanate emissions from a nearby industrial door manufacturing facility. Mobile source pollutants, such as benzene and acrolein, were also monitored, as this school is also located in an area surrounded by two state highways and a railroad line.
- Air monitoring for 1,6-HDI; MDI; and 2,4-TDI as well as benzene, acrolein, and other volatile organic compounds (VOCs), was performed from September 4, 2009 through November 9, 2009.
- The three diisocyanate chemicals (1,6-HDI; MDI; and 2,4-TDI) were not detected in the air samples, thereby suggesting no influence of a nearby source.
- Measured benzene levels and associated longer-term concentration estimates at this school, while below levels of significant concern, do indicate the ubiquitous nature and influence of mobile source pollutants of concern that are the focus of EPA actions nationwide.
- Benzene is common in the outdoor air in urban areas where many sources occur together, particularly mobile sources such as cars and other motor vehicles and off-road machinery. Concentrations of benzene and other air toxics released by mobile sources are often elevated in locations close to roadways and railroad lines, such as at Elm Street Elementary School. EPA remains concerned about mobile source emissions and continues to work to reduce those emissions across the country through national rules, and by providing information and suggestions to assist with reductions in local areas (http://www.epa.gov/schoolair/mobile.html).
- EPA will not use the acrolein data in evaluating the potential for health concerns from exposure to air toxics in outdoor air as part of the School Air Toxics Monitoring project (SAT). The Agency made this determination after results of a short-term laboratory study raised questions about the consistency and reliability of monitoring results of acrolein. (More information is available at http://www.epa.gov/schoolair/acrolein.html).

- Based on the analysis described here, EPA will not extend air toxics monitoring at this school. However, EPA's ongoing research and national air toxics monitoring programs will continue to collect information on mobile source impacts on outdoor air nationally.
- The Ohio Environmental Protection Agency (OEPA) will continue to oversee industrial facilities in the area through air permits and other programs. Additionally, OEPA will also continue to implement reductions in mobile sources through implementation of national programs and its own programs.

II. Background on this Initiative

As part of an EPA initiative to implement Administrator Lisa Jackson's commitment to assess potentially elevated air toxics levels at some of our nation's schools, EPA and state and local air pollution control agencies are monitoring specific (key) air toxics in the outdoor air around priority schools in 22 states and 2 tribal areas (<u>http://www.epa.gov/schoolair/schools.html</u>).

- The schools selected for monitoring include some schools that are near large industries that are sources of air toxics, and some schools that are in urban areas, where emissions of air toxics come from a mix of large and small industries, cars, trucks, buses and other sources.
- EPA selected schools based on information available to us about air pollution in the vicinity of the school, including results of the 2002 National-Scale Air Toxics Assessment (NATA), results from a 2008 USA Today analysis on air toxics at schools, and information from state and local air agencies. The analysis by USA Today involved use of EPA's Risk Screening Environmental Indicators tool and Toxics Release Inventory (TRI) for 2005.
 - Available information had raised some questions about air quality near these schools that EPA concluded merited investigation. In many cases, the information indicated that estimated long-term average concentrations of one or more air toxics were above the upper end of the range that EPA generally considers as acceptable (e.g., above 1-in-10,000 cancer risk for carcinogens).
- Monitors are placed at each school for approximately 60 days, and take air samples on at least 10 different days during that time. The samples are analyzed for specific air toxics identified for monitoring at the school (i.e., key pollutants).¹
- These monitoring results and other information collected at each school during this initiative allow us to:
 - assess specific air toxics levels occurring at these sites and associated estimates of longer-term concentrations in light of health risk-based criteria for long-term exposures,
 - better understand, in many cases, potential contributions from nearby sources to key air toxics concentrations at the schools,
 - consider what next steps might be appropriate to better understand and address air toxics at the school, and

¹ In analyzing air samples for these key pollutants, samples are also being analyzed for some additional pollutants that are routinely included in the analytical methods for the key pollutants.

- improve the information and methods we will use in the future (e.g., NATA) for estimating air toxics concentrations in communities across the U.S.

Assessment of air quality under this initiative is specific to the air toxics identified for monitoring at each school. This initiative is being implemented in addition to ongoing state, local and national air quality monitoring and assessment activities, including those focused on criteria pollutants (e.g., ozone and particulate matter) or existing, more extensive, air toxics programs.

Several technical documents prepared for this project provide further details on aspects of monitoring and data interpretation and are available on the EPA website (e.g., www.epa.gov/schoolair/techinfo.html). The full titles of these documents are provided here:

- School Air Toxics Ambient Monitoring Plan
- Quality Assurance Project Plan For the EPA School Air Toxics Monitoring Program
- Schools Air Toxics Monitoring Activity (2009), Uses of Health Effects Information in Evaluating Sample Results

Information on health effects of air toxics being monitored² and educational materials describing risk concepts³ are also available from EPA's website.

III. Basis for Selecting this School and the Air Monitoring Conducted

This school was selected for monitoring in consultation with the State air agency, the Ohio Environmental Protection Agency (OEPA). We were interested in evaluating the ambient concentrations of 1,6-HDI; MDI; and 2,4-TDI in air outside the school, as the school was ranked in the top 25 on the USA Today list due to 2005 Toxics Release Inventory (TRI) estimates of diisocyanate emissions for a nearby industrial door manufacturing facility. In addition, mobile source pollutants, such as benzene, acrolein, and other volatile organic compounds (VOCs), were monitored, as this school is also located in an area surrounded by two state highways and a railroad line (Figure 1). More information on mobile sources of air toxics can be found on EPA's website (<u>http://www.epa.gov/schoolair/mobile.html</u>).

Monitoring commenced at this school on September 4, 2009, and continued through November 9, 2009. During this period, 10 VOC samples and 10 diisocyanate samples were collected and analyzed for the key pollutants (benzene; 1,6-HDI; MDI; and 2,4-TDI) and other air toxics at this school.

All VOC results, with the exception of acrolein, were evaluated for health concerns. Results of a recent short-term laboratory study have raised questions about the consistency and reliability of monitoring results of acrolein. As a result, EPA will not use these acrolein data in evaluating the potential for health concerns from exposure to air toxics in outdoor air as part of the SAT Monitoring project (<u>http://www.epa.gov/schoolair/acrolein.html</u>). All sampling methodologies

² For example, <u>http://www.epa.gov/schoolair/pollutants.html</u>, <u>http://www.epa.gov/ttn/fera/risk_atoxic.html</u>.

³ For example, <u>http://www.epa.gov/ttn/atw/3_90_022.html</u>, <u>http://www.epa.gov/ttn/atw/3_90_024.html</u>.

are described in EPA's schools air toxics monitoring plan (http://www.epa.gov/schoolair/acrolein.html).⁴

IV. Monitoring Results and Analysis

A. Background for the SAT Analysis

The majority of schools being monitored in this initiative were selected based on modeling analyses that indicated the potential for annual average air concentrations of some specific (key) hazardous air pollutants (HAPs or air toxics)⁵ to be of particular concern based on approaches that are commonly used in the air toxics program for considering potential for long-term risk. For example, such analyses suggested annual average concentrations of some air toxics were greater than long-term risk-based concentrations associated with an additional cancer risk greater than 10-in-10,000 or a hazard index on the order of or above 10. To make projections of air concentrations, the modeling analyses combined estimates of air toxics emissions from industrial, motor vehicle and other sources, with past measurements of winds, and other meteorological factors that can influence air concentrations, from a weather station in the general area. In some cases, the weather station was very close (within a few miles), but in other cases, it was much further away (e.g., up to 60 miles), which may contribute to quite different conditions being modeled than actually exist at the school. The modeling analyses are intended to be used to prioritize locations for further investigation.

The primary objective of this initiative is to investigate - through monitoring air concentrations of key air toxics at each school over a 2-3 month period - whether levels measured and associated longer-term concentration estimates are of a magnitude, in light of health risk-based criteria, for which follow-up activities may need to be considered. To evaluate the monitoring results consistent with this objective, we developed health risk-based air concentrations (the long-term comparison levels summarized in Appendix A) for the monitored air toxics using established EPA methodology and practices for health risk assessment⁶ and, in the case of cancer risk, consistent with the implied level of risk considered in identifying schools for monitoring. Consistent with the long-term or chronic focus of the modeling analyses, based on which these schools were selected for monitoring, we have analyzed the full record of concentrations of air toxics measured at this school, using routine statistical tools, to derive a 95 percent confidence

⁴ OEPA staff operated the monitors and sent the canisters and filters to the analytical laboratory under contract to EPA.

⁵ The term hazardous air pollutants (commonly called HAPs or air toxics) refers to pollutants identified in section 112(b) of the Clean Air Act which are the focus of regulatory actions involving stationary sources described by CAA section 112 and are distinguished from the six pollutants for which criteria and national ambient air quality standards (NAAQS) are developed as described in section 108. One of the criteria pollutants, lead, is also represented, as lead compounds, on the HAP list.

⁶ While this EPA initiative will rely on EPA methodology, practices, assessments and risk policy considerations, we recognize that individual state methods, practices and policies may differ and subsequent analyses of the monitoring data by state agencies may draw additional or varying conclusions.

interval⁷ for the estimate of the longer-term average concentration of each of these pollutants. In this project, we are reporting all actual numerical values for pollutant concentrations including any values below method detection limit (MDL).⁸ Additionally, a value of 0.0 is used when a measured pollutant has no value detected (ND). The projected range for the longer-term concentration estimate for each chemical (most particularly the upper end of the range) is compared to the long-term comparison levels. These long-term comparison levels conservatively presume continuous (all-day, all-year) exposure over a lifetime. The analysis of the air concentrations also includes a consideration of the potential for cumulative multiple pollutant impacts.⁹ In general, where the monitoring results indicate estimates of longer-term average concentrations that are above the comparison levels - i.e., above the cancer-based comparison levels or notably above the noncancer-based comparison levels - we will consider the need for follow-up actions such as:

- \rightarrow Additional monitoring of air concentrations and/or meteorology in the area,
- → Evaluation of potentially contributing sources to help us confirm their emissions and identify what options (regulatory and otherwise) may be available to us to achieve emissions reductions, and
- → Evaluation of actions being taken or planned nationally, regionally or locally that may achieve emission and/or exposure reductions. An example of this would be the actions taken to address the type of ubiquitous emissions that come from mobile sources.

We have further analyzed the dataset to describe what it indicates in light of some other criteria and information commonly used in prioritizing state, local and national air toxics program activities. State, local and national programs often develop long-term monitoring datasets in order to better characterize pollutants near particular sources. The 2-3 month dataset developed under this initiative will be helpful to those programs in setting priorities for longer-term monitoring dataset as useful as possible to state, local and national air toxics programs in their longer-term efforts to improve air quality nationally. To that end, this analysis:

⁷ When data are available for only a portion of the period of interest (e.g., samples not collected on every day during this period), statisticians commonly calculate the 95% confidence interval around the dataset mean (or average) in order to have a conservative idea of how high or low the "true" mean may be. More specifically, this interval is the range in which the mean for the complete period of interest is expected to fall 95% of the time (95% probability is commonly used by statisticians). The interval includes an equal amount of quantities above and below the sample dataset mean. The interval that includes these quantities is calculated using a formula that takes into account the size of the dataset (i.e., the 'n') as well as the amount by which the individual data values vary from the dataset mean (i.e., the "standard deviation"). This calculation yields larger confidence intervals for smaller datasets as well as ones with more variable data points. For example, a dataset including {1.0, 3.0, and 5.0}, results in a mean of 3.0 and a 95% confidence interval of 3.0 +/- ~5 (or -2.0 to 8.0). For comparison purposes, a dataset including {2.5, 3 and 3.5} results in a mean of 3.0 and a 95% confidence interval of 3.0 +/- ~1.2 (or 1.8 to 4.2). The smaller variation within the data in the second set of values causes the second confidence interval to be smaller.

⁸ Method detection limit (MDL) is the minimum concentration of a substance that can be measured and reported with 99% confidence that the pollutant concentration is greater than zero and is determined from the analysis of a sample in a given matrix containing the pollutant.

⁹ As this analysis of a 2-3 month monitoring dataset is not intended to be a full risk assessment, consideration of potential multiple pollutant impacts may differ among sites. For example, in instances where no individual pollutant appears to be present above its comparison level, we will also check for the presence of multiple pollutants at levels just below their respective comparison levels (giving a higher priority to such instances).

- → Describes the air toxics measurements in terms of potential longer-term concentrations, and, as available, compares the measurements at this school to monitoring data from national monitoring programs.
- → Describes the meteorological data by considering conditions on sampling days as compared to those over all the days within the 2-3 month monitoring period and what conditions might be expected over the longer-term (as indicated, for example, by information from a nearby weather station).
- → Describes available information regarding activities and emissions at the nearby source(s) of interest, such as that obtained from public databases such as TRI and/or consultation with the local air pollution authority.

B. Chemical Concentrations

We developed two types of long-term health risk-related comparison levels (summarized in Appendix A below) to address our primary objective. The primary objective is to investigate through the monitoring data collected for key pollutants at the school, whether pollutant levels measured and associated longer-term concentration estimates are elevated enough in comparison with health risk-based criteria to indicate that follow-up activities be considered. These comparison levels conservatively presume continuous (all-day, all-year) exposure over a lifetime.

In developing or identifying these comparison levels, we have given priority to use of relevant and appropriate air standards and EPA risk assessment guidance and precedents.¹⁰ These levels are based upon health effects information, exposure concentrations and risk estimates developed and assessed by EPA, the U.S. Agency for Toxic Substances and Disease Registry, and the California EPA. These agencies recognize the need to account for potential differences in sensitivity or susceptibility of different groups (e.g., asthmatics) or lifestages/ages (e.g., young children or the elderly) to a particular pollutant's effects so that the resulting comparison levels are relevant for these potentially sensitive groups as well as the broader population.

In addition to evaluating individual pollutants with regard to their corresponding comparison levels, we also considered the potential for cumulative impacts from multiple pollutants in cases where individual pollutant levels fall below the comparison levels but where multiple pollutant mean concentrations are within an order of magnitude of their comparison levels.

Using the analysis approach described above, we analyzed the chemical concentration data (Table 1 and Figure 2) with regard to areas of interest identified below.

¹⁰ This is described in detail in Schools Air Toxics Monitoring Activity (2009), Uses of Health Effects Information in Evaluating Sample Results

Key findings drawn from the information on chemical concentrations and the considerations discussed below include:

- None of the three diisocyanate compounds were detected in any of the samples collected.
- Benzene levels measured over the 2-month sampling period and associated longer-term concentration estimates at this school, while below levels of significant concern do indicate the influence of mobile source pollutants of concern that are the focus of EPA actions nationwide.

1,6-HDI; MDI; and 2,4-TDI key pollutants:

- Do the monitoring data indicate influence from a nearby source?
 - \rightarrow There were no detections of these pollutants in any of the samples collected.
- Do the monitoring data indicate elevated levels that pose significant long-term health concerns?
 - \rightarrow There were no detections of these pollutants in any of the samples collected.

Benzene, key pollutant:

Benzene is one of several air toxics that EPA recognizes as a key pollutant nationally. A large number of people live in areas across the U.S. with elevated ambient concentrations of this pollutant due to mobile sources.¹¹

- Do the monitoring data indicate elevated levels that pose significant long-term health concerns?
 - → Measured benzene levels and associated longer-term concentration estimates at this school, while below the levels of significant concern, do indicate the ubiquitous nature and influence of mobile source pollutants of concern that are the focus of EPA actions nationwide.
 - The estimate of longer-term benzene concentration (i.e., the upper bound of the 95 percent confidence interval on the mean of the dataset) is well below the long-term comparison levels (Table 1).¹² These comparison levels are continuous exposure concentrations (24 hours a day, all year, over a lifetime).
 - Further, the longer-term concentration estimate is more than tenfold lower than the cancer-based comparison level, indicating the longer-term estimate is well below a continuous (24 hours a day, 7 days a week) lifetime exposure concentration associated with 1-in-100,000 additional cancer risk.

¹¹ Additional information on mobile sources of air toxics is available at http://www.epa.gov/schoolair/mobile.html

¹² The upper end of the interval is less than 1.3 times the mean of the monitoring data and less than 7% of the long-term cancer-based comparison level.

 \rightarrow Additionally, we did not identify any concerns regarding short-term exposures as each individual measurement is below the individual sample screening level for benzene (which is based on consideration of exposure all day, every day over a period from a couple of weeks to longer for some pollutants).¹⁰

Other Air Toxics:

- Do the monitoring data indicate elevated levels of any other air toxics (or HAPs) that pose significant long-term health concerns?
 - → The monitoring data show low levels of the other HAPs monitored, with longer-term concentration estimates for these HAPs below their long-term comparison levels (Appendix C). Additionally each individual measurement for these pollutants is below the individual sample screening level¹⁰ for that pollutant.

Multiple Pollutants:

- Do the data collected for the air toxics monitored indicate the potential for other monitored pollutants to be present at levels that in combination with the key pollutant levels indicate an increased potential for cumulative impacts of significant concern (e.g., that might warrant further investigation)?
 - \rightarrow Although the multiple air toxics monitored at this site were below the levels of significant concern for multi-pollutant cumulative risk that had been suggested by the modeling information, these results indicate the influence of multiple mobile source pollutants of concern that are the focus of EPA actions nationwide (Appendix C).¹³

C. Wind and Other Meteorological Data

At each school monitored as part of this initiative, we are collecting meteorological data, minimally for wind speed and direction, during the sampling period. Additionally, we have identified the nearest National Weather Service (NWS) station at which a longer record is available.

In reviewing these data at each school in this initiative, we are considering if these data indicate that the general pattern of winds on our sampling dates are significantly different from those occurring across the full sampling period or from those expected over the longer-term. Additionally, we are noting, particularly for school sites where the measured chemical concentrations show little indication of influence from a nearby source, whether wind conditions on some portion of the sampling dates were indicative of a potential to capture contributions from the nearby "key" source in the air sample collected.

¹³ We note that this initiative is focused on investigation for a school-specific set of key pollutants indicated by previous analyses (and a small set of others for which measurements are obtained in the same analysis). Combined impacts of pollutants or stressors other than those monitored in this project is a broader area of consideration in other EPA activities. General information on additional air pollutants is available at http://www.epa.gov/air/airpollutants.html.

The meteorological station at Elm Street Elementary School collected wind speed and wind direction measurements beginning on September 3, 2009, continuing through the sampling period (September 4, 2009-November 9, 2009), and ending on January 20, 2010. As a result, on-site data for these meteorological parameters are available for all dates of sample collection, and also a period before and after the sampling period, producing a continuous record of over four months of on-site meteorological data. The meteorological data collected on sampling days are presented in Figures 3a-3b and Tables 2a-2b.

The nearest NWS station is at Toledo Express Airport in Toledo, Ohio. This station is approximately 17 miles east-northeast of the school. Measurements taken at that station include wind, temperature, and precipitation. These are presented in Tables 2a-2b and Appendix E.

Key findings drawn from this information and the considerations discussed below include:

- Both the sampling results for the diisocyanates and the on-site wind data indicate that the nearby key source was not contributing to conditions at the school location.
- The key sources for benzene were identified as nearby roadway mobile sources surrounding the school and a railroad line to the south of the school. Therefore, wind from any direction may be considered as from the direction of a key source.
- The wind patterns at the monitoring site across sampling dates are somewhat similar to those observed across the record of on-site meteorological data during the sampling period.
- Our ability to provide a confident characterization of the wind flow patterns at the monitoring site over the long-term is somewhat limited, as the NWS station at Toledo Express Airport somewhat appears to represent the specific wind flow patterns at the school location.
- Although we lack long-term wind data at the monitoring site, the wind pattern at the NWS station during the sampling period is generally similar to the historical long-term wind flow pattern at that same NWS station. This suggests that, on a regional scale, the 2-month sampling period may be representative of year-round wind patterns.
- What are the directions of the key sources of 1,6-HDI; MDI; 2,4-TDI, and benzene emissions in relation to the school location?
 - → The nearby source emitting 1,6-HDI; MDI; and 2,4-TDI into the air (described in section III above) lies less than one mile northwest of the school.
 - → Using the property boundaries of the full facility (in lieu of information regarding the location of specific source of 1,6-HDI; MDI; and 2,4-TDI emissions at the facility), we have identified an approximate range of wind directions to use in considering the potential influence of this facility on air concentrations at the school.

- → This general range of wind directions, from approximately 281 to 304 degrees, is referred to here as the expected zone of source influence (ZOI) when referring to the diisocyanate compounds.
- → The key sources for benzene were identified as nearby roadway mobile sources surrounding the school and a railroad line to the south of the school. Therefore, wind from any direction may be considered as from the direction of a key source.
- On days the air samples were collected, how often did wind come from direction of the key source?
 - → For the diisocyanate sampling, there were five sampling days in which a portion of the winds were from the expected ZOI (Figure 3a, Table 2a).
 - → For benzene sampling, since any wind direction may be considered as the direction of a key source, all ten sampling days for which on-site wind data are available were from the direction of the key sources (Figure 3b, Table 2b).
- How do wind patterns on the air monitoring days compare to those across the complete monitoring period and what might be expected over the longer-term at the school location?
 - \rightarrow Wind patterns across the air monitoring days appear somewhat similar to those observed over the record of on-site meteorological data during the sampling period.
 - → We note that wind patterns at the nearest NWS station (at Toledo Express Airport) during the sampling period are similar to those recorded at the NWS station over the long-term (2002-2007 period; Appendix E), supporting the idea that regional meteorological patterns in the area during the sampling period were consistent with long-term patterns. There is some uncertainty as to whether the general wind patterns at the school location for longer periods would be similar to the general wind patterns at the Toledo Express Airport (see below).
- How do wind patterns at the school compare to those at the Toledo Express Airport NWS station, particularly with regard to prevalent wind directions and the direction of the key source?
 - → During the sampling period for which data are available both at the school site and at the reference NWS station (approximately 2 months), prevalent winds at the school site are predominantly from the southwest to northwest and northeast to east, while those at the NWS station are more from the south, southwest, and west. The windroses for the two sites during the sampling period (Figures 3a-3b and Appendix E) show some differences in wind flow patterns.
- Are there other meteorological patterns that may influence the measured concentrations at the school monitoring site?
 - \rightarrow No, we did not observe other meteorological patterns that may influence the measured concentrations at the school monitoring site.

V. Key Source Information

- Was the source operating as usual during the monitoring period?
 - The nearby source of diisocyanates (described in section III above) has an operating permit issued by OEPA that includes operating requirements.¹⁴
 - Information from the nearby source indicates that the door manufacturing facility was operating at a typical rate of 80% during the sampling period.
 - This facility did not report any diisocyanate emissions to the 2008 TRI and revised their 2007 TRI emissions downward from 0.55 tons per year (tpy) to 0.004 tpy. The last reported diisocyanate emissions (unspeciated from TRI) for this source (2007 TRI) were lower than those relied upon in previous modeling analysis for this area (e.g., 2002 and 2005 TRI).
- Was mobile source activity typical during the monitoring period?
 - The most recently available county-level benzene emissions for on-road mobile and locomotive sources (2005 NATA) are lower than those reported for this area in the 2002 NATA.

VI. Integrated Summary and Next Steps

A. Summary of Key Findings

- 1. What are the key HAPs for this school?
 - → 1,6-HDI; MDI; and 2,4-TDI are key HAPs for this school, identified based on emissions information considered in identifying the school for monitoring. There were no detections of diisocyanates monitored during the sampling period.
 - → Benzene and acrolein are also key HAPs, as this school is located in an area surrounded by two state highways and a railroad line. Acrolein was not evaluated due to reasons discussed in Section III.
- 2. Do the data collected at this school indicate an elevated level of concern, as implied by information that led to identifying this school for monitoring?
 - \rightarrow There were no detections of diisocyanates in any of the samples collected.
 - \rightarrow Although measured benzene levels and associated long-term concentrations were below levels of significant concern, these results do indicate the influence of mobile source pollutants of concern that are the focus of EPA actions nationwide.
 - → EPA will not use the acrolein data in evaluating the potential for health concerns from exposure to air toxics in outdoor air as part of the SAT Monitoring project. The Agency made this determination after results of a short-term laboratory study raised questions about the consistency and

¹⁴ Operating permits, which are issued to air pollution sources under the Clean Air Act, are described at: <u>http://www.epa.gov/air/oaqps/permits/</u>

reliability of monitoring results of acrolein. (More information is available at <u>http://www.epa.gov/schoolair/acrolein.html</u>).

- 3. Are there indications, e.g., from the meteorological or other data, that the sample set may not be indicative of longer-term air concentrations? Would we expect higher (or lower) concentrations at other times of year?
 - \rightarrow The data we have collected appear to reflect air concentrations during the entire sampling period, with no indications from the on-site meteorological data that the sampling day conditions were inconsistent with conditions overall during this period.
 - \rightarrow Among the data collected for this site, we have none that would indicate generally higher (or lower) concentrations during other times of year. The wind flow patterns at the nearest NWS station during the sampling period appear to be representative of long-term wind flow at that site. The lack of long-term meteorological data at the school location, along with our finding that the wind patterns from the nearest NWS station differ slightly from those at the school, limits somewhat our ability to confidently predict longer-term wind patterns at the school (which might provide further evidence relevant to concentrations during other times).

B. Next Steps for Key Pollutants

- 1. Based on the analysis described here, EPA will not extend air toxics monitoring at this school.
- 2. EPA's ongoing research and national air toxics monitoring programs will continue to collect information on mobile source impacts on outdoor air nationally. EPA will also continue to work toward reductions in mobile source emissions nationally and to facilitate reductions in local areas (http://www.epa.gov/schoolair/mobile.html).
- 3. OEPA will continue to oversee industrial facilities in the area through air permits and other programs. OEPA will continue to implement reductions in mobile sources through implementation of national programs and its own programs.

VII. Figures and Tables

A. Tables

- 1. Elm Street Elementary School Key Pollutant Analysis.
- 2a. Elm Street Elementary School Key Pollutant (Diisocyanates) Concentrations and Meteorological Data.
- 2b. Elm Street Elementary School Key Pollutant (Benzene) Concentrations and Meteorological Data.

B. Figures

- 1. Elm Street Elementary School and Sources of Interest.
- 2. Elm Street Elementary School Key Pollutant (Benzene) Analysis.
- 3a. Elm Street Elementary School (Wauseon, OH) 1,6-HDI; MDI; and 2,4-TDI Concentration and Wind Information.
- 3b. Elm Street Elementary School (Wauseon, OH) Benzene Concentration and Wind Information.

VIII. Appendices

- A. Summary Description of Long-term Comparison Levels.
- B. National Air Toxics Trends Stations Measurements (2004-2008).
- C. Analysis of Other (non-key) Air Toxics Monitored at the School and Multiplepollutant Considerations.
- D. Elm Street Elementary School Pollutant Concentrations.
- E. Windroses for Toledo Express Airport NWS Station.



Distance from School
(mi)
0.03
0.06
0.03
0.59

Table 1. Elm Street Elementary School - Key Pollutant Analysis.

			95% Confidence	Long-term Co	omparison Level ^a
		Mean of	Interval on the		
Parameter	Units	Measurements	Mean	Cancer-Based ^b	Noncancer-Based ^c
Hexamethylene Diisocyanate (1,6-HDI)	µg/m ³	1	ND ^d	NA	0.01
Methylenediphenyl Diisocyanate, 4,4- (MDI)	µg/m ³	I	ND ^d	NA	0.6
Toluene Diisocyanate, 2,4- (2,4-TDI)	µg/m ³	I	ND ^d	9.1	0.07
Benzene	µg/m ³	0.62 ^e	0.42 - 0.83	13	30

 $\mu g/m^3$ micrograms per cubic meter

NA Not applicable

^a Details regarding these values are in the technical report, Schools Air Toxics Monitoring Activity (2009) Uses of Health Effects Information.

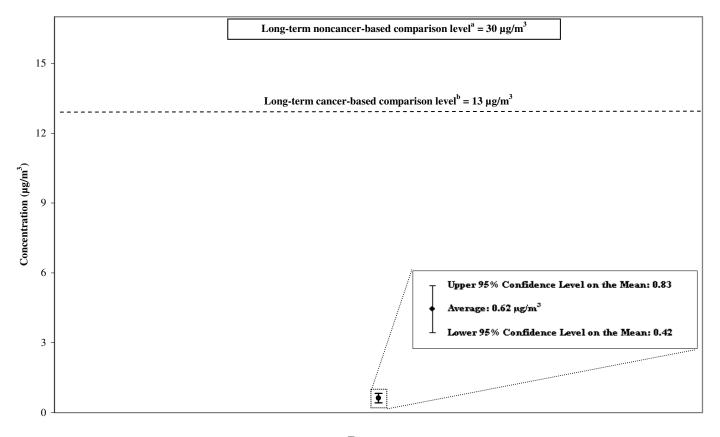
^b Air toxics for which the upper 95% confidence limit on the mean concentration is above this level will be fully discussed in the text and may be considered a priority for potential follow-up activities, if indicated in light of the full set of information available for the site. Findings of the upper 95% confidence limit below 1% of the comparison level (i.e., where the upper 95% confidence limit is below the corresponding 1-in-1-million cancer risk based concentration) are generally considered a low priority for follow-up activity. Situations where the summary statistics for a pollutant are below this comparison level but above 1% of this level are fully discussed in the text of the report.

^c Air toxics for which the upper 95% confidence limit on the mean concentration are near or below the noncancer-based comparison level are generally of low concern and will generally be considered a low priority for follow-up activity. Pollutants for which the 95% confidence limits extend appreciably above the noncancer-based comparison level are fully discussed in the school-specific report and may be considered a priority for follow-up activity, if indicated in light of the full set of information available for the site.

^d There were no detections of 1,6-HDI, MDI, or 2,4-TDI during the sampling period.

^e The mean of measurements for benzene is the average of all sample results, which include ten detections that ranged from 0.23 to 1.15 µg/m³.

Figure 2. Elm Street Elementary School - Key Pollutant (Benzene) Analysis.





^a Air toxics for which the upper 95% confidence limit on the mean concentration are near or below the noncancer-based comparison level are generally of low concern and will generally be considered a low priority for follow-up activity. Pollutants for which the 95% confidence limits extend appreciably above the noncancer-based comparison level are fully discussed in the school-specific report and may be considered a priority for follow-up activity, if indicated in light of the full set of information available for the site.

^b Air toxics for which the upper 95% confidence limit on the mean concentration is above this cancer-based comparison level will be fully discussed in the text and may be considered a priority for potential follow-up activities, if indicated in light of the full set of information available for the site. Findings of the upper 95% confidence limit below 1% of the comparison level (i.e., where the upper 95% confidence limit is below the corresponding 1-in-1-million cancer risk based concentration) are generally considered a low priority for follow-up activity. Situations where the summary statistics for a pollutant are below this comparison level but above 1% of this level are fully discussed in the text of the report.

Parameter	Units	9/4/2009	9/10/2009	9/16/2009	9/22/2009	10/4/2009	10/10/2009	10/16/2009	10/28/2009	11/3/2009	11/9/2009
Hexamethylene Diisocyanate (1,6-HDI)	µg/m ³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
Methylenediphenyl Diisocyanate, 4,4- (MDI)	µg/m ³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
Toluene Diisocyanate, 2,4- (2,4-TDI)	µg/m ³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
% Hours w/Wind Direction from Expected ZOI ^a	%	0.0	0.0	0.0	0.0	4.2	45.8	0.0	12.5	16.7	8.3
Wind Speed (avg. of hourly speeds)	mph	2.0	3.4	5.1	3.1	5.1	4.0	5.0	3.2	4.4	4.0
Wind Direction (avg. of unitized vector) ^b	deg.	33.9	73.6	63.3	184.1	258.5	288.4	30.5	10.8	284.3	257.5
% of Hours with Speed below 2 knots	%	75.0	37.5	16.7	33.3	20.8	20.8	4.2	16.7	16.7	20.8
Daily Average Temperature	°F	65.3	69.5	62.2	72.3	50.9	47.0	40.0	53.9	39.1	58.0
Daily Precipitation	inches	0.00	0.00	0.00	0.19	0.04	0.00	0.01	0.00	0.04	0.00

All precipitation and temperature data were from the Toledo Express Airport NWS Station.

^a Based on count of hours for which vector wind direction is from expected zone of influence.

^b Wind direction for each day is represented by values derived by scalar averaging of hourly estimates that were produced (by wind instrumentation's logger) as unitized vectors (specified as degrees from due north).

ND No detection of this chemical was registered by the laboratory analytical equipment. The method detection limit for MDI and 2,4-TDI is $0.190 \,\mu\text{g/m}^3$, while the method detection limit for 1,6-HDI is $0.380 \,\mu\text{g/m}^3$.

Parameter	Units	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/16/2009	10/22/2009	10/28/2009	11/3/2009 ^a	11/9/2009
Benzene	$\mu g/m^3$	0.978	0.627	0.416	0.521	0.23	0.339	0.515	0.729	1.15	0.716
% Hours w/Wind Direction from Expected ZOI ^b	%	100	100	100	100	100	100	100	100	100	100
Wind Speed (avg. of hourly speeds)	mph	2.0	3.4	5.1	3.1	10.0	5.0	5.5	3.2	3.5	4.0
Wind Direction (avg. of unitized vector) ^c	deg.	33.9	73.6	63.3	184.1	261.5	30.5	239.3	10.8	228.2	257.5
% of Hours with Speed below 2 knots	%	75.0	37.5	16.7	33.3	0.0	4.2	8.3	16.7	50.0	20.8
Daily Average Temperature	°F	65.3	69.5	62.2	72.3	56.8	40.0	55.5	53.9	36.9	58.0
Daily Precipitation	inches	0.00	0.00	0.00	0.19	0.11	0.01	0.03	0.04	0.00	0.00

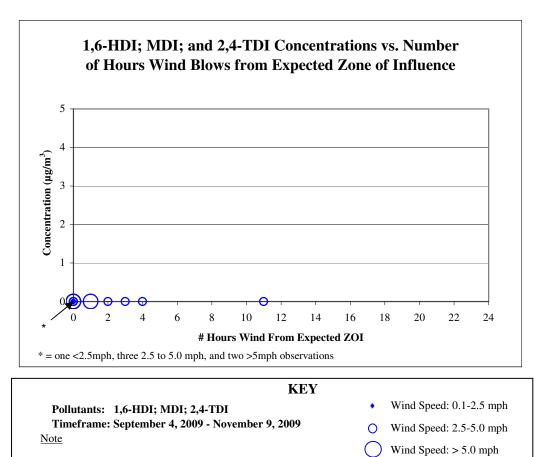
There was no hourly observation for one hour on this day. As a surrogate, the wind speed and wind direction was taken from the Toledo Express Airport NWS station.

All precipitation and temperature data were from the Toledo Express Airport NWS Station.

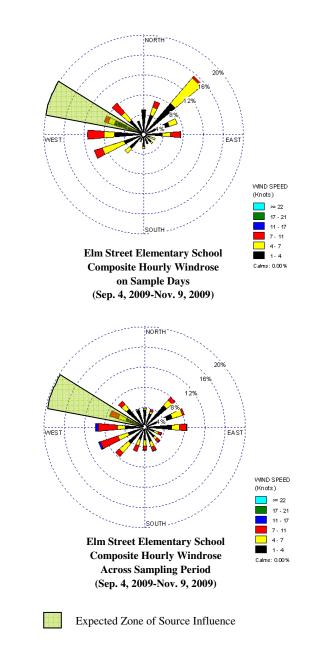
Due to issues with the timer, manual sampling was conducted on 11/3/2009. The sample start time was 11:15am, and continued for 24 hours into the following day. As such, the hourly meteorological data used to correlate the sample was adjusted accordingly.

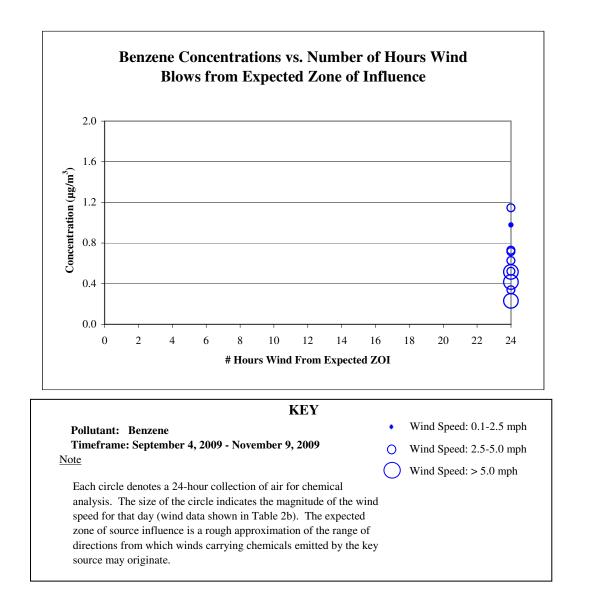
^b Based on count of hours for which vector wind direction is from expected zone of influence.

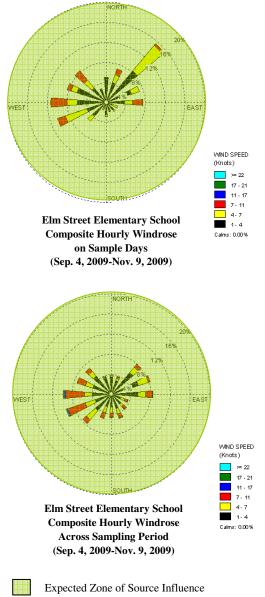
^c Wind direction for each day is represented by values derived by scalar averaging of hourly estimates that were produced (by wind instrumentation's logger) as unitized vectors (specified as degrees from due north).



Each circle denotes a 24-hour collection of air for chemical analysis. The size of the circle indicates the magnitude of the wind speed for that day (wind data shown in Table 2a). The expected zone of source influence is a rough approximation of the range of directions from which winds carrying chemicals emitted by the key source may originate.







Appendix A. Summary Description of Long-term Comparison Levels

In addressing the primary objective identified above, to investigate through the monitoring data collected for key pollutants at the school whether levels are of a magnitude, in light of health risk-based criteria, to indicate that follow-up activities be considered, we developed two types of long-term health risk-related comparison levels. These two types of levels are summarized below.¹⁵

Cancer-based Comparison Levels

- For air toxics where applicable, we developed cancer risk-based comparison levels to help us consider whether the monitoring data collected at the school indicate the potential for concentrations to pose incremental cancer risk above the range that EPA generally considers acceptable in regulatory decision-making to someone exposed to those concentrations continuously (24 hours a day, 7 days a week) over an entire lifetime.¹⁶ This general range is from 1 to 100 in a million.
- Air toxics with long-term mean concentrations below one one-hundredth of this comparison level would be below a comparably developed level for 1-ina-million risk (which is the lower bound of EPA's traditional acceptable risk range). Such pollutants, with long-term mean concentrations below the Agency's traditional acceptable risk range, are generally considered to pose negligible risk.
- Air toxics with long-term mean concentrations above the acceptable risk range would generally be a priority for follow-up activities. In this evaluation, we compare the upper 95% confidence limit on the mean concentration to the comparison level. Pollutants for which this upper limit falls above the comparison level are fully discussed in the school monitoring report and may be considered a priority for potential follow-up activities in light of the full set of information available for that site.
- Situations where the summary statistics for a pollutant are below the cancerbased comparison level but above 1% of that level are fully discussed in Appendix C.

¹⁵ These comparison levels are described in more detail *Schools Air Toxics Monitoring Activity (2009), Uses of Health Effects Information in Evaluating Sample Results.*

¹⁶ While no one would be exposed at a school for 24 hours a day, every day for an entire lifetime, we chose this worst-case exposure period as a simplification for the basis of the comparison level in recognition of other uncertainties in the analysis. Use of continuous lifetime exposure yields a lower, more conservative, comparison level than would use of a characterization more specific to the school population (e.g., 5 days a week, 8-10 hours a day for a limited number of years).

Noncancer-based Comparison Levels

- To consider concentrations of air toxics other than lead (for which we have a national ambient air quality standard) with regard to potential for health effects other than cancer, we derived noncancer-based comparison levels using EPA chronic reference concentrations (or similar values). A chronic reference concentration (RfC) is an estimate of a long-term continuous exposure concentration (24 hours a day, every day) without appreciable risk of adverse effect over a lifetime.¹⁷ This differs from the cancer risk-based comparison level in that it represents a concentration without appreciable risk *vs* a risk-based concentration.
- In using this comparison level in this initiative, the upper end of the 95% confidence limit on the mean is compared to the comparison level. Air toxics for which this upper confidence limit is near or below the noncancer-based comparison level (i.e., those for which longer-term average concentration estimates are below a long-term health-related reference concentration) are generally of low concern and will generally be considered a low priority for follow-up activity. Pollutants for which the 95% confidence limits extend appreciably above the noncancer-based comparison level are fully discussed below and may be considered a priority for follow-up activity if indicated in light of the full set of information available for the pollutant and the site.
- For lead, we set the noncancer-based comparison level equal to the level of the recently revised national ambient air quality standard (NAAQS). It is important to note that the NAAQS for lead is a 3-month rolling average of lead in total suspended particles. Mean levels for the monitoring data collected in this initiative that indicate the potential for a 3-month average above the level of the standard will be considered a priority for consideration of follow-up actions such as siting of a NAAQS monitor in the area.

In developing or identifying these comparison levels, we have given priority to use of relevant and appropriate air standards and EPA risk assessment guidance and precedents. These levels are based upon health effects information, exposure concentrations and risk estimates developed and assessed by EPA, the U.S. Agency for Toxic Substances and Disease Registry, and the California EPA. These agencies recognize the need to account for potential differences in sensitivity or susceptibility of different groups (e.g., asthmatics) or lifestages/ages (e.g., young children or the elderly) to a particular pollutant's effects so that the resulting comparison levels are relevant for these potentially sensitive groups as well as the broader population.

¹⁷ EPA defines the RfC as "an estimate (with uncertainty spanning perhaps an order of magnitude) of a continuous inhalation exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. It can be derived from a NOAEL, LOAEL, or benchmark concentration, with uncertainty factors generally applied to reflect limitations of the data used. Generally used in EPA's noncancer health assessments." <u>http://www.epa.gov/ncea/iris/help_gloss.htm#r</u>

		# Samples	%		Arithmetic	Geometric	5th	25th	50th	75th	95th
Pollutant	Units	Analyzed	Detections	Maximum	Mean ^b	Mean	Percentile	Percentile	Percentile	Percentile	Percentile
Acetonitrile	$\mu g/m^3$	1,804	69%	542.30	3.55	0.72	ND	ND	0.27	0.76	8.60
Acrylonitrile	$\mu g/m^3$	3,673	31%	5.51	0.06	0.10	ND	ND	ND	0.03	0.33
Benzene	$\mu g/m^3$	6,313	94%	10.19	1.03	0.84	ND	0.48	0.80	1.31	2.81
Benzyl chloride	$\mu g/m^3$	3,046	9%	2.49	0.01	0.05	ND	ND	ND	ND	0.05
Bromoform	$\mu g/m^3$	2,946	4%	1.18	0.01	0.16	ND	ND	ND	ND	ND
Bromomethane	$\mu g/m^3$	5,376	61%	120.76	0.11	0.05	ND	ND	0.03	0.05	0.12
Butadiene, 1,3-	$\mu g/m^3$	6,427	67%	15.55	0.10	0.09	ND	ND	0.05	0.13	0.38
Carbon disulfide	$\mu g/m^3$	1,925	91%	46.71	2.32	0.25	ND	0.03	0.09	0.96	12.65
Carbon tetrachloride	$\mu g/m^3$	6,218	86%	1.76	0.52	0.58	ND	0.47	0.57	0.65	0.87
Chlorobenzene	$\mu g/m^3$	5,763	30%	1.10	0.02	0.04	ND	ND	ND	0.01	0.11
Chloroethane	$\mu g/m^3$	4,625	37%	0.58	0.02	0.04	ND	ND	ND	0.03	0.08
Chloroform	$\mu g/m^3$	6,432	73%	48.05	0.17	0.14	ND	ND	0.10	0.17	0.61
Chloromethane	$\mu g/m^3$	5,573	95%	19.70	1.17	1.20	ND	1.03	1.18	1.36	1.68
Chloroprene	$\mu g/m^3$	2,341	11%	0.17	< 0.01	0.03	ND	ND	ND	ND	0.02
Dichlorobenzene, p-	$\mu g/m^3$	5,409	60%	13.65	0.19	0.16	ND	ND	ND	0.18	0.90
Dichloroethane, 1,1-	$\mu g/m^3$	5,670	16%	0.36	0.01	0.02	ND	ND	ND	ND	0.02
Dichloroethylene, 1,1-	$\mu g/m^3$	5,480	19%	0.44	0.01	0.02	ND	ND	ND	ND	0.04
Dichloromethane	$\mu g/m^3$	6,206	82%	214.67	0.59	0.34	ND	0.14	0.28	0.49	1.35
Dichloropropane,1,2-	µg/m ³	6,225	17%	1.80	0.01	0.03	ND	ND	ND	ND	0.04
Dichloropropylene, cis-1,3-	$\mu g/m^3$	4,705	18%	0.80	0.01	0.05	ND	ND	ND	ND	0.11
Dichloropropylene, trans -1,3-	$\mu g/m^3$	4,678	18%	1.13	0.02	0.05	ND	ND	ND	ND	0.11
Ethyl acrylate	µg/m ³	1,917	1%	0.08	<0.01	0.04	ND	ND	ND	ND	ND
Ethylbenzene	$\mu g/m^3$	6,120	84%	8.84	0.42	0.32	ND	0.10	0.29	0.53	1.33
Ethylene dibromide	µg/m ³	5,646	19%	4.15	0.01	0.05	ND	ND	ND	ND	0.05
Ethylene dichloride	µg/m ³	6,143	38%	4.49	0.03	0.05	ND	ND	ND	0.04	0.09
Hexachlorobutadiene	$\mu g/m^3$	3,727	20%	0.97	0.03	0.10	ND	ND	ND	ND	0.18
Methyl chloroform	$\mu g/m^3$	5,944	73%	3.17	0.09	0.10	ND	ND	0.08	0.11	0.20
Methyl isobutyl ketone	$\mu g/m^3$	2,936	60%	2.95	0.11	0.09	ND	ND	0.02	0.12	0.49
Methyl methacrylate	$\mu g/m^3$	1,917	9%	14.05	0.13	0.49	ND	ND	ND	ND	0.53
Methyl tert- butyl ether	$\mu g/m^3$	4,370	41%	20.50	0.28	0.12	ND	ND	ND	0.04	1.53
Styrene	$\mu g/m^3$	6,080	70%	27.22	0.16	0.11	ND	ND	0.05	0.16	0.60
Tetrachloroethane, 1,1,2,2-	$\mu g/m^3$	5,952	20%	2.47	0.02	0.04	ND	ND	ND	ND	0.07

Appendix B. National Air Toxics Trends Stations Measurements (2004-2008).^a

Pollutant	Units	# Samples Analyzed	% Detections		Arithmetic Mean ^b	Geometric Mean	5th Percentile	25th Percentile	50th Percentile	75th Percentile	95th Percentile
Tetrachloroethylene	µg/m ³	6,423	71%	42.12	0.28	0.20	ND	ND	0.13	0.27	0.88
Toluene	$\mu g/m^3$	5,947	95%	482.53	2.46	1.54	0.01	0.70	1.51	3.05	7.42
Trichlorobenzene, 1,2,4-	$\mu g/m^3$	4,301	21%	45.27	0.07	0.10	ND	ND	ND	ND	0.16
Trichloroethane,1,1,2-	µg/m ³	5,210	19%	5.89	0.01	0.04	ND	ND	ND	ND	0.05
Trichloroethylene	$\mu g/m^3$	6,410	46%	6.50	0.05	0.07	ND	ND	ND	0.05	0.22
Vinyl chloride	$\mu g/m^3$	6,284	18%	1.61	0.01	0.02	ND	ND	ND	ND	0.03
Xylene, <i>m/p</i> -	µg/m ³	4,260	90%	21.41	1.12	0.71	ND	0.26	0.69	1.43	3.65
Xylene, o-	$\mu g/m^3$	6,108	83%	9.21	0.41	0.30	ND	0.09	0.24	0.52	1.39

Appendix B. National Air Toxics Trends Stations Measurements (2004-2008).^a

Key Pollutant

ND No results of this chemical were registered by the laboratory analytical equipment.

^a The summary statistics in this table represent the range of actual daily HAP measurement values taken at NATTS sites from 2004 through 2008. These data were extracted from AQS in summer 2008 and 2009. During the time period of interest, there were 28 sites measuring VOCs, carbonyls, metals, and hexavalent chromium. We note that some sites did not sample for particular pollutant types during the initial year of the NATTS Program, which was 2004. Most of the monitoring stations in the NATTS network are located such that they are not expected to be impacted by single industrial sources. The concentrations typically measured at NATTS sites can thus provide a comparison point useful to considering whether concentrations measured at a school are likely to have been influenced by a significant nearby industrial source, or are more likely to be attributable to emissions from many small sources or to transported pollution from another area. For example, concentrations at a school above the 75th percentile may suggest that a nearby industrial source is affecting air quality at the school.

^b In calculations involving non-detects (ND), a value of zero is used.

Appendix C. Analysis of Other (non-key) Air Toxics Monitored at the School and Multiple-pollutant Considerations.

At each school, monitoring has been targeted to get information on a limited set of key hazardous air pollutants (HAPs).¹⁸ These pollutants are the primary focus of the monitoring activities at a school and a priority for us based on our emissions, modeling and other information. In analyzing air samples for these key pollutants, we have also obtained results for some other pollutants that are routinely included with the same test method. Our consideration of the data collected for these additional HAPs is described in the first section below. In addition to evaluating monitoring results for individual pollutants, we also considered the potential for cumulative impacts from multiple pollutants as described in the second section below (see Table C-1).

Other Air Toxics (HAPs)

- Do the monitoring data indicate elevated levels of any other air toxics or hazardous air pollutant (HAPs) that pose significant long-term health concerns?
 - \rightarrow The longer-term concentration estimates for the other HAPs monitored are below their long-term comparison levels.
 - → Further, for pollutants with cancer-based comparison levels, longer-term concentration estimates for all but two (carbon tetrachloride and 1,3-butadiene) are more than 100-fold lower.¹⁹
 - \rightarrow Additionally, each individual measurement for these pollutants is below the individual sample screening level developed for considering potential short-term exposures for that pollutant.²⁰

Additional Information on Two HAPs:

• The first of the two HAPs mentioned above is carbon tetrachloride. The mean and 95 percent upper bound on the mean for carbon tetrachloride are approximately 5% of the cancer-based comparison level. Additionally, a review of information available at other sites nationally shows that the mean concentration of carbon tetrachloride at this site is between the 75th and 95th percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B). Carbon tetrachloride is found globally as a result of its significant past uses in refrigerants and propellants for aerosol cans and its chemical persistence. Virtually all uses have been discontinued.

¹⁸ Section 112(b) of the Clean Air Act identifies 189 hazardous air pollutants, three of which have subsequently been removed from this list. These pollutants are the focus of regulatory actions involving stationary sources described by CAA section 112 and are distinguished from the six pollutants for which criteria and national ambient air quality standards (NAAQS) are developed as described in section 108. One of the criteria pollutants, lead, is also represented as lead compounds on the HAP list.

¹⁹ For pollutants with cancer-based comparison levels, this would indicate longer-term estimates below continuous (24 hours a day, 7 days a week) lifetime exposure concentrations associated with 10⁻⁶ excess cancer risk, respectively.

²⁰ The individual sample screening levels and their use is summarized on the website and described in detail in *Schools Air Toxics Monitoring Activity (2009), Uses of Health Effects Information in Evaluating Sample Results.*

However, it is still measured throughout the world as a result of its slow rate of degradation in the environment and global distribution in the atmosphere.

• The second HAP mentioned above is 1,3-butadiene. The mean and 95 percent upper bound on the mean for 1,3-butadiene are approximately 2-3% of the cancer-based comparison level. A review of information available at other sites nationally shows that the mean concentration of 1,3-butadiene at this site is between the 50th and 75th percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).

Multiple Pollutants

As described in the main body of the report and background materials, this initiative and the associated analyses are focused on investigation of key pollutants for each school that were identified by previous analyses. This focused design does not provide for the consideration of combined impacts of pollutants or stressors other than those monitored in this project. Broader analyses and those involving other pollutants may be the focus of other EPA activities.²¹

In our consideration of the potential for impacts from key pollutants at the monitored schools, we have also considered the potential for other monitored pollutants to be present at levels that in combination with the key pollutant levels contribute to an increased potential for cumulative impacts. This was done in cases where estimates of longer-term concentrations for any non-key HAPs are within an order of magnitude of their comparison levels even if these pollutant levels fall below the comparison levels. This analysis is summarized below.

- Do the data collected for the air toxics monitored indicate the potential for other monitored pollutants to be present at levels that in combination with the key pollutant levels indicate an increased potential for cumulative impacts of significant concern (e.g., that might warrant further investigation)?
 - → Although the multiple air toxics monitored at this site were below the levels of significant concern for multi-pollutant cumulative risk that had been suggested by the information available prior to monitoring, these results do indicate the influence of multiple moble source pollutants of concern that are the focus of EPA actions nationwide.
 - There were no HAPs monitored for which the longer-term concentration estimate was within an order of magnitude of their comparison levels.

²¹ General information on additional air pollutants is available at <u>http://www.epa.gov/air/airpollutants.html</u>.

		Mean of	95% Confidence		mparison Level ^b
Parameter	Units	Measurements ^a	Interval on the Mean	Cancer-Based ^c	Noncancer-Based ^d
No	on-Key H	APs with mean lowe	r than 10% of the lowest	comparison level	
Carbon Tetrachloride	$\mu g/m^3$	0.78	0.68 - 0.88	17	100
Butadiene, 1,3-	$\mu g/m^3$	0.06	0.03 - 0.09	3.3	2
Chloromethane	$\mu g/m^3$	1.18	1.01 - 1.35	NA	90
Bromomethane	µg/m ³	0.04	0.03 - 0.05	NA	5
Tetrachloroethylene	$\mu g/m^3$	0.09	0.06 - 0.11	17	270
Ethylbenzene	$\mu g/m^3$	0.17	0.10 - 0.23	40	1,000
Xylene, <i>m/p</i> -	µg/m ³	0.43	0.26 - 0.61	NA	100
Acetonitrile	µg/m ³	0.22	0.17 - 0.26	NA	60
Dichloromethane	$\mu g/m^3$	0.56	0.33 - 0.79	210	1,000
Xylene, o-	$\mu g/m^3$	0.17	0.10 - 0.23	NA	100
Chloroform	$\mu g/m^3$	0.13	0.11 - 0.15	NA	98
Methyl isobutyl ketone	µg/m ³	0.61	0.28 - 0.93	NA	3,000
Toluene	µg/m ³	0.95	0.53 - 1.38	NA	5,000
Carbon Disulfide	µg/m ³	0.07	0.05 - 0.08	NA	700
Styrene	µg/m ³	0.05	0.03 - 0.06	NA	1,000
Methyl Chloroform	$\mu g/m^3$	0.08	0.07 - 0.09	NA	5,000
Chloroethane	µg/m ³	0.02	0.01 - 0.03	NA	10,000
Dichlorobenzene, p-	µg/m ³	0.03 ^e	0.01 - 0.04 ^e	9.1	800
		Non-Key HAPs wit	h more than 50% ND Re	sults	
Trichloroethylene	$\mu g/m^3$	70% of the r	results were ND ^f	50	600
Trichloroethane, 1,1,2-	$\mu g/m^3$	90% of the r	esults were ND ^g	6.3	400
Vinyl chloride	$\mu g/m^3$	90% of the r	esults were ND ^h	11	100
		No other HAPs w	ere detected in any samp	oles	

Table C-1. Elm Street Elementary School - Other Monitored Pollutant Analysis.

 $\mu g/m^3$ micrograms per cubic meter

NA Not applicable

ND No detection of this chemical was registered by the laboratory analytical equipment.

^a Mean of measurements is the average of all sample results which include actual measured values. If no chemical was registered, then a value of zero is used when calculating the mean.

- ^b Details regarding these values are in the technical report, Schools Air Toxics Monitoring Activity (2009) Uses of Health Effects Information value is assumed to be zero in Evaluating Sample Results.
- ^c Air toxics for which the upper 95% confidence limit on the mean concentration is above this cancer-based cancer level will be fully discussed in the text and may be considered a priority for potential follow-up activities, if indicated in light of the full set of information available for the site. Findings of the upper 95% confidence limit below 1% of the comparison level (i.e., where the upper 95% confidence limit is below the corresponding 1-in-1-million cancer risk based concentration) are generally considered a low priority for follow-up activity. Situations where the summary statistics for a pollutant are below this comparison level but above 1% of this level are fully discussed in the text of the report.
- ^d Air toxics for which the upper 95% confidence limit on the mean concentration are near or below the noncancer-based comparison level are generally of low concern and will generally be considered a low priority for follow-up activity. Pollutants for which the 95% confidence limits extend appreciably above the noncancer-based comparison level are fully discussed in the school-specific report and may be considered a priority for follow-up activity, if indicated in light of the full set of information available for the site.

^e p-Dichlorobenzene was detected in 6 of 10 samples, ranging from 0.04 to 0.05 μ g/m³. The MDL is 0.024 μ g/m³.

^f Trichloroethylene was detected in only 3 of 10 samples, ranging from 0.086 to 0.19 μ g/m³. The MDL is 0.011 μ g/m³.

^g 1,1,2-Trichloroethane was detected in only 1 sample, with a value equal to 0.03 μ g/m³. The MDL is 0.016 μ g/m³.

^h Vinyl Chloride was detected in only 1 sample, with a value equal to 0.01 μ g/m³. The MDL is 0.005 μ g/m³.

Parameter	Units	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/4/2009	10/10/2009	10/16/2009	10/22/2009	10/28/2009	11/3/2009	11/9/2009	Sample Screening Level ^a
Hexamethylene Diisocyanate (1,6-HDI)	μg/m ³	ND	ND	ND	ND		ND	ND	ND		ND	ND	ND	0.2
Methylenediphenyl Diisocyanate, 4,4- (MDI)	µg/m ³	ND	ND	ND	ND		ND	ND	ND		ND	ND	ND	6
Toluene Diisocyanate, 2,4- (2,4-TDI)	μg/m ³	ND	ND	ND	ND		ND	ND	ND		ND	ND	ND	0.7
Benzene	μg/m ³	0.978	0.627	0.416	0.521	0.23			0.339	0.515	0.729	1.15	0.716	30
Carbon Tetrachloride	$\mu g/m^3$	0.787	0.774	0.781	0.951	1.01			0.724	0.875	0.62	0.724	0.53	200
Butadiene, 1,3-	µg/m ³	0.097	0.027	0.038	0.02	0.01			0.031	0.038	0.066	0.15	0.082	20
Chloromethane	µg/m ³	1.17	1.38	1.35	1.54	1.25			0.738	1.13	1.29	1.03	0.905	1,000
Bromomethane	$\mu g/m^3$	0.043	0.051	0.043	0.054	0.043			0.03	0.043	0.039	0.03	0.03	200
Tetrachloroethylene	μg/m ³	0.12	0.12	0.081	0.075	ND			0.06	0.081	0.12	0.12	0.095	1,400
Ethylbenzene	$\mu g/m^3$	0.23	0.10	0.13	0.078	0.03			0.32	0.14	0.17	0.27	0.20	40,000
Xylene, <i>m/p</i> -	$\mu g/m^3$	0.74	0.28	0.36	0.17	0.08			0.86	0.32	0.38	0.65	0.47	3,000
Acetonitrile	µg/m ³	0.264	0.328	0.197	0.316	0.15			0.217	0.15	0.203	0.15	0.178	600
Dichloromethane	$\mu g/m^3$	0.820	0.539	0.358	0.27	0.25			0.487	0.33	0.386	1.03	1.15	2,000
Xylene, o-	µg/m ³	0.32	0.11	0.15	0.074	0.043			0.24	0.13	0.14	0.27	0.18	9,000
Chloroform	µg/m ³	0.16	0.19	0.15	0.13	0.10			0.11	0.11	0.098	0.15	0.10	500
Methyl isobutyl ketone	μg/m ³	0.799	1.60	1.22	0.500	0.34			0.26	0.33	0.20	0.36	0.471	30,000
Toluene	μg/m ³	1.89	0.705	0.909	0.430	0.21			0.452	0.649	1.02	1.97	1.31	4,000
Carbon Disulfide	μg/m ³	0.075	0.093	0.059	0.069	0.056			0.090	0.059	0.044	0.02	0.10	7,000
Styrene	µg/m ³	0.081	0.03	0.051	0.043	0.03			ND	0.060	0.060	0.043	0.064	9,000
Methyl Chloroform	µg/m ³	0.076	0.093	0.071	0.10	0.082			0.093	0.076	0.066	0.066	0.055	10,000
Chloroethane	μg/m ³	0.029	0.040	0.02	0.02	ND			0.02	0.02	0.02	ND	0.02	40,000
Dichlorobenzene, p-	µg/m ³	0.05	0.05	0.04	ND	ND			ND	ND	0.04	0.05	0.04	10,000
Trichloroethylene	µg/m ³	0.11	ND	0.086	ND	ND			ND	ND	ND	0.19	ND	10,000
Trichloroethane, 1,1,2-	µg/m ³	0.03	ND	ND	ND	ND			ND	ND	ND	ND	ND	440
Vinyl chloride	$\mu g/m^3$	ND	ND	ND	ND	ND			0.01	ND	ND	ND	ND	1,000
Acrylonitrile	µg/m ³	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	200
Benzyl Chloride	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	140

Parameter	Units	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/4/2009	10/10/2009	10/16/2009	10/22/2009	10/28/2009	11/3/2009	11/9/2009	Sample Screening Level ^a
Bromoform	$\mu g/m^3$	ND	ND	ND	ND	ND	-	-	ND	ND	ND	ND	ND	6,400
Chlorobenzene	µg/m ³	ND	ND	ND	ND	ND	-	-	ND	ND	ND	ND	ND	10,000
Chloroprene	$\mu g/m^3$	ND	ND	ND	ND	ND	-		ND	ND	ND	ND	ND	70
Ethylene dibromide	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	12
Dichloroethane, 1,1-	$\mu g/m^3$	ND	ND	ND	ND	ND	1	1	ND	ND	ND	ND	ND	4,400
Dichloroethylene, 1,1-	$\mu g/m^3$	ND	ND	ND	ND	ND	-		ND	ND	ND	ND	ND	80
Dichloropropane, 1,2-	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	200
Dichloropropylene, cis-1,3-	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	14
Dichloropropylene, trans-1,3-	$\mu g/m^3$	ND	ND	ND	ND	ND	-		ND	ND	ND	ND	ND	14
Ethyl Acrylate	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	20,000
Ethylene dichloride	$\mu g/m^3$	ND	ND	ND	ND	ND	1	-	ND	ND	ND	ND	ND	270
Hexachlorobutadiene	$\mu g/m^3$	ND	ND	ND	ND	ND	-	-	ND	ND	ND	ND	ND	320
Methyl Methacrylate	$\mu g/m^3$	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	7,000
Methyl tert -Butyl Ether	μg/m ³	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	7,000
Tetrachloroethane, 1,1,2,2-	μg/m ³	ND	ND	ND	ND	ND	-	-	ND	ND	ND	ND	ND	120
Trichlorobenzene, 1,2,4-	μg/m ³	ND	ND	ND	ND	ND			ND	ND	ND	ND	ND	2,000

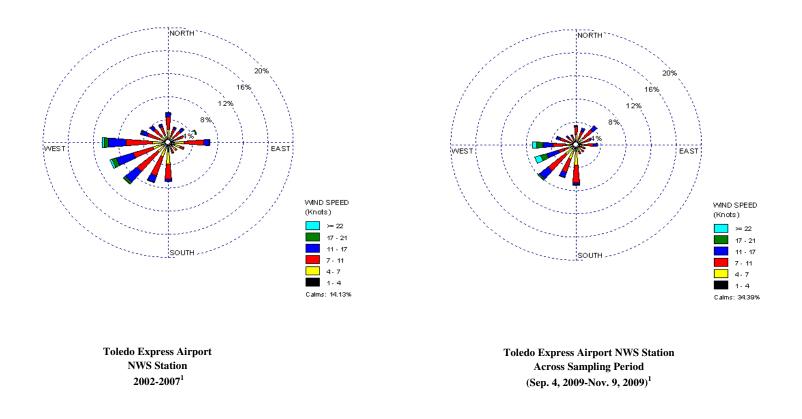
Key Pollutant

 $\mu g/m^3$ micrograms per cubic meter

-- No sample was collected for this pollutant on this day or the sample was invalidated.

ND No detection of this chemical was registered by the laboratory analytical equipment.

^a The individual sample screening levels and their use is summarized on the web site and described in detail in Schools Air Toxics Monitoring Activity (2009), "Uses of Health Effects Information in Evaluating Sample Results", see http://www.epa.gov/schoolair/pdfs/UsesOfHealthEffectsInfoinEvalSampleResults.pdf. These screening levels are based on consideration of exposure all day, every day over a period ranging up to at least a couple of weeks, and longer for some pollutants.



¹ Toledo Express Airport NWS Station (WBAN 94830) is 16.55 miles from Elm Street Elementary School.