## **SAT Initiative: Lincoln Elementary School (Warsaw, Indiana)**

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 Lincoln 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 manganese and nickel metals in air outside the school. The Indiana Department of Environmental Management recommended this school because it is near a large foundry and previous monitoring indicated potentially elevated levels of these pollutants.
- Air monitoring for manganese, nickel, and other metals in particulate matter less than 10 microns (PM<sub>10</sub>) was performed from August 23, 2009 through October 16, 2009.
- Measured levels of manganese (PM<sub>10</sub>) and nickel (PM<sub>10</sub>) and associated longer-term concentration estimates are below levels of significant concern for short- and long-term exposures. They are not as high as suggested by the information available prior to monitoring.
- The levels of manganese (PM<sub>10</sub>) measured in the outdoor air at this school do indicate influence of a nearby source. The levels of nickel (PM<sub>10</sub>) measured in the outdoor air at this school do not indicate influence of a nearby source.
- Based on the analysis described here, EPA will not extend air toxics monitoring at this school.
- IDEM will continue to oversee industrial facilities in the area through air permits and other 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 (http://www.epa.gov/schoolair/schools.html).

• 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).<sup>1</sup>
- 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
  - 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

<sup>1</sup> 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.

2

Information on health effects of air toxics being monitored<sup>2</sup> and educational materials describing risk concepts<sup>3</sup> 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 Indiana state air agency, the Indiana Department of Environmental Management (IDEM). We were interested in evaluating the ambient concentrations of manganese and nickel in air outside Lincoln Elementary. IDEM recommended this school because it is near a large foundry and previous monitoring indicated potentially elevated levels of these pollutants.

Monitoring commenced at this school on August 23, 2009 and continued through October 16, 2009. During this period, ten samples of airborne particles were collected using a PM<sub>10</sub> sampler.<sup>4</sup> The samples were analyzed for manganese and nickel (the key pollutants at this school) and a standardized set of additional metals that are routinely included in the analytical methods for the key pollutants. Additionally, ten VOC samples were collected and analyzed for 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 School Air Toxics Monitoring project (SAT) (<a href="http://www.epa.gov/schoolair/acrolein.html">http://www.epa.gov/schoolair/acrolein.html</a>). Sampling methodologies are described in EPA's schools air toxics monitoring plan (<a href="http://www.epa.gov/schoolair/techinfo.html">http://www.epa.gov/schoolair/techinfo.html</a>).

## 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)<sup>6</sup> 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

\_

<sup>&</sup>lt;sup>2</sup> For example, http://www.epa.gov/schoolair/pollutants.html, http://www.epa.gov/ttn/fera/risk atoxic.html.

<sup>&</sup>lt;sup>3</sup> For example, http://www.epa.gov/ttn/atw/3\_90\_022.html, http://www.epa.gov/ttn/atw/3\_90\_024.html.

<sup>&</sup>lt;sup>4</sup> In general this sampler collects airborne particles with a diameter of 10 microns or smaller, more of which would be considered to be in the respirable range which are what the health-based comparison levels for manganese and nickel are based on.

<sup>&</sup>lt;sup>5</sup> IDEM staff operated the monitors and sent the filters to the analytical laboratory under contract to EPA.

<sup>&</sup>lt;sup>6</sup> 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.

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 assessment 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 interval<sup>8</sup> 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). 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

<sup>&</sup>lt;sup>7</sup> 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.

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.

<sup>&</sup>lt;sup>9</sup> 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.

pollutant impacts.<sup>10</sup> 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:

- → 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 projects. The intent of this analysis is to make this 2-3 month 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:

- → 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.

-

<sup>&</sup>lt;sup>10</sup> 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).

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.<sup>11</sup> 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 Figures 1a-1b) with regard to the areas of interest identified below.

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

• The sampling data are below concentrations of significant concern for short-term exposures, and the related longer-term concentration estimates are below concentrations of significant concern for long-term exposures. The air sampling data indicate influence from a nearby source of manganese emissions, but do not indicate the influence of a nearby source of nickel emissions.

## Manganese, key pollutant:

- Do the monitoring data indicate influence from a nearby source?
  - $\rightarrow$  The monitoring data include some manganese (PM<sub>10</sub>) concentrations that are higher than concentrations commonly observed in other locations nationally. <sup>12</sup>
- Do the monitoring data indicate elevated levels that pose significant long-term health concerns?
  - → The monitoring data for manganese do not indicate levels of health concern for long-term exposures.
    - The estimate of longer-term manganese (PM<sub>10</sub>) concentration (i.e., the upper bound of the 95 percent confidence interval on the mean of the dataset) is well

\_

<sup>&</sup>lt;sup>11</sup> This is described in detail in *Schools Air Toxics Monitoring Activity (2009), Uses of Health Effects Information in Evaluating Sample Results.* 

<sup>&</sup>lt;sup>12</sup> For example, two of the manganese concentrations at this site (Table 2) were higher than 75 percent of samples collected at the National Air Toxics Trends Stations (NATTS) from 2004-2008 (Appendix B). Because the NATTS sites are generally sited so as not to be influenced by specific nearby sources, EPA is using the 75th percentile point of concentrations at these sites as a benchmark for indicating potential influence from a source nearby to this school.

below the long-term comparison level (Table 1). <sup>13</sup> This comparison level is a continuous exposure concentration (24 hours a day, all year, over a lifetime) associated with little risk of adverse effect; it is not an exposure concentration at which effects have been observed or are predicted to occur. <sup>14</sup>

- As manganese has not been found to be carcinogenic, it has no cancer-based comparison level.<sup>15</sup>
- → Additionally, we did not identify any concerns regarding short-term exposures as each individual measurement is below the individual sample screening level for manganese (which is based on consideration of exposure all day, every day over a period ranging up from a couple of weeks to longer for some pollutants). 11
- → In summary, none of the individual measurements indicate concentrations of concern for short-term exposures, and the combined contributions of all individual measurements in the estimate of longer-term concentration do not indicate a level of concern for long-term exposure.

## Nickel, key pollutant:

- Do the monitoring data indicate influence from a nearby source?
  - $\rightarrow$  The monitoring data do not include nickel (PM<sub>10</sub>) concentrations that are higher than concentrations commonly observed in other locations nationally. <sup>16</sup>
- Do the monitoring data indicate elevated levels that pose significant long-term health concerns?
  - → The monitoring data for nickel do not indicate levels of significant health concern for long-term exposures.
    - The estimate of longer-term nickel (PM<sub>10</sub>) concentration (i.e., the upper bound of the 95 percent confidence interval on the mean of the dataset) is well below both of the long-term comparison levels (Table 1). These comparison levels are based on consideration of continuous exposure concentrations (24 hours a day, all year, over a lifetime).
    - Further, the longer-term concentration estimate is more than 100-fold lower than the cancer-based comparison level, indicating the longer-term estimate is

1

<sup>&</sup>lt;sup>13</sup> The upper end of the interval is about twice the mean of the monitoring data, but less than 40% of the noncancerbased long-term comparison level.

<sup>&</sup>lt;sup>14</sup> The comparison level for manganese is based on the RfC. Manganese concentrations at which health effects have been documented are higher than the RfC (http://www.atsdr.cdc.gov/tfacts151.html, http://www.epa.gov/ttn/atw/hlthef/manganes.html#conversion)

<sup>15</sup> www.epa.gov/iris

<sup>&</sup>lt;sup>16</sup> For example, none of the nickel concentrations at this site (Table 2) were higher than 75 percent of samples collected at the National Air Toxics Trends Stations (NATTS) from 2004-2008 (Appendix B). Because the NATTS sites are generally sited so as not to be influenced by specific nearby sources, EPA is using the 75th percentile point of concentrations at these sites as a benchmark for indicating potential influence from a source nearby to this school.

<sup>&</sup>lt;sup>17</sup> The upper end of the interval is 1.3 times the mean of the monitoring data, but less than 1% of the noncancerbased long-term comparison level.

below a continuous (24 hours a day, 7 days a week) lifetime exposure concentration associated with 1-in-1 million additional cancer risk.

- → Additionally, we did not identify any concerns regarding short-term exposures as each individual measurement is below the individual sample screening level for nickel (which is based on consideration of exposure all day, every day over a period ranging from a couple of weeks to longer for some pollutants). <sup>11</sup>
- → In summary, none of the individual measurements indicate concentrations of concern for short-term exposures, and the combined contributions of all individual measurements in the estimate of longer-term concentration do not indicate a level of significant concern for long-term exposure.

## 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 level<sup>11</sup> 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)?
  - → The data collected for the key pollutants and other air toxics and the associated longer-term concentration estimates do not pose significant concerns for cumulative health risk from these pollutants (Appendix C). <sup>18</sup>

## 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

<sup>&</sup>lt;sup>18</sup> 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

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.

The meteorological station at Lincoln Elementary School collected wind speed and wind direction measurements beginning August 7, 2009, continuing through the sampling period (August 23, 2009-October 16, 2009), and ending May 18, 2010. As a result, on-site data for these meteorological parameters are available for all dates of sample collection, and also for a period before and after the sampling period, producing a continuous record of over nine months of on-site meteorological data. The meteorological data collected at the school on sampling days are presented in Figures 2a-2b and Table 2.

The nearest NWS station is at Goshen Municipal Airport in Goshen, IN. This station is approximately 20 miles north of the school. Measurements taken at that station include wind speed and direction, temperature, and precipitation. These are presented in Table 2 and Appendix E.

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

- Both the sampling results and the on-site wind data indicate that some of the air samples were collected on days when the nearby key source was contributing to conditions at the school location.
- The wind patterns at the monitoring site across sampling dates are similar to those observed across the record of on-site meteorological data during the sampling period, with respect to the expected zone of source influence.
- Our ability to provide a confident characterization of the wind flow patterns at the
  monitoring site over the long-term is limited, as the NWS station in Goshen Municipal
  Airport does not appear to represent the specific wind flow patterns at the school
  location.
- We lack long-term wind data at the monitoring site, and the wind pattern at the NWS station during the sampling period is generally not similar to the historical long-term wind flow pattern at that same NWS station. Therefore, the 2-month sampling period may not be representative of year-round wind patterns.
- What is the direction of the key source of manganese and nickel emissions in relation to the school location?
  - → The nearby industrial facility emitting the key pollutants into the air (described in section III above) lies about 0.2 miles south of the school.
  - → Using the property boundaries of the full facility (in lieu of information regarding the location of specific sources of manganese and nickel 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 146 to 214 degrees, is referred to here as the expected zone of source influence (ZOI).
- On days the air samples were collected, how often did wind come from direction of the key source?
  - → There were four sampling days in which a portion of the winds were from the expected ZOI (Table 2, Figures 2a-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?
  - → Wind patterns across the air monitoring days appear similar to those observed over the record of on-site meteorological data during the sampling period, with respect to the expected zone of source influence.
  - → We note that wind patterns at the nearest NWS station at Goshen Municipal Airport during the sampling period are not similar those recorded at the NWS station over the long-term (2002-2007 period; Appendix E). 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 Goshen Municipal Airport (see below).
- How do wind patterns at the school compare to those at the Goshen Municipal Airport 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 northeast, while those at the NWS station are somewhat more from the east. The windroses for the two sites during the sampling period (Figures 2a-2b and Appendix E) show differences in wind flow patterns.
- Are there other meteorological patterns that may influence the measured concentrations at the school monitoring site?
  - → 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 manganese and nickel (described in section III above) has an operating permit issued by IDEM that includes operating requirements.

<sup>19</sup> Operating permits, which are issued to air pollution sources under the Clean Air Act, are described at: http://www.epa.gov/air/oaqps/permits/

10

- Information from the nearby source indicates that this facility was operating at 36 percent of maximum permitted levels during the sampling period, in comparison to its typical 51 percent.
- The most recently available manganese and nickel emissions for this source (2008 TRI) are lower than those relied upon in previous modeling analysis for this area (e.g., 2002 NATA, 2005 TRI).

# VI. Integrated Summary and Next Steps

## A. Summary of Key Findings

- 1. What are the key HAPs for this school?
  - → Manganese and nickel are the key HAPs for this school, identified based on emissions information considered in identifying the school for monitoring. The ambient air concentrations on two days during the monitoring period indicate contributions from a source of manganese in the area.
- 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?
  - → The measured levels and associated longer-term concentration estimates are not as high as suggested by the information available prior to monitoring and are below levels of concern for long-term exposures.
- 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?
  - → The data we have collected appear to reflect air concentrations during the entire monitoring period, with no indications from the on-site meteorological data that the sampling day conditions were inconsistent with conditions overall during this period.
  - → Among the data collected for this site, we have none that would indicate generally higher (or lower) concentrations during other times of year.

#### **B.** Next Steps for Key Pollutants

- 1. Based on the analysis described here, EPA will not extend air toxics monitoring at this school.
- 2. IDEM will continue to oversee industrial facilities in the area through air permits and other programs.

#### VII. Figures and Tables

#### A. Tables

1. Lincoln Elementary School – Key Pollutant Analysis.

2. Lincoln Elementary School Key Pollutant Concentrations and Meteorological Data.

## **B.** Figures

- 1a. Lincoln Elementary School Key Pollutant (Manganese (PM<sub>10</sub>)) Analysis.
- 1b. Lincoln Elementary School Key Pollutant (Nickel (PM<sub>10</sub>)) Analysis.
- 2a. Lincoln Elementary School (Warsaw, IN) Manganese (PM<sub>10</sub>) Concentration and Wind Information.
- 2b. Lincoln Elementary School (Warsaw, IN) Nickel (PM<sub>10</sub>) 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 Multiple-pollutant Considerations.
- D. Lincoln Elementary School Pollutant Concentrations.
- E. Windroses for Goshen Municipal Airport NWS Station.

Table 1. Lincoln Elementary School - Key Pollutant Analysis.

			95% Confidence	Long-term Comparison Level <sup>a</sup>			
		Mean of	Interval on the				
Parameter	Units	Measurements	Mean	Cancer-Based <sup>b</sup>	Noncancer-Based <sup>c</sup>		
Manganese (PM <sub>10</sub> )	ng/m <sup>3</sup>	9.14 <sup>d</sup>	0 - 19.2	NA	50		
Nickel (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.35 <sup>e</sup>	0.24 - 0.47	420	90		

ng/m³ nanograms per cubic meter NA Not applicable

<sup>&</sup>lt;sup>a</sup> Details regarding these values are in the technical report, Schools Air Toxics Monitoring Activity (2009) Uses of Health Effects Information.

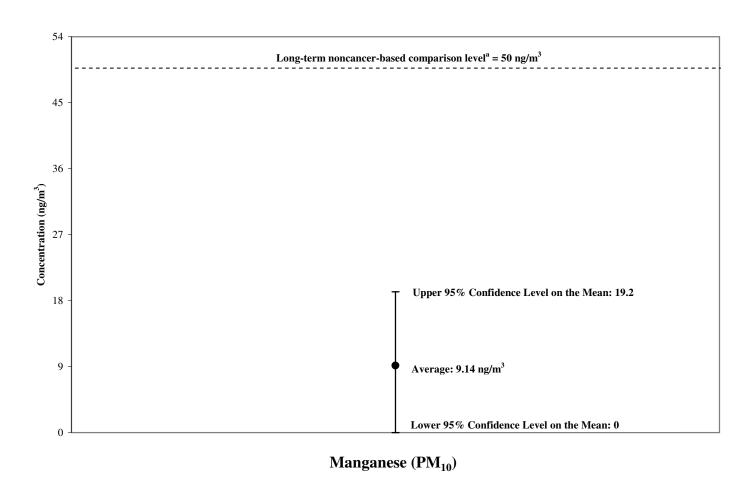
b Air toxics for which the upper 95% confidence limit on the mean concentration is above this level will be fully discussed in the text and may be considered a priority for potential follow-up activities, if indicated in light of the full set of information available for the site. Findings of the upper 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.

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.

<sup>&</sup>lt;sup>d</sup> The mean of measurements for manganese (PM<sub>10</sub>) is the average of all sample results, which include ten detections that ranged from 0.62 to 48.3 ng/m<sup>3</sup>.

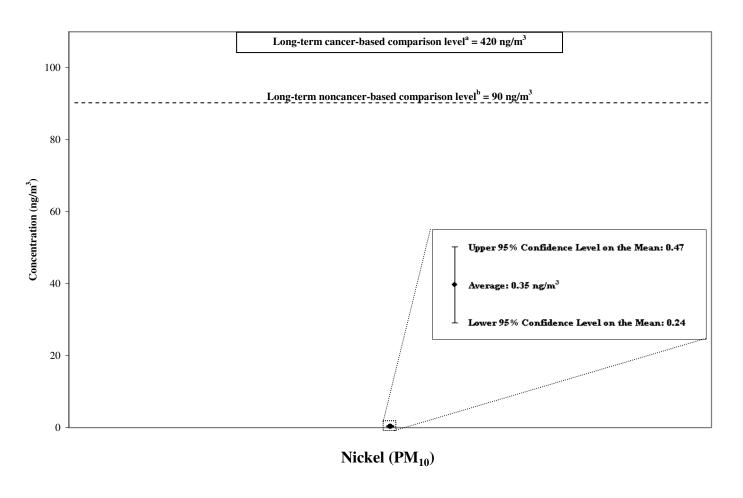
<sup>&</sup>lt;sup>e</sup> The mean of measurements for nickel (PM<sub>10</sub>) is the average of all sample results, which include ten detections that ranged from 0.09 to 0.58 ng/m<sup>3</sup>.

Figure 1a. Lincoln Elementary School - Key Pollutant (Manganese (PM<sub>10</sub>)) Analysis.



<sup>a</sup> 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.

Figure 1b. Lincoln Elementary School - Key Pollutant (Nickel (PM<sub>10</sub>)) Analysis.



<sup>&</sup>lt;sup>a</sup> 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.

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.

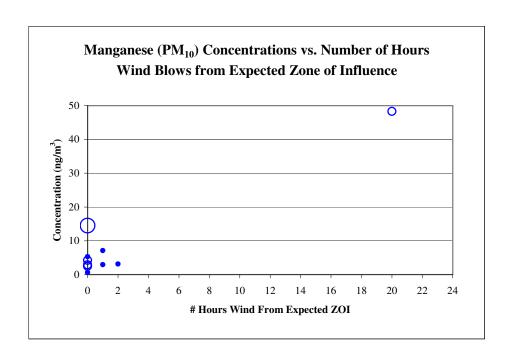
Table 2. Lincoln Elementary School Key Pollutant Concentrations and Meteorological Data.

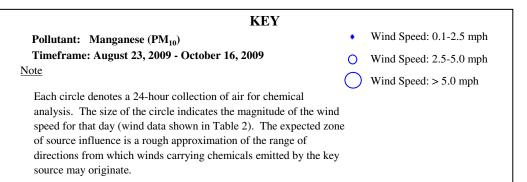
Parameter	Units	8/23/2009	8/29/2009	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/4/2009	10/10/2009	10/16/2009
Manganese (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.62	2.89	7.08	5.30	4.21	48.3	14.5	2.94	3.12	2.44
Nickel (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.09	0.36	0.58	0.41	0.41	0.58	0.20	0.32	0.39	0.17
% Hours w/Wind Direction from Expected ZOI <sup>a</sup>	%	0.0	0.0	4.2	0.0	0.0	83.3	0.0	4.2	8.3	0.0
Wind Speed (avg. of hourly speeds)	mph	2.5	3.3	1.7	2.0	2.7	3.2	6.0	2.4	2.3	4.7
Wind Direction (avg. of unitized vector) <sup>b</sup>	deg.	348.7	289.2	50.1	88.0	60.7	172.4	290.7	282.1	295.4	60.4
% of Hours with Speed below 2 knots	%	37.5	33.3	66.7	66.7	41.7	25.0	0.0	41.7	58.3	0.0
Daily Average Temperature	° F	62.3	60.9	63.7	67.1	61.0	70.1	54.8	49.4	42.5	39.3
Daily Precipitation	inches	0.01	0.00	0.00	0.00	0.00	1.08	0.21	0.01	0.00	0.02

All precipitation and temperature data were from the Goshen Municipal Airport NWS Station.

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

Figure 2a. Lincoln Elementary School (Warsaw, IN) Manganese (PM<sub>10</sub>) Concentration and Wind Information.





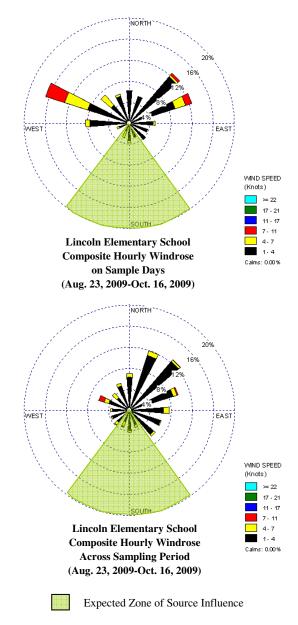
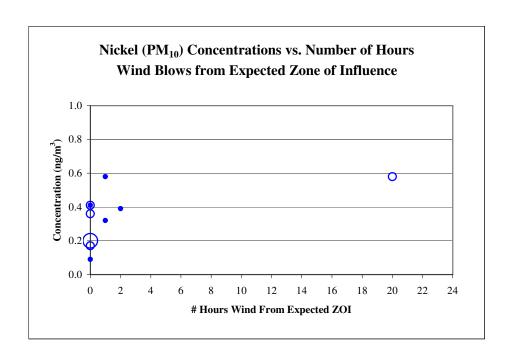
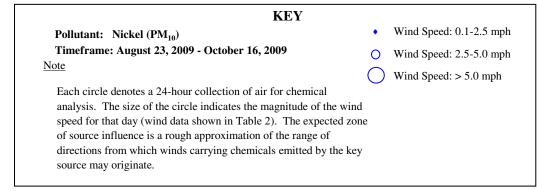
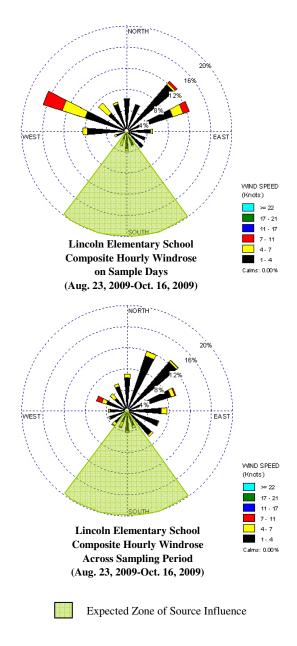


Figure 2b. Lincoln Elementary School (Warsaw, IN) Nickel (PM<sub>10</sub>) Concentration and Wind Information.







## 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.<sup>20</sup>

#### 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.<sup>21</sup> 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 cancer-based comparison level but above 1% of that level are fully discussed in Appendix C.

-

<sup>&</sup>lt;sup>20</sup> These comparison levels are described in more detail *Schools Air Toxics Monitoring Activity* (2009), *Uses of Health Effects Information in Evaluating Sample Results*.

<sup>&</sup>lt;sup>21</sup> 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.<sup>22</sup> 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.

<sup>22</sup> 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

 $EPA's \ noncancer \ health \ assessments." \ http://www.epa.gov/ncea/iris/help\_gloss.htm \#r$ 

\_

Appendix B. National Air Toxics Trends Stations Measurements (2004-2008).<sup>a</sup>

		# Samples	%		Arithmetic	Geometric	5th	25th	50th	75th	95th
Pollutant	Units	Analyzed	Detections	Maximum	Mean <sup>b</sup>	Mean	Percentile	Percentile	Percentile	Percentile	Percentile
Antimony (PM <sub>10</sub> )	ng/m <sup>3</sup>	2,372	94%	43.30	1.71	1.21	ND	0.60	1.13	2.17	4.33
Arsenic (PM <sub>10</sub> )	ng/m <sup>3</sup>	5,076	86%	47.70	0.93	0.70	ND	0.29	0.56	1.02	2.89
Beryllium (PM <sub>10</sub> )	ng/m <sup>3</sup>	4,771	64%	1.97	0.05	0.02	ND	ND	< 0.01	0.02	0.50
Cadmium (PM <sub>10</sub> )	ng/m <sup>3</sup>	4,793	85%	15.30	0.27	0.17	ND	0.05	0.13	0.29	0.94
Chromium (PM <sub>10</sub> )	ng/m <sup>3</sup>	5,094	92%	172.06	2.71	1.66	ND	0.93	1.98	2.85	7.10
Cobalt (PM <sub>10</sub> )	ng/m <sup>3</sup>	2,614	91%	20.30	0.28	0.18	ND	0.08	0.15	0.27	1.00
Manganese (PM <sub>10</sub> )	ng/m <sup>3</sup>	4,793	99%	734.00	10.39	5.20	< 0.01	2.41	4.49	9.96	33.78
Mercury (PM <sub>10</sub> )	ng/m <sup>3</sup>	1,167	81%	2.07	0.07	0.04	ND	0.01	0.02	0.06	0.32
Nickel (PM <sub>10</sub> )	ng/m <sup>3</sup>	4,815	90%	110.10	2.05	1.49	ND	0.74	1.44	2.50	5.74
Selenium (PM <sub>10</sub> )	ng/m <sup>3</sup>	2,382	96%	13.00	1.10	0.53	< 0.01	0.24	0.53	1.07	5.50
Acetonitrile	μg/m <sup>3</sup>	1,804	69%	542.30	3.55	0.72	ND	ND	0.27	0.76	8.60
Acrylonitrile	μg/m <sup>3</sup>	3,673	31%	5.51	0.06	0.10	ND	ND	ND	0.03	0.33
Benzene	μg/m <sup>3</sup>	6,313	94%	10.19	1.03	0.84	ND	0.48	0.80	1.31	2.81
Bromomethane	μg/m <sup>3</sup>	5,376	61%	120.76	0.11	0.05	ND	ND	0.03	0.05	0.12
Butadiene, 1,3-	μg/m <sup>3</sup>	6,427	67%	15.55	0.10	0.09	ND	ND	0.05	0.13	0.38
Carbon disulfide	μg/m <sup>3</sup>	1,925	91%	46.71	2.32	0.25	ND	0.03	0.09	0.96	12.65
Carbon tetrachloride	μg/m <sup>3</sup>	6,218	86%	1.76	0.52	0.58	ND	0.47	0.57	0.65	0.87
Chloro-1,3-butadiene, 2-	μg/m <sup>3</sup>	2,341	11%	0.17	< 0.01	0.03	ND	ND	ND	ND	0.02
Chlorobenzene	μg/m <sup>3</sup>	5,763	30%	1.10	0.02	0.04	ND	ND	ND	0.01	0.11
Chloroethane	μg/m <sup>3</sup>	4,625	37%	0.58	0.02	0.04	ND	ND	ND	0.03	0.08
Chloroform	μg/m <sup>3</sup>	6,432	73%	48.05	0.17	0.14	ND	ND	0.10	0.17	0.61
Chloromethane	μg/m <sup>3</sup>	5,573	95%	19.70	1.17	1.20	ND	1.03	1.18	1.36	1.68
Chlorotoluene, alpha-	μg/m <sup>3</sup>	3,046	9%	2.49	0.01	0.05	ND	ND	ND	ND	0.05
Dibromoethane, 1,2-	μg/m <sup>3</sup>	5,646	19%	4.15	0.01	0.05	ND	ND	ND	ND	0.05
Dichlorobenzene, p-	μg/m <sup>3</sup>	5,409	60%	13.65	0.19	0.16	ND	ND	ND	0.18	0.90
Dichloroethane, 1,1-	μg/m <sup>3</sup>	5,670	16%	0.36	0.01	0.02	ND	ND	ND	ND	0.02
Dichloroethylene, 1,1-	μg/m <sup>3</sup>	5,480	19%	0.44	0.01	0.02	ND	ND	ND	ND	0.04
Dichloropropane,1,2-	μg/m <sup>3</sup>	6,225	17%	1.80	0.01	0.03	ND	ND	ND	ND	0.04
Dichloropropylene, Cis-1,3-	μg/m <sup>3</sup>	4,705	18%	0.80	0.01	0.05	ND	ND	ND	ND	0.11
Dichloropropylene, Trans -1,3-	μg/m <sup>3</sup>	4,678	18%	1.13	0.02	0.05	ND	ND	ND	ND	0.11
Ethyl acrylate	μg/m <sup>3</sup>	1,917	1%	0.08	< 0.01	0.04	ND	ND	ND	ND	ND

Appendix B. National Air Toxics Trends Stations Measurements (2004-2008).<sup>a</sup>

Pollutant	Units	# Samples Analyzed	% Detections	Maximum	Arithmetic Mean <sup>b</sup>	Geometric Mean	5th Percentile	25th Percentile	50th Percentile	75th Percentile	95th Percentile
Ethylbenzene	μg/m <sup>3</sup>	6,120	84%	8.84	0.42	0.32	ND	0.10	0.29	0.53	1.33
Ethylene dichloride	μg/m <sup>3</sup>	6,143	38%	4.49	0.03	0.05	ND	ND	ND	0.04	0.09
Hexachloro-1,3-butadiene	μg/m <sup>3</sup>	3,727	20%	0.97	0.03	0.10	ND	ND	ND	ND	0.18
	μg/m <sup>3</sup>		9%	14.05	0.13	0.49	ND	ND	ND	ND	0.53
Methyl tert -butyl ether	μg/m <sup>3</sup>	4,370	41%	20.50	0.28	0.12	ND	ND	ND	0.04	1.53
Methyl-2-pentanone, 4-	μg/m <sup>3</sup>	2,936	60%	2.95	0.11	0.09	ND	ND	0.02	0.12	0.49
Methylene chloride	μg/m <sup>3</sup>	6,206	82%	214.67	0.59	0.34	ND	0.14	0.28	0.49	1.35
	μg/m <sup>3</sup>		70%	27.22	0.16	0.11	ND	ND	0.05	0.16	0.60
	μg/m <sup>3</sup>	5,952	20%	2.47	0.02	0.04	ND	ND	ND	ND	0.07
Tetrachloroethylene	μg/m <sup>3</sup>	6,423	71%	42.12	0.28	0.20	ND	ND	0.13	0.27	0.88
	μg/m <sup>3</sup>	5,947	95%	482.53	2.46	1.54	0.01	0.70	1.51	3.05	7.42
	μg/m <sup>3</sup>	2,946	4%	1.18	0.01	0.16	ND	ND	ND	ND	ND
Trichlorobenzene, 1,2,4-	μg/m <sup>3</sup>	4,301	21%	45.27	0.07	0.10	ND	ND	ND	ND	0.16
Trichloroethane, 1,1,1-	μg/m <sup>3</sup>	5,944	73%	3.17	0.09	0.10	ND	ND	0.08	0.11	0.20
Trichloroethane,1,1,2-	μg/m <sup>3</sup>	5,210	19%	5.89	0.01	0.04	ND	ND	ND	ND	0.05
Trichloroethylene	μg/m <sup>3</sup>	6,410	46%	6.50	0.05	0.07	ND	ND	ND	0.05	0.22
Vinyl chloride	μg/m <sup>3</sup>	6,284	18%	1.61	0.01	0.02	ND	ND	ND	ND	0.03
Xylene, m/p-	μg/m <sup>3</sup>	4,260	90%	21.41	1.12	0.71	ND	0.26	0.69	1.43	3.65
Xylene, o-	μg/m <sup>3</sup>	6,108	83%	9.21	0.41	0.30	ND	0.09	0.24	0.52	1.39

**Key Pollutant** 

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

<sup>&</sup>lt;sup>a</sup> 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.

<sup>&</sup>lt;sup>b</sup> 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).<sup>23</sup> 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?
  - → 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 one of these (chromium) is more than tenfold lower and all but six (also arsenic, benzene, carbon tetrachloride, 1,3-butadiene, and tetrachloroethylene) are more than 100-fold lower. <sup>24</sup>
  - → Additionally each individual measurement for these pollutants is below the individual sample (short-term) screening level developed for considering potential short-term exposures for that pollutant.<sup>25</sup>

#### Additional Information on Six HAPs

• The first HAP mentioned above is chromium. The comparison values for chromium are conservatively based on the most toxic form of chromium (hexavalent chromium,  $Cr^{+6}$ ) which is only a fraction of the chromium. Nonetheless, the longer-term concentration estimate for chromium (PM<sub>10</sub>) is well below even these very restrictive comparison values. The mean and 95 percent upper bound on the mean for chromium (PM<sub>10</sub>) are approximately 39-45% of the lowest comparison level. Further, as  $Cr^{+6}$  is commonly

\_

<sup>&</sup>lt;sup>23</sup> 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.

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<sup>-5</sup> and 10<sup>-6</sup> excess cancer risk, respectively.

<sup>&</sup>lt;sup>25</sup> 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*.

only a small fraction of the total,  $^{26}$  the levels of  $Cr^{+6}$  in these samples would be expected to be appreciably lower than this. A review of information available at other sites nationally shows that the mean concentration of chromium ( $PM_{10}$ ) at this site is between the  $75^{th}$  and  $95^{th}$  percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).

- The second HAP mentioned above is arsenic. The mean and 95% percent upper bound on the mean for arsenic (PM<sub>10</sub>) are approximately 4-6% of the lowest comparison level. A review of information available at other sites nationally shows that the mean concentration of arsenic (PM<sub>10</sub>) at this site is between the 50<sup>th</sup> and 75<sup>th</sup> percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).
- The third HAP mentioned above is benzene. The mean and 95% percent upper bound on the mean for benzene are approximately 4-7% of the lowest comparison level. A review of information available at other sites nationally shows that the mean concentration of benzene at this site is between the 25<sup>th</sup> and 50<sup>th</sup> percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).
- The fourth HAP mentioned above is carbon tetrachloride. The mean and 95 percent upper bound on the mean for carbon tetrachloride are approximately 4-5% of the lowest comparison level. A review of information available at other sites nationally shows that the mean concentration of carbon tetrachloride at this site is between the 75<sup>th</sup> and 95<sup>th</sup> 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. 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 fifth HAP mentioned above is 1,3-butadiene. The mean and 95% percent upper bound on the mean for 1,3-butadiene are approximately 2% of the lowest 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 50<sup>th</sup> and 75<sup>th</sup> percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).
- The final HAP mentioned above is tetrachloroethylene. The mean and 95% percent upper bound on the mean for tetrachloroethylene are approximately 1% of the lowest comparison level. A review of information available at other sites nationally shows that the mean concentration of for tetrachloroethylene at this site is between the 25<sup>th</sup> and 50<sup>th</sup> percentile of samples collected from 2004 to 2008 (the most recently compiled period) at the NATTS sites (Appendix B).

<sup>26</sup> Data in EPA's Air Quality System for locations that are not near a facility emitting hexavalent chromium indicate hexavalent chromium concentrations to comprise less than approximately 10% of total chromium concentrations.

C-2

#### **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.<sup>27</sup>

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)?
  - → The data collected for the key and other air toxics and the associated longer-term concentration estimates do not together pose significant concerns for cumulative health risk from these pollutants.
    - In addition to the key pollutant manganese, the only other HAP monitored whose longer-term concentration estimates is more than ten percent of its lowest comparison level is chromium. The lowest comparison level for chromium (conservatively based on the most toxic form of chromium, hexavalent chromium)<sup>28</sup> is based on carcinogenic risk to the respiratory system; however, hexavalent chromium is commonly a small fraction of the total chromium reported. The comparison level for manganese is based on non-carcinogenic effects to the nervous system. Taken together, these considerations reduce any concerns for cumulative health risk from these pollutants.

<sup>27</sup> General information on additional air pollutants is available at http://www.epa.gov/air/airpollutants.html.

C-3

The noncancer-based comparison level for chromium is much higher than the cancer-based level and is based on risk of other effects posed to the respiratory system by hexavalent chromium in particulate form.

 Table C-1. Lincoln Elementary School - Other Monitored Pollutant Analysis.

			95% Confidence	Long-term Co	omparison Level <sup>b</sup>	
		Mean of	Interval on the		d	
Parameter		Measurements <sup>a</sup>		Cancer-Based <sup>c</sup>	Noncancer-Based <sup>d</sup>	
			an 10% of the lowes			
Chromium (PM <sub>10</sub> )	ng/m <sup>3</sup>		2.75 - 3.77	8.3 e	100 <sup>e</sup>	
Non-K			an 10% of the lowest	_		
Arsenic (PM <sub>10</sub> )	ng/m <sup>3</sup>		0.43 - 1.42	23	15	
Benzene	$\mu g/m^3$	0.58	0.28 - 0.88	13	30	
Carbon Tetrachloride	$\mu g/m^3$	0.72	0.51 - 0.93	17	100	
Butadiene, 1,3-	$\mu g/m^3$	0.06	0.03 - 0.08	3.3	2	
Chloromethane	$\mu g/m^3$	1.26	1.03 - 1.49	NA	90	
Cadmium (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.13	0.02 - 0.25	56	10	
Bromomethane	μg/m <sup>3</sup>	0.05	0.04 - 0.06	NA	5	
Tetrachloroethylene	μg/m <sup>3</sup>	0.12	0.06 - 0.18	17	270	
Dichlorobenzene, p-	μg/m <sup>3</sup>	0.05	0.03 - 0.08	9.1	800	
Acetonitrile	$\mu g/m^3$	0.31	0.18 - 0.43	NA	60	
Antimony (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.76	0.44 - 1.09	NA	200	
Vinyl chloride	μg/m <sup>3</sup>	0.04	0.03 - 0.05	11	100	
Ethylbenzene	μg/m <sup>3</sup>	0.11	0.08 - 0.15	40	1000	
Xylene, <i>m/p</i> -	$\mu g/m^3$	0.26	0.17 - 0.34	NA	100	
Dichloromethane	μg/m <sup>3</sup>	0.33	0.26 - 0.39	210	1000	
Chloroform	μg/m <sup>3</sup>	0.13	0.11 - 0.15	NA	98	
Xylene, o-	μg/m <sup>3</sup>	0.11	0.08 - 0.15	NA	100	
Cobalt (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.05	0.02 - 0.07	NA	100	
Toluene	μg/m <sup>3</sup>	1.36	0.71 - 2.01	NA	5000	
Methyl isobutyl ketone	μg/m <sup>3</sup>	0.75	0.13 - 1.36	NA	3000	
Carbon Disulfide	μg/m <sup>3</sup>	0.14	0.08 - 0.21	NA	700	
Styrene	μg/m <sup>3</sup>	0.06	0.02 - 0.10	NA	1000	
Mercury (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.011	0.001 - 0.022	NA	300 <sup>f</sup>	
Selenium (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.59	0.21 - 0.97	NA	20000	
Methyl chloroform	μg/m <sup>3</sup>	0.08	0.05 - 0.10	NA	5000	
Chloroethane	μg/m <sup>3</sup>	0.05	0.04 - 0.06	NA	10000	

Table C-1. Lincoln Elementary School - Other Monitored Pollutant Analysis.

			95% Confidence	Long-term Comparison Level <sup>b</sup>								
Parameter	Units	Mean of Measurements <sup>a</sup>	Interval on the Mean	Cancer-Based <sup>c</sup>	Noncancer-Based <sup>d</sup>							
	Non-Key HAPs with more than 50% ND Results											
Trichloroethylene	μg/m <sup>3</sup>	80% of the re	esults were ND <sup>g</sup>	NA	5000							
Beryllium (PM <sub>10</sub> )	ng/m <sup>3</sup>	60% of the re	esults were ND <sup>h</sup>	42	20							
No other HAPs were detected in any samples												

μg/m<sup>3</sup> micrograms per cubic meter

ng/m<sup>3</sup> micrograms per cubic meter

NA Not applicable

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

- <sup>c</sup> 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.
- <sup>d</sup> 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.

<sup>&</sup>lt;sup>a</sup> 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

<sup>&</sup>lt;sup>b</sup> Details regarding these values are in the technical report, Schools Air Toxics Monitoring Activity (2009) Uses of Health Effects Information.

<sup>&</sup>lt;sup>e</sup> The comparison levels are specific to hexavalent chromium (recognized as the most toxic form) which is a fraction of the total chromium reported.

<sup>&</sup>lt;sup>f</sup> The comparison level is specific to elemental mercury, which is more readily and completely absorbed into the body than mercury conveyed on particles (e.g., divalent species).

 $<sup>^</sup>g$  Trichloroethylene was detected in only 2 of 10 samples, ranging from 0.054 to 0.17 ng/m3. The MDL is 0.011  $\mu$ g/m3.

<sup>&</sup>lt;sup>h</sup> Beryllium (PM<sub>10</sub>) was detected in only 4 of 10 samples, ranging from 0.004 to 0.05 ng/m3. The MDL is 0.03 ng/m3.

Appendix D. Lincoln Elementary School Pollutant Concentrations.

	rr.											
Parameter	Units	8/23/2009	8/29/2009	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/4/2009	10/10/2009	10/16/2009	Sample Screening Level <sup>a</sup>
Manganese (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.62	2.89	7.08	5.30	4.21	48.3	14.5	2.94	3.12	2.44	500
Nickel (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.09	0.36	0.58	0.41	0.41	0.58	0.20	0.32	0.39	0.17	200
Chromium (PM <sub>10</sub> )	ng/m <sup>3</sup>	2.29	3.21	3.83	4.29	3.78	4.08	3.41	2.74	2.46	2.54	580 <sup>b</sup>
Arsenic (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.62	1.04	1.39	0.79	0.22	0.74	0.18	0.22	1.73	2.29	150
Benzene	μg/m <sup>3</sup>	0.329	0.403	0.857	0.706	0.643	1.59	ND	0.419	0.422	0.400	30
Carbon Tetrachloride	μg/m <sup>3</sup>	0.755	0.680	0.730	0.793	1.22	0.837	ND	0.774	0.736	0.674	200
Butadiene, 1,3-	μg/m³	0.035	0.038	0.093	0.044	0.055	0.12	0.027	0.060	0.044	0.040	20
Chloromethane	μg/m³	1.13	1.06	1.15	1.41	1.90	1.53	1.20	1.01	0.723	1.50	1,000
Cadmium (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.04	0.09	0.17	0.14	0.06	0.57	0.05	0.05	0.12	0.05	30
Bromomethane	μg/m <sup>3</sup>	0.047	0.051	0.062	0.058	0.062	0.070	0.047	0.039	0.03	0.03	200
Tetrachloroethylene	μg/m <sup>3</sup>	0.068	0.10	0.22	0.10	0.075	0.12	0.06	0.081	0.31	0.06	1,400
Dichlorobenzene, p-	μg/m <sup>3</sup>	0.066	0.060	0.11	0.060	0.05	0.11	0.04	0.04	ND	ND	10,000
Acetonitrile	μg/m³	0.018	0.482	0.457	0.433	0.368	0.553	0.292	0.200	0.178	0.11	600
Antimony (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.87	0.39	1.48	1.31	0.44	1.32	0.15	0.42	0.61	0.65	2,000
Vinyl chloride	μg/m <sup>3</sup>	0.033	0.031	0.041	0.046	0.072	0.046	0.033	0.02	0.02	0.02	1,000
Ethylbenzene	μg/m <sup>3</sup>	0.070	0.087	0.23	0.12	0.100	0.15	0.052	0.11	0.096	0.091	40,000
Xylene, <i>m/p</i> -	μg/m <sup>3</sup>	0.195	0.208	0.51	0.27	0.20	0.44	0.113	0.24	0.204	0.182	3,000
Dichloromethane	μg/m³	0.30	0.26	0.535	0.28	0.382	0.32	0.24	0.27	0.428	0.24	2,000
Chloroform	μg/m³	0.13	0.10	0.19	0.14	0.15	0.14	0.11	0.13	0.12	0.11	500
Xylene, o-	μg/m <sup>3</sup>	0.070	0.087	0.22	0.13	0.11	0.19	0.048	0.11	0.091	0.078	9,000
Cobalt (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.05	0.03	0.11	0.05	0.05	0.08	0.05	0.005	0.02	0.02	100
Toluene	μg/m <sup>3</sup>	0.682	1.44	1.69	0.777	0.656	1.26	1.49	1.35	3.71	0.524	4,000
Methyl isobutyl ketone	μg/m³	2.16	2.34	0.14	1.21	0.516	0.685	ND	0.21	0.16	0.04	30,000
Carbon Disulfide	μg/m³	0.062	0.18	0.21	0.13	0.361	0.12	0.14	0.034	0.15	0.047	7,000
Styrene	μg/m³	0.03	0.043	0.22	0.064	0.047	0.060	0.04	0.03	ND	0.055	9,000
Mercury (PM <sub>10</sub> )	ng/m <sup>3</sup>	ND	0.005	0.02	ND	0.006	0.01	0.01	ND	0.01	0.05	3,000°
Selenium (PM <sub>10</sub> )	ng/m <sup>3</sup>	0.17	0.32	1.22	1.13	0.67	1.61	0.17	0.20	0.29	0.11	20,000
Methyl Chloroform	μg/m <sup>3</sup>	0.066	0.082	0.087	0.087	0.10	0.098	ND	0.087	0.071	0.071	10,000
Chloroethane	μg/m <sup>3</sup>	0.045	0.045	0.092	0.058	0.050	0.045	0.050	0.034	0.029	0.055	40,000
Trichloroethylene	μg/m <sup>3</sup>	ND	ND	0.17	ND	ND	0.054	ND	ND	ND	ND	10,000

Appendix D. Lincoln Elementary School Pollutant Concentrations.

Parameter	Units	8/23/2009	8/29/2009	9/4/2009	9/10/2009	9/16/2009	9/22/2009	9/28/2009	10/4/2009	10/10/2009	10/16/2009	Sample Screening Level <sup>a</sup>
Beryllium (PM <sub>10</sub> )	ng/m <sup>3</sup>	ND	ND	0.05	ND	ND	ND	0.004	ND	0.007	0.007	20
Acrylonitrile	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	200
Benzyl Chloride	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	140
Bromoform	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	6,400
Chlorobenzene	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	10,000
Chloroprene	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	70
Dichloroethane, 1,1-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	4,400
Dichloroethylene, 1,1-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	80
Dichloropropane, 1,2-	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	200
Dichloropropylene, cis-1,3-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	14
Dichloropropylene, trans -1,3-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	14
Ethyl Acrylate	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	20,000
Ethylene dibromide	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	12
Ethylene dichloride	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	270
Hexachloro-1,3-butadiene	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	320
Methyl Methacrylate	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	7,000
Methyl tert -Butyl Ether	μg/m³	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	7,000
Tetrachloroethane, 1,1,2,2-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	120
Trichlorobenzene, 1,2,4-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	2,000
Trichloroethane, 1,1,2-	μg/m <sup>3</sup>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	440

Key Pollutant

ng/m<sup>3</sup> nanograms per cubic meter

ug/m<sup>3</sup> micrograms per cubic meter

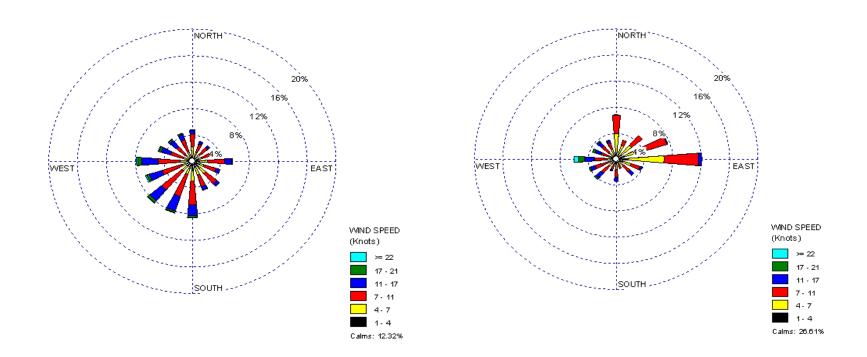
- -- No sample was conducted for this pollutant on this day or the sample was invalid.
- ND No detection of this chemical was registered by the laboratory analytical equipment.

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 <a href="http://www.epa.gov/schoolair/pdfs/UsesOfHealthEffectsInfoinEvalSampleResults.pdf">http://www.epa.gov/schoolair/pdfs/UsesOfHealthEffectsInfoinEvalSampleResults.pdf</a>. 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.

b The sample screening levels are specific to hexavalent chromium (recognized as the most toxic form) which is a fraction of the total chromium reported.

<sup>&</sup>lt;sup>c</sup> The sample screening level is specific to elemental mercury, which is more readily and completely absorbed into the body than mercury conveyed on particles (e.g., divalent species).

# Appendix E. Windroses for Goshen Municipal Airport NWS Station.



Goshen Municipal Airport NWS Station 2002-2007

Goshen Municipal Airport NWS Station Across Sampling Period (Aug. 23, 2009-Oct. 16, 2009)<sup>1</sup>

 $<sup>^{1}</sup>$  Goshen Municipal Airport NWS Station (WBAN 14829) is 19.97 miles from Lincoln Elementary School.