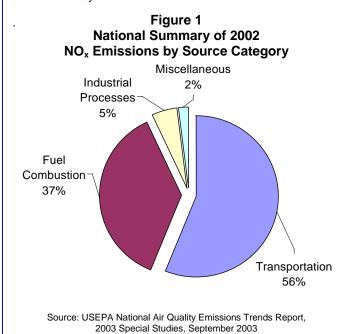


### **2004 Nitrogen Dioxide Summary**

**New Jersey Department of Environmental Protection** 

#### **NATURE AND SOURCES**

Nitrogen Dioxide (NO<sub>2</sub>) is a reddish-brown, highly reactive gas that is formed in the air through the oxidation of Nitric Oxide (NO). When NO<sub>2</sub> reacts with other chemicals, it can form ozone, particulate matter, and other compounds which can contribute to regional haze and acid rain. Nitrogen Oxides (NO<sub>x</sub>) is a mixture of gases which is mostly comprised of NO and NO2. These gases are emitted from the exhaust of motor vehicles, the burning of coal, oil or natural gas, and during industrial processes such as welding, electroplating, and dynamite blasting. Although most NO<sub>x</sub> is emitted as NO, it is readily converted to NO2 in the atmosphere. In the home, gas stoves and heaters produce substantial amounts of nitrogen dioxide. A pie chart summarizing the major sources of NO<sub>x</sub> is shown below (Figure 1). As much of the NO<sub>x</sub> in the air is emitted by motor vehicles, concentrations tend to peak during the morning and afternoon rush hours. This is shown in the graph in Figure 2 (page 2), which also indicates that concentrations tend to be higher in the winter than the summer. This is due in part to poorer local dispersion conditions caused by light winds and other weather conditions that are more prevalent in the colder months of the year.



## HEALTH AND ENVIRONMENTAL EFFECTS

Short-term exposures (less than 3 hours) to low levels of nitrogen dioxide may aggravate pre-existing respiratory illnesses, and can cause respiratory illnesses, particularly in children ages 5-12. Symptoms of low level exposure to NO and NO2 include irritation to eyes, nose, throat and lungs, coughing, shortness of breath, tiredness and nausea. Longterm exposures to NO2 may increase susceptibility to respiratory infection and may cause permanent damage to the lung. NO and NO2 are found in tobacco smoke, so people who smoke or breathe in second-hand smoke may be exposed to NO<sub>x</sub>. The U.S. Department of Health and Human Services (DHHS), the International Agency for Research on Cancer (IARC), and the U.S. Environmental Protection Agency (EPA) have determined that, with the available information, no conclusion can be made as to the carcinogenicity of NO or NO2 to human beings.

Nitrogen Oxides contribute to a wide range of environmental problems. These include potential changes in the composition of some plants in wetland and terrestrial ecosystems, acidification of freshwater bodies, eutrophication of estuarine and coastal waters, increases in levels of toxins harmful to fish and other aquatic life, and visibility impairment.

#### **STANDARDS**

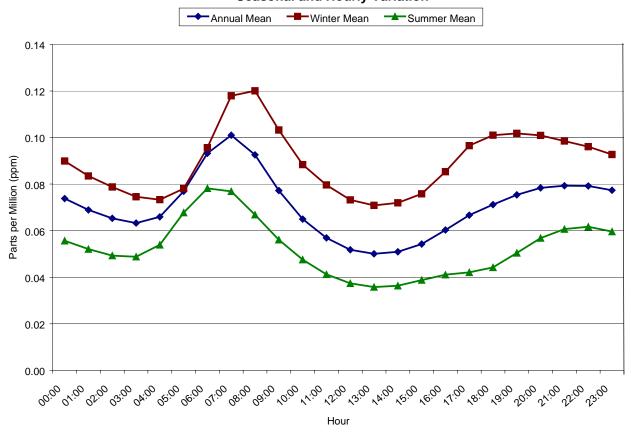
The primary (health based) and secondary (welfare based) National Ambient Air Quality Standards (NAAQS) for NO $_2$  are the same. They are set at a calendar year average concentration of 0.053 parts per million (ppm). The New Jersey Ambient Air Quality Standards (NJAAQS) are identical to the NAAQS except micrograms per cubic meter ( $\mu$ g/m $^3$ ) are the standard units and the state standard applies to any 12-month period, not just the calendar year. The state of California has a one-hour average standard of 470  $\mu$ g/m $^3$  that New Jersey uses as a guideline in assessing short-term impacts from specific sources. Table 1 provides a summary of the NO $_2$  standards.

Table 1
National and New Jersey Ambient Air Quality Standards for Nitrogen Dioxide

Parts Per Million (ppm)
Micrograms Per Cubic Meter (µg/m³)

micrograms for Subjective (µg/m)								
Averaging Period	Туре	New Jersey	National	California				
12-month average	Primary	100 μg/m <sup>3</sup> (0.05 ppm)						
Annual average	Primary		0.053 ppm (100 μg/m <sup>3</sup> )					
12-month average	Secondary	100 μg/m <sup>3</sup> (0.05 ppm)						
Annual average	Secondary		0.053 ppm (100 μg/m <sup>3</sup> )					
1-hour average	Primary			470 μg/m <sup>3</sup> (0.25 ppm)				

Figure 2
Nitrogen Dioxide & Nitric Oxide Concentrations – New Jersey 1967-1999
Seasonal and Hourly Variation



#### **MONITORING LOCATIONS**

The state monitored  $NO_2$  levels at 9 locations in 2004. These sites are shown in the map to the right. The Camden Lab monitoring station was temporarily discontinued on September 22, 2003, and resumed operation on January 8, 2004.

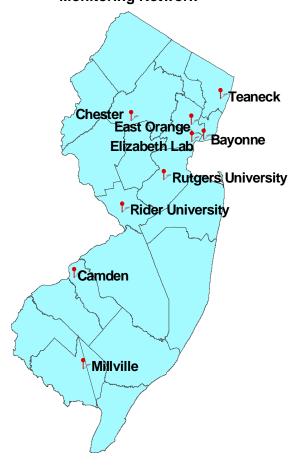
#### NO<sub>2</sub> Levels in 2004

None of the monitoring sites recorded exceedances of either the National or New Jersey Air Quality Standards for  $NO_2$  during 2004. The maximum annual average concentration recorded was 0.030 ppm at the Elizabeth Lab site located at Exit 13 of the New Jersey Turnpike. While national health and welfare standards have not been established for Nitric Oxide (NO), it is considered to be an important pollutant that contributes to the formation of ozone, fine particles and acid rain. The maximum annual average concentration of NO recorded in 2004 was 0.044 ppm, also at the Elizabeth Lab site (see Table 2 and Figure 4, page 4).

#### **TRENDS**

Since routine monitoring for NO<sub>2</sub> began in 1966, concentrations have never exceeded the NAAQS in New Jersey. A graph of NO<sub>2</sub> levels provided in Figure 5

#### Figure 3 2004 Oxides of Nitrogen Monitoring Network

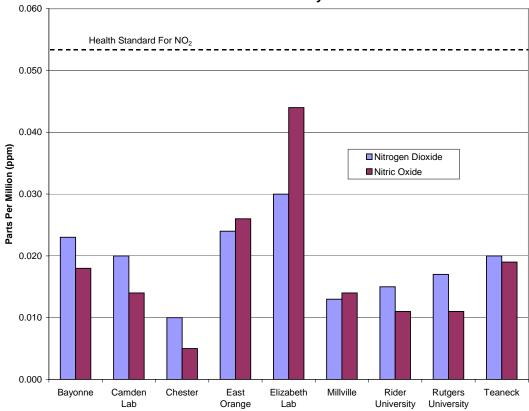


Tabe 2
Nitrogen Dioxide and Nitric Oxide Data-2004
1-Hour and 12-Month Averages

Parts Per Million (ppm)
California 1-Hour Standard = 0.25 ppm
National 12-Month Standard = 0.053 ppm

	Nitroge	n Dioxide	Nitrogen Dioxide		Nitric Oxides
	1-Hour Ave	1-Hour Average (ppm)		12-Month Average (ppm)	
Monitoring Sites	Maximum	2nd Highest	Maximum	Calendar year	Average(ppm)
Bayonne	0.115	0.084	0.024	0.023	0.018
Camden Lab	0.078	0.074	0.020	0.020	0.014
Chester	0.063	0.062	0.012	0.010	0.005
East Orange	0.150	0.143	0.027	0.024	0.026
Elizabeth Lab	0.156	0.125	0.032	0.030	0.044
Millville	0.057	0.053	0.013	0.013	0.014
Rider University	0.054	0.054	0.015	0.015	0.011
Rutgers University	0.077	0.076	0.018	0.017	0.011
Teaneck	0.083	0.078	0.020	0.020	0.019

Figure 4
Annual Average NO and NO₂ Concentrations in New Jersey - 2004



shows the statewide average annual mean concentrations recorded from 1975 to 2004 in the form of a trendline. The graph also includes the levels of the sites that measured the highest annual mean and lowest annual mean in each year as points above and below this trendline. Although NO<sub>2</sub> concentrations are well within the NAAQS, there is still a great deal of interest in oxides of nitrogen because of their role in the formation of other pollutants – most notably ozone and fine particles. Both these pollutants are of concern over much of the northeastern United States and efforts to reduce levels of ozone and fine particles are likely to require reductions in NO emissions.

## TOTAL REACTIVE OXIDES OF NITROGEN (NO<sub>y</sub>)

Although not specifically defined, there is a broad group of nitroxyl compounds in the ambient air that react in the troposphere and contribute to the formation of ozone. These compounds, called Total Reactive Oxides of Nitrogen (NO<sub>y</sub>), include nitrogen oxides (NO<sub>x</sub>), peroxyacyl

nitrates (RC(O)OONO $_2$  or PAN), peroxynitric acid (HO $_2$ NO $_2$ ), nitrous acid (HONO), nitric acid (HNO $_3$ ), dinitrogen pentoxide (N $_2$ O $_5$ ) and nitrate radicals (•NO $_3$ ). NO $_y$  can also be described as the sum of the nitrogen oxides (NO $_x$ ) and the atmospheric NO $_x$  oxidation products. Although measuring NO $_y$  is not required by the federal regulations, it is strongly recommended by the EPA to supplement the data collected by Photochemical Assessment Monitoring Stations (PAMS) Network. NO $_y$  measurements may provide valuable information for evaluating chemical mechanisms in ozone (O $_3$ ) prediction models, indicate NO and NO $_2$  emission trends, and assist in developing regional control strategies for O $_3$ .

The identification and measurement of individual NO<sub>y</sub> compounds is technically difficult and expensive, however, an analyzer that measures total NO<sub>y</sub> concentrations is commercially available and New Jersey has been evaluating its performance. The NJDEP began

monitoring for  $NO_y$  at the Rider University station in March 2002. Nitrogen oxides ( $NO_x$ ) and a group of volatile organic compounds (VOCs) known as ground-level ozone precursors are also measured at this station.

The monthly average  $NO_y$  concentrations for 2004 are presented in Table 3 along with the corresponding  $NO_x$  concentrations measured by the  $NO_2$  analyzer at Rider University. Theoretically, concentrations of  $NO_y$  should be greater than  $NO_x$  when both are measured at the same monitoring location. Since several monthly average  $NO_y$  concentrations in 2004 are actually less than their corresponding  $NO_x$  monthly averages, and since there have been some technical difficulties with the method which indicate a lot of uncertainty in its measurements, the usefulness of the  $NO_y$  data is limited.

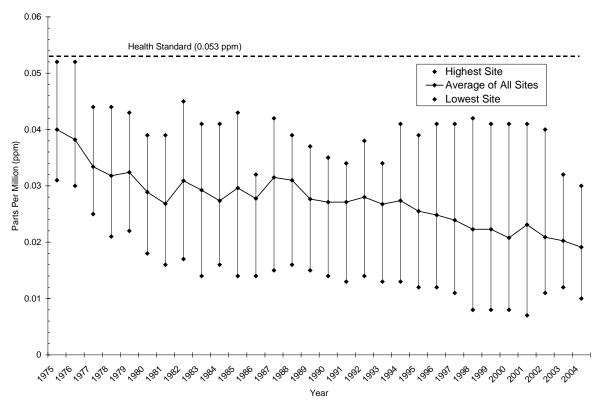
# Table 3 Nitrogen Oxides (NO<sub>x</sub>) and Total Reactive Oxides of Nitrogen (NO<sub>y</sub>) Data Rider University - 2004 Monthly Average

Parts Per Million (ppm)

	NO <sub>x</sub> Monthly Average (ppm)	NO <sub>y</sub> 1 Monthly Average (ppm)
January	0.027	0.024
February	0.041	0.038
March	0.028	0.028
April	0.020	0.021
May	0.015	0.015
June	0.012	0.011
July	0.013	0.012
August	0.015	0.012
September	0.018	0.018
October	0.029	0.029
November	0.040	0.039
December	0.043	0.032

See text for explanation of NO<sub>y</sub> measurement issues. It is presented here to show that the data are available and the range of concentrations recorded.

Figure 5
Nitrogen Dioxide Concentrations in New Jersey
1975-2004
12-Month (Calendar Year) Average



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