#### Year One Project Report To Meadowlands Environmental Research Institute The NJ Meadowlands Commission

(07/27/07)

### TITLE: Characterization of Atmospheric Nitrogen Oxides over the Meadowlands

PI: Yuan Gao Department of Earth and Environmental Sciences Rutgers University 101 Warren Street Newark, NJ 07102

> Phone: (973) 353-1139 Fax: (973) 353-1965 E-mail: yuangaoh@andromeda.rutgers.edu

- **PERIOD:** Year1: 11/01/06 08/31/07 (Approved Project Period: 11/01/06 – 10/31/08)
- BUDGET: Year1: \$40,500 (2 Year Total: \$81,000)

## STATEMENT OF THE PROBLEMS

Nitrogen oxides  $(NO_x)$  are highly reactive gases in the ambient air, such as NO and NO<sub>2</sub> which are precursor molecules for the production of ground-level ozone  $(O_3)$ . The O<sub>3</sub> production rate strongly depends on the concentrations of NO and NO<sub>2</sub>. High concentrations of O<sub>3</sub> in the ambient air can trigger serious respiratory and other health-related problems. Therefore, an adequate knowledge of the characteristics of NO<sub>x</sub> in the ambient air is crucial for the assessment of O<sub>3</sub> pollution and air quality.

According to the EPA data, the major anthropogenic source of NO<sub>x</sub> in the ambient air in the United States is motor vehicles emissions, accounting for ~55% of the total emissions. Additional sources of NO<sub>x</sub> include electric utilities (~22%) and other industrial, commercial, and residential sources. Nitrogen oxides form from fossil fuel burning at high temperature, as in a combustion process. Therefore, high concentrations of NO<sub>x</sub> are often found in the areas heavily impacted by traffic and power plants, although they can also come from natural sources and be transported over long distances by prevailing winds.

Despite the critical role of  $NO_x$  in air quality, the concentration levels and temporal variations of  $NO_x$  over the Meadowlands District have not been monitored, and this represents a serious data gap in air quality assessment in the District. With the heavy transportation and the existence of power plants and landfills in the region, the emissions of  $NO_x$  and the production of  $O_3$  in the

District could be substantial. Therefore, the measurements of  $NO_x$  in the ambient air in the region are critically needed.

## **GOAL AND OBJECTIVES**

The research effort under this plan aims at generating new data on the ambient levels of  $NO_x$  over the Meadowlands District, and this new research will complement the on-going air quality monitoring at Meadowlands Environmental Research Institute (MERI). The ultimate goal of this project is to contribute to the understanding and control of air pollution in the District, maintaining the health of the Meadowlands. Within this context, the specific objectives are:

- (1) To install a new NO<sub>x</sub> analyzer at MERI and to initiate atmospheric nitrogen monitoring at this location,
- (2) To measure the concentrations of atmospheric  $NO_x$  in order to assess the current levels of  $NO_x$  in the ambient air,
- (3) To generate temporal variation, in particular seasonal trends of NO<sub>x</sub> under ambient environmental conditions.

# **ACTIVITIES ACCOMPLISHED DURING YEAR ONE PERIOD**

A brief summary of the accomplishment toward the goals in the year1 period is as follows:

- (1) A new NO<sub>x</sub> analyzer (Model 42i, Thermal Electron Corp.) was purchased based on the selection made by the PI, and it was installed on site in a laboratory at MERI in December 2006. The instrument calibration was completed in January 2007 with the assistance from MERI staff.
- (2) Routinely measuring NO<sub>x</sub> in the ambient air at the MERI site has been on since February 2007.

Besides the above targeted gas-phase measurement activities, additional efforts that are not in the original research plan were made on aerosol nitrate during this period to strengthen the project, which includes:

- (1) Particulate N sampling was made for a period of three months from February to May 2007 on a daily basis. Sample collection was carried out through the use of an inhouse-built aerosol sampler assembled at the PI's lab that is primarily composed of a ChemComb cartridge, mass flow meter, and vacuum pump.
- (2) Aerosol sampling structure was constructed on the roof of MERI building to support daily aerosol sample collection.
- (3) Analyses of all air samples for particulate nitrate and nitric acid vapor have been performed through a DIONEX ion chromatograph analyzer at the PI's lab.

In addition to the efforts on aerosol nitrate, a new ozone analyzer was installed at the MERI site and has been up running for data collection, which is also not in the original research plan. Although this  $O_3$  analyzer has generated tremendous data that requires more time for data process, the  $O_3$  data will strengthen this project and help interpret  $NO_x$  data. Therefore we have included extra  $O_3$  data processing in our current efforts.

#### **PRELIMINARY RESULTS**

**Summary:** Table 1 shows a brief summary of the concentrations of the gas-phase NO,  $NO_x$ ,  $O_3$ , aerosol nitrate and HNO<sub>3</sub> vapor observed at the MERI site. During the measurement period of February – May 2007, the concentrations of NO,  $NO_x$  and  $NO_3^-$  in the ambient air decreased with time, while the O<sub>3</sub> concentration increased with time. The HNO<sub>3</sub> gas concentration showed less variation.

**Diurnal Patterns:** Figure 1 shows the diurnal patterns of the concentrations of NO,  $NO_x$ , and  $O_3$  based on the hourly averages of data collected on the minute basis. The  $O_3$  concentration showed an opposite variation compared with those of NO and  $NO_x$ . The concentration peaks of NO and  $NO_x$  appeared at 6:00am-8:00am, while the  $O_3$  concentration peak appeared in early afternoon hours. There are about 8 hours of lag between the  $NO_x$  peak and  $O_3$  peak.

**Daily Patterns:** The daily variation of  $O_3$  was less than those of NO and  $NO_x$  and mostly opposite to those of NO and  $NO_x$ , as shown in Figure 2. These plots were made based on daily average concentrations that were converted from minute/hourly data. Figure 3 shows the daily concentration patterns of aerosol nitrate and HNO<sub>3</sub> vapor. Compared to the HNO<sub>3</sub> gas, aerosol nitrate showed greater daily variation. In earlier spring,  $NO_3^-$  also showed more variation and higher values in the ambient concentration; however, HNO<sub>3</sub> gas showed less variation in earlier spring compared with the late spring.

#### **YEAR-2 EFFORTS**

The efforts under this project in Year 2 starting in September 2007 should include:

- To continue the measurements of NO<sub>x</sub> and O<sub>3</sub> for a full year in order to get the seasonal patterns. By now, the measurements of NO<sub>x</sub> and O<sub>3</sub> have been made for 7 months (Feb July, 2007). Therefore, the measurements should be continued to February 2008;
- (2) To continue and finish the remaining IC chemical analyses for aerosol samples;
- (3) To perform data analyses and interpretation of all species (NO, NO<sub>2</sub>, O<sub>3</sub>, NO<sub>3</sub>, HNO<sub>3</sub>);
- (4) To generate a publication on atmospheric nitrogen over the Meadowlands and final project report.

Month 2007	NO (ppb)	NO <sub>x</sub> (ppb)	O <sub>3</sub> (ppb)	$NO_3^{-1}$ (µg m <sup>-3</sup> )	HNO <sub>3</sub> (µg m <sup>-3</sup> )
February	25.4	50.7	20.5	3.08	1.28
March	16.8	41.2	25.6	2.42	1.10
April	9.36	25.1	31.8	1.39	1.16
May	5.26	19.1	34.4	1.06	1.50
Average	16.4	34.0	28.1	1.26	1.99

Table 1. Springtime Concentrations of Atmospheric NO, NO<sub>x</sub>, O<sub>3</sub>, Aerosol Nitrate and HNO<sub>3</sub> Vapor over the Meadowland in Central New Jersey



Figure 1. Averaged diurnal variations of NO, NO<sub>x</sub> and O<sub>3</sub> during Feb – May 2007.



Figure 2. Daily Variations of NO, NO<sub>x</sub> and O<sub>3</sub> during Feb – May 2007.



Figure 3. Daily Variations of NO<sub>3</sub><sup>-</sup> and HNO<sub>3</sub> vapor from February to May 2007.