

#### **NOAA Platforms and Facilities Available in ICARTT 2004**

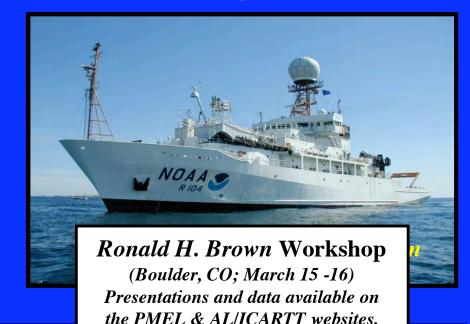


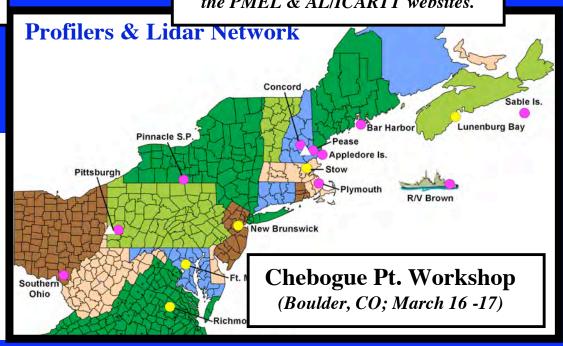
#### WP-3D/DC3 Workshop

(*Boulder*, *CO*; *April 11 - 13*) Data available on AL/ICARTT website. Requires ICARTT username and password.









### **NOAA** and **NSF**

**Chebogue Point: Location and site** 



**Pico: Location and site** 





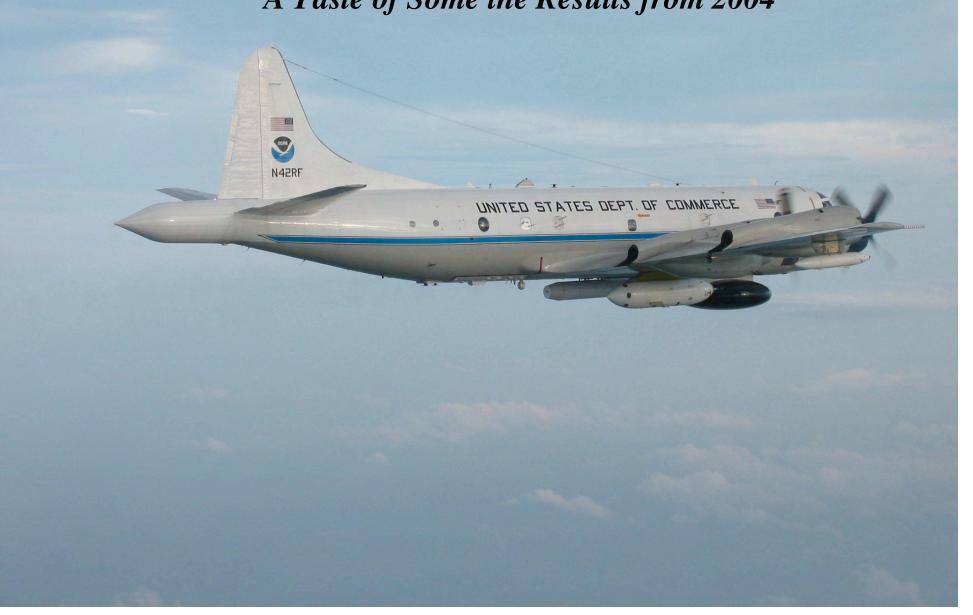
(March 10 and 11)

The Chebogue Point presentations (in pdf format) and data are available at Chebogue Pt. ftp site:

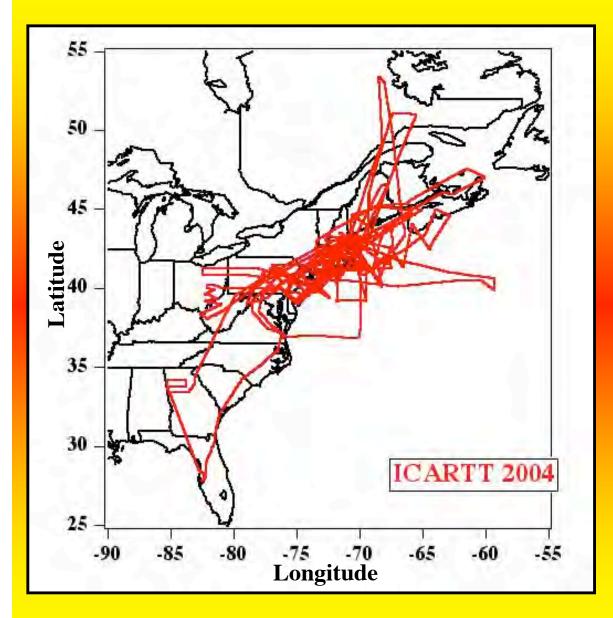
- Host: ftp.al.noaa.gov
- User ID: chebpt

## NOAA WP-3D Research Aircraft

A Taste of Some the Results from 2004

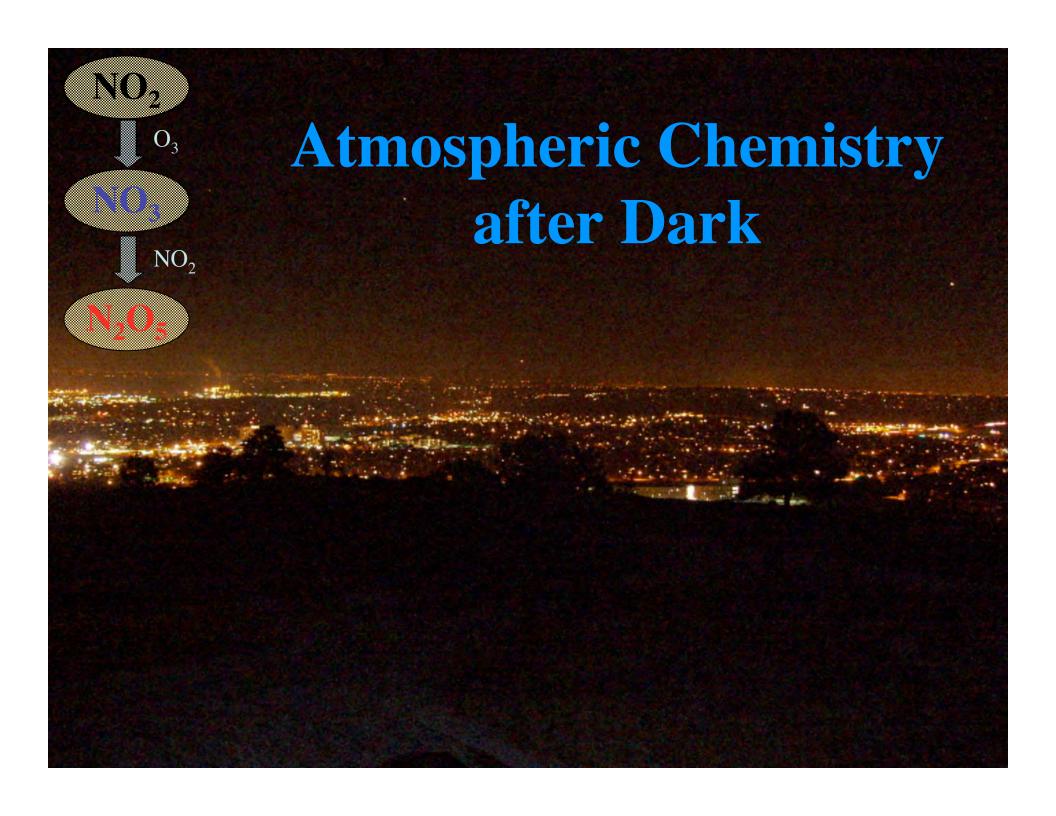


#### A Taste of Some the Results from 2004

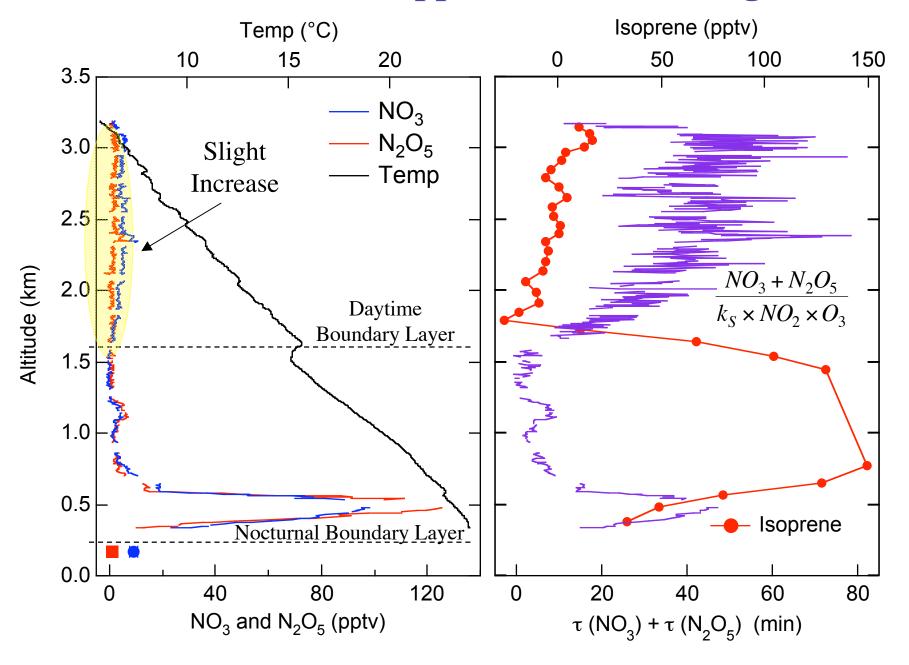


# **During ICARRT Eighteen Flights (~ 130 hrs.) Studying:**

- Urban plumes (day and night)
- Power plant plumes
- Forest fire plumes
- Regional composition and chemistry
- Forecast model evaluation
- Long-range transport
- Intercomparison activities

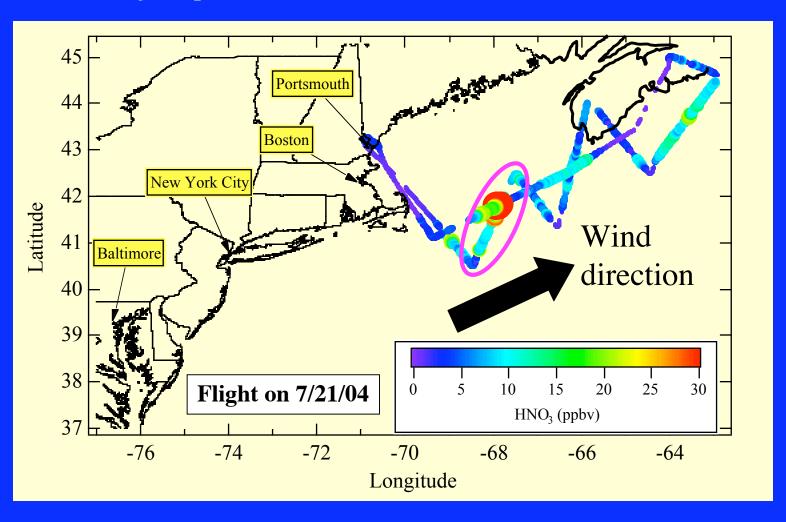


### Vertical Profile Over Appledore Island - Aug 3, 2004

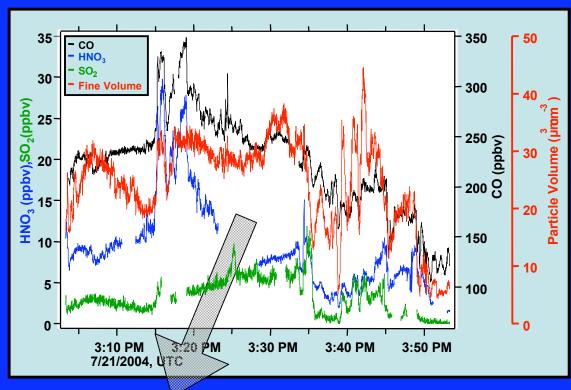


### **Tracking East Coast Oxidation Products over the Atlantic**

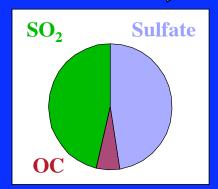
Data from flights on July 20, 21, 22.
CO, HNO<sub>3</sub>, SO<sub>2</sub>, aerosol measurements from NOAA WP-3D.



### Pollution plumes over the Atlantic



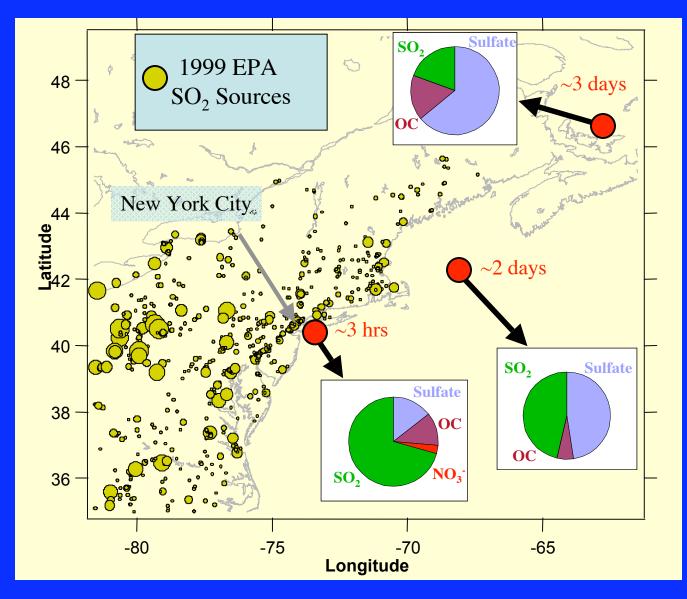
- The CO reflects crossing a mixture of East Coast sources.
- HNO<sub>3</sub> was the most abundant reactive nitrogen species and correlated with CO. (flexpart retro-plume analysis indicates ~ 2 day transport time).
- SO<sub>2</sub> not highly correlated with CO.
- SO<sub>2</sub> and fine particle volume are well correlated and greatly enhanced in plume.



- Most of the fine particle mass is sulfate (PILS, R. Weber, GIT).
- Water-soluble organic carbon (PILS, R. Weber, GIT) is generally < 15% of the fine mass; suggests < 30% total organic contribution. (OC estimated from WSOC\*2.25.)

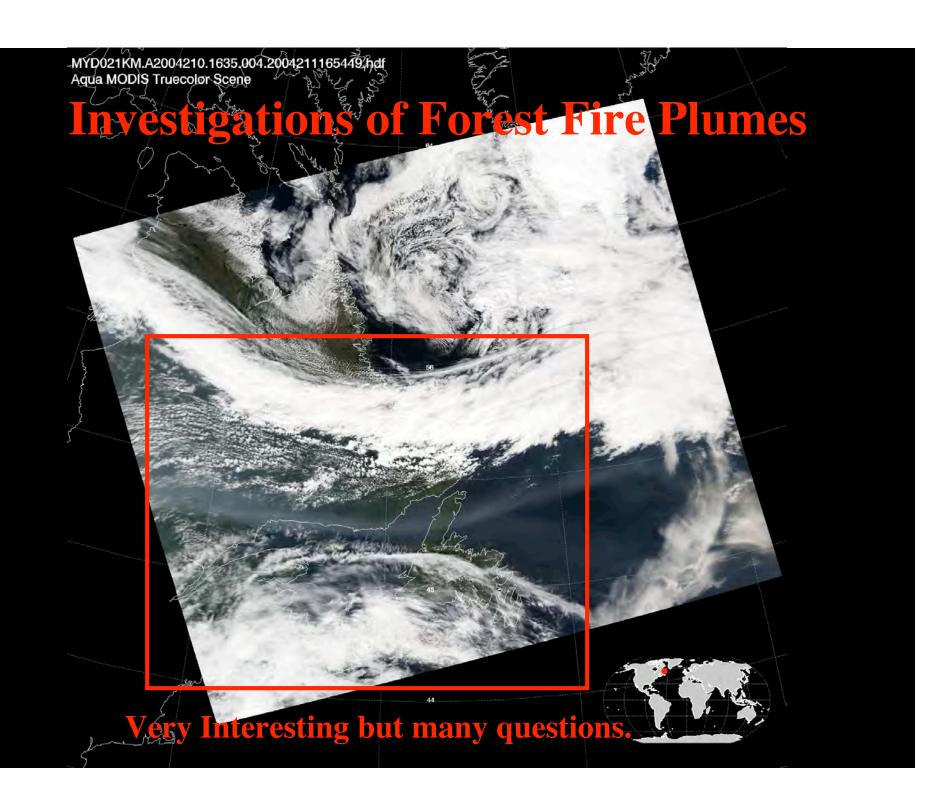
Particulate sulfate and potential particulate sulfate (from  $SO_2$ ) dominate aerosol mass.

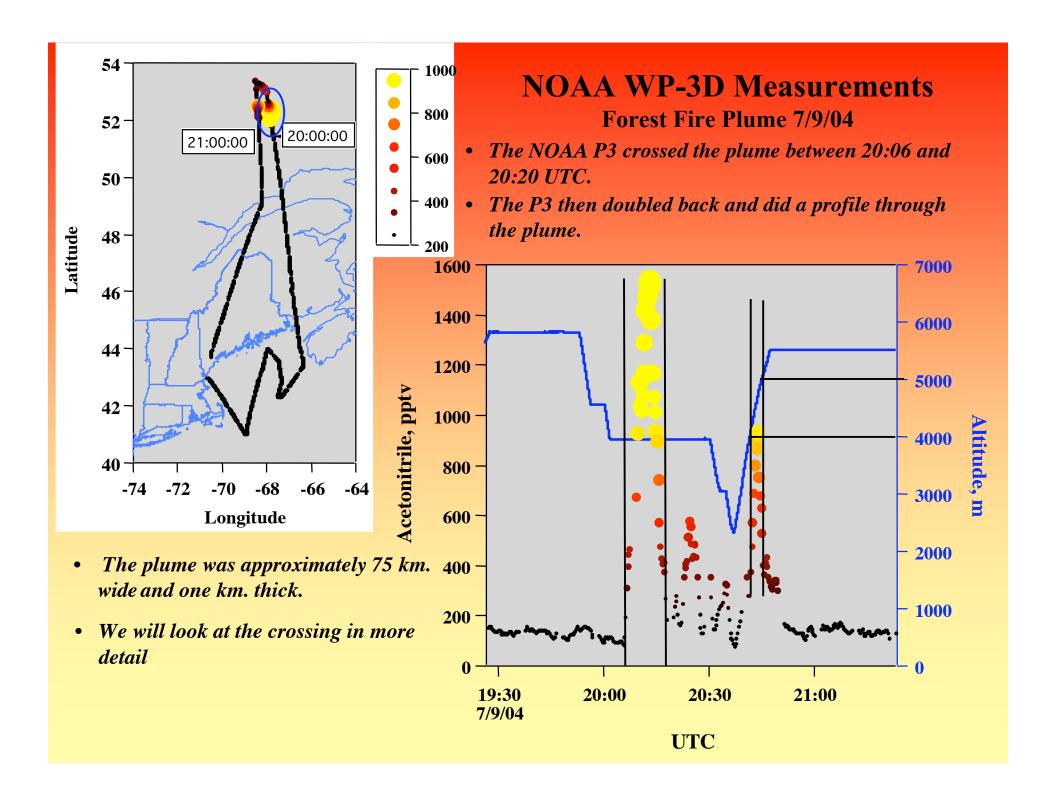
## How does aerosol composition change with aging plume?



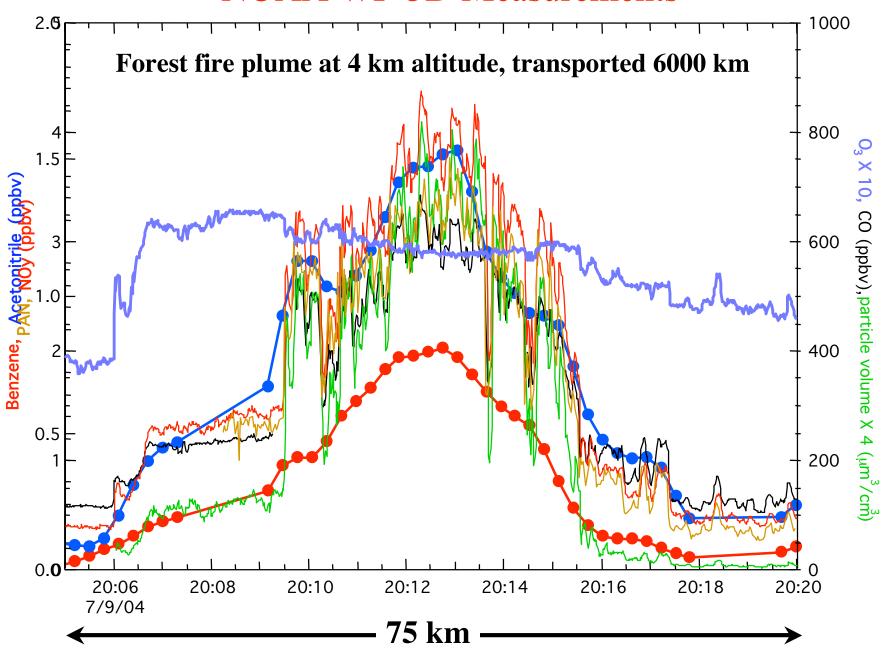
- Many SO<sub>2</sub> sources embedded along urban corridor.
- Plumes crossed on 7/20-21-22.
- Flexpart retro-plume analysis indicates E.
   Coast sources
- Total OC fraction <
  30% fine particle mass
  (OC fraction estimated
  from WSOC).</li>

Sulfate and Potential Sulfate (SO<sub>2</sub>) Dominate Exported Fine Particle Mass

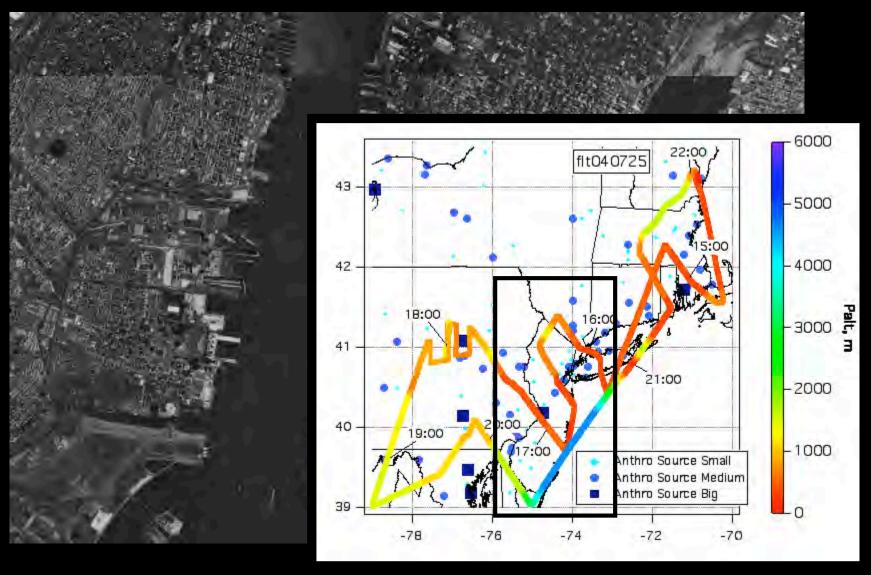




### NOAA WP-3D Measurements

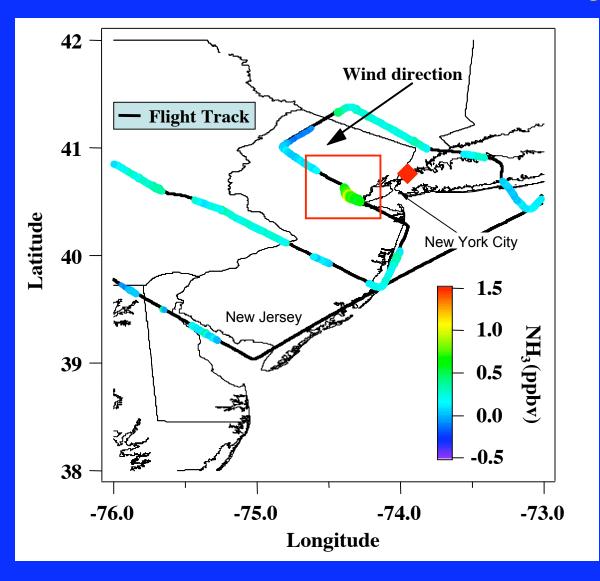


### NH<sub>3</sub> in the New York City Plume



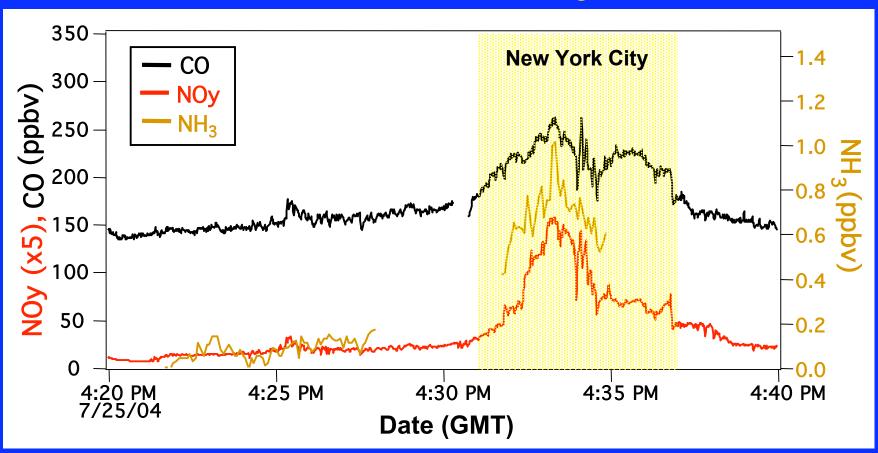
Out first results on the P3 using a recently developed CIMS chemistry

# New York NH<sub>3</sub>



- July 25th flight downwind of New York City
- Flight track colored and sized by NH<sub>3</sub> mixing ratios.

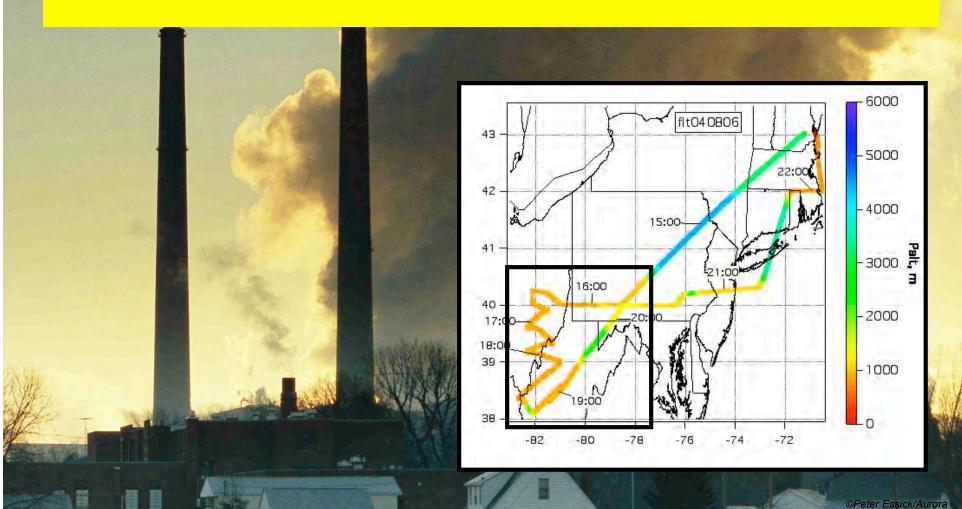
# New York NH<sub>3</sub>



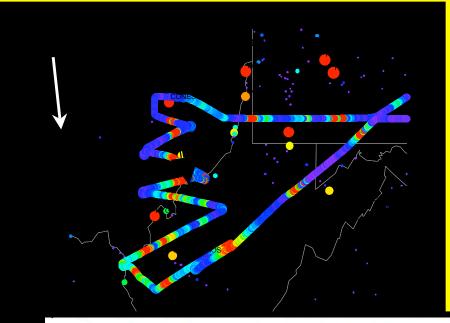
- Time series shows NH<sub>3</sub> correlated with CO and NOy in New York city outflow.
- NH<sub>3</sub> observed in forest fire, urban, and some power plant plumes.

## **Investigating Power Plant Emissions**

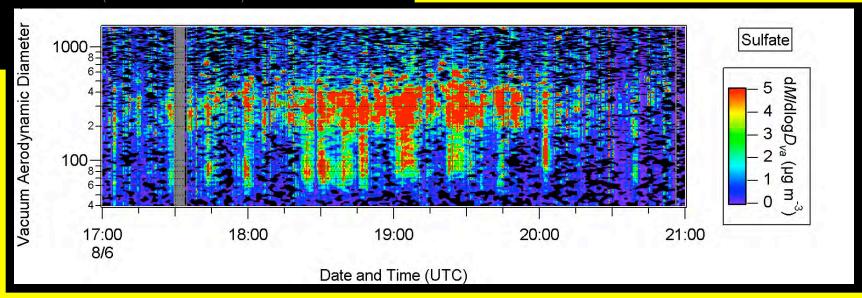
The emission from seventeen different power plants were studied.



## Power Plants Study: Flight on 8/6/04



- Daytime flight
- Wind from the north-north-west
- Power plants and flight tracks are colorized by SO<sub>2</sub> emissions.



### Conclusions from Power Plant Studies

During ICARTT, we obtained the plumes from several power plant on different days plumes at different stages of chemical evolution. Near the plant:

- High levels of aerosol sulfate were measured (often more than 15 μg/m3).
- Most (>90%) of the total sulfur was still in SO<sub>2</sub>.
- Very little ammonium -> acidic particles
- Very little organic mass -> probably not enough precursor.

#### As the plumes aged:

- Particle sizes increase, due initially to sulfate increases.
- Further downwind, aerosol ammonium and organic mass increases.
- When clouds were present, cloud processing rapidly converted SO<sub>2</sub> into aerosol sulfate.

Findings: We made measurement in plumes downwind of several power plants that had revised their emission inventories (EI) since 1999. The result:

- 2003  $E(NO_x) \sim 50\%$  1999  $E(NO_x)$
- 2003  $E(SO_2) \sim 85\% 1999 E(SO_2)$
- Updated EI agrees with observations

#### **SUMMARY**

We established broad goals for the study that we were able to accomplish. Some important finding are:

**NOAA:ICARTT-2004)** 

- It's a whole new day for the night!
- Sulfate dominated the exported fine particle mass.
- The aerosols measured after long-range transport were largely neutralized.
- We have obtained a wealth of new information about the transport and chemical transformation processes that are occurring in forest fire emissions.
- Nitric acid and aerosols can be efficiently scavenged by clouds.
- NH<sub>3</sub> was measured (a first for the P3) in urban, forest fire and some power plant plumes.
- Air quality has significantly reduced the emission of  $NO_x$  and  $SO_2$  from power plants in the Northeast.



We stand ready to share our results with our study partners.