# **Evaluating the HYSPLIT-Hg Atmospheric Mercury Model Using Ambient Monitoring Data**

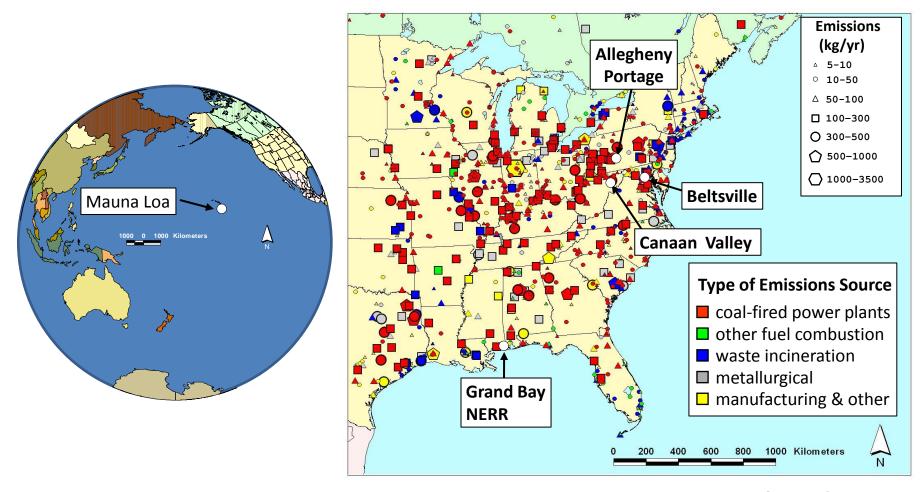
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10th International Conference on Mercury as a Global Pollutant July 23-29, 2011, Halifax, Nova Scotia

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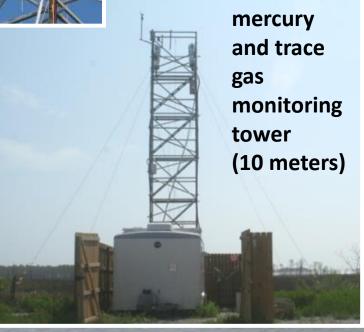
# NOAA Air Resources Laboratory: Five Collaborative Speciated Atmospheric Mercury Measurement Sites

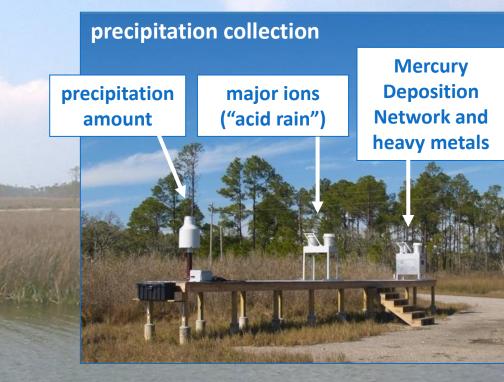


2002 mercury emissions sources based on data from USEPA, Envr. Canada and the CEC

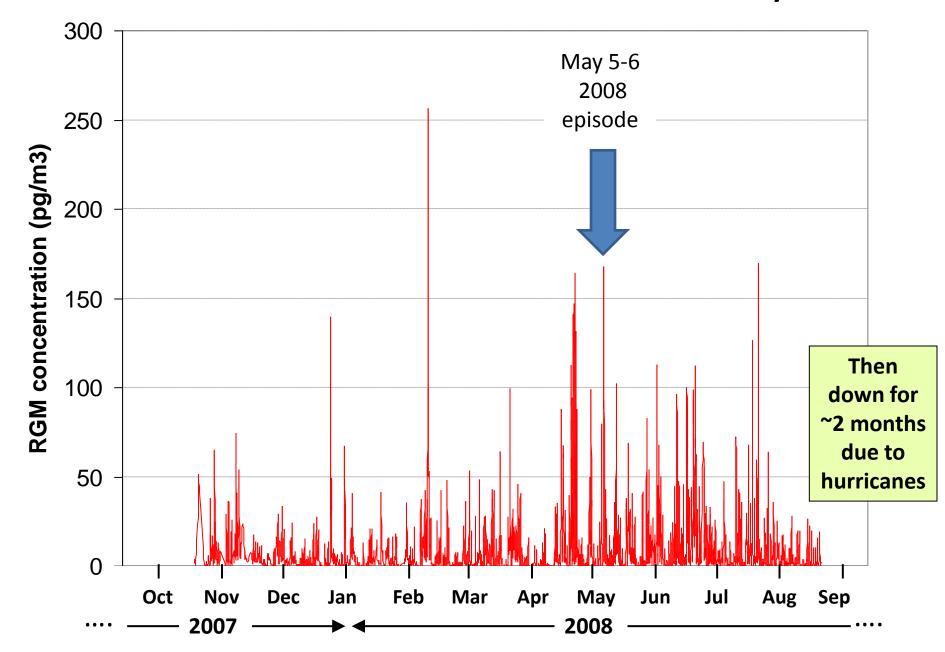


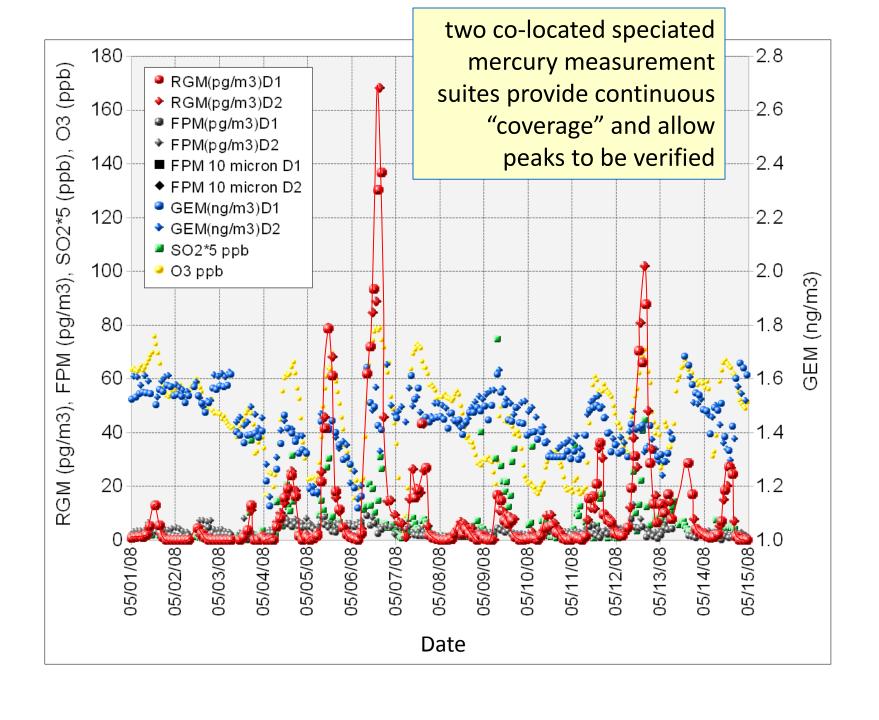
# Atmospheric Mercury Monitoring Station at the Grand Bay NERR





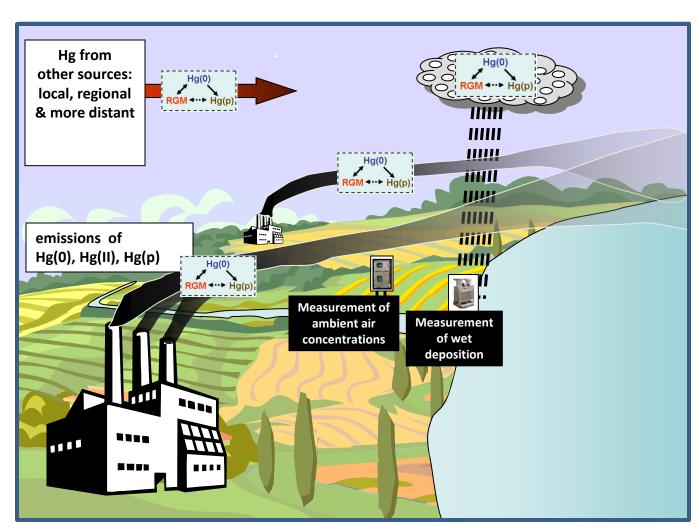
#### time series of RGM concentrations measured at Grand Bay



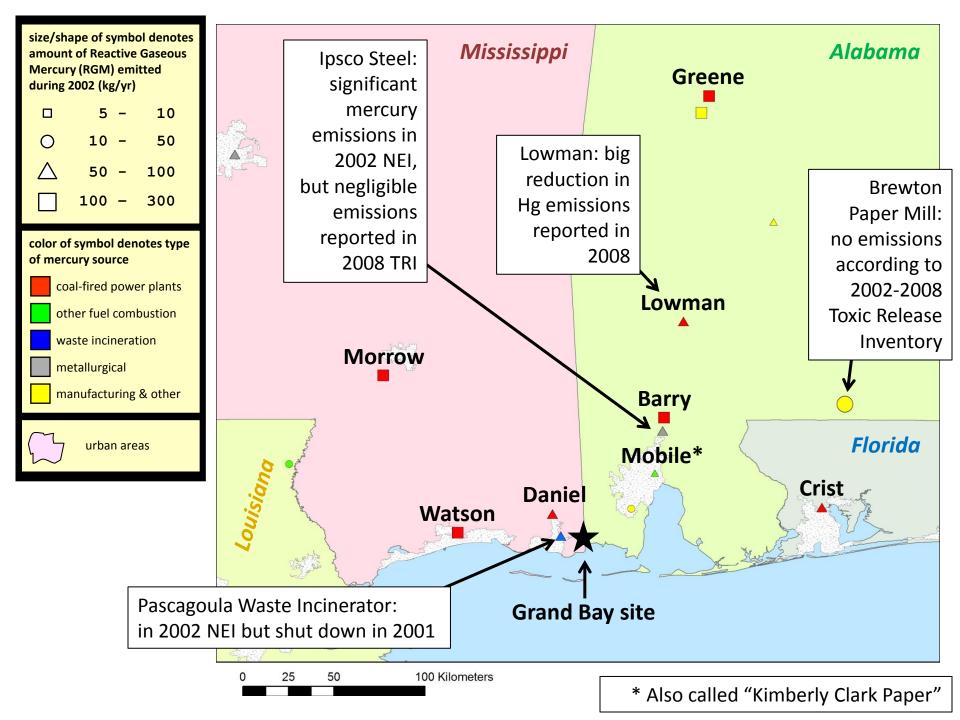


# Can we reproduce this episode with an atmospheric mercury model?

- Peaks are important to understand
- Opportunity for model evaluation
- Just one episode of many



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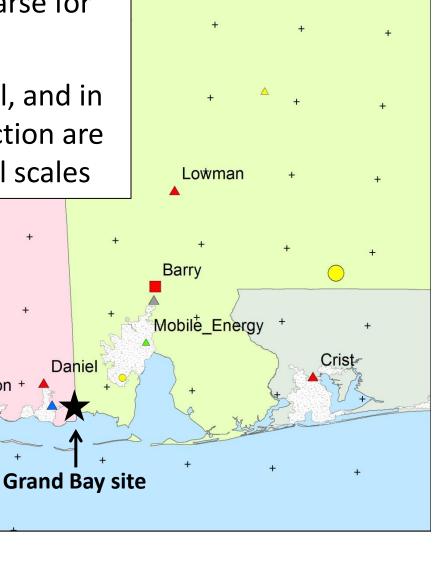


- NOAA "EDAS 40km" met data (grid points shown here)
- Ok for regional analysis, but too coarse for local analysis
- Unlikely to simulate sea-breeze well, and in general, wind speed and wind direction are unlikely to be very accurate on local scales

+ Morrow <sub>+</sub>

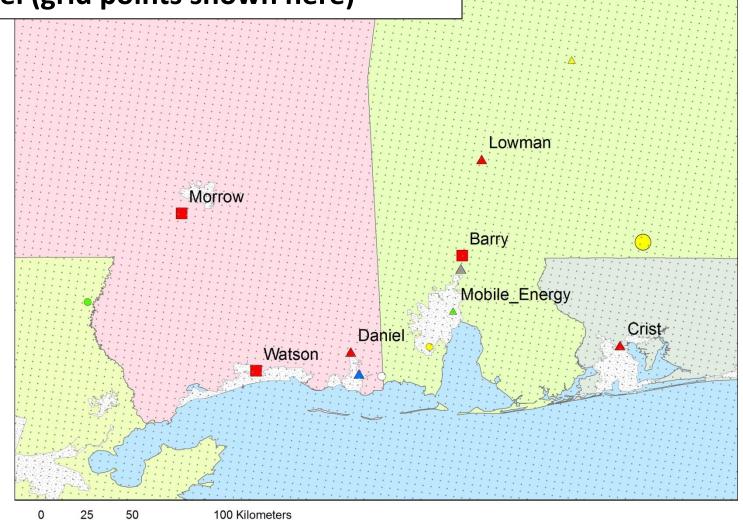
Watson +

100 Kilometers

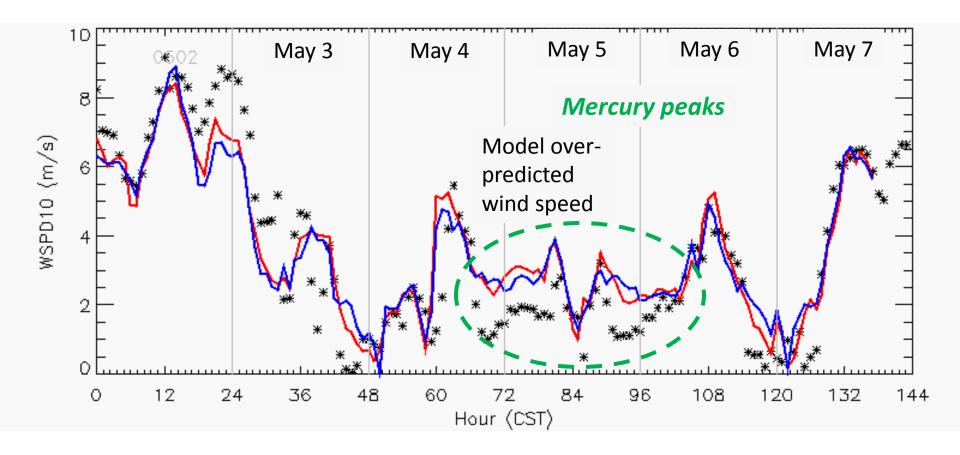


Greene County

We created a 4km resolution met data set – also with enhanced vertical resolution – using the Weather Research and Forecasting (WRF) model (grid points shown here)



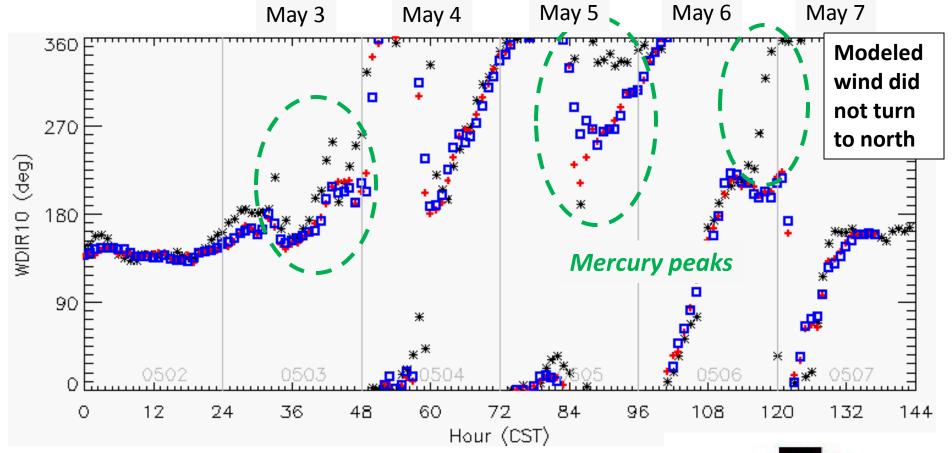
Greene County



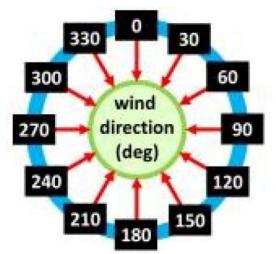
\* Measurements at the Grand Bay NERR site

Model: WRF(a) (NOAH Land Surface Model + nudging)

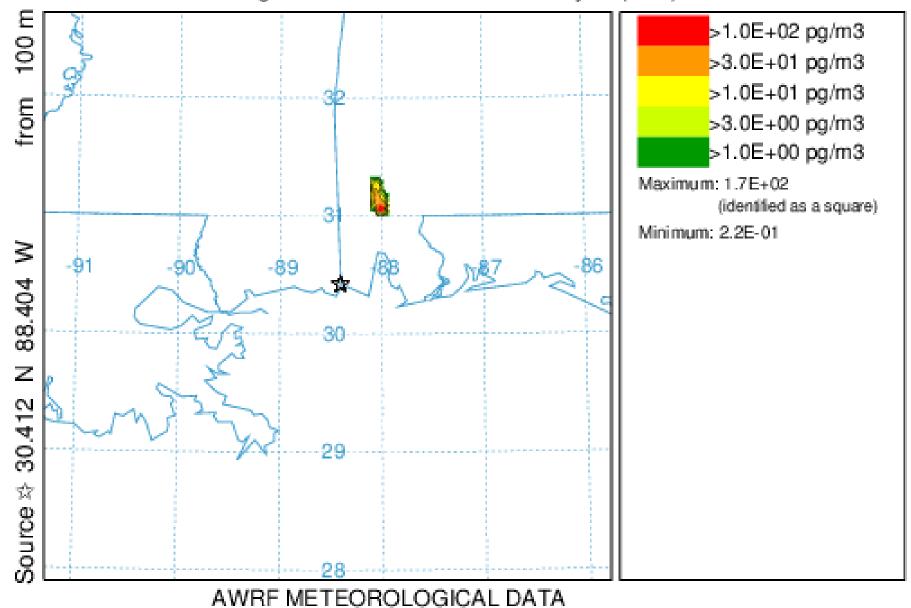
Model: WRF(b) (PX Land Surface Model + nudging)



- \* Measurements at the Grand Bay NERR site
- Model: WRF(a)(NOAH Land Surface Model + nudging)
- Model: WRF(b)(PX Land Surface Model + nudging)

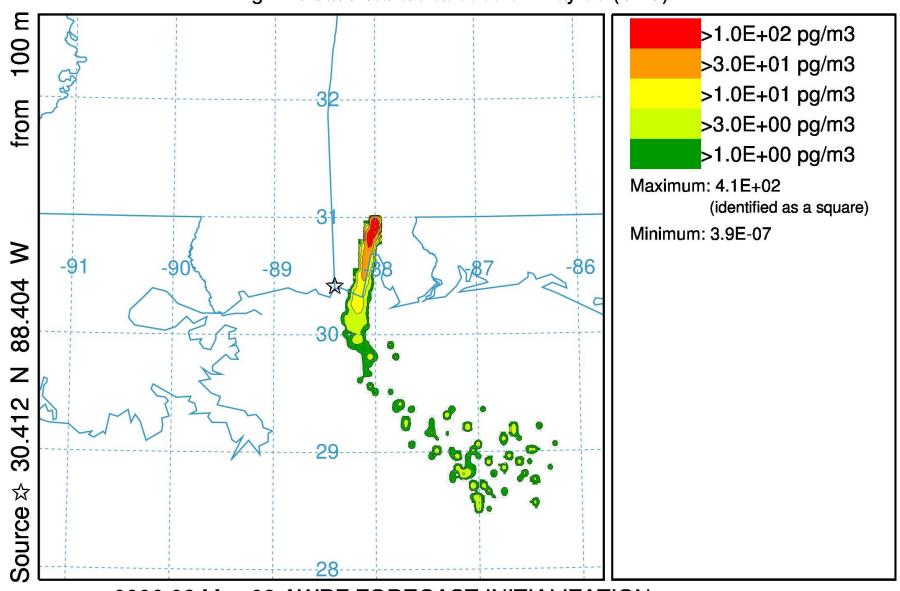


Concentration (pg/m3) averaged between 0 m and 50 m Integrated from 0000 02 May to 0100 02 May 08 (UTC) HgII Release started at 0000 02 May 08 (UTC)



Concentration (pg/m3) averaged between 0 m and 50 m Integrated from 1000 05 May to 1100 05 May 08 (UTC)

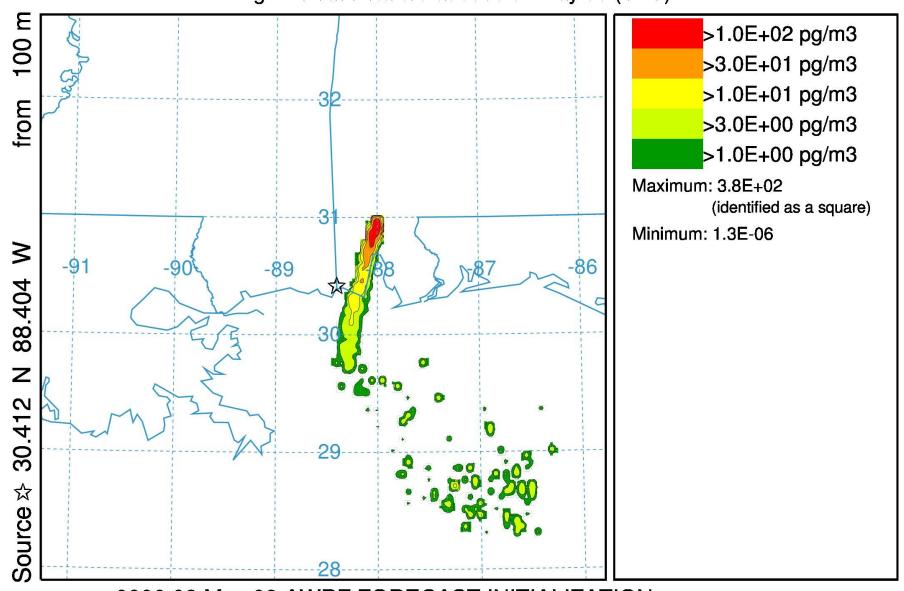
Hgll Release started at 0000 02 May 08 (UTC)



0000 02 May 08 AWRF FORECAST INITIALIZATION

Concentration (pg/m3) averaged between 0 m and 50 m Integrated from 1100 05 May to 1200 05 May 08 (UTC)

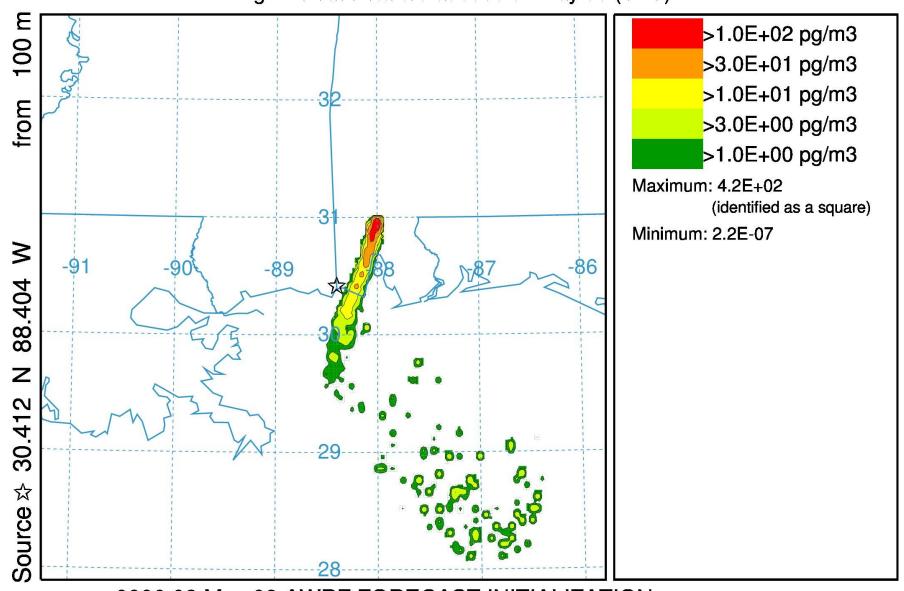
Hgll Release started at 0000 02 May 08 (UTC)



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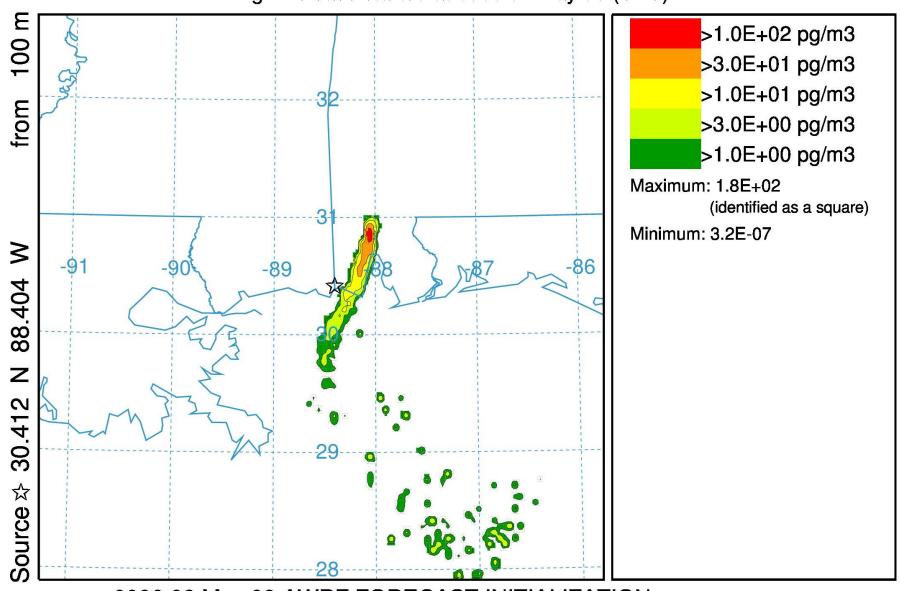
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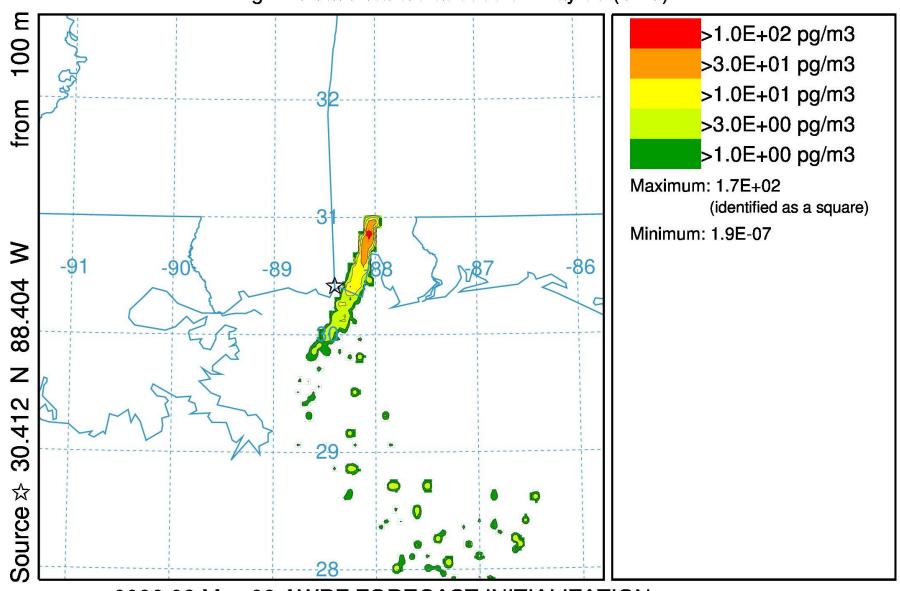
HgII Release started at 0000 02 May 08 (UTC)



0000 02 May 08 AWRF FORECAST INITIALIZATION

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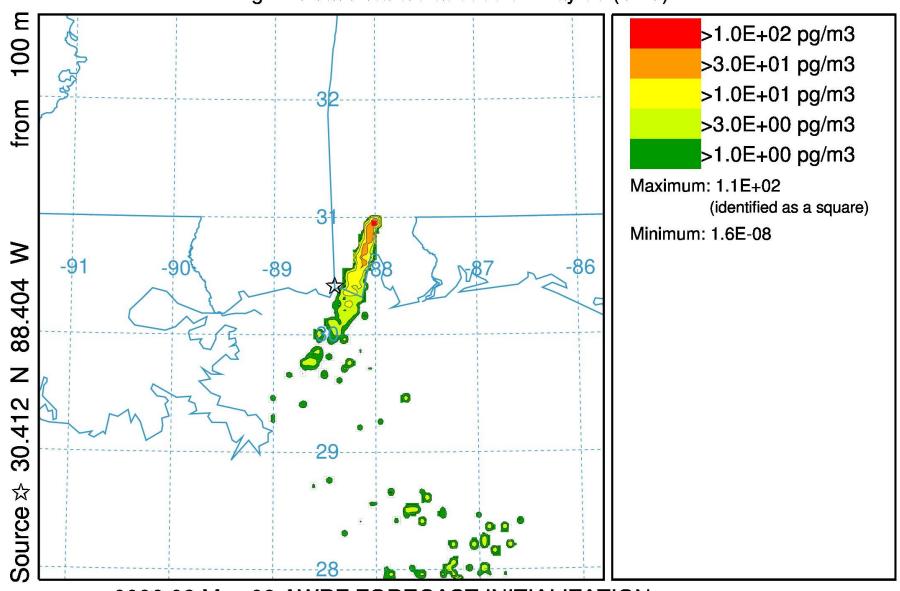
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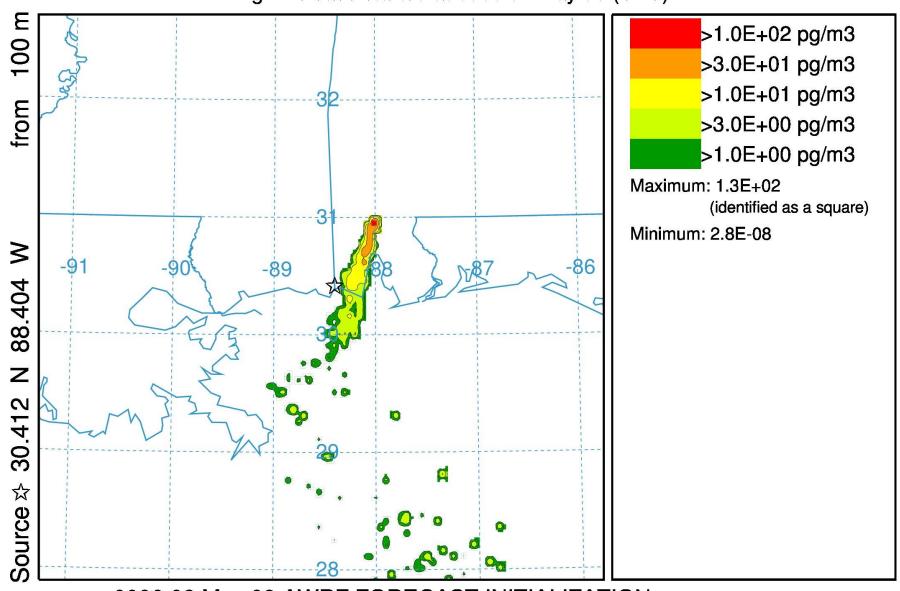
HgII Release started at 0000 02 May 08 (UTC)



0000 02 May 08 AWRF FORECAST INITIALIZATION

Concentration (pg/m3) averaged between 0 m and 50 m Integrated from 1600 05 May to 1700 05 May 08 (UTC)

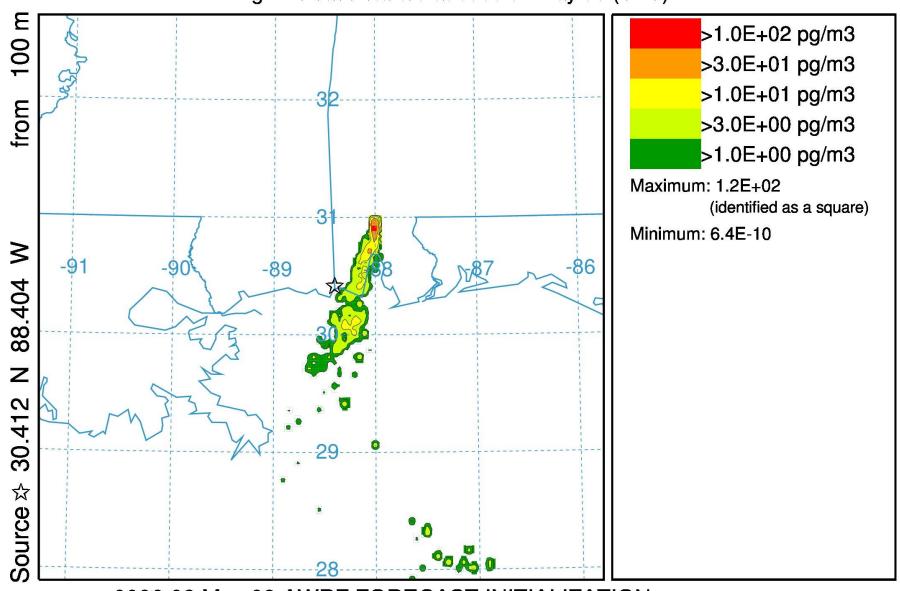
HgII Release started at 0000 02 May 08 (UTC)



0000 02 May 08 AWRF FORECAST INITIALIZATION

Concentration (pg/m3) averaged between 0 m and 50 m Integrated from 1700 05 May to 1800 05 May 08 (UTC)

Hgll Release started at 0000 02 May 08 (UTC)

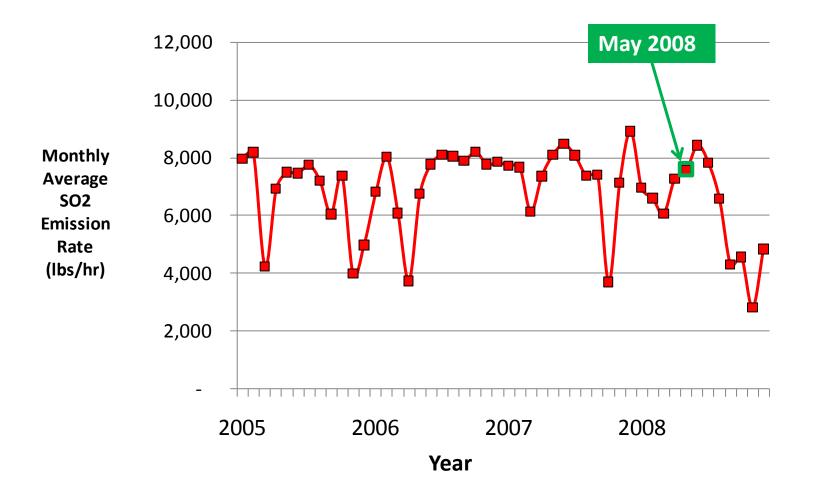


0000 02 May 08 AWRF FORECAST INITIALIZATION

This is the period that you just saw – a measured peak, but not a modeled peak (BARRY - blue line) **RGM** Measurements #2 100 RGM Concentration (pg/m3) One set of model results **BARRY** 10 **DANIEL CRIST GREENE** 12 AM **12 PM** 12 PM 12 AM 12 PM 12 PM 12 PM 12 AM 12 PM 12 AM 12 PM 12 PM 5/4/08 12 AM 5/7/08 12 AM 5/8/08 12 AM **LOWMAN** 5/3/08 5/4/08 2/5/08 2/6/08 2/6/08 5/2/08 2/1/08 5/3/08 2/2/08 5/8/08 5/2/08 **WATSON MOBILE Central Standard Time** 

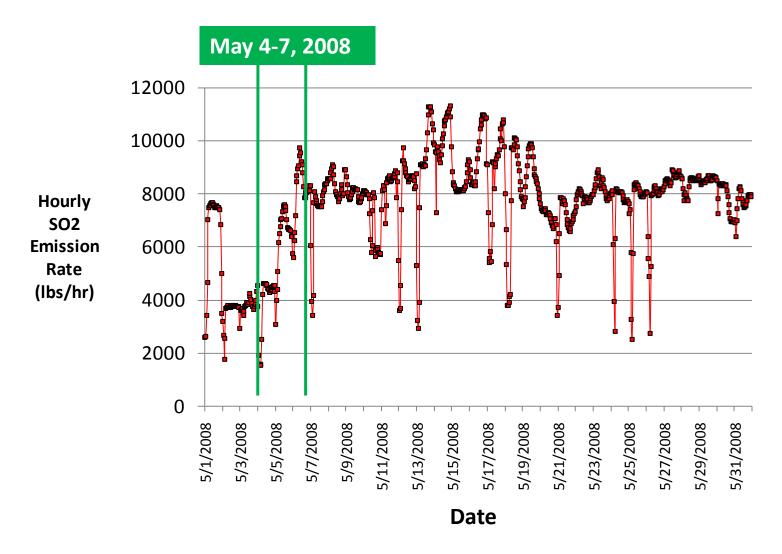
☐ We typically assume emissions are constant throughout the year

☐ May 2008: "typical" month for SO2 emissions at the Daniel plant



Source of data: USEPA Clean Air Markets Division

### But, hourly SO2 emissions at the Daniel plant were not constant... and so Hg emissions were also likely variable



Source of data: USEPA Clean Air Markets Division

#### **SOME CONCLUDING THOUGHTS (and questions):**

In order to "use" the AMNet data, we need

- Emissions information with higher temporal resolution (better than annual once every 3 years)
- Meteorological data with higher resolution

...no matter what kind of analyses we are doing

Comparing Trends; Trajectories; Dispersion

Justification for the network depends on being able to use the data meaningfully!

How can we improve this situation?

☐ This model evaluation exercise will be more fully characterized for this one episode at this one site ☐ Of course, must look at other episodes, other sites ■ Variations and uncertainties in meteorology and emissions make local plumes "stochastic"... ☐ Even if a model was "perfect" it would not be generally possible to reproduce a local plume hit at a monitoring site. ■ What is the best way to evaluate (and improve) models?

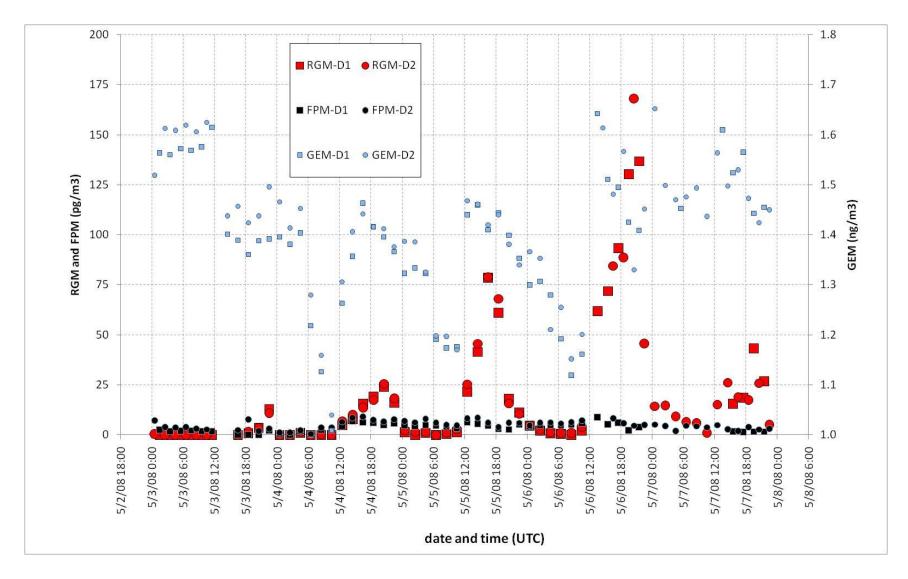


# EXTRA SLIDES

#### Modeling – Comprehensive Fate and Transport Simulations

- Start with an emissions inventory
- Use gridded meteorological data
- Simulate the dispersion, chemical transformation, and wet and dry deposition of mercury emitted to the air
- Source-attribution information needed at the end, so optimize modeling system and approach to allow source-receptor information to be captured
- HYSPLIT-Hg developed over the last ~10 years with specialized algorithms for simulation of atmospheric mercury





(Proposed Alternative) Figure 3. Time series of Reactive Gaseous Mercury (RGM), Fine Particulate Mercury (FPM) and Gaseous Elemental Mercury (GEM) from two co-located instruments (D1 and D2) (top graph) and of SO2, O3, NO, NOy, and CO (bottom graph) measured at the Grand Bay NERR from May 3-8, 2008

