Utilizing @MAQP Forcess A Aalysisishtan Attemptstandhtlerstanactseolmpiactseof Climatea Opange OnaOd and 25M2 5 C. Hogrefe<sup>1</sup>, B. Lynn<sup>2</sup>, C. Rosenzweig<sup>3</sup>, R. Goldberg<sup>3</sup>, K. Civerolo<sup>4</sup>, J.-Y. Ku<sup>4</sup>, J. Rosenthal<sup>2</sup>, K. Knowlton<sup>2</sup>, and P.L. Kinney<sup>2</sup> <sup>1</sup>Atmospheric Sciences Research Center, University at Albany <sup>2</sup>Columbia University <sup>3</sup>NASA-Goddard Institute for Space Studies <sup>4</sup>New York State Department of Environmental Conservation Models-3 Users' Workshop, September 26-28, 2005 This project was supported by the U.S. **Environmental Protection Agency under STAR** New York Climate & Health

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Proiect

# Previously shown at CMAS ...



 CMAQ simulates an increase in average (~3-5 ppb) and 4<sup>th</sup>-highest (~5-7 ppb) summertime daily maximum 8-hr ozone concentrations for future decades as a result of climate change alone

## And now the Continuation:

- Include simulation of aerosols
- Utilize regional climate simulations from two configurations of MM5
- Include process analysis

# GCM/MM5/CMAQ Model Setup

- GISS coupled global ocean/atmosphere model driven by IPCC "A2" greenhouse gas scenario
- MM5 was run on 2 nested domains of 108km and 36km over the U.S. with two cumulus parameterizations:
  - Betts-Miller (MM5-BM)
  - Grell (MM5-G)
- Simulations periods :

June – August 1993-1997 June – August 2053-2057

- 1996 U.S. Emissions processed by SMOKE
- BEIS2 for biogenic emissions and Mobile5b for mobile source emissions
- CMAQ 4.4 was run at 36km to simulate ozone
- CB-IV mechanism, aerosols, process analysis
- Note: No coupling to global chemistry model, no feedback from aerosols to climate simulations

## Effect of MM5 Cumulus Parameterization on Regional Climate Fields

Summertime Average Temperatures Simulated by MM5-BM (left) and MM5-G (right) for the 1990s (top), 2050s (center), and 2050s-1990s (bottom)



## Changes in Summertime Average Species Concentrations, 2050s – 1990s, MM5-BM/CMAQ (left), MM5-G/CMAQ (right)



Changes in Summertime Average PM<sub>2.5</sub> Species Between the 2050s and 1990s for both the MM5-BM / CMAQ and MM5-G / CMAQ Simulations



- Increase in total  $PM_{2.5}$ ,  $SO_4$ , EC, and other primary particles, decreases in  $NO_3$  and OC
- Direction of change consistent for both MM5 regional climate scenarios

 Could one parameterize these changes in concentration fields based on the changes in regional climate parameters?

## Changes in Summertime Average Ozone, HO<sub>x</sub> and Meteorology (2050s – 1990s, MM5-BM / CMAQ)



No strong relationship between patterns of changes in meteorological parameters, summertime average O concentrations, and summertime average HO, concentrations is evident

## How Are Changes In O<sub>3</sub> Related To Changes in Climate Parameters?

#### MM5-BM / CMAQ

#### MM5-G / CMAQ



Little relationship between changes in individual meteorological parameters at a given location and average O<sub>3</sub> changes at the same location for either MM5-BM / CMAQ or MM5-G / CMAQ (2050 A2 scenario)

## Correlations Between the Spatial Patterns of Changes in Summertime Average O<sub>3</sub> and Meteorology (MM5-BM)

	∆CloudFr	∆PBL	∆WaVap	ΔΤ	∆Windsp
ΔCloudFr		-0.06	0.01	0.06	0.33
ΔPBL			0.42	0.83	0.47
∆WaVap				0.25	0.44
ΔΤ					0.38
ΔCO	0.12	0.09	0.60	0.08	0.13
$\Delta O_3$	-0.10	0.04	0.48	0.01	0.07
ΔEC	-0.12	-0.32	0.00	-0.35	-0.35
$\Delta NO_3$	0.10	0.01	-0.19	0.00	0.03
ΔOC	-0.08	-0.63	-0.52	-0.53	-0.49
$\Delta SO_4$	0.20	-0.21	0.10	-0.28	-0.02

## **Process Analysis**

- Goal: Keep track of the contributions of different science processes to the changes in species concentrations
- In this analysis, Integrated Process Rates (IPR) were used and four processes were defined:
  - Vertical: Advection + diffusion + mass adjustment + dry deposition (+ emissions)
  - Horizontal: advection + diffusion
  - Clouds (includes aqueous chemistry, scavenging, cloud vertical mixing)
  - Chemistry/Aerosol Module:
- Analysis is presented for the first model layer and for the MM5-BM / CMAQ simulations only

## Temporal and Spatial Patterns of O<sub>3</sub> IPR Factors for the 1990s

#### Summertime Average of IPR Terms

#### Average Diurnal Cycles of IPR Terms





Vertical



Clouds



#### Chemistry



#### Spatially and Temporally Averaged O<sub>3</sub> Process Rates for the 1990s and 2050s



 Increase in the strength of the net chemical production rates for the future climate scenario

• Increase in the net loss due to vertical processes

#### Spatially and Temporally Averaged EC Process Rates for the 1990s and 2050s



- For EC as a primary aerosol, the vertical term (which includes emissions) is the largest net source while cloud processes are the largest net sink in the surface level
- Minor changes in strength between the horizontal and vertical components are seen between the two decades

## Changes in Summertime Average O<sub>3</sub>(left) and IPR Categories (center, right) 2050s – 1990s, MM5-BM / CMAQ



## **Correlations Between Spatial Patterns of Changes**

	ΔCO	<b>ΔΟ</b> <sub>3</sub>	ΔΕС	ΔSO <sub>4</sub>
<b>∆IPR(EC, Clouds)</b>	-0.19	-0.01	0.09	-0.02
<b>∆IPR(EC, Horizontal)</b>	0.17	0.06	0.17	0.08
<b>∆IPR(EC, Vertical)</b>	-0.11	-0.10	-0.05	-0.09
<b>∆IPR(CO, Chemistry)</b>	0.69	0.49	0.00	-0.02
<b>∆IPR(CO, Clouds)</b>	-0.26	-0.19	-0.18	-0.16
<b>∆IPR(CO</b> , Horizontal)	0.19	0.08	0.08	0.15
<b>∆IPR(CO, Vertical)</b>	-0.17	-0.11	0.05	-0.14
$\Delta$ IPR(O <sub>3</sub> , Chemistry)	0.57	0.70	0.37	0.31
ΔIPR(O <sub>3</sub> , Clouds)	0.09	-0.05	0.01	0.35
$\Delta$ IPR(O <sub>3</sub> , Horizontal)	0.10	-0.03	-0.10	0.09
ΔIPR(O <sub>3</sub> , Vertical)	-0.24	-0.21	-0.01	-0.24

## **Correlations Between Spatial Patterns of Changes**

	<b>∆CloudFr</b>	APBL	∆WaVap	ΔΤ	∆Windsp
ΔCΟ	0.12	0.09	0.60	0.08	0.13
ΔΕС	-0.12	-0.32	0.00	-0.35	-0.35
∆NO <sub>3</sub>	0.10	0.01	-0.19	0.00	0.03
∆SO <sub>4</sub>	0.20	-0.21	0.10	-0.28	-0.02
<b>∆IPR(EC, Clouds)</b>	-0.29	0.02	-0.27	0.02	-0.31
<b>∆IPR(EC, Horizontal)</b>	0.19	0.07	0.11	0.05	0.14
<b>∆IPR(EC, Vertical)</b>	-0.20	-0.01	-0.14	0.05	-0.11
$\Delta$ IPR(O <sub>3</sub> , Chemistry)	-0.17	0.05	0.37	-0.02	-0.04
∆IPR(O <sub>3</sub> , Clouds)	0.39	-0.17	0.04	-0.26	0.21
$\Delta$ IPR(O <sub>3</sub> , Horizontal)	0.34	0.13	0.00	0.14	0.09
$\Delta$ IPR(O <sub>3</sub> , Vertical)	-0.19	0.03	-0.25	0.10	-0.08

## Summary

- CMAQ simulations with regional climate change under the IPCC A2 scenario for the 2050s shows an increase of up to  $1 \mu g/m^3$  in summertime average total PM<sub>2.5</sub> concentrations, mostly driven by increases in sulfate
  - Decreases in the volatile species nitrate and organic carbon are more than offset by increases in sulfate and primary PM<sub>2.5</sub> species
  - The directionality of changes is consistent for two different MM5 configurations
  - $\Rightarrow$  Performing regional climate ensemble modeling studies could help to quantify the uncertainty around simulated pollutant changes as a result of climate change
- Process analysis: strongest link between climate change and changes in pollutant concentrations is through chemical production rates for reactive gas-phase species (via water vapor / radical chemistry?)
- But: Even the strongest linear regression associations explain less than half of the concentration changes simulated by CMAQ
- This implies that the simulated changes in pollutant concentrations stemming from climate change are the result of a complex interaction between changes in transport, mixing and chemistry that cannot be parameterized by spatially uniform linear regression relationships
- Therefore, full-science photochemical modeling systems such as CMAQ are the tool of choice for quantitatively studying the impact of climate change on regional-scale air pollution.
- Need to include global chemistry models and aerosol/climate feedback