

# Task Force on Hemispheric Transport of Air Pollution

Work Package outline for:

# Task Force on Hemispheric Transport of Air Pollution Work Plan 2012-2016

Work Package 3.2: Inflow processes influencing air quality over western North America

Lead: Owen R. Cooper<sup>1,2</sup>: owen.r.cooper@noaa.gov Contributors: Greg Carmichael<sup>3</sup>, Joshua Fu<sup>4</sup>, Tracy Holloway<sup>5</sup>, Meiyun Lin<sup>6,7</sup>

<sup>1</sup>Cooperative Institute for Research in Environmental Sciences, University of Colorado, Boulder

<sup>5</sup>Nelson Institute for Environmental Studies, University of Wisconsin-Madison

<sup>6</sup>Atmospheric and Oceanic Sciences, Princeton University, Princeton, NJ

<sup>7</sup>NOAA Geophysical Fluid Dynamics Laboratory, Princeton, NJ

**Updated:** June 28, 2012

# 1. Purpose

In recognition of an increasing body of scientific evidence suggesting the potential importance of intercontinental transport of air pollutants, the UNECE (United Nations Economic Commission for Europe) Convention on Longrange Transboundary Air Pollution (CLRTAP) created the Task Force on Hemispheric Transport of Air Pollution (TF HTAP). Established in 2004, this international team of scientists was charged with improving our understanding of the intercontinental transport of air pollutants across the Northern Hemisphere. TF HTAP completed its first assessment report in 2010, covering ozone, particulate matter, persistent organic pollutants and mercury [*Dentener et al.*, 2010]. Later that year the LRTAP Executive Body revised and expanded the TF HTAP mandate to continue to examine the transport of air pollution across the Northern Hemisphere, identify emission mitigation opportunities, and assess the impacts of those mitigation opportunities on regional and global air quality, public health, ecosystems and climate change. (Further background information on TF HTAP and its activities can be found at http://www.htap.org)

In response to this new mandate a TF HTAP meeting was held in Pasadena, California, Feb 1-3, 2012, where the TF HTAP Work Plan 2012-2016 was developed around the following 6 themes:

Theme 1: Emission inventories and projections

<sup>&</sup>lt;sup>2</sup>NOAA Earth System Research Laboratory, Boulder, CO

<sup>&</sup>lt;sup>3</sup>Department of Chemical & Biochemical Engineering, The University of Iowa, Iowa City

<sup>&</sup>lt;sup>4</sup>Department of Civil and Environmental Engineering, The University of Tennessee, Knoxville

Theme 2: Source Apportionment and Source/Receptor Analysis Theme 3: Model-Observation Evaluation and Process Diagnosis Theme 4: Assessment of Health, Ecosystem, and Climate Impacts Theme 5: Assessment of Climate Change impact on Pollution Theme 6: Expansion of the Data Network and Analysis Tool

The work under each of the 6 themes will contribute to the achievement of three overarching goals:

- 1) deliver policy-relevant products
- 2) improve scientific understanding, and
- 3) reach-out beyond CLRTAP to create a common understanding of hemispheric air pollution issues.

One specific science question to be answered under Theme 3 is: How well do current global and regional chemical transport models (CTMs) simulate the atmospheric boundary layer (ABL) processes that transport intercontinental pollution plumes from the free troposphere down to the surface of a receptor region? To address this question, TF HTAP participants developed Work Package 3.2: Inflow processes influencing air quality over western North America. This paper provides an experimental outline to explore these continental inflow processes using global and regional scale CTM simulations of ozone and particulate matter (PM), and it also serves as a call for modeling groups to participate in the experiment. Details of the proposed study are provided in Sections 2-6 below, while a summary and dates for deliverables are provided in Table 1.

#### 2. Background

TF HTAP's 2010 assessment report, *Hemispheric Transport of Air Pollution, Part A: Ozone and Particulate Matter* [*Dentener et al.*, 2010] reported several findings and recommendations regarding continental inflow processes, paraphrased as follows.

The most intense pollution plumes are found in the mid- and upper troposphere but to be relevant to air quality in a downwind continent, the pollution must descend to the surface. As the plumes descend they are diluted and can be difficult to distinguish from local pollution, especially in receptor regions with relatively high emissions. While great progress has been made in forecasting and intercepting intercontinental pollution plumes in the free troposphere, to date there has been little focus on measuring the composition of these plumes as they descend into the boundary layer of the receptor continent (Chapter 1.4). Existing studies show that onshore marine airflow above the marine boundary layer can carry ozone concentrations that approach urban air quality standards, and that air can mix to the surface and contribute substantially to air quality standard violations. This is particularly noticeable in low emission regions. The impact depends upon vertical mixing of air into the ABL, which is enhanced by complex topography (Chapter 2.2).

Resolution determines a model's ability to resolve local features that influence import processes and chemical transformation. In particular, topography affects the mixing of free tropospheric pollutants into the ABL, and sharp gradients in the chemical environment (especially in urban regions) affect the strength of surface ozone responses to emission changes. Model resolution also affects the response of pollutants at specific receptors of interest, e.g., Megacities, and is an important source of uncertainty in the analysis of human health impacts in urban areas, as urban areas are not adequately resolved in global models (Chapter 4.5.2). Many urban areas have strong gradients in emissions and topography, and this can lead to imported ozone and precursors having qualitatively different impacts on urban areas and their surroundings due to the strong dependency of ozone chemistry on the ratio of NO<sub>x</sub> to VOCs (Chapter 4.5.3).

In response to these findings the TF HTAP 2010 assessment report recommended support for the application of high-resolution global and regional models to investigate the effect of smaller-scale processes on continental import and export budgets (Chapter 4.5.2).

## 3. Continental Inflow Processes

The 21 CTMs used for the experiments in *Hemispheric Transport of Air Pollution, Part A: Ozone and Particulate Matter* were run at various horizontal resolutions, with 18 being coarser than 2x2 degrees, and none being finer than 1x1 degree. These resolutions were adequate for quantifying broad impacts of upwind emission reductions on downwind receptor regions as large as the USA or Europe. However, these relatively coarse simulations are not able to demonstrate the impact of upwind emissions on scales less than 100 km especially in regions with variable terrain [*Huang et al.*, 2010]. Key import processes to be explored with finer scale models are:

- Quantification of low altitude vs. high altitude import Transport of upwind pollution to a surface receptor can either occur entirely within the ABL, or it can occur in the free troposphere followed by descent into the ABL at the receptor. Previous studies indicate descent from the free troposphere may be the most effective route, however the resulting subsidence inversion of the descending air mass creates a barrier that must be overcome if the pollution is to reach the surface. How much of the upwind pollution that impacts the receptor region is imported entirely within the ABL and how much descends into the ABL?
- 2) Influence of boundary layer depth What is the difference in the quantity of pollution transported to the surface on days with deep mixed layers vs. days with shallow mixed layers? Nighttime and daytime conditions can also be compared.
- 3) Influence of terrain Higher terrain has a greater impact from intercontinental plumes because at night these regions can lie above the nighttime temperature inversion, and during daytime high elevation regions have mixed layers that penetrate further into the free troposphere allowing plumes at higher altitudes to be mixed down to the surface. To what degree do high elevation sites experience greater impact from upwind sources?
- 4) Impact of upwind pollution on rural and urban regions How do imported pollution plumes impact the chemistry and pollutant loading in rural areas and in nearby urban areas [*Lin et al.*, 2010]? In which cases do imported plumes increase the probability of air quality standard exceedances? In which cases do strongly descending pollution plumes, already diluted after long transport times, merely displace greater quantities of local pollutants, lowering the probability of air quality exceedances?
- 5) The same processes that bring pollution plumes to the surface also affect stratospheric intrusions that reach the lower troposphere. What is the impact of intrusions on surface ozone [*Zhang et al.*, 2011; *Langford et al.*, 2012; *Emery et al.*, 2012]?

#### 4. Western North America study region

Western North America is an ideal region for studying continental inflow processes due to its location downwind of Asia, its varied terrain and boundary layer heights, its mix of rural and urban areas as well as the availability of extensive meteorological and chemical measurements for model evaluation.

Modelling the intercontinental transport of air pollution is most straight forward when there is a strong pollution signal. Western North America is downwind of East Asia, the region of the world with the fastest growing emissions, with NO<sub>x</sub> emissions exceeding all other regions in 2010 (Figures 1 and 2). Large East Asian pollution plumes are frequently exported across the North Pacific Ocean and reach western North America several days later [*Hudman et al.*, 2004; *Wang et al.*, 2006; *Lin et al.*, 2012].

Western North America's terrain allows for a variety of inflow processes to be explored as higher elevation regions tend to have greater impact from free-tropospheric air masses transported into North America [Zhang et al., 2011; Cooper et al., 2011]. The major terrain features are distinct enough that regional scale models run at resolutions as course as 50 km can resolve the largest valleys and mountain ranges (Figure 3). The region's well defined coastline allows for clear comparisons to be made between marine boundary layer and continental boundary layer transport process. The marine boundary layer meets the land in areas with both high coastal mountains and low lying terrain. The Central Valley of California is an extensive intermontane basin close to sea level that interfaces with both the marine boundary layer and the surrounding mountains. Mountain ranges near the coast or the Central Valley can rise abruptly from the marine boundary layer or low elevation valleys to heights 2000-3000+ m above sea level. Further inland, regions such as Nevada, Arizona, Utah and Colorado contain mountain ranges situated on high plateaus increasing the likelihood of interaction with free tropospheric air. The varied terrain also influences the maximum height of the daytime mixed layer with depths less than 1 km over the cool eastern North Pacific Ocean, while mixed layers over high mountainous terrain can reach several km. High surface temperatures across the elevated deserts of the southwestern USA can routinely push the mixed layer to 4 km above sea level.

Demographically, western North America is home to 57 million people living within 350 km of the coast, from southern British Columbia to northern Baja California. Anthropogenic emissions closely match the population distribution (Figure 3) with major urban emission regions being Vancouver/Seattle, Portland, San Francisco, the Central Valley, Los Angeles/San Diego/Tijuana, Phoenix and Salt Lake City. Western North America also contains extensive low-emission rural regions, and there are often abrupt spatial transitions between the high and low emission regions.

Meteorological and chemical measurements for model evaluation are widely available across western North America, especially during the CALNEX, CalMex and CARES field experiments across California and northwest Mexico in May-June 2010. Routine surface hourly meteorological observations are made at dozens of National Weather Service locations across the region. Important for evaluating model simulations of boundary layer depth, the National Weather Service launches rawinsondes from 18 sites across the region (Figure 4) at 4:00 and 16:00 local time every day, which provides the ABL depth close to the times of its daily minimum and maximum. During May-June 2010 additional daytime temperature and humidity profiles are available from 6 ozonesondes sites, two ground sites near Sacramento and a ground site on the US-Mexico border (Figure 4). Boundary layer depth and wind vectors are also available from this period as measured by 11 NOAA radar wind profilers.

Figure 5 shows the locations of 18 rural ozone monitors across the western United States, maintained by the National Park Service, the EPA CASTNET program, the University of Washington and NOAA. While these rural sites are best for evaluating the influence of modeled pollution plumes, the models can also be compared to dozens of urban ozone monitors throughout the western USA (archived by the EPA), mainly in central and southern California as well as the urban regions of Seattle, Salt Lake City, Las Vegas, Phoenix and Tucson. Many rural and urban ozone monitors are also located further east in Wyoming and Colorado. Dozens of IMPROVE aerosol composition monitoring sites are also spread

across the western USA (Figure 6), measuring PM2.5 and PM10 mass (also mass of constituents such as Cl-, NO3-, SO4=, OC) as 24-hour samples, once every 3 days. Other routine chemical measurements include:

- hourly observations in the marine boundary layer at Trinidad Head, CA, of ozone, aerosol back scatter, absorption, and total particle number (data collected by NOAA).
- hourly measurements of ozone, CO, total Hg, and aerosol scattering, size and chemistry at Mt. Bachelor, Oregon (2.7 km a.s.l.), (collected by U. of Washington, *Fischer et al.*, 2011a).
- weekly ozonesondes at Trinidad Head; continuous aerosol lidar at Trinidad Head (operated by the NASA Micro-Pulse Lidar Network )
- tropospheric ozone lidar at Table Mountain, southern California, operated by NASA JPL.

During May-June 2010 the CALNEX experiment brought extra measurements to California. Six ozonesondes sites were implemented with near daily ozonesondes [*Cooper et al.*, 2011]. The NOAA WP-3D aircraft sampled the chemistry of the lower troposphere above central and southern California during 18 flights. The NOAA Twin Otter flew 46 flights across California with a downward looking ozone lidar. The NASA King Air made 31 flights equipped with a downward looking high spectral resolution lidar (HSRL) for measuring aerosol extinction, backscatter and optical thickness as well as planetary boundary layer heights. The CIRPAS Twin Otter and the DOE G-1 aircraft made extensive aerosol measurements across the Los Angeles Basin and Sacramento, respectively. Ground sites for measuring trace gases and aerosols were located in Los Angeles, Bakersfield, Sacramento and Tijuana, with two tall towers in central California for measuring carbon cycle gases. The R/V Atlantis made trace gas and aerosol measurements in the marine boundary layer along the coast of central and southern California. Three UC Davis rotating drum impactors provided continuous, 3-hr resolution PM mass at 8 stages from 10  $\mu$ m to 90 nm. Finally, continuous PAN measurements were made from Mt. Bachelor during May, 2010 [*Fischer et al.*, 2011b].

Figure 7 shows the distribution of the CALNEX ozonesonde network, the design of which was inspired by the research needs outlined in the TF HTAP 2010 assessment report. Four of the ozonesonde sites were located on the coast to measure baseline ozone, while ozone was measured within the polluted regions of California by two additional ozonesonde sites and the NOAA WP-3D aircraft. Comparison of the baseline profiles to the inland profiles shows that average daytime baseline ozone throughout the depth of the daytime boundary layer was equal to more than 80% of the ozone measured above the Central Valley, and 63-76% of the measured ozone above Joshua Tree National Park and the Los Angeles Basin [*Cooper et al.*, 2011]. Two recently published papers have identified three major Asian plumes that impacted the western USA during CALNEX (all sampled by the ozonesondes or NOAA WP-3D) [*Lin et al.*, 2012], as well as stratospheric intrusions that impacted the Los Angeles region [*Langford et al.*, 2012]. *Lin et al.* [2012] show that satellite retrievals of tropospheric trace gases can be used to constrain model simulations of trans-Pacific pollution plume transport (Figure 8).

#### 5. Design of regional scale model experiment

The objective of Work Package 3.2 is to evaluate the ability of global and regional scale CTMs to represent processes driving the import of intercontinental air pollution into western North America and its influence on surface air quality using observations from intensive field campaigns in May-June 2010. The goal is not to determine which model is most accurate but to evaluate how the range of regional scale models in use today simulate pollution inflow processes and to gauge the improvement in accuracy afforded by regional models in comparison to the range of global scale model simulations. To meet this

objective several global and regional-scale CTMs (five or more in each category) will be used to study the impact of terrain (large mountain chains vs. broad valleys) and ABL depth (marine boundary layer, daytime mixed layer, nighttime inversion) on the transport of intercontinental pollution plumes down to the surface. Another important goal is to quantify the impact of intercontinental plumes on surface chemistry in rural and urban areas. These processes will be studied with global scale models at 1-2.5 degree horizontal resolutions and with regional scale models that operate on horizontal scales of 50 - 12 km. The regional models also need a vertical resolution fine enough to resolve the transition between the ABL and the free troposphere.

The time period for the model intercomparison is May-June 2010 due to the extensive chemical and boundary layer measurements made across California. The model domain should cover all of California but it would be helpful if the domain could also include Vancouver and Tijuana so that air quality issues in countries other than the USA (i.e. Canada and Mexico) are also addressed. Ideally the eastern boundary of the intercomparison should extend to central Arizona to take advantage of several rural ozone and CASTNET monitoring sites. Models that can also cover Colorado will provide extra information on the impact of long range transport on the highest terrain of the western USA as well as the impact on the Denver region located eastward of, and 2 km below, the Continental Divide.

The regional scale models will require realistic boundary conditions of the chemical and particulate matter composition of the baseline air flowing into western North America; climatological boundary conditions will not suffice. The recommendation is that the boundary conditions be provided by global scale models that are run at half degree resolution or better (such as the newly developed GFDL AM3 model) to limit the influence of numerical diffusion that artificially dilutes plumes transported over long distances. The latest methods in downscaling between boundary conditions generated by global models to finer resolution regional models should also be considered [*Lam and Fu*, 2010]. Because the experiment is focused on assessing the range of simulations produced by models in use today, regional scale modeling groups are free to use any global scale model for their boundary conditions. The experimental design accepts that part of the variability between regional scale model simulations will be due to variation in model boundary conditions. To produce boundary conditions as accurate as possible, the global scale model strengths and weaknesses regarding air quality prediction and assessments in Asia. MICS-III simulations for spring 2010 will improve our understanding of the Asian pollution events that impact western North America.

A new global emission inventory will be compiled for the latest round of HTAP model studies, based on EDGAR v4.2 (for the year 2008) with 2010 regional updates for North America, Europe and East Asia. The inventory is due for release on January 1, 2013. The emission inventory and observations for model comparison will be made available to TF HTAP participants through centralized data bases in standard formats.

The model simulations and measurement comparisons will proceed as follows:

 Several global CTMs will be run for spring and summer 2010 to quantify the impact of Asian emissions on surface ozone and PM in western North America. Requirements are a base run for 2010 with full global emissions, and a run with zero East Asian anthropogenic emissions to quantify the impact of East Asia on western North America. In addition 2 perturbation runs will be are conducted, with East Asian anthropogenic emissions increased and decreased by 20%.

- 2) Output from the global models will be used as boundary conditions for the regional models. Regional models are free to use any global scale model as a source for boundary conditions. The experimental design accepts that part of the variability between regional scale model simulations will be due to variation in boundary conditions.
- 3) Output from each global and regional scale model will be compared to observations:

A) Compare model daytime boundary layer depth to temperature and humidity profiles from the rawinsonde network. Do the models accurately simulate the entrainment of free tropospheric air into the ABL? Comparisons will be made on a monthly basis (May and June), as well as on a daily basis for case studies.

B) Use the rawinsonde network to identify the general structure of the subsidence inversion associated with the eastern North Pacific anticyclone. How well do the models simulate this inversion and what effect does it have on the quantity of pollutants entrained into the ABL? How much of the upwind pollution that impacts the receptor region is imported entirely within the ABL and how much descends into the ABL?

C) Compare modeled ozone and PM to the ozonesondes and to the measured values at high and low elevation surface sites across western North America. What is the influence of terrain on the quantity of pollution transported to the surface? Are impacts greatest in the high elevation regions of Nevada, Utah, Arizona and Colorado?

D) Perturbation experiments: What is the impact of 20% reduction and 20% increases in East Asian anthropogenic emissions on ozone and PM in western N. America? Are impacts proportional at both high and low elevation sites? Are impacts proportional in rural and urban areas?

E) When assessing the impacts of foreign pollution at the surface, attention will be paid to the absolute contribution to surface pollution and to the number of exceedances of the U. S. National Ambient Air Quality Standards for ozone and PM (as well as exceedances of air quality standards in the neighboring regions of Canada and Mexico).

4) Finally, the ensemble of regional scale model results will be compared to the ensemble of global scale model results. To what degree do regional-scale models improve our understanding of the impacts of intercontinental transport on surface air quality in the receptor region?

Results from the model intercomparison will be documented in a manuscript for submission to a peerreviewed journal such as *Journal of Geophysical Research* or *Atmospheric Chemistry and Physics*. The results will also be made available to the TF HTAP community for inclusion in any subsequent assessment reports.

# 6. Participating Modeling groups

Now that the outline of Work Package 3.2 has been developed the organizers are placing a call to ask for the participation of regional and global-scale modeling groups. Participation is open to any modeling group that agrees to follow forthcoming TF HTAP protocols for data sharing and archiving, output format, and participation in the publication of the results in the peer-reviewed literature. The timeline for the experiment and deliverables is given in Table 1. Participation in Work Package 3.2 is entirely voluntary and modeling groups must provide their own funding. Interested participants should contact Owen Cooper (owen.r.cooper@noaa.gov). Following is a list of groups that have so far agreed to participate in Work Package 3.2.

Regional-scale (half degree and finder resolutions) Greg Carmichael and Min Huang, U. of Iowa STEM CTM, 12 km resolution Tracy Holloway, U. of Wisconsin CMAQ CTM Meiyun Lin, Princeton/NOAA GFDL NOAA FDL AM3 global chemistry-climate model, half degree resolution Joshua Fu and Xinyi Dong, U. of Tennessee-Knoxville CMAQ CTM, two ways including direct and indirect in CMAQ, down to 4x4 km Owen Cooper, U. of Colorado/NOAA ESRL: FLEXPART Lagrangian Particle Dispersion Model, half degree resolution, passive tracers only *Global scale (resolutions coarser than half degree)* Greg Carmichael and Min Huang, U. of Iowa STEM CTM, 2 degree resolution??? Meiyun Lin, Princeton/NOAA GFDL NOAA FDL AM3 global chemistry-climate model, 2 degree resolution??? Joshua Fu and Xinyi Dong, U. of Tennessee-Knoxville

Hemispheric CMAQ, 1 degree resolution

*Acknowledgements:* We acknowledge the free use of tropospheric NO<sub>2</sub> column data from the GOME and SCIAMACHY sensors from <u>www.temis.nl</u>. Global population data for the year 2010 were made available by the Center for International Earth Science Information Network (CIESIN), Columbia University, the United Nations Food and Agriculture Programme (FAO) and Centro Internacional de Agricultura Tropical (CIAT); see full reference below. EDGAR v4.1 data for the year 2005 were downloaded from <u>http://edgar.jrc.ec.europa.eu/index.php</u>; see full reference below. The global topography data at 4 minute resolution data were downloaded from: NOAA National Geophysical Data Center, Boulder, <u>www.ngdc.noaa.gov/mgg/global/global.html</u>.

#### **Table 1.** Summary of Work Package 3.2

Work Package	3.2	
Title	Inflow processes influencing air quality over the	
	Western North America	
Lead	Owen Cooper	
Contributor	Greg Carmichael, Joshua Fu, Tracey Holloway, Meiyun Lin	
Start date	01.06.2012	
End date	01.05.2014	

## **Objectives:**

To evaluate the ability of global and regional models to represent processes driving the import of intercontinental air pollution into western North America and its influence on surface air quality using observations from intensive field campaigns in May-June 2010.

# **Task Descriptions:**

- Task 3.2.1 Contribute to specifications document of model experiments (WP 2.2) Identify the model outputs needed to perform the model to observation comparisons needed in Task 3.2.3.
- Task 3.2.2 Collect relevant measurement data and make available in networked archive

Identify the available observational data sets for the time and spatial domain of interest: Western North America, for May-June 2010. Make data sets available through one of the distributed archives participating in Theme 6 (AeroCom, NILU, ENSEMBLE, ...) or create a new node in the data network. Note that much of the relevant observational data from national networks will already be available through ENSEMBLE for purposes of AQMEII.

Relevant observational data include:

- CALNEX, CalMex and CARES field experiments
- Routine surface hourly meteorological observations from NWS
- National Weather Service rawinsondes
- NPS, IMPROVE
- CASTNET
- NOAA Trinidad Head surface, ozonesondes, lidar
- NASA Table Mountain ozone lidar
- University of Washington Mt. Bachelor Observations
- Task 3.2.3 Conduct analysis of 2010 model simulations in comparison to observations Specific analyses that will be conducted include:
  - Compare model daytime boundary layer depth to temperature and humidity profiles from the rawinsonde network. Do the models accurately simulate the entrainment of free tropospheric air into the ABL? Comparisons will be made on a monthly basis (May and June), as well as on a daily basis for case studies.
  - Use the rawinsonde network to identify the general structure of the subsidence inversion associated with the eastern North Pacific anticyclone. How do the models simulate this inversion and what effect

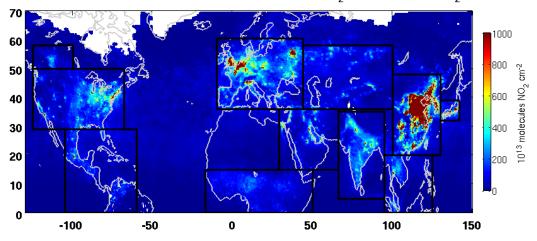
	<ul> <li>does it have on the quantity of pollutants entrained into the ABL? How much of the upwind pollution that impacts the receptor region is imported entirely within the ABL and how much descends into the ABL?</li> <li>3) Compare modeled ozone and PM to the ozonesondes and to the measured values at high and low elevation surface sites across western North America. What is the influence of terrain on the quantity of pollution transported to the surface? Are impacts greatest in the high</li> </ul>
	<ul> <li>elevation regions of Nevada, Utah, Arizona and Colorado?</li> <li>4) Perturbation experiments: What is the impact of 20% reduction and 20% increases in East Asian emissions on ozone and PM in western N. America? Are impacts proportional at both high and low elevation sites?</li> </ul>
	<ul> <li>Are impacts proportional in rural and urban areas?</li> <li>5) When assessing the impacts of foreign pollution at the surface, attention will be paid to the absolute contribution to surface pollution and to the number of exceedances of the U. S. National Ambient Air Quality Standards for ozone and PM (as well as exceedances of air quality</li> </ul>
	<ul> <li>standards in the neighboring regions of Canada and Mexico).</li> <li>6) This model intercomparison will benefit greatly from coordination with MICS-III, a multi-scale model intercomparison to evaluate model strengths and weaknesses regarding air quality prediction and</li> </ul>
	<ul> <li>assessments in Asia. MICS-III simulations for spring 2010 will improve our understanding of the Asian pollution events that impact western North America.</li> <li>7) Finally, the results from this regional-scale model inter-comparison will</li> </ul>
	be compared to coarser scale simulations by global scale models participating in the HTAP model experiments. To what degree do regional-scale models improve our understanding of the impacts of intercontinental transport on surface air quality in the receptor region?
	Results of analyses will be circulated via the HTAP wiki for comments by Task Force participants.
Task 3.2.4	Publish paper based on analysis.

Deliverables:			
3.2.1 Model output specification	1-Sep-2012		
3.2.2 Observational data available for model evaluation	1-Jan-2013		
3.2.3 Draft analysis results	1-Oct-2013		
3.2.4 Submit Publication	1-Jan-2014		

#### **References:**

- Boersma, K. F., H. J. Eskes and E. J. Brinksma (2004), Error analysis for tropospheric NO<sub>2</sub> retrieval from space, J. Geophys. Res., 109, D04311, doi:10.1029/2003JD003962.
- Center for International Earth Science Information Network (CIESIN), Columbia University; United Nations Food and Agriculture Programme (FAO); and Centro Internacional de Agricultura Tropical (CIAT). 2005. Gridded Population of the World: Future Estimates (GPWFE). Palisades, NY: Socioeconomic Data and Applications Center (SEDAC), Columbia University. Available at http://sedac.ciesin.columbia.edu/gpw
- Cooper, O. R., S. J. Oltmans, B. J. Johnson, J. Brioude, W. Angevine, M. Trainer, D. D. Parrish, T. R. Ryerson, I. Pollack, P. D. Cullis, M. A. Ives, D. W. Tarasick, J. Al-Saadi, and I. Stajner (2011), Measurement of western U.S. baseline ozone from the surface to the tropopause and assessment of downwind impact regions, *J. Geophys. Res.*, *116*, D00V03, doi:10.1029/2011JD016095.
- Dentener F., T. Keating and H. Akimoto (eds.) Hemispheric Transport of Air Pollution 2010, Part A: ozone and Particulate Matter, Air Pollution Studies No. 17, United Nations, New York and Geneva, ISSN 1014-4625, ISBN 978-92-1-117043-6.
- Emery et al. (2012), Regional and global modeling estimates of policy relevant background ozone over the United States, *Atmos. Environ.*, 47, 206-217.
- European Commission, Joint Research Centre (JRC)/Netherlands Environmental Agency (PBL). Emission Database for Global Atmospheric Research (EDGAR), release version 4.1. http://edgar.jrc.ec.europa.eu, 2010.
- Fischer, E. V., K. D. Perry, and D. A. Jaffe (2011a), Optical and chemical properties of aerosols transported to Mount Bachelor during spring 2010, J. Geophys. Res., 116, D18202, doi:10.1029/2011JD015932.
- Fischer, E. V., D. A. Jaffe, and E. C. Weatherhead (2011b), Free tropospheric peroxyacetyl nitrate (PAN) and ozone at Mount Bachelor: potential causes of variability and timescale for trend detection, *Atmos. Chem. Phys.*, *11*, 5641-5654.
- Huang, M., G. R. Carmichael, B. Adhikary, S. N. Spak, S. Kulkarni, Y. F. Cheng, C. Wei, Y. Tang, D. D. Parrish, S. J. Oltmans, A. D'Allura, A. Kaduwela, C. Cai, A. J. Weinheimer, M. Wong, R. B. Pierce, J. A. Al-Saadi, D. G. Streets, and Q. Zhang (2010), Impacts of transported background ozone on California air quality during the ARCTAS-CARB period a multi-scale modeling study, *Atmos. Chem. Phys.*, *10*, 6947-6968.
- Hudman, R. C., D. J. Jacob, O.R. Cooper, M. J. Evans, C. L. Heald, R. J. Park, F. Fehsenfeld, F. Flocke, J. Holloway, G. Hübler, K. Kita, M. Koike, Y. Kondo, A. Neuman, J. Nowak, S. Oltmans, D. Parrish, J. M. Roberts, and T. Ryerson, Ozone production in transpacific Asian pollution plumes and implications for ozone air quality in California, *J. Geophys. Res.*, 109, D23S10, doi:10.1029/2004JD004974, 2004.
- Lam, Y. F. and J. S. Fu (2010), Corrigendum to "A novel downscaling technique for the linkage of global and regional air quality modeling", *Atmos. Chem. Phys.*, 10, 4013-4031.
- Langford, A. O., J. Brioude, O.R. Cooper, C.J Senff, R.J. Alvarez II, R.M. Hardesty, B.J. Johnson, and S.J. Oltmans (2012), Stratospheric influence on surface ozone in the Los Angeles area during late spring and early summer of 2010, *J. Geophys. Res.*, 117, D00V06, doi:10.1029/2011JD016766.

- Lin, M., Holloway, T., Carmichael, G. R., and Fiore, A. M.: Quantifying pollution inflow and outflow over East Asia in spring with regional and global models, *Atmos. Chem. Phys.*, 10, 4221-4239, doi:10.5194/acp-10-4221-2010, 2010.
- Lin, M., A. M. Fiore, L. W. Horowitz, O. R. Cooper, V. Naik, J. Holloway, B. J. Johnson, A. Middlebrook, S. J. Oltmans, I. B. Pollack, T. B. Ryerson, J. X. Warner, C. Wiedinmyer, J. Wilson, B. Wyman (2012), Transport of Asian ozone pollution into surface air over the western United States in spring, J. Geophys. Res., in-press.
- Wang, Y., Y. Choi, T. Zeng, B. Ridley, N. Blake, D. Blake, and F. Flocke (2006), Late-spring increase of trans-Pacific pollution transport in the upper troposphere, *Geophys. Res. Lett.*, 33, L01811, doi:10.1029/2005GL024975.
- Zhang, L., D. J. Jacob, N. V. Downey, D. A. Wood, D. B., C. C. Carouge, A. van Donkelaar, D.B.A. Jones, L. T. Murray, Y. Wang (2011), Improved estimate of the policy-relevant background ozone in the United States using the GEOS-Chem global model with ½ x 2/3 horizontal resolution over North America, *Atmos. Environ.*, 45, 6769-6776.



April-May 2009-2011 average SCIAMACHY tropospheric NO $_2$ , 10<sup>13</sup> molecules NO $_2$  cm<sup>-2</sup>

Figure 1. Tropospheric NO<sub>2</sub> column retrievals from the SCIAMACHY sensor, averaged over April-May 2009-2011. Gridded data used in this plot were freely downloaded from: www.temis.nl

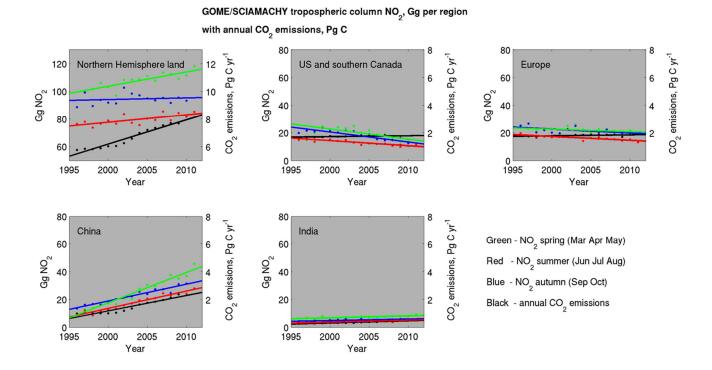
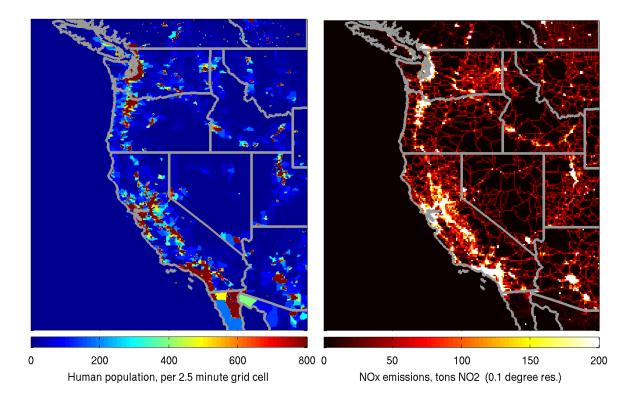


Figure 2. Trends in tropospheric column NO<sub>2</sub>, as detected by GOME/SCIAMACHY (gridded data used in these plots were freely downloaded from <u>www.temis.nl</u>), and anthropogenic CO<sub>2</sub> emissions (U.S. Energy Information Agency).



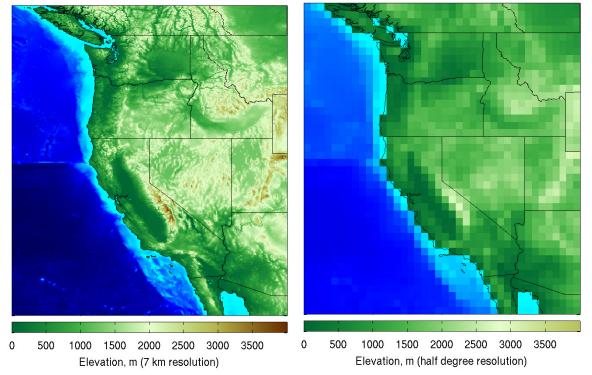
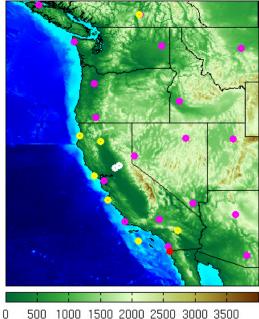


Figure 3. The human population of mid-latitude western North America in 2010, and the land-based anthropogenic  $NO_x$  emissions in 2005 (EDGAR v4.1). Also shown is the corresponding topography at 7 km resolution, and half degree resolution.



) 500 1000 1500 2000 2500 3000 3500 Elevation, m (7 km resolution)

Figure 4. Locations of the NWS daily rawinsondes (magenta circles), the May-June 2010 daily ozonesondes (yellow), and the additional May-June 2010 daily rawinsondes launched from the CARES sites (white) and the CalMex site (red). NOAA radar wind profilers (not shown) were also situated in the LA Basin (5 profilers), the Central Valley (5 profilers) and on the coast north of San Francisco (1).

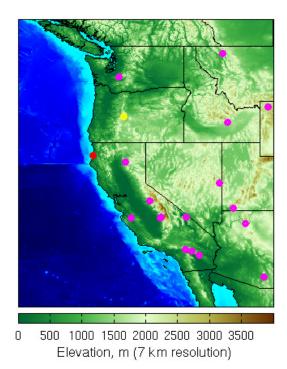


Figure 5. Locations of the CASTNET and National Park Service rural ozone monitors (magenta circles) and the University of Washington Mt. Bachelor (yellow) and the NOAA Trinidad Head (red) monitors.



Figure 6. Map showing the locations of the IMPROVE particulate matter monitoring network. IMPROVE data are available for download from http://views.cira.colostate.edu/fed/.

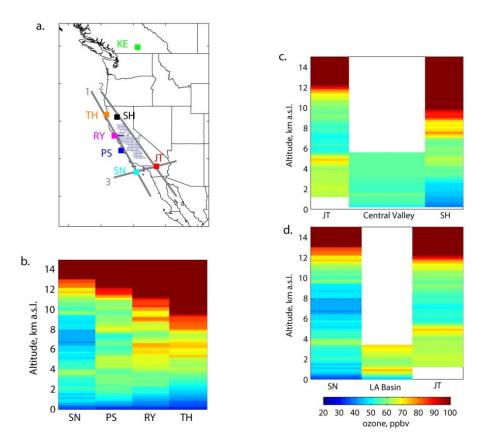


Figure 7. a.) Locations of the daily ozonesonde sites in May-June 2010, and the regions of the Central Valley and the LA Basin sampled by the NOAA P3 aircraft (blue shading). The numbered transects correspond to the ozone curtains in b. (Transect 1), c. (Transect 2) and d. (Transect 3). The O<sub>3</sub> curtains are median profiles using all available ozonesonde measurements at each site, or all available NOAA P3 measurements above the Central Valley and the LA Basin. Figures are from *Cooper et al.*, 2011.

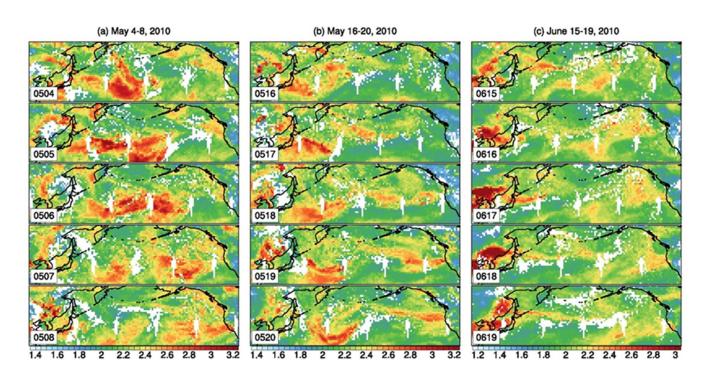


Figure 8. Major trans-Pacific Asian pollution events in May-June 2010 as seen from AIRs retrievals of CO total columns (1018 molecules cm-2; level 3 daily 1 degree gridded products). Reprint from *Lin et al.*, 2012.