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**Spring 2006 INTEX-B
Campaign**

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Chemistry and transport of pollution over the Gulf of Mexico and the Pacific: Spring 2006 INTEX-B Campaign overview and first results

H. B. Singh¹, W. H. Brune², J. H. Crawford³, F. Flocke⁴, and D. J. Jacob⁵

¹NASA Ames Research Center, Moffett Field, CA 94035, USA

²Pennsylvania State University, PA 16902, USA

³NASA Langley Research Center, Hampton, VA 23665, USA

⁴National Center for Atmospheric Research, Boulder, CO 80301, USA

⁵Harvard University, MA 02138, USA

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Correspondence to: H. B. Singh (hanwant.b.singh@nasa.gov)

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Abstract

Intercontinental Chemical Transport Experiment-B (INTEX-B) was a major NASA¹ led multi-partner atmospheric field campaign completed in the spring of 2006 (<http://cloud1.arc.nasa.gov/intex-b/>). Its major objectives aimed at (i) investigating the extent and persistence of the outflow of pollution from Mexico; (ii) understanding transport and evolution of Asian pollution and implications for air quality and climate across western North America; and (iii) validating space-borne observations of tropospheric composition. INTEX-B was performed in two phases. In its first phase (1–21 March), INTEX-B operated as part of the MILAGRO campaign with a focus on observations over Mexico and the Gulf of Mexico. In the second phase (17 April–15 May), the main INTEX-B focus was on the trans-Pacific Asian pollution transport. Multiple airborne platforms carrying state of the art chemistry and radiation payloads were flown in concert with satellites and ground stations during the two phases of INTEX-B. Validation of Aura satellite instruments (TES, OMI, MLS, HIRDLS) was a key objective within INTEX-B. Satellite products along with meteorological and 3-D chemical transport model forecasts were integrated into the flight planning process to allow targeted sampling of air parcels. Inter-comparisons were performed among and between aircraft payloads to quantify the accuracy of data and to create a unified data set. Pollution plumes were sampled over the Gulf of Mexico and the Pacific several days after downwind transport from source regions. Signatures of Asian pollution were routinely detected by INTEX-B aircraft, providing a comprehensive data set on gas and aerosol composition to test models and evaluate pathways of pollution transport and their impact on air quality and climate. This overview provides details about campaign implementation and a context within which the present and future INTEX-B/MILAGRO publications can be understood.

¹Acronyms are provided in Appendix A.

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1 Introduction

Air pollution is one of the most important environmental challenges of this century. This challenge is particularly acute in the developing world where the rapid growth of megacities (>10 million population) is producing atmospheric pollution of unprecedented severity and extent (Molina and Molina, 2004; Chan and Yao, 2008). In recent decades, a mounting body of atmospheric data have demonstrated that gas and aerosol emissions from major urban and industrial centers can influence both air quality and climate on scales ranging from regional to continental and global (Holloway et al., 2003; Stohl, 2004; Oltmans et al., 2006; EPA, 2007 and references there in). During the last decade, the NASA Tropospheric Chemistry Program has organized several field campaigns to investigate the outflow of pollution from Asia and North America with the goal of understanding the transport and transformation of gases and aerosols on transcontinental/intercontinental scales and their impact on air quality and climate (Hoell et al., 1996, 1997; Raper et al., 2001; Jacob et al., 2003; Singh et al., 2006). INTEX-NA was a major NASA led integrated field experiment that focused on the inflow and outflow of pollution over the North American continent. The first phase of INTEX-NA (INTEX-A) was completed in the summer of 2004 and results widely disseminated (Singh et al., 2006 and references there in). The second phase (INTEX-B) was completed in the spring of 2006 and forms the basis of the MILAGRO/INTEX-B Special Issue in ACP.

INTEX-B/MILAGRO was an integrated mission performed principally over Mexico, the Gulf of Mexico, and the northern Pacific Ocean during the spring of 2006 in cooperation with multiple national and international partners (Molina et al., 2008). Major objectives for INTEX-B were aimed at (i) investigating the extent and persistence of the outflow of pollution from Mexico; (ii) understanding the transport and evolution of Asian pollution and implications for air quality and climate across western North America; (iii) relating atmospheric composition to sources and sinks; (iv) characterizing the effects of aerosols on solar radiation; and (v) validating space-borne observations of tropo-

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spheric composition. This effort had a broad scope to investigate the chemistry and transport of long-lived greenhouse gases, oxidants and their precursors, aerosols and their precursors, as well their relationship with radiation and climate.

2 Mission design and implementation

2.1 Measurement platforms

The principal mobile platforms that participated in the INTEX-B/MILAGRO campaign, their bases of operation, campaign duration, and specifications are summarized in Table 1. These airborne platforms had the capability to sample much of the troposphere over long distances. Complementing these airborne platforms were several ground stations that operated during this study. An ozonesonde-radiosonde network (IONS-06) operated over 16 North American sites. Satellite observations both guided flight planning and helped to extend the range of airborne observations. Description of Mexico based platforms and ground stations focused exclusively on the MILAGRO part of the campaign are being summarized in an overview paper by L. Molina and S. Madronich (private communication). Additional details about these platforms are available from <http://cloud1.arc.nasa.gov/intex-b/> and <http://www.eol.ucar.edu/projects/milagro>.

2.2 Instrument payload and measurement capability

The payloads of a majority of airborne platforms in Table 1 were selected to measure major greenhouse gases, ozone and precursors, aerosols and precursors, and a large number of tracers. Table 2a shows the DC-8 payload of in situ and remote sensors for measurements of both gases and aerosols. A nadir and zenith viewing UV lidar measured tropospheric O₃ and aerosols remotely. A second zenith viewing lidar was tuned to stratospheric ozone and temperature measurements for HIRDLS validation. Spectral radiometers allowed direct measurement of actinic flux to derive key photolysis

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frequencies. Multiple instruments measured physical, chemical, and optical properties of fine and coarse aerosol. The DC-8 flew 145 hs from 1 March–15 May 2006 in 18 science/transit flights of 8–10 h duration from its three bases of operation (Houston, Hawaii, Anchorage). Six DC-8 sorties were undertaken from Houston targeted on Mexico and the Gulf of Mexico with three sorties from Hawaii and four from Anchorage targeted on Asian pollution over the Pacific. The C-130 payload was similar to the in situ payload of the DC-8 and is described in more detail by L. Molina and S. Madronich (private communication). The C-130 performed 13 flights over Mexico and the Gulf and undertook 11 sorties from Seattle that sampled pollution over western North America and the eastern Pacific.

An instrumented Environment Canada Cessna 207 performed some 33 short flights (approx 2-h duration) from 22 April to 17 May 2006 from a base in western Canada in close coordination with the C-130, and with measurements at the Whistler Peak Station (50.1° N, 122.9° W, 2182 m) that has provided continuous measurements of meteorological parameters, CO, and O₃ since its establishment in 2002 (Leaitch et al., 2008). The Cessna 207 measured CO, O₃, aerosol size, number density, and composition (Table 2b). A Raman lidar was operated during 11–17 May from the base of Whistler Mountain to provide profiles of aerosol backscatter in clear skies. University of Washington Beechcraft Duchess 76 performed eight flights between 12 April and 15 May 2006 out of Paine Field in Everett Washington. The somewhat limited payload (Table 2c) included measurements of CO, O₃, Hg, and aerosols. Complementing the airborne observations were more extensive measurements of speciated Hg, PANs, CO, O₃, NO_x, and Radon made at the Mt. Bachelor Observatory (43.98° N 121.69° W; 2.7 km a.s.l.). The DLR Falcon-20, with a payload summarized in Table 2d, operated out of Oberpfaffenhofen, Germany and performed some 15 flights from 24 March to 3 May 2006. The objectives of the DLR Falcon flights during INTEX-B included the chemical and aerosol characterization of pollution plumes over the eastern Atlantic and Europe originating from Asia and North America. The INTEX-B Ozonesonde Network Study (IONS-06) regularly launched ozonesondes from over 16 North American

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sites (Table 3). These launches were coordinated with the daytime Aura overpass (Thompson et al., 2008).

2.3 Flight planning and execution

Satellite observations, along with meteorological and chemical forecasts from global and regional chemical transport models (CTMs), were extensively used to plan, coordinate, and implement the INTEX-B field mission. Table 4 lists the major CTMs in use for planning this experiment. Meteorological products were derived from the NCEP reanalysis data. Daily meteorological and chemical forecasts guided the design and execution of the mission. Typical model products included source-tagged CO tracers, aerosol tracers, O₃, and PAN distributions. Meteorological products included satellite imagery (visible, infrared, and water channels), precipitation and cloud fields, backward and forward trajectories, and multi-day synoptic weather forecasts. Principal satellite instruments that provided near real-time data to guide flight planning were AIRS and MOPITT (CO columns), MODIS (aerosol, fire counts), and OMI (total column O₃, tropospheric NO₂, absorbing aerosols). These data were combined with forward trajectories for flight planning purposes.

The NSF/NCAR C-130 coordinated activities with the DC-8 with bases in Mexico and Seattle. This coordination was necessary to accomplish inter-comparisons among instruments, to achieve several objectives that were unattainable by a single platform, and to collect complementary data where appropriate. This also permitted quasi-Lagrangian investigations when pollution plumes sampled over Mexico and the Pacific were sampled again after several days of transport.

2.4 Flight tracks and sampled air masses

Figures 1 and 2a–e show flight tracks of several major airborne platforms (DC-8, C-130, Cessna-207, Beechcraft 76, Falcon-20) and their spatial coverage. Table 5 provides a brief summary of the flight objectives of each of the DC-8 flights with more

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details available at <http://cloud1.arc.nasa.gov/intex-b/>. Air masses influenced by anthropogenic pollution, forest fires, and stratosphere were sampled during INTEX-B. It was possible to fly extremely low (100 m a.g.l.) for source characterization and spiral up to 12 km for purposes of satellite validation. Many of the flight planning procedures used in INTEX-B were similar to INTEX-A and have been previously described (Singh et al., 2006).

2.5 Inter-comparisons

Intra- and inter-platform comparisons of measurements were extensively performed to establish accuracies and precisions. On the DC-8 itself, several chemical measurements were duplicated using a variety of different techniques. Salient among these were measurements of O₃, H₂O, HCHO, and jNO₂. Inter-comparisons were also performed with other aircraft by flying in formation with a separation of less than 300 m in the horizontal and 100 m in the vertical. Two to three levels were chosen between 0 and 6 km altitudes, and the entire inter-comparison including descent and ascent typically lasted one hour. Table 6 provides a summary of the inter-comparisons carried out by the airborne platforms, which included species such as O₃, OH, HO₂, CO, SO₂, PAN, NO₂, HNO₃, VOC, and aerosol composition and properties. Comparisons were conducted in two phases: a blind phase occurring in the field shortly after completion of the flight, and a final phase after data was fully calibrated and approved by the investigators. The actual inter-comparison data and results are available at <http://www-air.larc.nasa.gov/cgi-bin/ic>. While the agreement among instruments on various platforms was generally good for molecules such as O₃, NO₂, CO, H₂O, CO₂, HCHO, and aerosol physical properties, there were large differences in measurements of HO_x, OVOC, HNO₃, PAN, NO, and H₂O₂ that suggest a need for concerted efforts to reduce these errors. Specific details on the inter-comparison results will be published in a forthcoming paper to the MILAGRO/INTEX-B Special Issue by Chen et al. (private communication, 2008).

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2.6 Satellite validation

Satellite sensors measure radiances from which information on atmospheric composition is retrieved using radiative transfer algorithms with substantial and often poorly characterized uncertainties. Validation of satellite derived tropospheric composition and optical properties were a primary goal and received a great deal of attention from the NASA DC-8, J-31, and to a lesser extent, the C-130 platforms. The principal focus of INTEX-B was on the Aura satellite (Schoeberl et al., 2006), but validation studies for SCIAMACHY/Envisat, MODIS-Terra/Aqua, and MISR/Terra were also carried out (Table 7a,b). The DC-8 focused on several trace gases (O_3 , CO , NO_2 , HNO_3 , and $HCHO$) and together with the J-31 on aerosols and clouds. Opportunities to under-fly satellites for integrated science and validation were examined and planned for each flight. Validations were performed to test retrievals with a variety of underlying surfaces. A typical DC-8 validation flight for nadir viewing instruments (e.g. OMI, TES) involved a profile from 0.2 to 11.5 km, spiraling with a 20 km radius within ± 30 min of the satellite overpass time under relatively cloud free conditions. Limb viewing instruments (e.g. MLS) frequently required sampling along the satellite track utilizing both in situ and remote sensing capabilities on the DC-8. Two flights were dedicated to validation of HIRDLS O_3 and temperature in the stratosphere using the AROTAL lidar, which required night flights for observations to reach maximum altitudes. In all cases, the DC-8 provided important chemical and physical data that could be used to improve satellite retrievals. While satellite under-flights were usually planned to focus on a specific satellite instrument, the proximity of A-Train observations and wide swaths for some instruments allowed these flights to routinely benefit multiple satellite sensors.

3 Overview of first results

We provide here a brief overview of the INTEX-B results presented in the first collection of papers assembled in this Special Issue of Atmospheric Chemistry and Physics

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and published elsewhere to date. We focus principally on the Pacific component assessing the influence of transpacific transport. Additional information on the INTEX-B/MILAGRO mission is available from Molina et al. (2008) and a complementary overview paper that focuses on activities from platforms based within Mexico is in preparation (L. Molina and S. Madronich, private communication, 2008). In the following sections we briefly discuss results based on (1) integrated analysis of data using observations and models; (2) chemistry and transport model evaluation; and (3) satellite validation results and analysis from satellite observations. The reader is referred to the specific papers for more information. We also note that the data continue to be further analyzed and new results are anticipated.

3.1 Pollution transport and chemistry: integrated analysis of data

Many studies being published in the ACP Special Issue analyzed observed data from aircraft, ground stations, and satellites with the help of models. Here we briefly discuss INTEX-B papers that have been submitted for publication. At this point a majority of the studies are focused on the Pacific component of INTEX-B.

3.1.1 Airborne observations and trans-pacific transport

There was extensive evidence for pollution transport from Asia to North America and all the way to Europe. Concurrent satellite CO and ozone observations (TES, AIRS, MOPITT) showed direct evidence for trans-Pacific pollution transport (Molina et al., 2008; Zhang et al., 2008). Fiedler et al. (private communication, 2008) intercepted pollution plumes transported all the way from Asia to the eastern Atlantic and investigated the SO₂ induced aerosol formation in these plumes. Zhang et al. (2008) and Walker et al. (2008) interpret INTEX-B and satellite observations to investigate the transport and evolution of pollution over the Pacific and its impact on surface air quality over western North America. Satellite derived tropospheric columns of NO₂ indicate an increase of some 8–10%/year from 2000–2006. The semi-permanent Pacific High and Aleutian

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Low cause splitting of trans-Pacific pollution plumes over the northeast Pacific. Both aircraft measurements and model results showed sustained ozone production driven by PAN decomposition in the southern branch, adding to ozone produced in the Asian continental boundary layer. Model simulation of ozone observations suggest that Asian pollution in spring 2006 enhanced surface O₃ concentrations by 5–7 ppb over western North America. Modeled sensitivities of Pacific tropospheric O₃ columns to Asian anthropogenic emissions and lightning are >3.7 DU and >2.1 DU respectively.

INTEX-B data over the Pacific clearly showed that a dominant fraction of reactive nitrogen was in the form of transported PAN. Comparison with previous observations suggests that the relative fraction of PAN has increased over time while that of HNO₃ has decreased (Molina et al., 2008). It is argued that these trends are largely driven by changes in surface emissions of VOC, NO_x and aerosol over Asia. Wolfe et al. (2008) utilize observations of PANs at the MBO to conclude that trans-Pacific transport of Asian pollution leads to substantial increases in PANs and ozone at this site. The ensemble of trajectories indicates that Asian-influenced free tropospheric air contained a median PAN mixing ratio double that of the full dataset. Walker et al. (2008) and Zhang et al. (2008) use observations and 3-D models to elucidate the important role PAN plays in O₃ formation resulting from its ability to redistribute NO_x. Measurements of CO at MBO and meteorological indices show that long-range transport of CO from the heavily industrialized region of East Asia was significantly lower in 2006 compared to 2005 (Reidmiller et al., 2008). In the western US both ground site data and MOPITT and TES satellite observations reveal a significant decrease (from 2–21%) in spring-time maximum CO between 2005 and 2006. They attribute this decrease to moderate biomass burning in Southeast Asia during 2006 and a transport pattern that limited the inflow of Asian pollution to the lower free troposphere over western North America.

Analysis of aircraft sulfate measurements from the DC-8 over the central Pacific, C-130 over the east Pacific, and the Cessna over British Columbia indicated that most Asian aerosol over the ocean was in the lower free troposphere (2–5 km). Large amounts of sulfate and little organic carbon were found in imported Asian pollution

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over western Canada (Fig. 3). Asian plumes were not only significantly reduced of fine particle organic material, but organic compounds were attached to coarse particles in significant quantities. Scavenging of organic aerosol precursors by dust near source regions is suggested, and any formation of secondary organic aerosol during transport from Asian source regions across the Pacific was principally associated with the coarse particles (Leaitch et al., 2008; McKendry et al., 2008). In agreement with Zhang et al. (2008), an average of profiles indicated that trans-Pacific transport between 2 and 5 km during this period increased ozone by about 10 ppb and fine particle sulfate by $0.2\text{--}0.5\ \mu\text{g m}^{-3}$. Campaign-average simulations showed that anthropogenic East Asian sulfur emissions increased mean springtime sulfate in western Canada at the surface by 25–30% and accounted for 40% of the overall regional sulfate burden between 1 and 5 km (van Donkelaar et al., 2008). McNaughton et al. (2008) analyze data from the eastern Pacific and deduce that the presence of CaCO_3 in Asian dust enhanced heterogeneous conversion of gas phase S and N into aerosol phase.

Fine particle median sulfate and water-soluble organic carbon (WSOC) concentrations in transported Asian air masses (2–5 km) were two to four times lower than North American air masses. In contrast, in air masses below 2 km median WSOC-sulfate ratios were consistently between one and two (Peltier et al., 2008). Sun et al. (2008) use measured aerosol mass spectra at Whistler peak to conclude that sulfate and OA were mostly present in external mixtures, indicating different origins, and OA was highly oxygenated, with an average organic-mass-to-organic-carbon ratio of 2.0 and an atomic ratio of oxygen-to-carbon of 0.8. The nominal formula for OA was $\text{C}_{1.52}\text{H}_{1.52}\text{N}_{0.03}\text{O}_{0.82}$ for the entire study. Heald et al. (2008) synthesize measurements of organic carbon compounds in both the gas and particle phases made upwind, over, and downwind of North America. The daytime mean total organic carbon ranges from 4 to $456\ \mu\text{g C m}^{-3}$ with organic aerosol making up 3–17% of this mean. Anthropogenic emissions account for much of this variability but correlation with tracers such as isoprene suggest that biogenic emissions are an important contributor. Dunlea et al. (2008) describe two case studies for pollution layers transported across the Pacific from the Asian continent, in-

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tercepted 3–4 days and 7–10 days downwind of Asia, respectively. These observations of sulfate and organic aerosol in aged Asian pollution layers are consistent with fast formation near the Asian continent, followed by washout during lofting and subsequent transformation during transport across the Pacific. No evidence is found to support significant secondary organic aerosol formation in the free troposphere.

A number of specialized measurements, new to the DC-8, were performed to define Pacific composition for the first time. Kim et al. (2008) provide specific gas phase HCl measurements from the marine boundary layer (MBL) to the lower stratosphere. They find that HCl mixing ratios are lower than previously believed and show evidence for its production in the mid troposphere by the de-chlorination of dust aerosols. Over the Pacific, mercury was weakly correlated with anthropogenic tracers (Swartzendruber et al., 2008). A prominent feature of the INTEX-B dataset was frequent total depletion of Hg⁰ in the upper troposphere when stratospherically influenced air was encountered (Talbot et al., 2007, 2008). Global 3-D chemical transport models have been used to interpret new INTEX-B observations of methanol to further constrain the atmospheric methanol budget (Millet et al., 2008).

INTEX-B provided a most detailed exploration of HO_x free radicals and its precursors. Simulation studies highlight many uncertainties in our quantitative understanding of the HO_x-NO_x-O₃ system. Mao et al. (2008) deployed a new airborne OH reactivity instrument for the first time on the NASA DC-8 during the INTEX-B campaign. From the median vertical profile obtained over the Pacific, the measured OH reactivity was 30% higher than the OH reactivity from steady state assumptions and some 200% higher than the OH reactivity calculated from the total measurements of all OH reactants. They attribute the missing OH reactivity to some highly reactive unmeasured VOCs that have HCHO as an oxidation product.

3.2 Model simulations and INTEX-B observations

INTEX-B provided a vast amount of observational data to both test CTMs and use them for further analysis. In general, the models in Table 4 captured the presence of pollution

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plumes and were an excellent aid in flight planning. However, significant quantitative differences remain. Figure 4a shows a comparison of selected observed constituents (OH, CO, and O₃) and results from post mission simulations over the Gulf of Mexico (Region 1) and Pacific (Region 4). As was previously observed in INTEX-A (Singh et al., 2007), model disagreements far exceed any potential error in observations and are also not consistent between missions (INTEX-A and INTEX-B). In case of GEOS-Chem, overestimation of OH appears to be a feature that is propagated throughout the INTEX-B Gulf and Pacific domain, and is principally responsible for CO underestimation. The “box model”, which is most constrained by observations, is consistent with measurements above ~4 km but significantly over predicts OH in the lower troposphere. Disagreements with observations are often much larger for more complex organic molecules such as acetaldehyde (Fig. 4b). Comparisons with CTMs of data from the entire campaign reveal an under-prediction of organic aerosol mass in the MOZART model, but much smaller discrepancies with the GEOS-Chem model than found in previous studies over the western Pacific (Dunlea et al., 2008). Model disagreements can be due to a large number of factors that include emissions, boundary conditions, chemistry, meteorology, and even computational errors that need to be fully investigated. Plans are presently underway to create a community model inter-comparison and test project within the IGAC umbrella.

3.3 Satellite derived results and analysis

3.3.1 Validation results

Providing coincident observations for satellite validation of Aura instruments was a central theme for the DC-8 in both phases of INTEX-B. A limited number of validation comparisons are summarized in Fig. 5 and more details referenced. Figure 5a,b show a correlation plot of MLS retrieved CO and O₃ in the UT (215 hPa) with the DC-8 data indicating large differences in CO and relatively good agreement with O₃ (Livesey et al., 2008). Figure 5c compares tropospheric NO₂ column measurements from OMI

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with airborne observations over the southern United States, Mexico, and the Gulf of Mexico during the INTEX-B campaign. Reasonably good correlation with no significant bias ($R^2=0.67$, slope=0.99) is found for the ensemble of comparisons when the aircraft could spiral sufficiently low to sample most of the NO_2 column (Boersma et al., 2008; Bucsela et al., 2008). TES CO data show good ($\pm 20\%$) overall agreement in the troposphere with the DC-8 in situ measurements (Fig. 5d) with some biases in the UT and LT (Luo et al., 2007). Comparisons of the TES O_3 with the DC-8 DIAL (Fig. 5e) and ozonesonde data show a somewhat positive bias for TES (Richards et al., 2008; Nassar et al., 2008). Santee et al. (2007) provide an in depth comparison of limb derived MLS HNO_3 data with that of INTEX-B and other observations.

Although INTEX-B primarily focused on the troposphere, one of the DC-8 remote sensing instruments (AROTAL) was targeted for stratospheric observations. Figure 5f shows a comparison of the DC-8 AROTAL and HIRDLS retrieved O_3 with good agreement above 20 km and poor agreement in the lower stratosphere. Nardi et al. (2008) provide details on these comparisons and the reasons for the disagreements. Froidevaux et al. (2008) investigate MLS O_3 retrievals in the stratosphere and compare them with the INTEX-B AROTAL data.

Extensive validation of CO from AIRS and MOPITT was also carried out. Emmons et al. (2008) use INTEX-B/MILAGRO and other in situ observations to conclude that a significant positive bias, around 20%, is present in MOPITT derived CO. Comparisons to the long-term records revealed that this bias is increasing with time. Possible causes for this drift and improvements in future retrievals have been suggested. Felker et al. (2008) blend GOES water vapor and TES ozone to derive a Multi-sensor UT Ozone Product (MUTOP) that has the advantage of producing the spatial coverage of a geostationary image while retaining TES's ability to resolve UT ozone. Results show 70–80% of TES-observed UT ozone variability can be explained by correlation with these two dynamical tracers. It is noted that satellite retrievals are continually being refined and it is possible that the comparisons shown here will improve over time.

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3.3.2 Satellite based analysis

OMI NO₂ satellite observations were used to constrain Asian anthropogenic NO_x emissions, indicating a factor of two increases in emissions from China from 2000 to 2006 (Walker et al., 2008; Zhang et al., 2008). Halland et al. (2008) quantify the vertical transport of CO by deep mesoscale convective systems and assess the ability of TES to detect the resulting enhanced CO in the upper atmosphere. A squall line similar to one in INTEX-B was simulated using the Goddard Cumulus Ensemble model. Results show that the simulated squall line transports the greatest mass of CO in the UT. They conclude that TES has sufficient sensitivity to resolve convectively lofted CO, as long as the retrieval scene is cloud-free. Arellano et al. (2007) present a global chemical data assimilation system and apply it to constrain global tropospheric CO by assimilating meteorological observations of temperature and horizontal wind velocity and satellite CO retrieved from MOPITT. They conclude that MOPITT data assimilation provides significant improvements in capturing the observed CO variability. Shim et al. (2008) use concurrent tropospheric O₃ and CO vertical profiles from TES over Mexico City to characterize mega-city pollution outflow on a regional scale. They find that TES captures much of the spatial and day-to-day variability seen in the in situ data for O₃, but not for CO and cautions against using such correlations. Shinozuka et al. (2008) report that number concentration of particles >100 nm are a proxy for CCN and can be stratified by two optical properties, the extinction coefficient and its wavelength dependence. They use data from INTEX-B/MILAGRO and INTEX-A to quantify these relationships and explore the feasibility of satellite retrieval of CCN concentration from optical property measurements.

It is increasingly possible to retrieve many organic and inorganic species from existing satellite obtained spectra (e.g. acetone, methanol, HCHO, PAN, CO₂, CH₄) (Beer et al., 2001; Allen et al., 2004; Frankenberg et al., 2005; Glatthor et al., 2007; Rinsland et al., 2007). In Table 8, we provide UT mixing ratios and tropospheric columns of key atmospheric constituents that are potentially retrievable from existing satellite

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data from the four selected regions of INTEX-B (Fig. 2a). The variability in the CO₂ column and UT mixing ratios is 2.7% (≈10 ppm) and 1.5% (≈6 ppm), respectively, irrespective of the nature of underlying sources. Filtering the data for only clean conditions (bottom two CO quartiles) had little effect (<5%) on the relative variability. Boundary layer (1–2 km) CO₂ elevation due to urban pollution and reductions due to biosphere uptake would need to be 40 ppm or more to be detected in the CO₂ column. Airborne observations indicate that these elevations are typically less than 15 ppm and only on rare occasions may exceed 40 ppm (Choi et al., 2008). Relating satellite derived atmospheric columns of CO₂ (Orbiting Carbon Observatory; <http://oco.jpl.nasa.gov/>; Miller et al., 2007) to its surface sources and sinks would be extremely challenging.

4 Conclusions

The INTEX-B/MILAGRO measurements provide a rich data set for describing the composition and chemistry of the troposphere over Mexico, the Gulf of Mexico, the Pacific, and western North America, for investigating the transformation of gases and aerosols during long-range transport, for the radiation balance of the troposphere, and for validating a variety of satellite observations as well as models of chemistry and transport. The results presented in this special issue of ACP represent only the initial analysis, and much more integrated analysis is expected in the near future. The data are available to all interested parties for further analysis.

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Appendix A

List of acronyms

ACP: Atmospheric Chemistry and Physics

5 AIRS: Atmospheric Infrared Sounder

AOD: Aerosol Optical Depth

AROTAL: Airborne Raman Ozone Temperature and Aerosol Lidar

CIMS: Chemical Ionization Mass Spectrometry

CTM: Chemical Transport Model

10 DLR: Deutsches Zentrum für Luft- und Raumfahrt

DU: Dobson Unit (2.7×10^{16} ozone molecules per cm^2)

GOES: Geostationary Operational Environmental Satellite

HIRDLS: High Resolution Dynamic Limb Sounder

IGAC: International Global Atmospheric Chemistry

15 INTEX-NA: Intercontinental Chemical Transport Experiment-North America

IONS: INTEX Ozoneprobe Network Study

LIF: Laser Induced Fluorescence

LIS: Lightning Imaging Sensor

LRLN: Long-Range Lightning Network

20 LS: Lower Stratosphere

MBO: Mount Bachelor Observatory

MILAGRO: Megacity Initiative: Local Global and Research Observations

MISR: Multiangle Imaging SpectroRadiometer

MLS: Microwave Limb Sounder

25 MOPITT: Measurement of Pollution in the Troposphere

MOZART: Model of Ozone and Related Tracers

MODIS: Moderate Resolution Imaging Spectroradiometer

NA: North America

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NASA: National Aeronautics and Space Administration

NCEP: National Centers for Environmental Prediction

NLDN: National Lightning Detection Network

OA: Organic Aerosol

5 OC: Organic Carbon

OMI: Ozone Monitoring Instrument

OVOC: Oxygenated Volatile Organic Chemicals

PANs: Peroxy Acetyl Nitrates

RAQMS: Real-time Air Quality Modeling System

10 SCIAMACHY: Scanning Imaging Absorption Spectrometer

for Atmospheric Chartography

SSFR: Solar Spectral Flux Radiometer

STEM: Sulfur Transport and Deposition Model

TES: Tropospheric Emission Spectrometer

15 TOMS: Total Ozone Measuring Spectrometer

UT: Upper Troposphere

WSOC: Water Soluble Organic Carbon

20 *Acknowledgement.* We thank all INTEX-B/MILAGRO participants and sponsoring agencies for making this project possible. DC-8 and C-130 activities were supported by the NASA Tropospheric Chemistry and NSF Atmospheric Chemistry Programs. We very much appreciate the dedicated efforts of all personnel associated with INTEX-B activities from NASA Dryden, NASA Wallops, Univ. of North Dakota in make this campaign a success.

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Table 1. Principal Mobile Platforms in Operation During the Spring 2006 MILAGRO/INTEX-B Study

Platform type ^a	Study name	Platforms (Country)	Dates of operation (2006)	Bases of Operation ^b	Nominal Specifications, km	
					Altitude	Range
Airborne	INTEX-B/MILAGRO	NASA DC-8 ^c (United States)	1 Mar to 15 May	HS, HI, AN	0.2–12.5	9000
Airborne	INTEX-B/MILAGRO	NASA/Sky Research J-31 ^d (United States)	1–21 Mar	VC	0.1–7.5	1500
Airborne	INTEX-B	DLR Falcon-20 ^e (Germany)	1–30 Apr	OB	0.1–12.8	3500
Airborne	INTEX-B	Cessna 207 ^f (Canada)	22 Apr to 17 May	PN	0.1–5.5	800
Airborne	INTEX-B	UW Beechcraft Duchess ^e (United States)	1–15 May	ST	0.1–6.0	1400
Airborne	Mirage-MEX/INTEX-B	NSF/NCAR C-130 ^e (United States)	1 Mar to 15 May	VC, ST	0.1–7.5	6000
Airborne	MAX-MEX/MILAGRO	DOE G-1 ^e (United States)	1–29 Mar	VC	0.3–7.5	2000
Airborne	MAX-MEX/MILAGRO	NASA B-200 ^g (United States)	1–29 Mar	VC	0.1–8.5	2000
Spaceborne ^f	Terra-NASA	Terra (MOPITT, MISR, MODIS) (United States)	continuous	LEO	all altitudes	global coverage
Spaceborne ^f	Aqua-NASA	Aqua (AIRS, MODIS) (United States)	continuous	LEO	all altitudes	global coverage
Spaceborne ^f	Aura-NASA	Aura (OMI, TES, MLS, HIRDLS) (United States)	continuous	LEO	all altitudes	global coverage
Spaceborne ^f	Envisat-ESA	Envisat (SCIAMACHY)(European Union)	continuous	LEO	All altitudes	Global coverage

^a A number of fixed surface stations and sonde release sites also operated during this experiment. See Molina et al. [2008] for details.

^b HS, Houston, Texas; HI, Honolulu, Hawaii; AN, Anchorage, Alaska; VC, Veracruz, Mexico; OB, Oberpfaffenhofen, Germany; ST, Seattle, Washington; PN, Pemberton, British Columbia, Canada; LEO, Low Earth Orbit.

^c Extensive payload principally designed to study atmospheric composition and chemistry. DC-8 also contained two remote sensors for ozone/aerosol/temperature.

^d J-31 payload was principally designed to study aerosols, Earth surfaces, and water vapor via their effects on solar radiation.

^e B-200 payload consisted of the High Spectral Resolution Lidar sensing aerosol extinction and backscatter.

^f Principal focus was on gas and aerosol observations in the troposphere.

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Table 2a. DC-8 Science Payload During INTEX-B^a.

Parameters	Method	Principal Investigators	Detection Limit/Response (Nominal Accuracy) ^b
O ₃	NO/O ₃ chemiluminescence	M. Avery, NASA LaRC	1 ppb/1 s (±5%)
O ₃ and aerosol profile	UV DIAL lidar	E. Browell, NASA LaRC	–/60 s (±10%)
O ₃ and temperature profile	Raman lidar	T. McGee, NASA GSFC	–/60 s (±10%)
NO, NO ₂ , HCHO	LIF	D. Tan, Georgia Institute of Technology	5 ppt/60 s (±10%), 15 ppt/60 s (±10%)
NO ₂ , NO _y	LIF and thermal dissociation	R. Cohen, University of California, Berkeley	1 ppt/60 s (±10%), 10 ppt/60 s (±10%)
PANs, OVOC, nitriles	GC-ECD/PID/RGD	H. Singh, NASA ARC	1 ppt/90 s (±15%), 10 ppt/120 s (±20%)
H ₂ O ₂ , CH ₃ OOH, HCHO	derivative HPLC and fluorescence	B. Heikes, University of Rhode Island	20 ppt/150 s (±20%)
HCHO	TDL absorption spectrometry	A. Fried, NCAR	20 ppt/60 s (±10%)
OH, HO ₂ , naphthalene	LIF	W. Brune, Pennsylvania State University	0.02 ppt/15 s (±15%), 10 ppt/15 s (±15%)
SO ₂ , HNO ₄ , HCl	CIMS	G. Huey, Georgia Institute of Technology	5 ppt/15 s (±15%)
NMHC, halocarbons, alkyl nitrates	whole air samples; GC-FID/EC/MS	D. Blake, University of California, Irvine, and E. Atlas, University of Miami	1 ppt/100 s (±2–10%)
CO, CH ₄ , N ₂ O	TDL absorption spectrometry	G. Sachse, NASA LaRC	1 ppb/5 s (±5%)
CO ₂	Non-dispersive infrared analyzer	S. Vay, NASA LaRC	0.1 ppm/1 s (±0.1%)
H ₂ O	open path TDL absorption spectrometry	G. Diskin, NASA LaRC, and J. Podolske, NASA ARC	5 ppm/10 s (±5%)
H ₂ O, J(NO ₂)	cryogenic hygrometer, filter radiometer	J. Barrick, NASA LaRC	5 ppm/10 s (±10%), 10 ⁻⁸ s/20 s (±10%) 10 ppt/300 s (±15%)
Bulk aerosol composition, HNO ₃ TGM (Total Gaseous Mercury)	Filter analysis, mist chamber/IC cold vapor atomic fluorescence spectrometry	J. Dibb, University of New Hampshire R. Talbot, University of New Hampshire	
Aerosol number, size, composition, microphysics, optical properties, light absorption & scattering, cloud water content	particle spectrometers, volatility TDMA, CN counters, nephelometer, soot photometer, f(RH), cloud and aerosol probes	A. Clarke, University of Hawaii, and B. Anderson, NASA LaRC	
Actinic fluxes and photolytic frequencies	spectral radiometry, zenith and nadir	R. Shetter, NCAR	10 ⁻⁸ s/15 s (±5%)

^a Instruments on board the DC-8 also measured pressure, temperature, dew point, aircraft position and orientation.

^b These are nominal values. Detailed measurement specifications, which may vary with altitude, are provided in the headers along with data.

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Table 2b. Canadian Cessna 207 Science Payload during INTEX-B^a.

Parameters	Method	Principal Investigator	Detection Limit/Response ^b
Aerosol composition	quadrupole aerosol mass spectrometer (Q-AMS)	Richard Leaitch, Environment Canada (EC)	SO ₄ ⁻ and NO ₃ ⁻ : 40 ng m ⁻³ /60 s organic: 400 ng m ⁻³ /60 s
Aerosol number > 15 nm	condensation particle counter	Richard Leaitch, EC	10 cm ⁻³ /1 s
Aerosol size distribution, 120 nm to 2.5 μm	optical particle counter (PMS PCASP-X)	Richard Leaitch, EC	1 cm ⁻³ /1 s
Aerosol size distribution, 300 nm to 20 μm	optical particle counter (PMS FSSP300)	Richard Leaitch, EC	0.3 cm ⁻³ /1 s
O ₃	UV absorption	Kurt Anlauf, EC	1 ppbv/1 s
CO	UV resonance fluorescence	Kurt Anlauf, EC	5 ppbv/1 s

^a Other measurements include pressure, temperature, relative humidity, and aircraft position.^b These are nominal values. Detailed measurement specifications, which may vary with altitude, are provided in the headers along with data.

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Table 2c. UW Beechcraft Duchess 76 aircraft payload during INTEX-B (12 Apr–15 May 2006).

Measurement	Method	Detection Limit ^a	Uncertainty ^a
Sub-micron aerosol scattering (Mm^{-1})	3 λ nephelometer (TSI Inc)	0.1 Mm^{-1}	10%
CO	vacuum UV fluorescence (AeroLaser)	2 ppb	5%
O ₃	UV absorption (2B Technologies)	4 ppb	6%
Hg ⁰ (ng/m^3)	pre-concentration on Au trap w/UV fluorescence (Tekran, Inc)	0.1 ng/m^3	10–15%
Standard met and GPS sensors	temperature, RH, wind speed, wind direction		

^a These are nominal values. Detailed measurement specifications, which may vary with altitude, are provided in the headers along with data.

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**Table 2d.** DLR Falcon science payload during INTEX-B^a.

Parameter	Method/Instrument	Principle Investigator	Detection Limit/Response ^b
O ₃	UV absorption/Thermo Electron TE49	Hans Schlager, DLR	1 ppb/1 s
CO	VUV fluorescence/Aerolaser AL5001		5 ppb/1 s
CO ₂	ND IR absorption/LI-COR LI-7000		0.2 ppm/1 s
HCHO	Hantzsch synthesis/Aerolaser AL 4021		50 ppt/90 s
NO	chemiluminescence/modified ECO-SR		5 ppt/1 s
NO _y	catalytic reduction + chemiluminescence/ modified ECO-SR		20 ppt/1 s
NO ₂ photolysis rate	filter radiometer		5 × 10 ⁻⁴ /1 s
SO ₂	chemical ionization ion trap mass spectrometer		20 ppt/1 s
Aerosol property	Method/Instrument	Principle Investigator	Size range (μm) ^b
Integral number concentration	condensation/CPC	Andreas Petzold, DLR	0.004–1
Nano particle size Distribution	multi-channel CPC system electrical mobility/DMA		0.004–0.1 0.01–0.2
Particle size distribution	light scattering/PCASP-100X		0.1–3
	light scattering/OPC Grimm 1.109		0.25–2.5
Cloud element size distribution	light scattering/FSSP-300		0.4–20
Integral absorption coefficient (& black carbon mass)	filter transmission photometry/PSAP		2–100

^a Other measurements include pressure, temperature, relative humidity, and aircraft position.^b These are nominal values. Detailed measurement specifications, which may vary with altitude, are provided in the headers along with data.

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Table 3. IONS-06 ozonesonde launch sites during INTEX-B.

Station	Lat (N), Lon (W)	Spring Period (SN)
Boulder, CO	40.3, 105.2	28 Mar–18 May
Bratt's Lake, SAS	50.2, 104.7	3 Mar–17 May
Egbert, ONT	44.2, 79.98	8 Mar–17 May
Houston, TX	29.7, 95.4	1 Mar–10 May
Huntsville, AL	35.3, 86.6	1 Mar–9 May
Kelowna, BC	49.9, 199.4	8 Mar–17 May
Mexico City	19.4, 98.6	7–20 Mar
Narragansett, RI	41.5, 71.4	8 Mar–12 May
Richland, WA	46.0, 119.0	21 Apr–15 May
Stonyplain, AB	53.6, 114.1	8 Mar–18 May
Trinidad Head, CA	40.9, 124.2	3 Mar–19 May
Table Mtn, CA	34.4, 117.7	10–26 Apr
Valparaiso, IN	41.5, 87.0	19 Apr–16 May
Wallops Is, VA	37.9, 75.5	1 Mar–17 May
Walsingham, On	42.6, 80.6	24 Apr–5 May
Yarmouth, NS	43.9, 66.1	8 Mar–16 May

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Table 4. Models used for flight forecasting and analysis in INTEX-B^a.

Model	Type	Resolution		Underlying Meteorological Model	Chemical Package	Originator (Principal Investigator)	Additional References
		Horizontal	Vertical levels				
GEOS-4 GMAO	global	1° × 1.25°	55	GEOS-4, on-line transport and clouds	tagged tracers	I. Stajner, GMAO (D. J. Jacob, Harvard)	
GEOS-Chem	global	2° × 2.5°	37	GEOS-4, archived winds, separate transport and cloud algorithms	ozone-NO _x -VOC-aerosol-Hg chemistry	R. Yantosca (D. J. Jacob, Harvard)	Hudman et al. [2007]
MOZART-4-NCAR	global	2.8° × 2.8°	28	NCAR/NCEP, reanalysis archived winds,	ozone-NO _x -VOC-aerosol chemistry; tagged tracers	L. K. Emmons, NCAR	Pfister et al. [2005]
RAQMS	global	2° × 2.5°	36	NCEP GFS, isentropic coordinates	ozone-NO _x -VOC-chemistry; tagged tracers	R. B. Pierce, NASA Langley	Pierce et al. [2003]
STEM	regional	4 × 4 km to 80 × 80 km	21	RAMS-WRF-based, archived winds NCEP, GFS convection	ozone-NO _x -VOC-aerosol chemistry; tagged tracers	G. R. Carmichael, University of Iowa	Tang et al. [2004]
FLEXPART	global	Lagrangian		NCEP GFS winds	tagged tracers	A. Stohl, NILU	Stohl et al. [1998]
Photochemical model/LaRC	0-d "box" model		ozone-NO _x -VOC-chemistry	J. H. Crawford, NASA Langley	Olson et al. [2004]

^a Flight planning process also used weather forecasts and FLEXPART trajectories based on NCEP analysis and near real-time satellite data from AIRS and MOPITT (CO columns), MODIS (aerosol, fire counts), OMI (NO₂), TOMS (tropospheric ozone, and absorbing aerosols), and LIS (lightning). Potential vorticity and convective influence plots were also available.

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Table 5. INTEX-B DC-8 missions and salient activity^a.

Flight	Date	Operational Base ^b	Flight Hours	Salient Mission Activity
3	4 Mar 2006	Houston, Texas	6.1	sampling of outflow from south eastern US along the satellite tracks, validation of Aura satellite instruments (TES and OMI), as well as Aqua and Envisat
4	9 Mar 2006	Houston, Texas	8.4	sampling of outflow from Mexico City to the Gulf of Mexico, coordination with C-130 to investigate the aging of the Mexico City pollution, validation of Aura satellite instruments (TES and OMI), as well as Envisat
5	11 Mar 2006	Houston, Texas	7.4	sampling of the aged Mexico City plume previously sampled by the C-130 near Mexico City on 3/10, sampling the Mexico City boundary layer both remotely (DIAL) and in situ as initial conditions for a C-130 flight on 3/12
6	12 Mar 2006	Houston, Texas	7.8	validation of Aura satellite instruments (TES and OMI), inter-comparison with the NSF/NCAR C-130, sampling of aged Mexico City pollution outflow and its near source characterization using both remote (DIAL) and in situ measurements
7	16 Mar 2006	Houston, Texas	7.1	sampling of aged pollution and plume scouting for the C-130 north of Mexico City, sampling fresh Mexico City pollution during weekday conditions, sampling convective outflow over the southern Gulf of Mexico, and satellite validation (TES, OMI, and AIRS) that included a profile under TES at the time of overpass followed by in-progress profiling along the TES special observation track
8	19 Mar 2006	Houston, Texas	6.5	inter-comparison with the NSF/NCAR C-130, coordinated spiral with the J-31, validation of the Aura satellite instruments (TES and OMI), sampling of aged Mexico City pollution outflow, and the near source characterization of Mexico City an Monterrey pollution
9	22 Mar 2006	Houston, Texas to NASA Ames Research Center (ARC)	7.4	validation of HIRDLS observations in the stratosphere and upper troposphere using the DC-8 AROTAL Lidar as the primary instrument
10	17 Apr 2006	NASA ARC to Honolulu, Hawaii	9.0	inter-comparison with the C-130, sampling of aged and fresh Asian pollution over the Pacific, and validation of TES and OMI observations
11	23 Apr 2006	Honolulu, Hawaii	8.3	sampling of aged and fresh Asian pollution over the Pacific and validation of TES and OMI observations
12	25 Apr 2006	Honolulu, Hawaii	8.9	sampling of aged and fresh Asian pollution over the Pacific and validation of MLS and TES observations
13	28 Apr 2006	Honolulu, Hawaii	8.1	sampling Asian pollution to the west of Hawaii, providing validation data for TES with two spirals along the orbit track
14	1 May 2006	Honolulu, Hawaii to Anchorage, Alaska	9.2	extensive validation of the HIRDLS instrument aboard the Aura satellite, sampling Asian pollution, TES validation, inter-comparisons with instrumentation at the Mauna Loa observatory, specialized channel run targeted to study air-sea exchange processes
15	4 May 2006	Anchorage, Alaska	7.7	validation for OMI especially behind a front, performing clear air sampling of Arctic air and recent Asian pollution, setting up coordination with the C-130 on 5/5
16	7 May 2006	Anchorage, Alaska	8.0	validation for MLS and TES along the limb track, vertical profiling through Asian plumes, setting up coordination for C-130 sampling the following day
17	9 May 2006	Anchorage, Alaska	8.4	providing validation for TES along its limb and nadir tracks, and vertical profiling through Asian plumes
18	12 May 2006	Anchorage, Alaska	9.2	sampling Asian pollution both ahead and behind the ridge associated with a large WCB, boundary sampling of the Bethel wetlands, validation spiral for TES
19	15 May 2006	Anchorage, Alaska to Grand Forks, North Dakota	8.8	inter-comparison with the C-130, spiral with the Duchess, boundary layer leg over Montana in the OMI view to seek apparent NO ₂ hotspots, interception of plumes of Asian outflow

^a More details on individual flights can be found at http://www-air.larc.nasa.gov/missions/intex-b/INTEX_B.Flight.times_DC8.htm

^b Houston, Texas, 29.6° N, 95.1° W; NASA ARC, 37.4° N, 122.0° W; Honolulu, Hawaii, 21.3° N, 157.9° W; Anchorage, Alaska, 61.2° N, 150.1° W; Grand Forks, North Dakota, 47.9° N, 97.4° W.

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Table 6. Principal DC-8 airborne intercomparisons.

Intercomparing Platforms	Dates (2006)	Approximate Location	Time, UT	Level Altitude	Legs, km
DC-8 and C-130	19 Mar	22° N, 98° W	18:34–19:12	0.6, 2.6	
DC-8 and J-31	19 Mar	20° N, 99° W	19:17–19:27	spiral	
DC-8 and C-130	17 Apr	43° N, 122° W	18:03–18:48	3.7, 6.1	
DC-8 and C-130	15 May	43° N, 126° W	18:18–19:23	5.2, 1.6, 0.3	
DC-8 and Beechcraft Duchess	15 May	47° N, 124° W	20:32–21:03	5.3, spiral	

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Table 7a. Satellite measurements targeted for validation in INTEX-B.

Satellite Instruments	Platform (Launch Date)	Validation Measurements	More Information
MISR	Terra (Dec 1999)	aerosol	http://terra.nasa.gov/About/MISR/index.php
MODIS	Terra (Dec 1999)	aerosol, H ₂ O	http://terra.nasa.gov/About/MODIS/index.php
MOPITT	Terra (Dec 1999)	CO	http://terra.nasa.gov/About/MOPITT/index.php
AIRS	Aqua (May 2002)	CO, H ₂ O	http://aqua.nasa.gov/about/instrument_airs.php
MODIS	Aqua (May 2002)	aerosol, H ₂ O	http://aqua.nasa.gov/about/instrument_modis_science.php
SCIAMACHY	Envisat (Mar 2002)	NO ₂ , HCHO	http://www.iup.physik.uni-bremen.de/sciamachy/
TES	Aura (July 2004)	O ₃ , CO, CH ₄	http://aura.gsfc.nasa.gov/
MLS	Aura (July 2004)	O ₃ , CO, HNO ₃	http://aura.gsfc.nasa.gov/
OMI	Aura (July 2004)	NO _x , HCHO, SO ₂	http://aura.gsfc.nasa.gov/
HIRDLS	Aura (July 2004)	O ₃ , CO (stratospheric)	http://aura.gsfc.nasa.gov/

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Table 7b. DC-8 satellite validation opportunities during INTEX-B.

Instrument	Flight	Dates (2006)	Approximate Location	Time, UT
TES/OMI/AIRS	3	4 Mar	33° N; 88° W	1754–1825
TES/OMI/AIRS/SCIAMACHY	3	4 Mar	24° N; 86° W	2016–2050
TES/OMI/SCIAMACHY	4	9 Mar	23° N; 91° W	1925–1958
TES/OMI	6	12 Mar	14–23° N; 97–99° W	1837–2107
TES/OMI/AIRS	7	16 Mar	20–28° N; 92–94° W	1914–2119
TES/OMI	8	19 Mar	19–25° N; 99–101° W	1942–2115
HIRDLS	9	22 Mar	20–40° N; 113–114° W	0619–0946
TES/OMI	10	17 Apr	32° N; 144° W	2252–2319
TES/OMI	11	23 Apr	25–40° N; 158–162° W	2225–0155
MLS	12	25 Apr	34–45° N; 155–156° W	2150–2318
TES/OMI	12	25 Apr	45° N; 160° W	2341–0007
TES/OMI	13	28 Apr	23–28° N; 162–163° W	2232–0027
TES/HIRDLS	14	1 May	19–55° N; 150–152° W	0751–1327
OMI	15	4 May	44° N; 139° W	2244–2248
MLS	16	7 May	51–59° N; 144–147° W	2100–2216
TES	16	7 May	40–50° N; 142–146° W	2238–0008
TES	17	9 May	42–61° N; 139–159° W	1957–2321
TES/OMI	17	9 May	42° N; 137° W	2345–0014
TES/OMI	18	12 May	49° N; 173° W	0016–0046

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Table 8. Upper tropospheric (7–12 km) mixing ratios and tropospheric columns (surface–12 km) of key atmospheric constituents that are potentially amenable to space-based observation*.

Region [†]		H ₂ O (ppm)	CO ₂ (ppm)	O ₃ (ppb)	CO (ppb)	CH ₄ (ppb)	N ₂ O (ppb)	OCS (ppt)	NO ₂ (ppt)	HNO ₃ (ppt)	PAN (ppt)
1	UT	617±478 [#]	382±4.9	38±10	79.1±11.6	1770±10	321±0.3	478±13	23±49	66±64	50±148
	Trop. Column (molec. cm ⁻²)	1.00E+23 ± 3.48E+22	6.55E+21 ± 1.80E+20	7.49E+17 ± 68.57E+16	1.84E+18 ± 2.04E+17	3.07E+19 ± 9.27E+17	5.02E+18 ± 1.28E+17	8.27E+15 ± 2.74E+14	1.78E+15 ± 2.13E+15	4.96E+15 ± 1.64E+15	2.09E+15 ± 1.79E+15
2	UT	368±328	382±5.1	39±8	75.3±7.8	1770±8	321±0.1	472±8	39±47	59±80	23±25
	Trop. Column (molec. cm ⁻²)	3.04E+22 ± 1.22E+22	4.83E+21 ± 1.20E+20	5.62E+17 ± 7.61E+16	1.32E+18 ± 5.46E+17	2.25E+19 ± 6.35E+17	3.53E+18 ± 8.92E+16	6.02E+15 ± 1.65E+14	4.91E+15 ± 1.60E+16	4.60E+15 ± 4.28E+15	3.91E+15 ± 5.33E+15
3	UT	315±300	385±8.4	74±14	134±30	1820±14	320±0.6	493±29	20±27	70±70	166±143
	Trop. Column (molec. cm ⁻²)	4.66E+22 ± 1.47E+22	6.92E+21 ± 1.95E+20	1.15E+18 ± 9.13E+16	2.62E+18 ± 1.71E+17	3.28E+19 ± 9.47E+17	5.74E+18 ± 1.58E+17	8.73E+15 ± 2.82E+14	9.13E+14 ± 2.64E+15	1.39E+15 ± 5.49E+14	3.26E+15 ± 1.64E+15
4	UT	208±258	385±7.3	64±13	130±19	1810±11	3.20±0.5	483±16	33±30	32±20	168±165
	Trop. Column (molec. cm ⁻²)	8.52E+22 ± 2.58E+22	6.81E+21 ± 1.78E+20	1.00E+18 ± 9.55E+16	2.30E+18 ± 1.62E+17	3.21E+19 ± 8.83E+17	3.48E+18 ± 7.08E+16	8.60E+15 ± 2.68E+14	1.04E+15 ± 1.93E+15	1.62E+15 ± 4.42E+14	1.88E+15 ± 8.91E+14
Region		CH ₂ O (ppt)	Acetone (ppt)	CH ₃ OH (ppt)	CH ₃ OOH (ppt)	H ₂ O ₂ (ppt)	HCN (ppt)	CH ₃ CN (ppt)	CH ₃ Cl (ppt)	C ₂ H ₆ (ppt)	C ₂ H ₂ (ppt)
1	UT	47±70	460±275	470±405	361±173	480±360	228±75	120±29	569±18	459±102	73±529
	Trop. Column (molec. cm ⁻²)	9.54E+15 ± 4.72E+15	1.42E+16 ± 3.47E+15	1.74E+16 ± 6.50E+15	8.13E+15 ± 1.63E+15	2.00E+16 ± 6.15E+15	5.49E+15 ± 1.13E+15	2.22E+15 ± 2.40E+14	1.05E+16 ± 7.20E+14	1.40E+16 ± 4.94E+15	3.25E+15 ± 1.01E+15
2	UT	33±56	401±126	437±220	285±83.3	373±216	191±55	129±44	556±9	406±68.6	54±19
	Trop. Column (molec. cm ⁻²)	1.69E+16 ± 9.41E+15	8.98E+15 ± 4.66E+15	1.18E+16 ± 8.60E+15	4.36E+15 ± 6.60E+14	1.06E+16 ± 2.64E+15	5.96E+15 ± 3.90E+15	1.75E+15 ± 3.46E+14	7.25E+15 ± 2.56E+14	7.16E+15 ± 2.29E+15	3.58E+15 ± 5.40E+15
3	UT	37±51	891±340	1540±1040	570±281	483±276	268±84	137±29	564±29	1030±237	227±106
	Trop. Column (molec. cm ⁻²)	1.90E+15 ± 6.63E+14	1.35E+16 ± 2.45E+15	2.21E+16 ± 5.70E+15	9.73E+15 ± 1.98E+15	1.11E+16 ± 4.25E+15	5.02E+15 ± 7.79E+14	2.11E+15 ± 3.16E+14	9.74E+15 ± 2.76E+14	2.34E+16 ± 1.73E+15	4.70E+15 ± 6.42E+14
4	UT	47±47	662±219	947±543	592±251	405±319	252±94	162±55	585±19	801±148	210±76
	Trop. Column (molec. cm ⁻²)	3.65E+15 ± 8.70E+14	1.02E+16 ± 1.87E+15	1.31E+16 ± 4.16E+15	1.28E+16 ± 2.44E+15	1.79E+16 ± 4.53E+15	5.58E+15 ± 1.01E+15	2.38E+15 ± 3.64E+14	9.99E+15 ± 2.91E+14	1.77E+16 ± 2.14E+15	3.43E+15 ± 6.52E+14

* Regions are as defined in Fig. 2a. Terrain effects in Region 2 cause the column to be 2–12 km above surface. Missing data in column calculation were interpolated.

[#] Mean ± one sigma.

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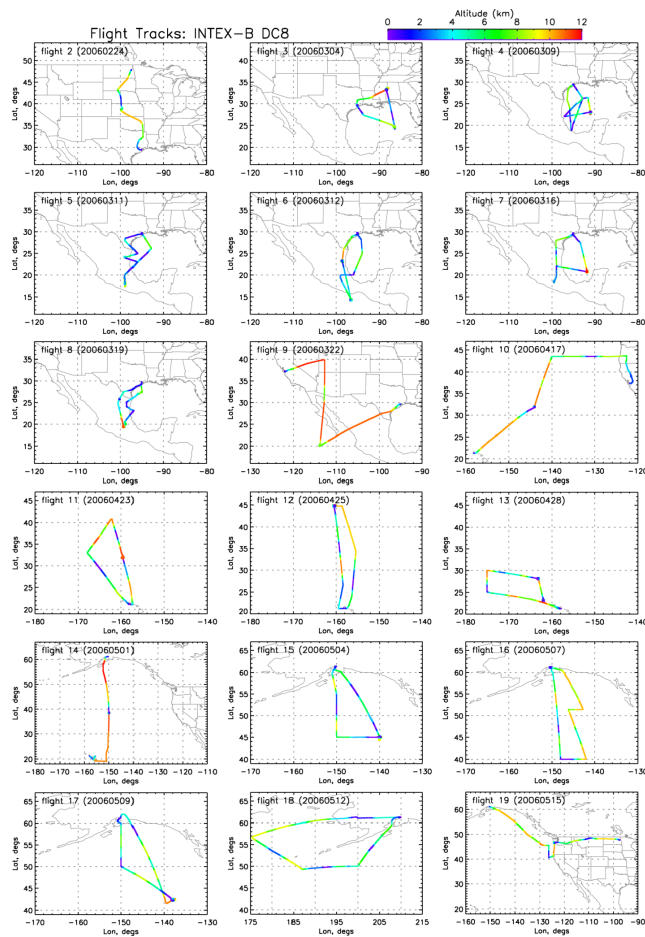


Fig. 1. DC-8 individual flight tracks in INTEX-B colored by altitude. 17 Science flights were performed from 4 March to 15 May 2008.

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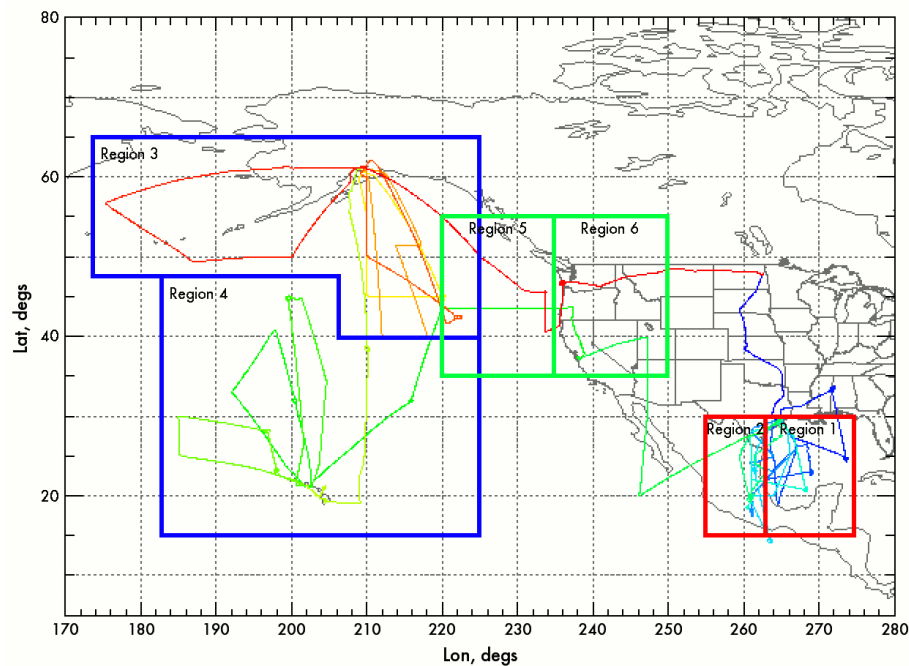


Fig. 2a. Composite of DC-8 flight tracks during INTEX-B. Boxes indicate select regions subjected to data analysis.

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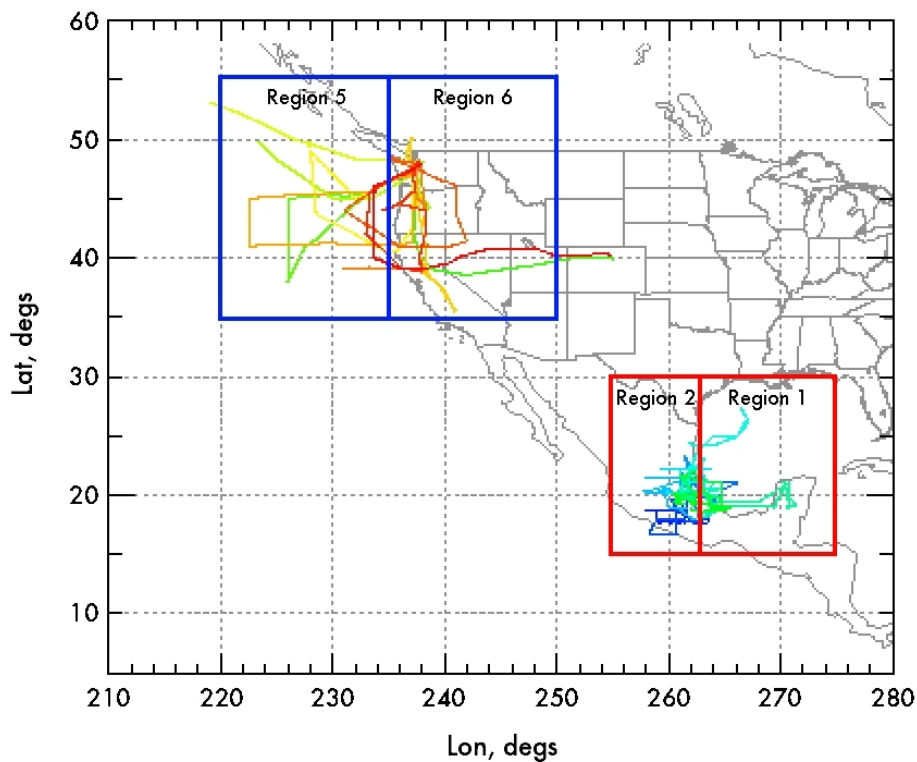


Fig. 2b. Composite of C-130 flight tracks during MILAGRO/INTEX-B.

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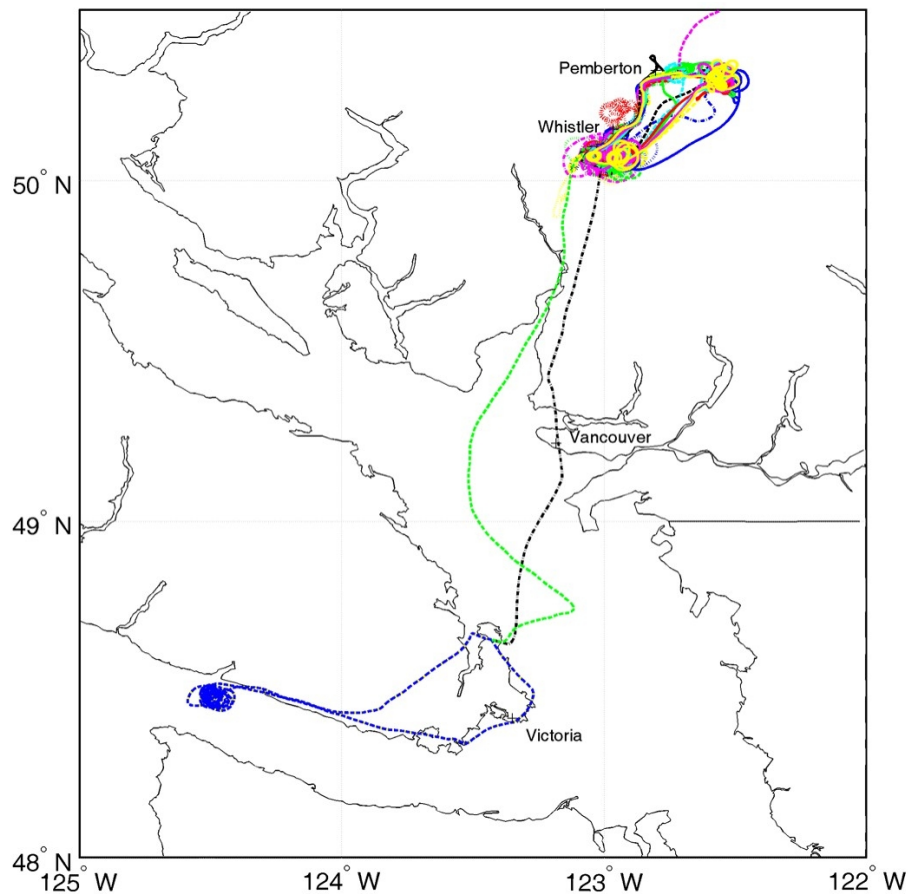


Fig. 2c. Flight paths of the Cessna 207 aircraft during the INTEX-B campaign over 22 April 2006 to 17 May 2006. Colors represent individual flights.

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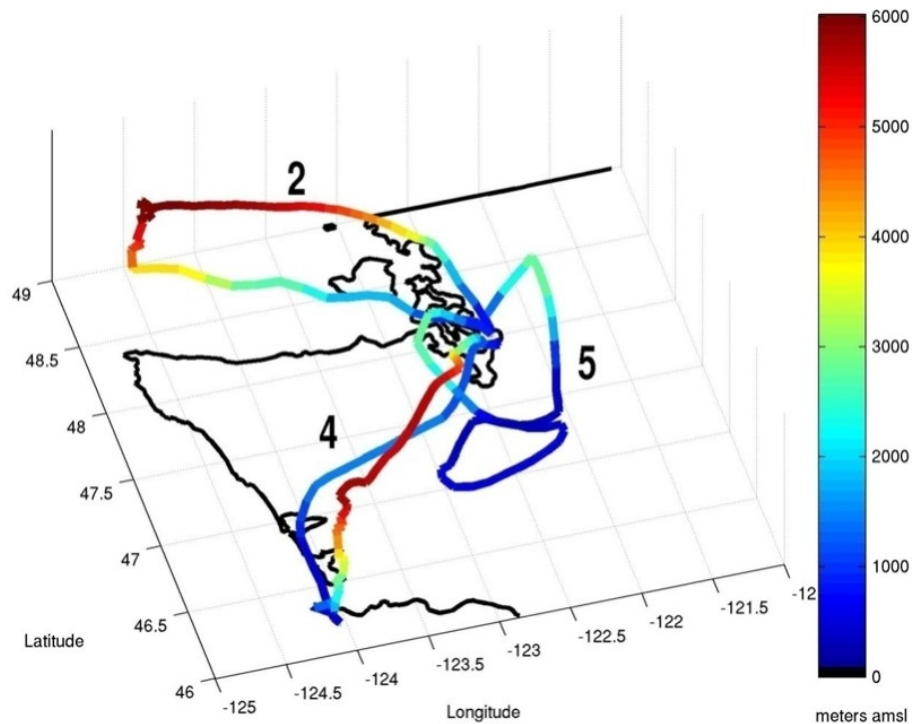


Fig. 2d. Sample flight tracks of the U. of Washington Beechcraft Duchess colored by altitude. Three representative flights are shown: flight 2 (18 April 2006), flight 4 (30 April 2006), and flight 5 (4 May 2006).

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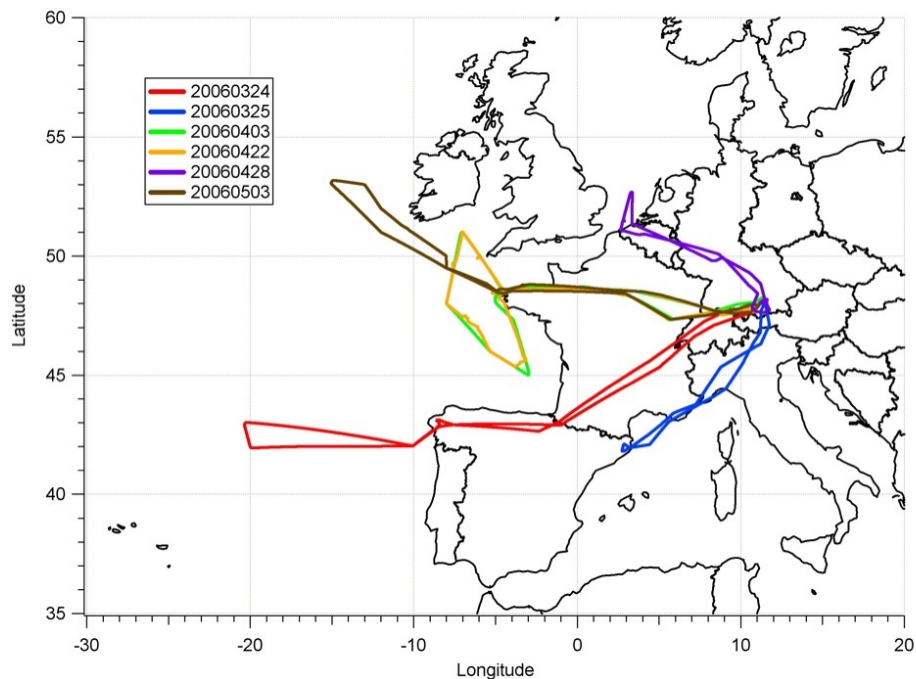


Fig. 2e. Flight paths of the DLR Falcon-20 during INTEX-B.

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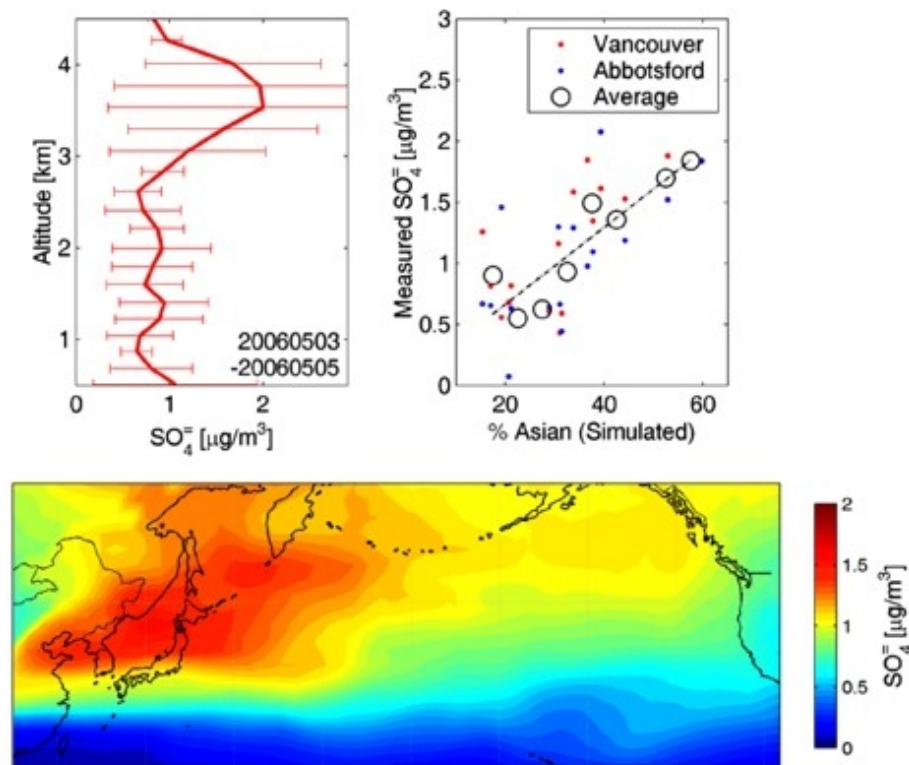


Fig. 3. Transpacific transport of sulfate. The top left panel shows average Cessna Q-AMS measured sulfate over Whistler, British Columbia during 3–5 May 2006. The top right panel shows surface filter pack measurements of sulfate from Canada's National Air Pollution Network versus simulated East Asian sulfate fraction during April–May 2006. Black circles denote 5% binned averages. Bottom panel shows simulated mean sulfate concentrations over April–May 2006 at 3.5 km from the GEOS-Chem model. Adapted from van Donkelaar et al. (2008).

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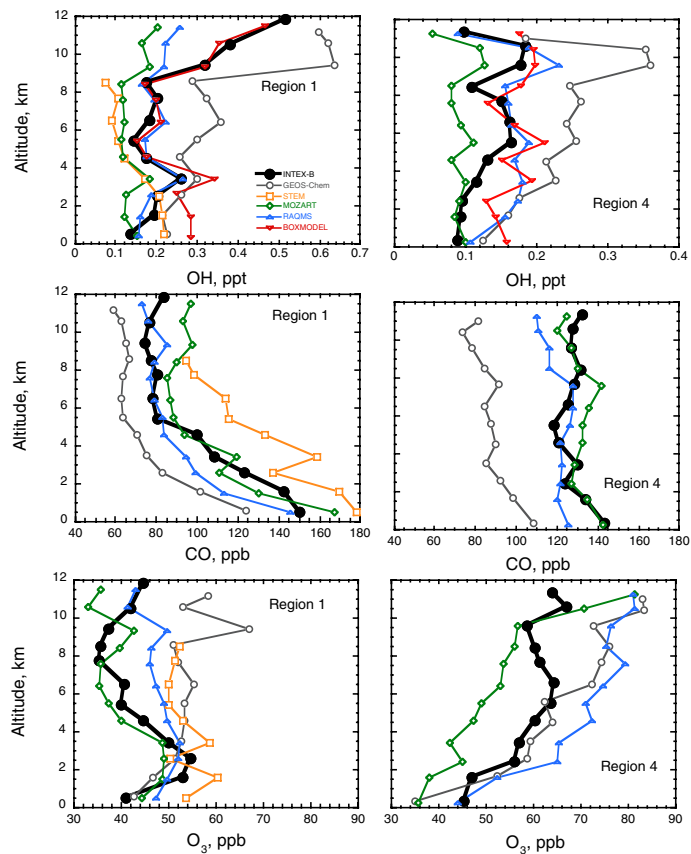


Fig. 4a. A comparison of INTEX-B observations over the Gulf of Mexico (Region 1) and subtropical Pacific (Region 4) with along track post mission simulations of select species. Regions 1 and 4 are as shown in Figure 2a. STEM simulations were not available for Region 4.

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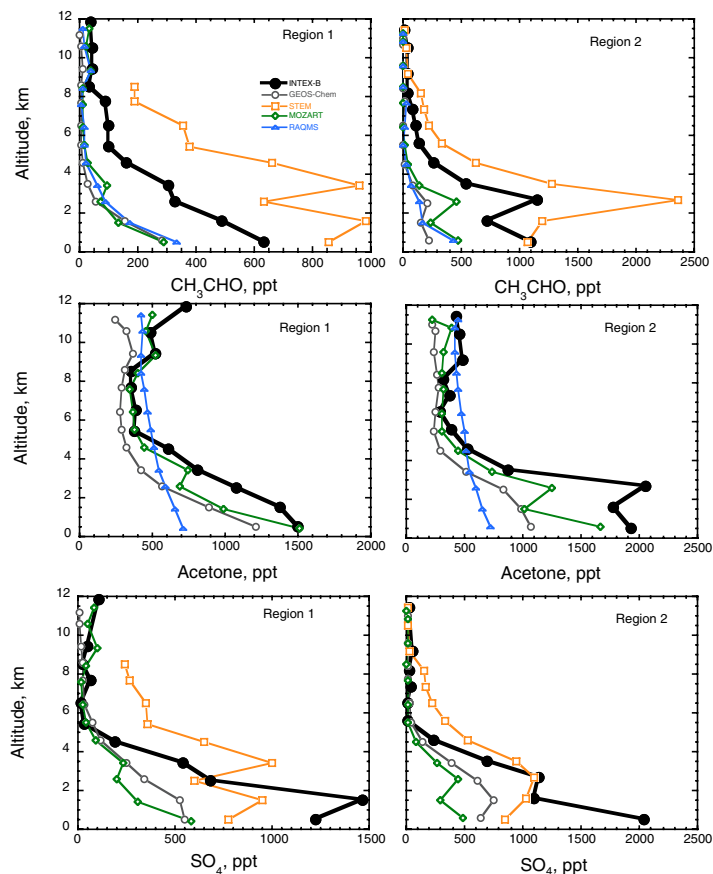


Fig. 4b. A comparison of INTEX-B observations over the Gulf of Mexico (Region 1) and Continental Mexico (Region 2) with along track post mission simulations of select species. Regions 1 and 2 are as shown in Fig. 2a.

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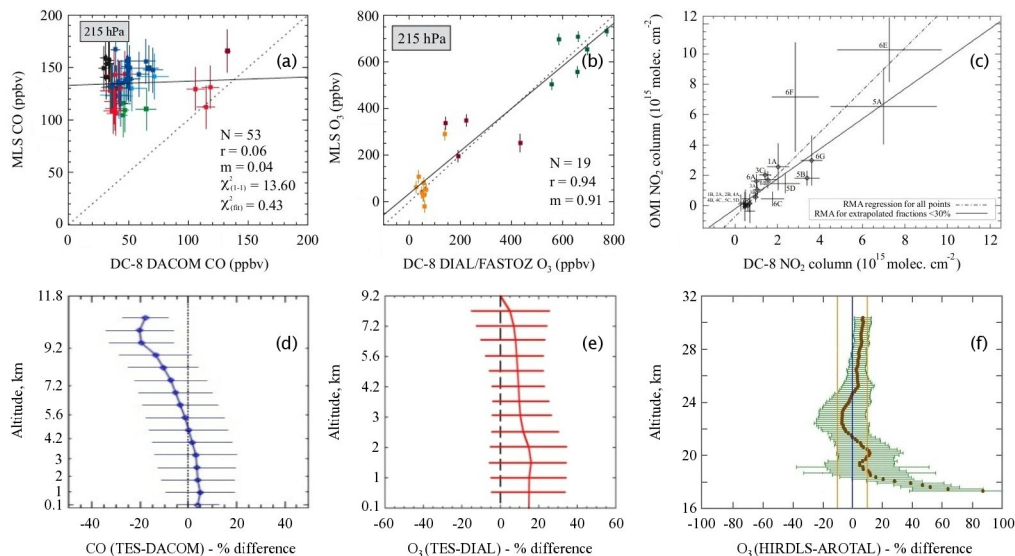


Fig. 5. Comparison of Aura satellite retrieved observations with DC-8 INTEX-B measurements. Adapted from Livesey et al. (2007), Boersma et al. (2008), Luo et al. (2007), and Nardi et al. (2008). See text for more details.

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