

# Ozone production in pollution plumes transported to the Arctic

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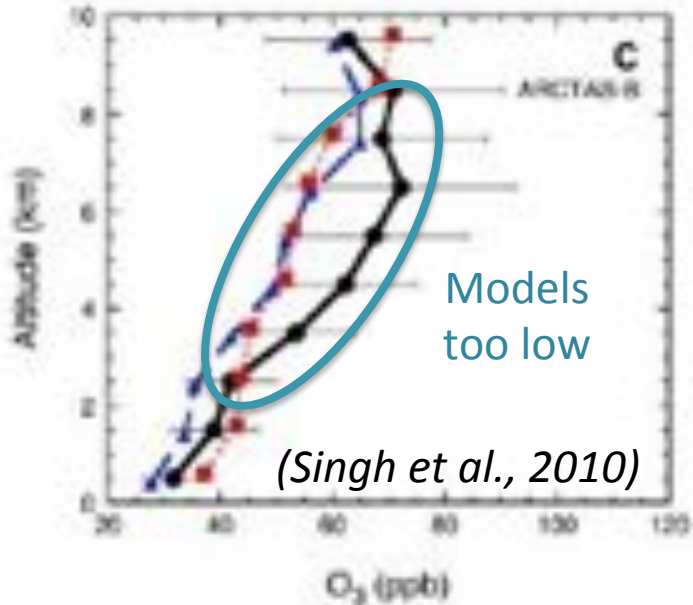
[5] Institut für Physik der Atmosphäre, Deutsches Zentrum für Luft- und Raumfahrt (DLR), Oberpfaffenhofen, Germany

# POLARCAT: Long-range transport of pollutants to the Arctic

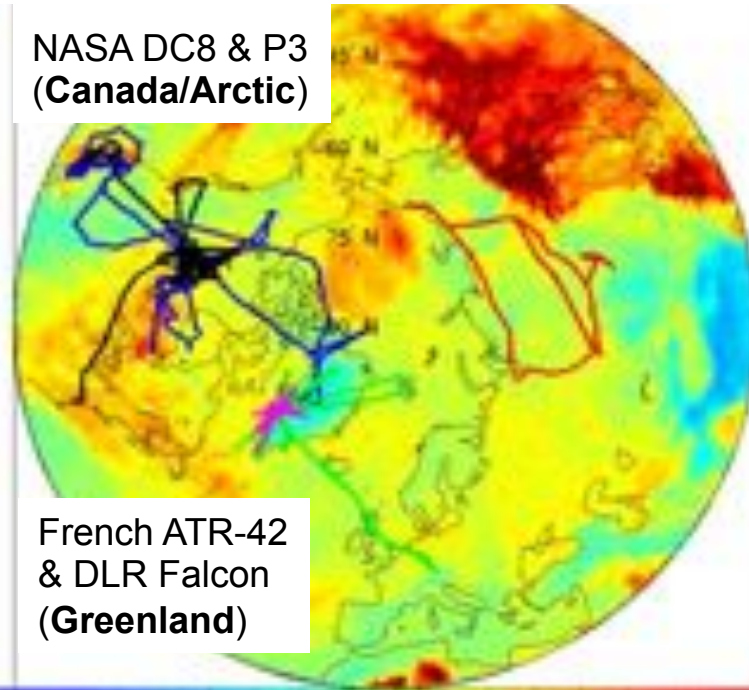
- Multiple aircraft campaigns spring/summer 2008
- Arctic ozone budget – how much  $O_3$  from mid-latitudes, formed (destroyed) during transport or from UTLS?



GEOS-Chem/MOZART vs ARCTAS-B aircraft data (DC8, summer 2008)



NASA DC8 & P3 (Canada/Arctic)



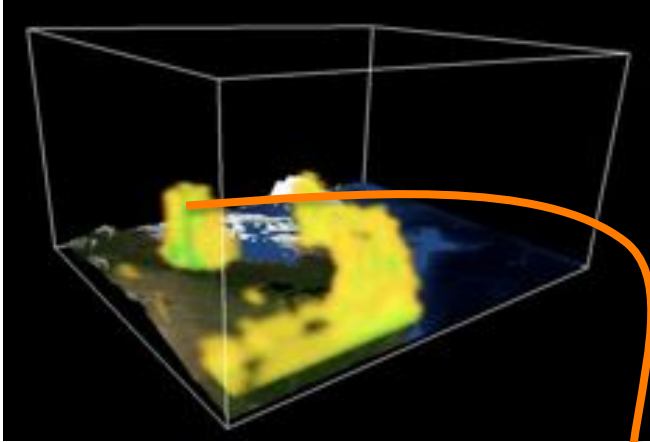
French ATR-42 & DLR Falcon (Greenland)

**POLARCAT  
summer 2008  
aircraft  
campaigns**

*IASI CO columns  
July 2008  
Pommier et al. (2010)*

0 0.2 0.4 0.6 0.8 1 1.2 1.4 1.6 1.8 2 2.2 2.4 2.6 2.8 3 3.2 3.4  
Total Column CO ( $10^{18}$  molecules/cm<sup>2</sup>)

# Using POLARCAT aircraft measurements to understand long range transport & chemistry

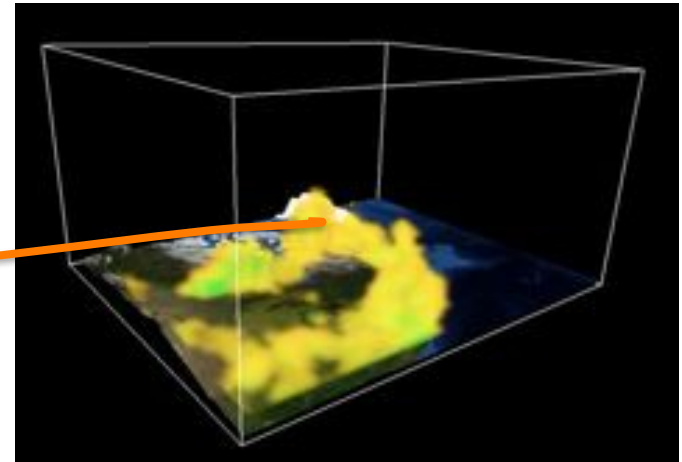


Use emissions with 3D chemistry and meteorology to predict measurements in fresh plumes and plumes after long range transport

**fresh emissions**

- **Chemical composition**
- **Injection height**

**transport &  
chemistry**



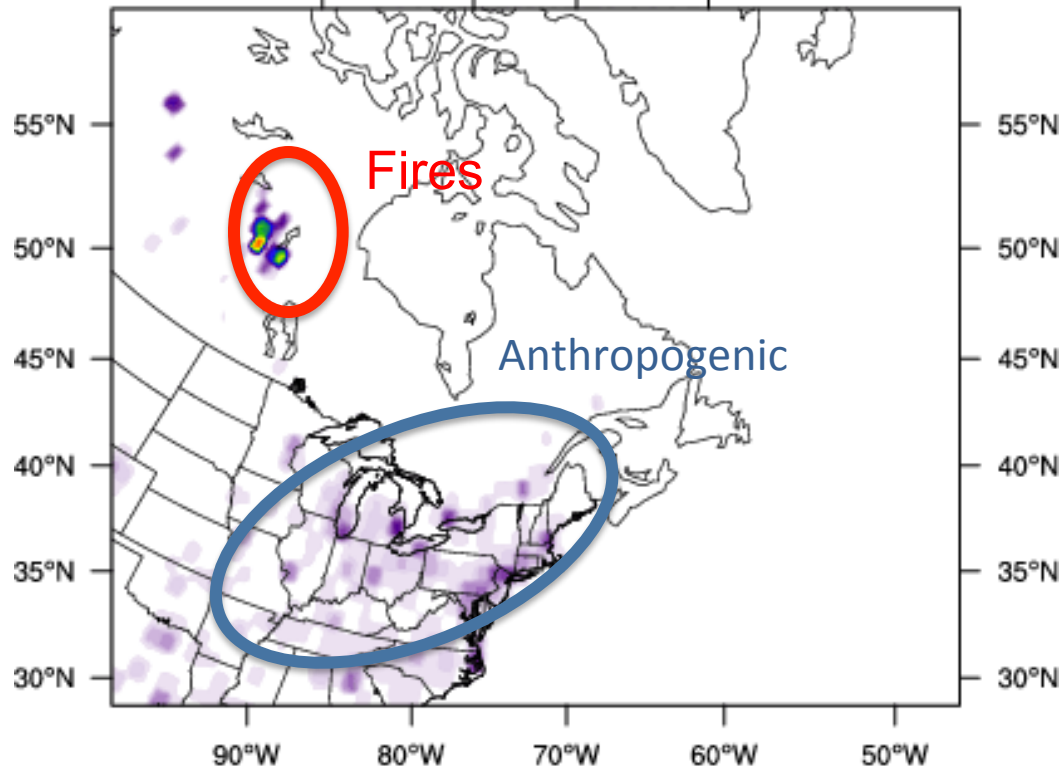
**predicted chemical  
composition and spatial  
extent of pollution plumes**

**Key issue: model resolution  
vs. plume spatial extent**

# WRF-Chem model setup and simulations

3 July 2008

CO EMISSIONS mole km<sup>-2</sup> hr<sup>-1</sup>



**Case study:**  
**28 June 2008 – 8 July 2008**

35 km x 35 km, 27 vertical levels

## **POLMIP emissions**

- **Daily Anthropogenic emissions** 2008 inventory for ARCTAS (D. Streets)
- **Daily Fire emissions** from FINN-v1 (Wiedinmyer et al., Geosci. Model Devel., 2011) + online plume rise model (Freitas et al., ACP, 2007)

Initial & boundary conditions from MOZART4 for chemistry & GFS for meteorology + nudging (6hr)

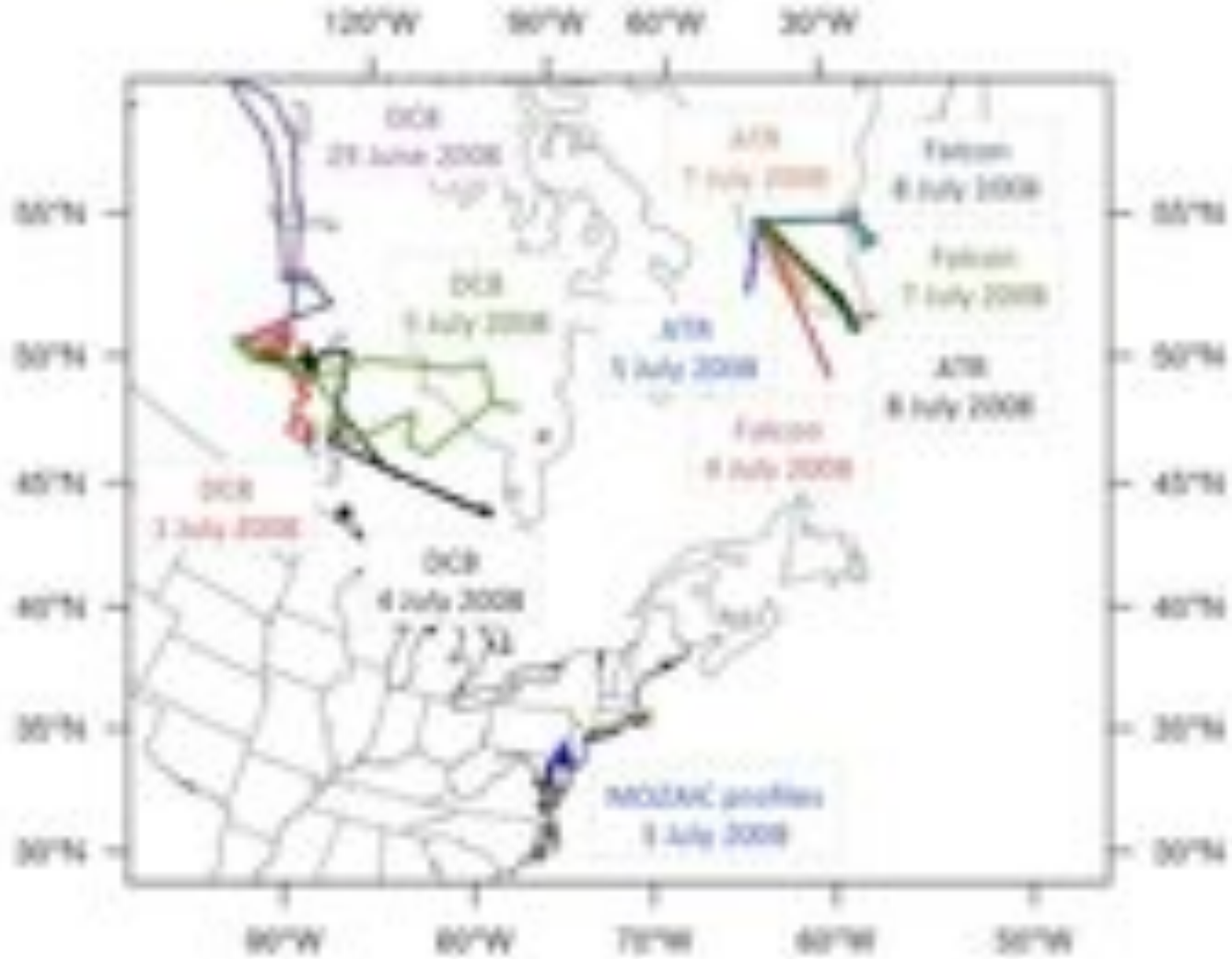
MOZART gas phase chemistry & GOCART aerosol scheme

Runs:

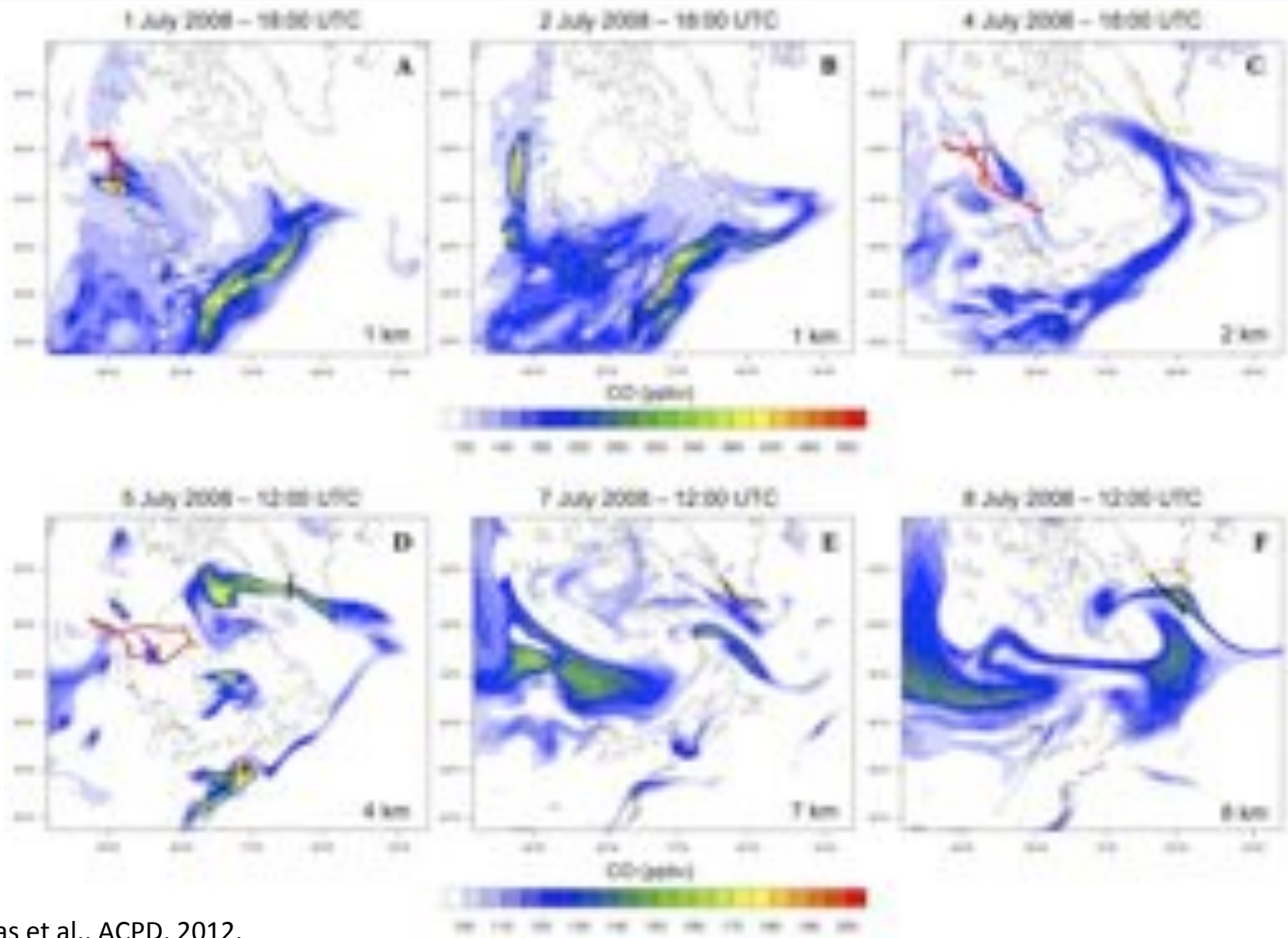
- Base case (all emissions included)
- No anthropogenic emissions
- No fire emissions

mol km<sup>-2</sup> hr<sup>-1</sup>

# POLARCAT flights used for evaluating model results

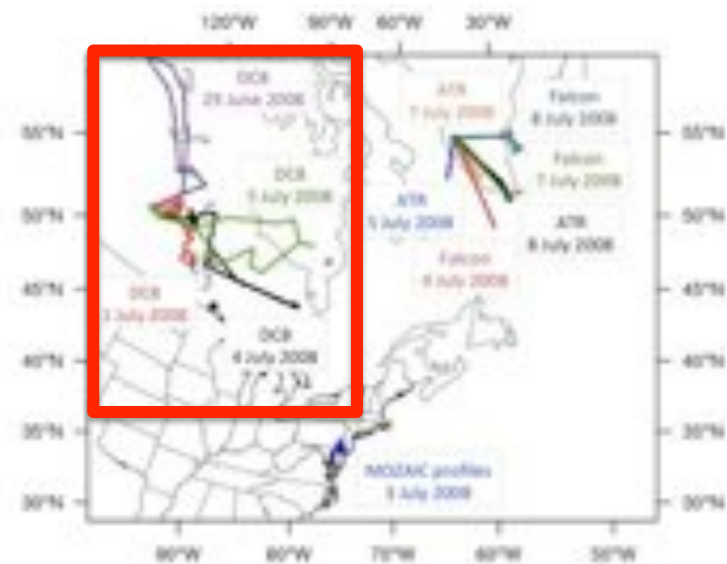
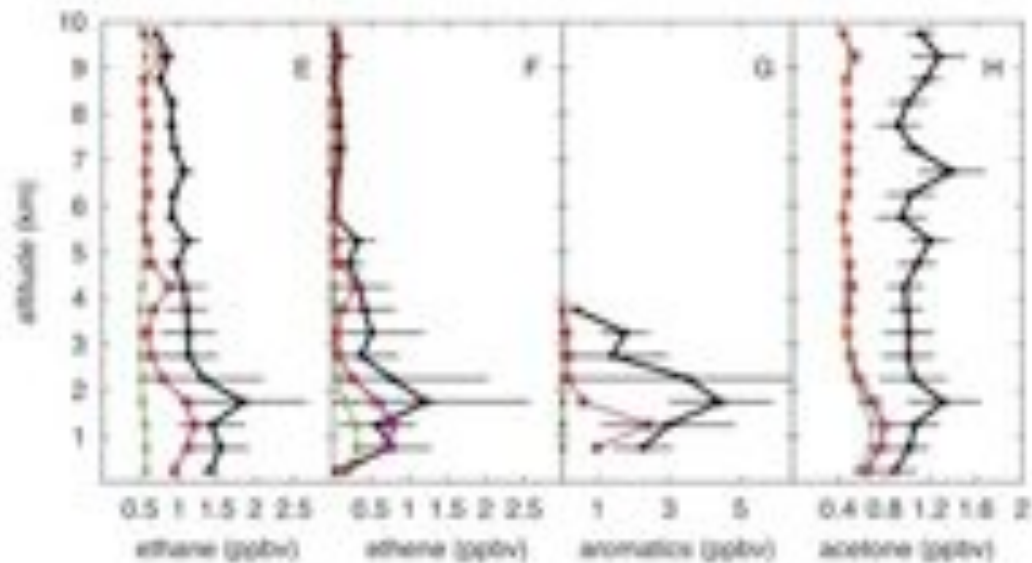
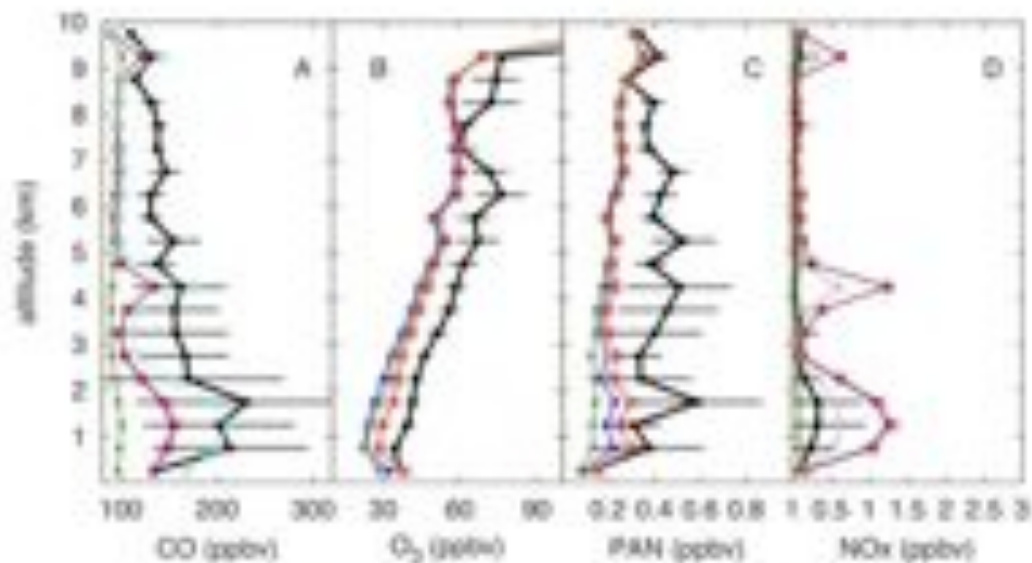


# Example pollution plumes – fresh emissions and uplift





# Comparison – fresh fire plumes in Canada



Measurements

Base model run

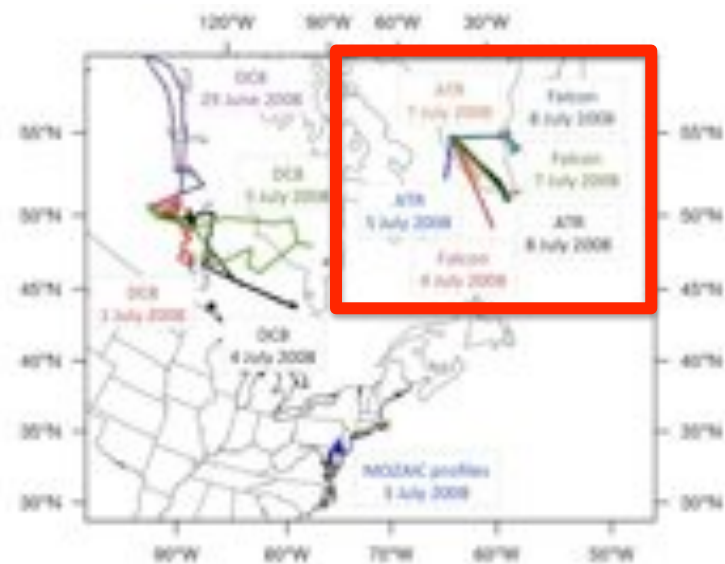
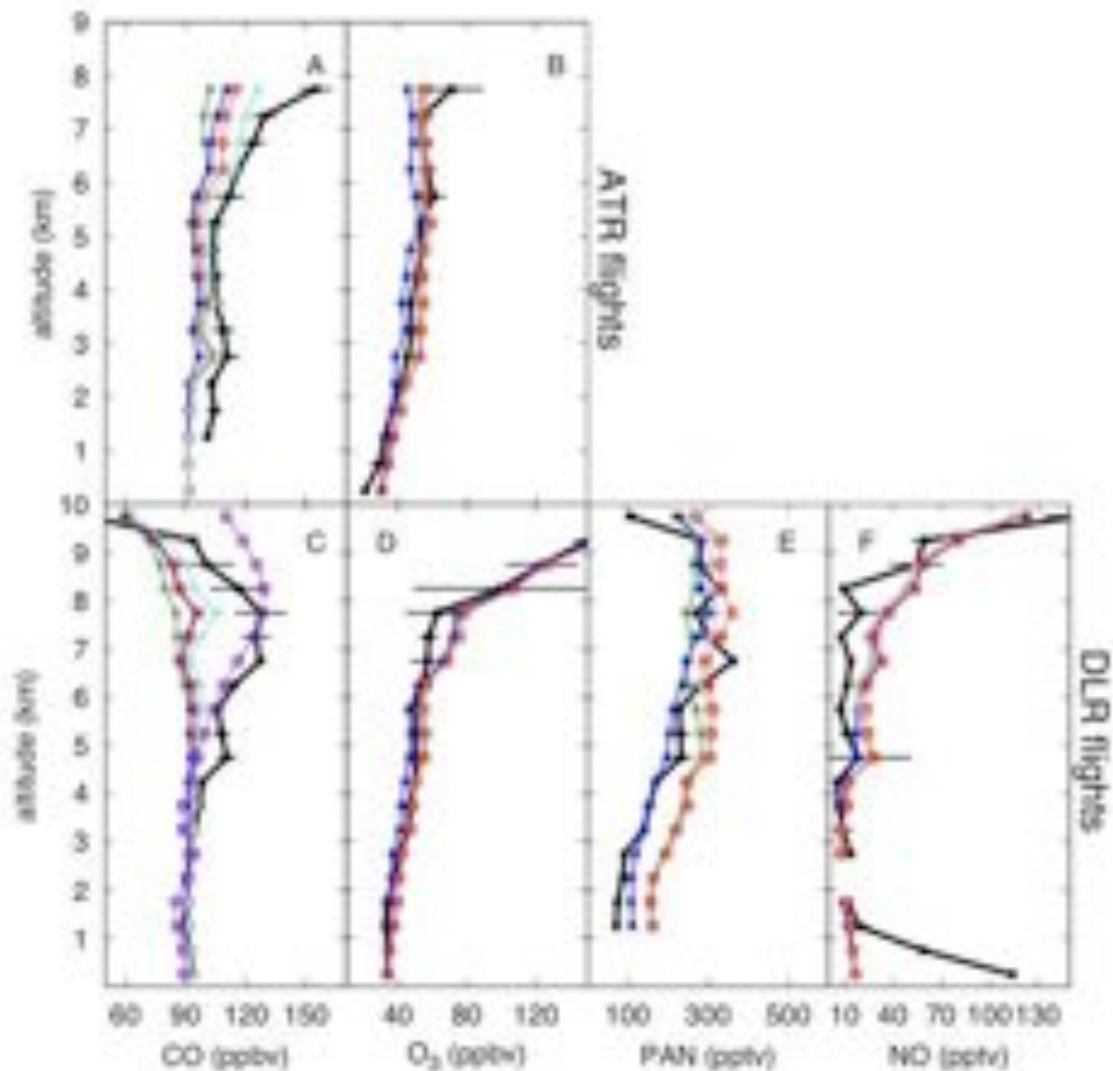
noAnthro emissions run

noFire emissions run

2x CO emissions from fires

½ NO<sub>x</sub> emissions from fires

# Comparison – aged plumes measured after long range transport (southern Greenland)



Measurements

Base model run

noAnthro emissions run

noFire emissions run

2x CO emissions from fires

½ NO<sub>x</sub> emissions from fires

CO profile near flight track



# Measurements of individual aged plumes – onboard the ATR-42 over S. Greenland

stratospheric  
airmass

pollution plume

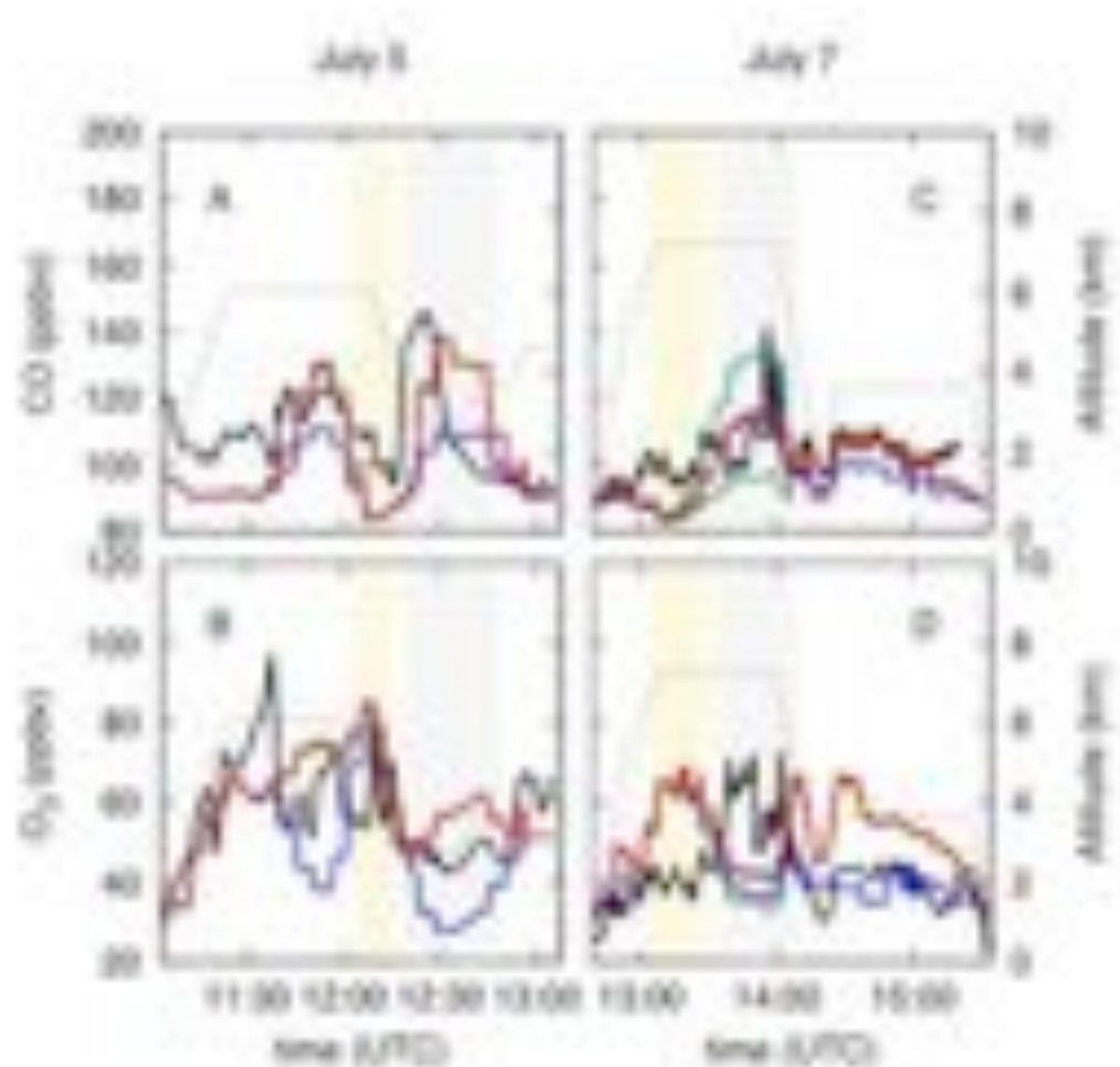
Measurements

Base model run

noFire emissions run

noAnthro emissions run

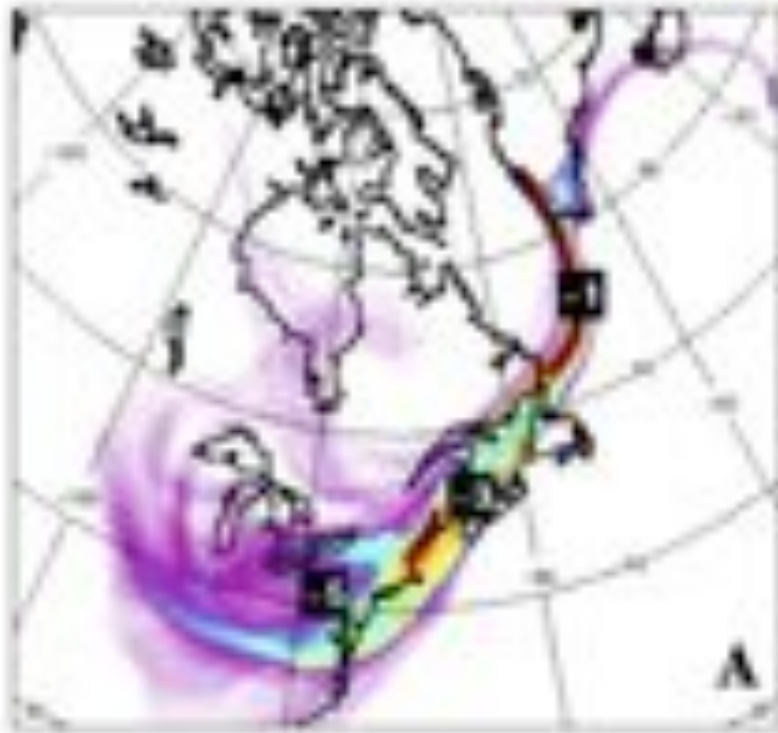
2x fire CO emissions



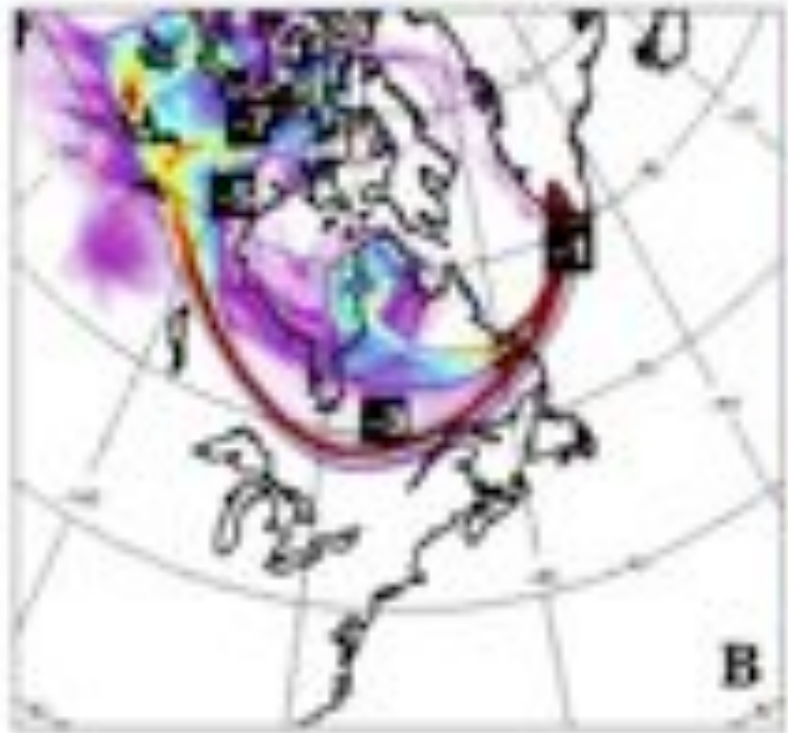
# FLEXPART-WRF used to identify air mass origins

## 5 July Plume (Anthropogenic) and 7 July Plume (BB)

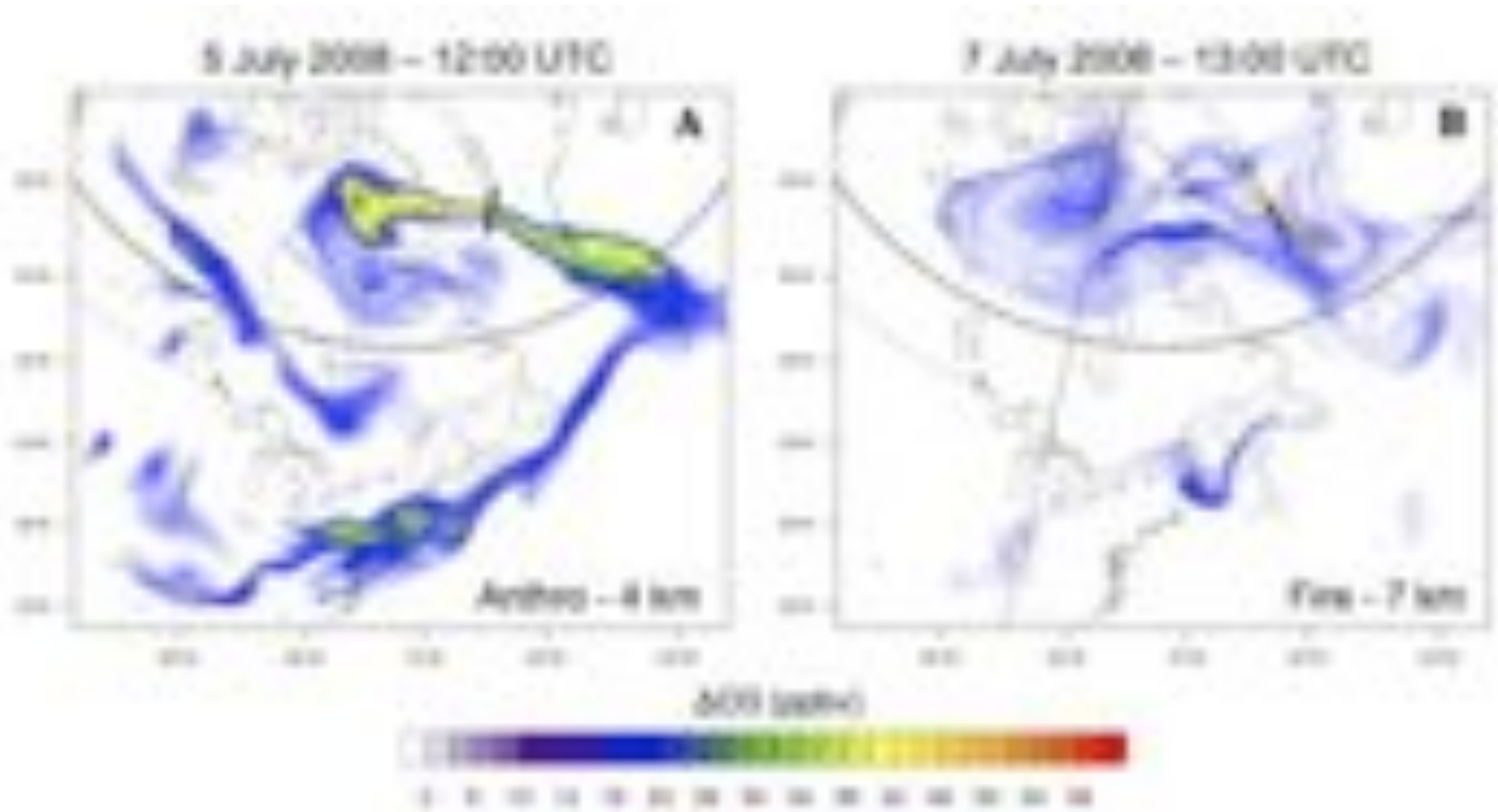
FLEXPART-WRF BACKWARDS MODE PER COLUMN  
Release July 5th 2008, 12:00 UTC



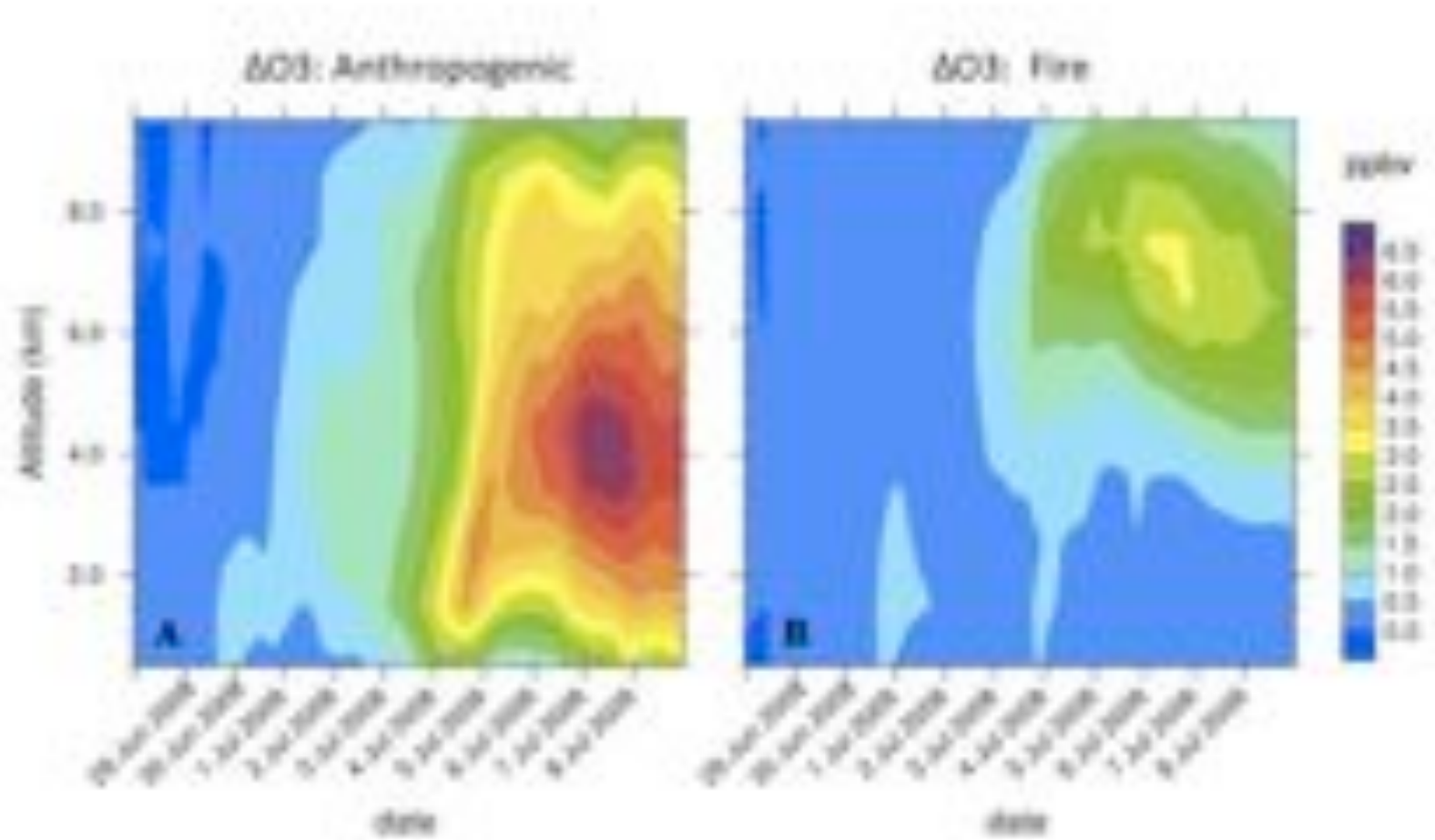
FLEXPART-WRF BACKWARDS MODE PER COLUMN  
Release July 7th 2008, 12:45 UTC



# Increase in ozone due to anthropogenic (left) and fire (right) emissions



# Quantifying the increase in ozone due to anthropogenic and fire pollution & chemistry (north of 55 °N)



# Summary & conclusions

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- Fresh and aged plumes correctly represented in the model
- The ozone produced due to anthropogenic and BB emissions has been quantified
- Air mass origin has been identified using FLEXPART-WRF PES
- Sensitivity to increased CO and decreased NO<sub>x</sub> emissions from fires has been studied
- Future work: focus on aerosol processes during long range transport to the Arctic



# Aircraft Campaign – July 2012

Jennie L. Thomas, Kathy Law, Claire Granier,  
Jean-Christophe Raut,  
Louis Marelle (LATMOS)

Hans Schlager, Anke Roiger, ACCESS Campaign team (DLR)

Ivar Isaksen, Stig Dalsoren (CICERO)



**ACCESS**  
Arctic Climate Change  
Economy and Society

# ACCESS aircraft campaign 2012

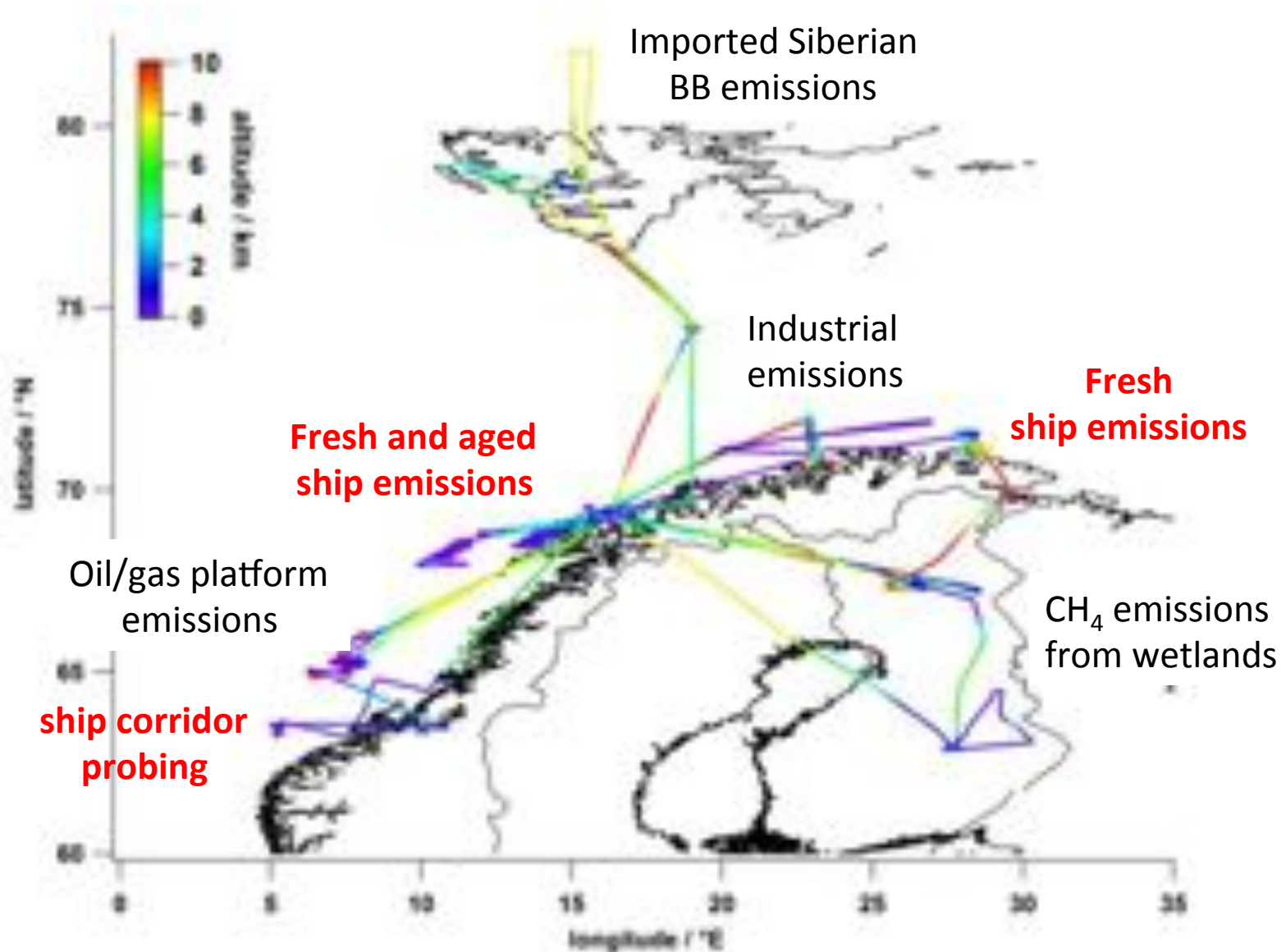
- DLR Falcon base: Andoya (Norway)
- Time period: 9th-27th July 2012
- 13 local flights performed in order to study Arctic pollution sources



# ACCESS campaign 2012 – flight overview

Date	Flight Objectives
11 July	F#1 Single ship plume study „Wilson Leer“ (and „Costa Delizioso“)
12 July	F#2 Single ship plume study „Wilson Nanjing“
13 July	F#3 Sampling of Kola Peninsula Plume
16 July	F#4 Low-level survey over southern Finland
17 July	F#5+6 Sampling of Siberian BB plume
19 July	F#7 Ship corridor probing F#8 Sampling of different oil/gas facility plumes
20 July	F#9 Single plume study at „Heidrun“ platform
22 July	F#10+11 Low-level survey over N/S Finland
25 July	F#12 Sampling of fresh ship emissions F#13 Vertical profile for Sodankylä FTS

# ACCESS campaign 2012 – flight overview

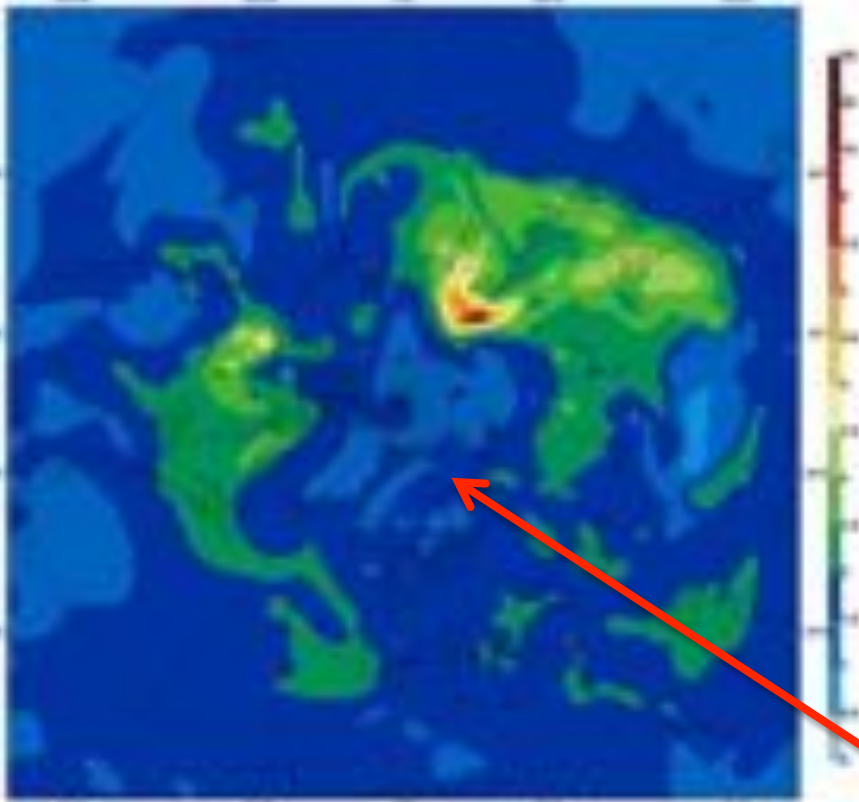


# Total column CO: IASI vs MACC Forecast

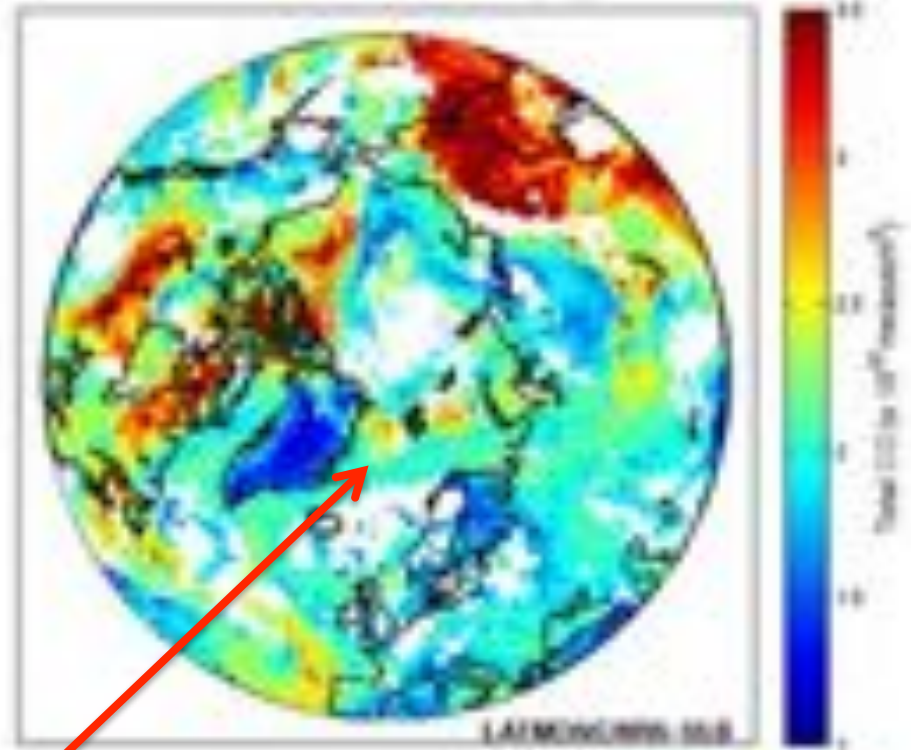
14 July MACC 12z

14 July

Satellite 14 July 2014 (00:00) (UTC) (UTC) (Forecast) (00:00) (UTC) (Saturday 14 July 2014 (00:00))  
North Hemisphere Total column Carbon monoxide (10<sup>-6</sup> molecules/cm<sup>2</sup>)



IASI Total CO - 20/12/2014 (00:00)



Different scales!

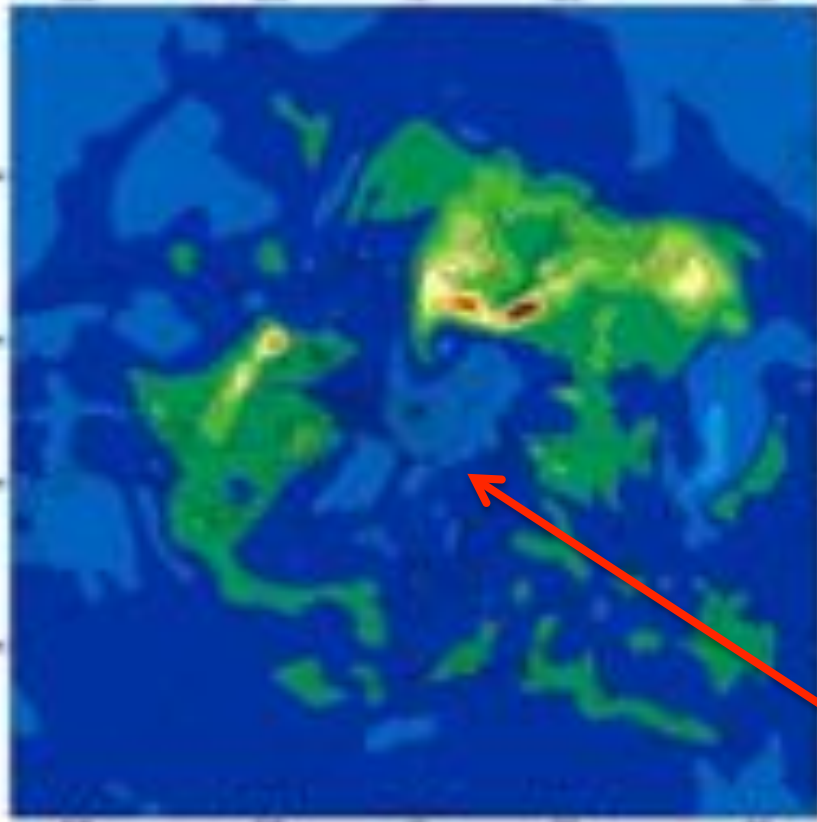


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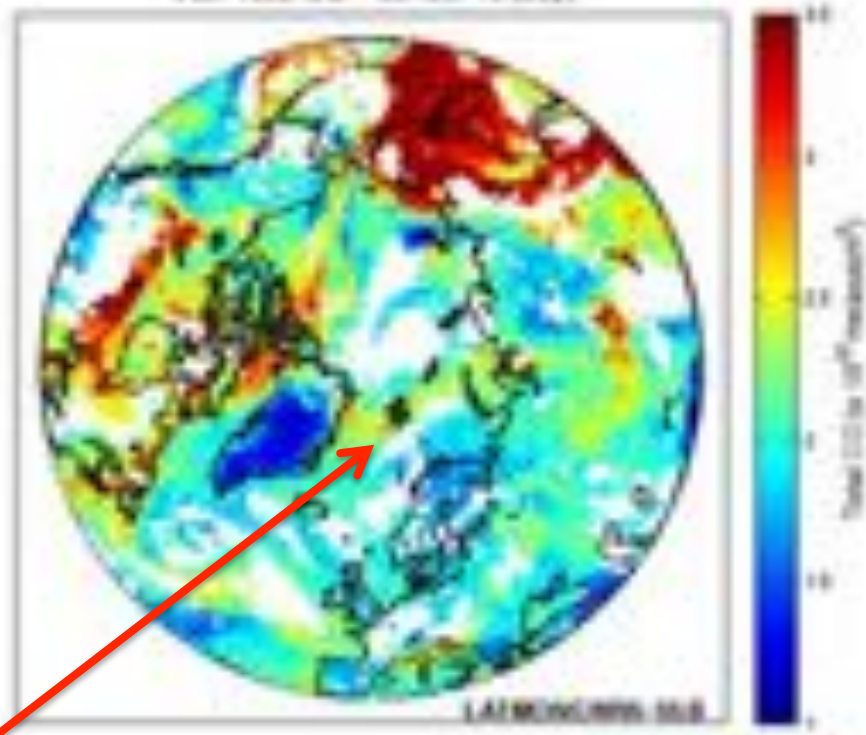
15 July MACC 12z

15 July

Sunday 15 July 2011 12:00 UTC (local) Forecast valid 15 July 2011 12:00 UTC  
North Hemisphere: Total column Carbon monoxide (10<sup>-6</sup> molecules/cm<sup>2</sup>)



IASI Total CO - 20110715 (12z)



Different scales!

# A multi-model evaluation of processes controlling spring and summer tropospheric ozone in the Arctic: The POLARCAT Model Intercomparison Project (POLMIP)

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S. Turquety<sup>4</sup>, S. A. Monks<sup>2</sup>, S. Tilmes<sup>1</sup>, B. Duncan<sup>5</sup>,  
Jennie L. Thomas<sup>3</sup>, I. Bouarar<sup>3</sup>, J. Mao<sup>6</sup>,  
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(1) Atmospheric Chemistry Division, NCAR, Boulder, CO, USA

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(5) Atmospheric Chemistry & Dynamics Laboratory, NASA Goddard, Greenbelt, MD, USA

(6) Atmospheric and Oceanic Sciences, Princeton University, USA

(7) ECMWF, Reading, England

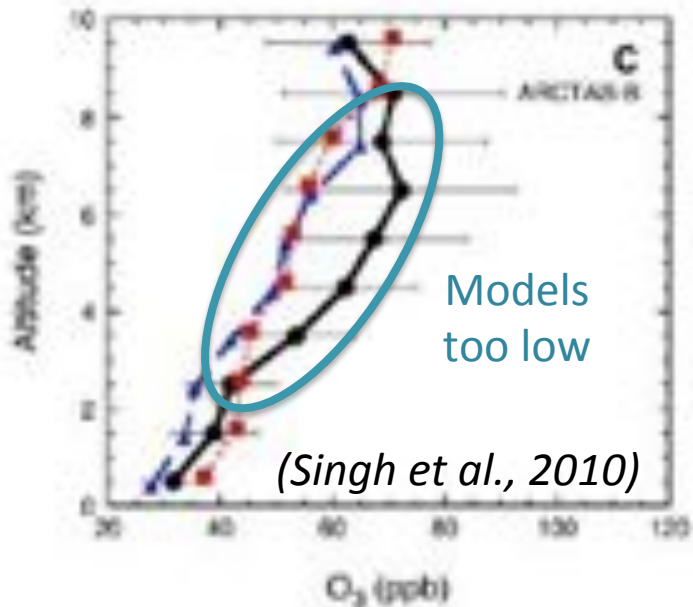
(8) Royal Netherlands Meteorological Institute (KNMI), De Bilt, Netherlands

# POLARCAT: Long-range transport of pollutants to the Arctic

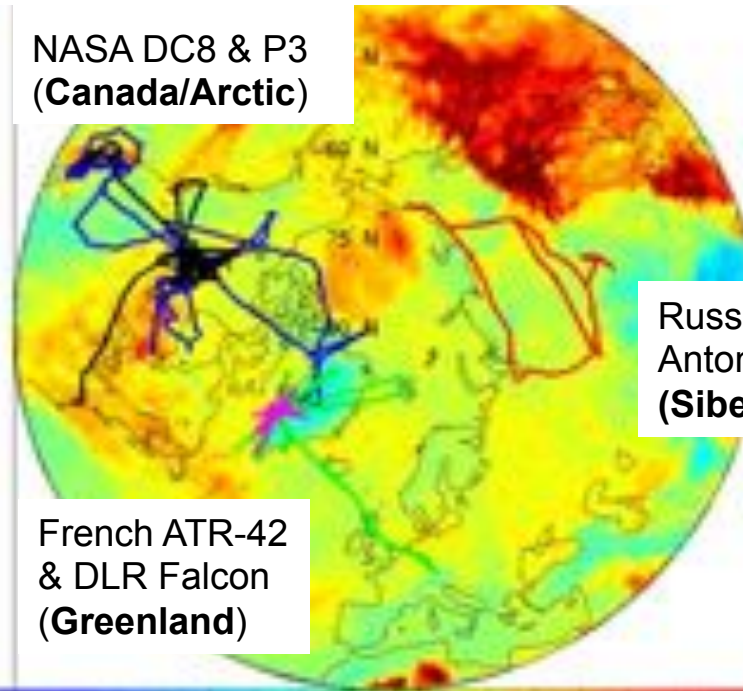
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GEOS-Chem/MOZART vs ARCTAS-B aircraft data (DC8, summer 2008)



NASA DC8 & P3 (Canada/Arctic)



**POLARCAT summer 2008 aircraft campaigns**

Russian-French Antonov-30 (Siberia)

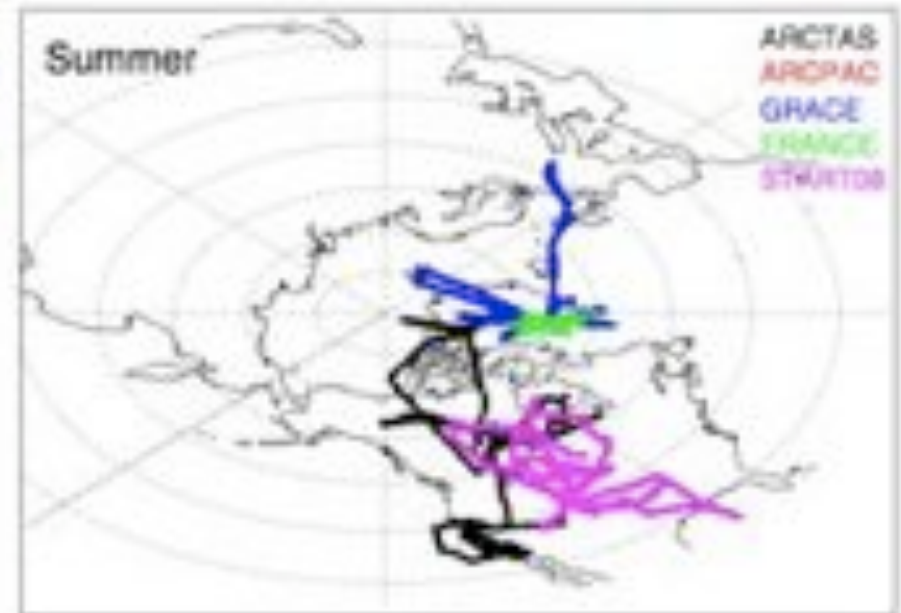
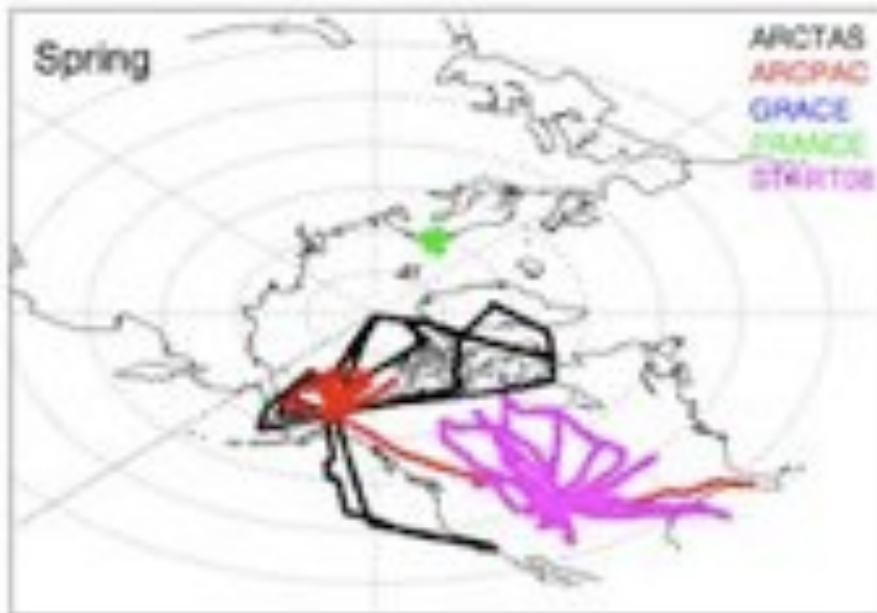
French ATR-42 & DLR Falcon (Greenland)

IASI CO columns July 2008  
Pommier et al. (2010)

# POLARCAT Aircraft Experiments

## April-July 2008

NASA - ARCTAS  
NOAA - ARCPAC  
DLR - GRACE  
POLARCAT-France



# Models

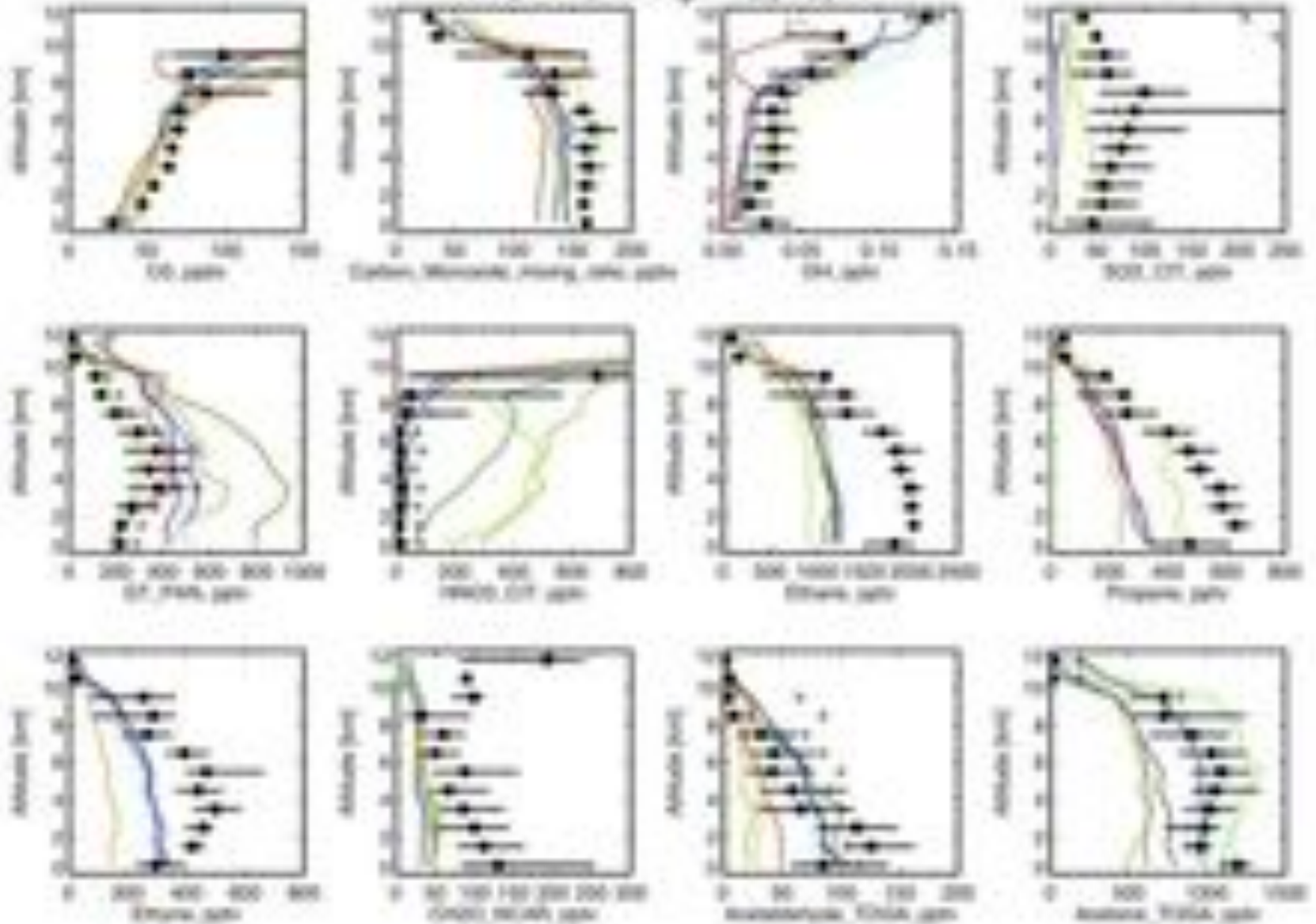
Emissions: Anthropogenic: Streets' ARCTAS-v1.2; Fires: FINN-v1; Other: MACCity

Model	Resolution	Meteorology	Chemistry
TOMCAT	2.8°x2.8°, 31 levels	ECMWF ERA-oper.	trop: 82 species
MOZART-4	1.9°x2.5°, 56 levels	GEOS-5	trop: 103 species
CAM-chem	1.9°x2.5°, 56 levels	GEOS-5	MOZART-4
LMDZ-INCA	1.9°x3.75°, 19 levels	ECMWF	trop: 89 species
C-IFS	T159 (~1°), 60 levels	ECMWF	trop: CB05, strat: linear. O3 (Cariolle)
TM5	2°x3°, 60 levels	ECMWF	trop: CB05
NASA GMI	2°x2.5°, 70 levels	GEOS-5	strat+trop (123 species)
GEOS-Chem	2°x2.5°, 47 levels	GEOS-5	trop: ~100 species
WRF-Chem	70, 35, 17.5 km		MOZART-GOCART

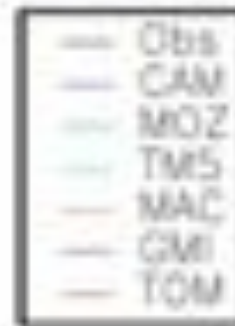
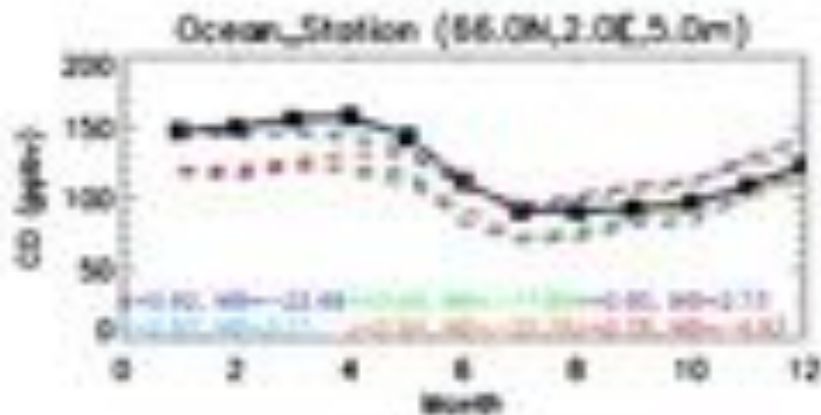
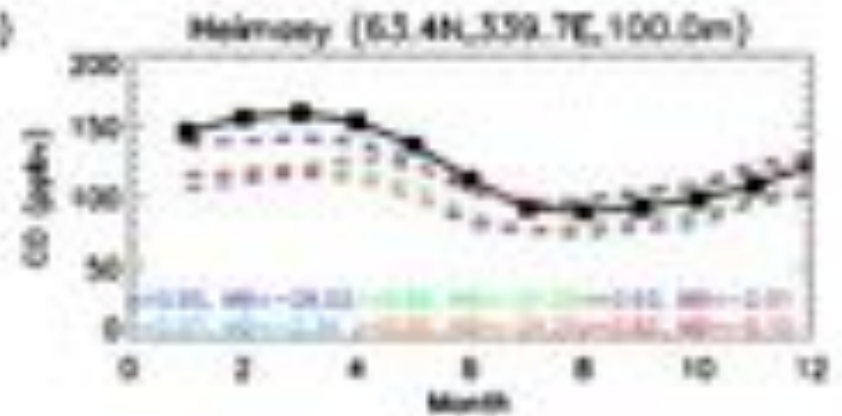
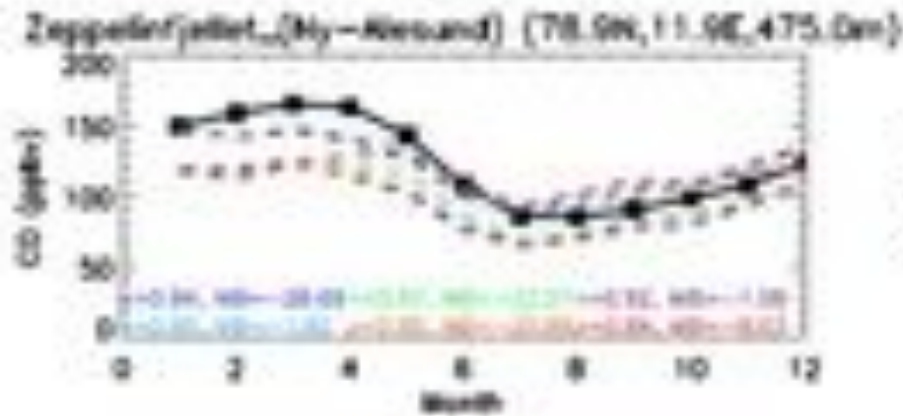
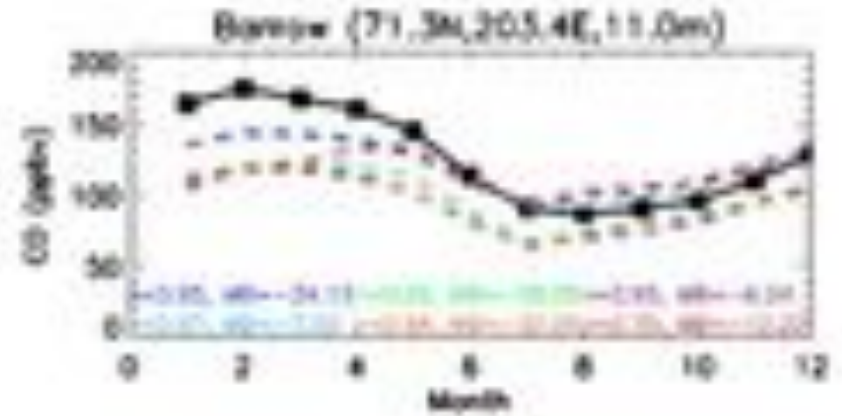
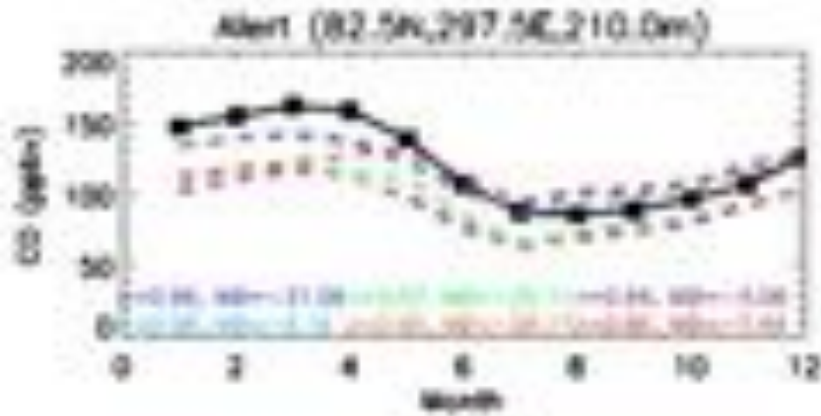


# Preliminary comparison: spring 2008 aircraft data

ARCTAS-A Apr 12,16,17

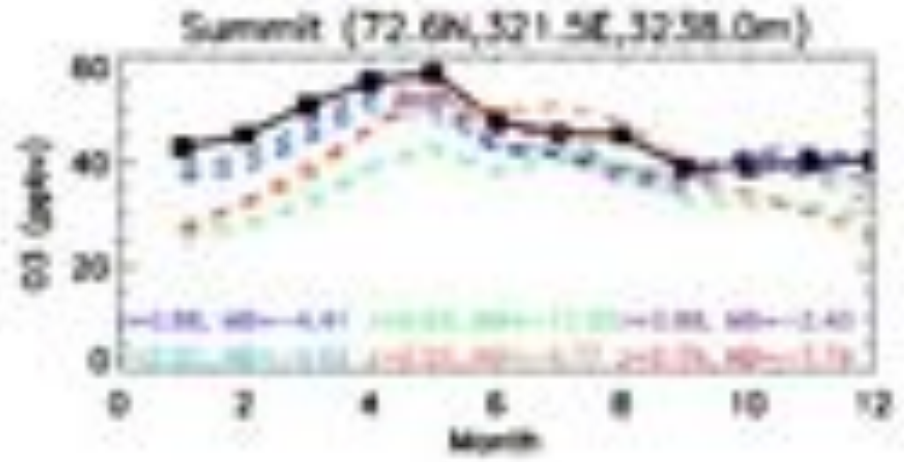
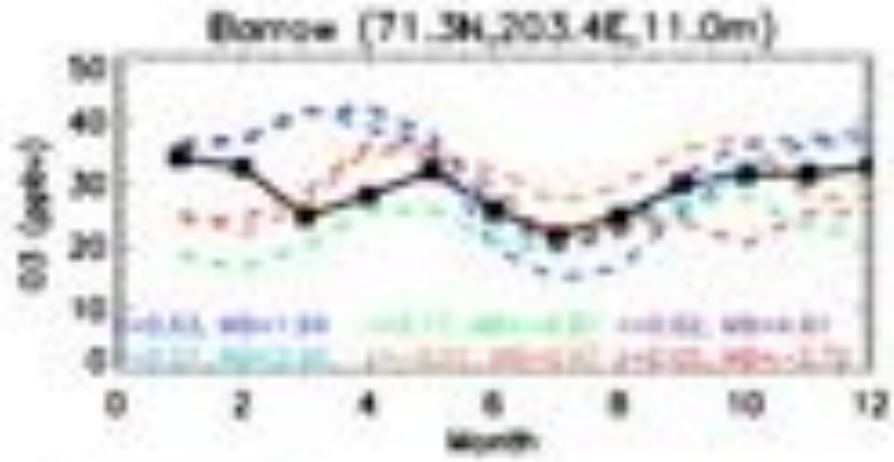


# Surface CO compared to Arctic Station data



S. Monks

# Surface ozone compared to Arctic Station data



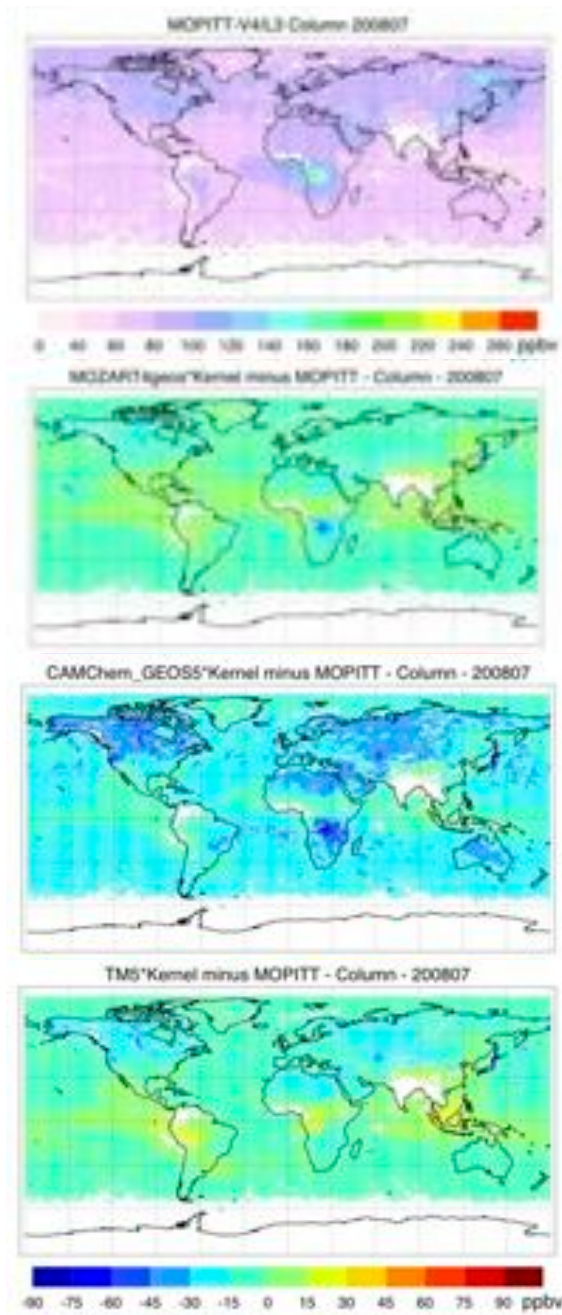
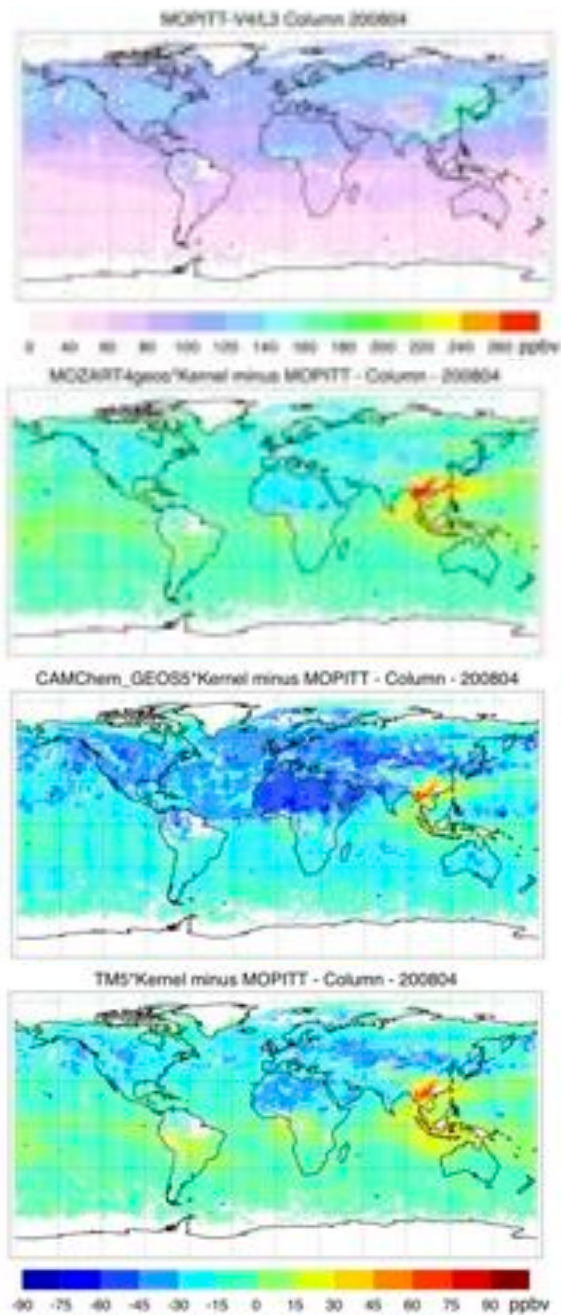
S. Monks



# Comparisons to MOPITT CO

April

July



# POLMIP Summary

- Models are being evaluated in the Arctic using POLARCAT data
- POLMIP will aim to understand the cause of the variability between models in the Arctic and the ability of models to predict the composition of the Arctic troposphere
- The correlation of model-observation differences with air mass origin will be used to identify poorly modeled transport processes
- Models vary in their simulation of ozone precursors in the Arctic
- Differences in chemistry are likely explained by differences in transport, efficiency of  $\text{NO}_y$  export, photolysis, and chemistry