



Estimation of SO_2 and NO_2 emissions from point sources using satellite retrievals

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with contribution from

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Outline

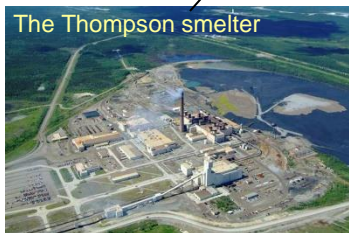
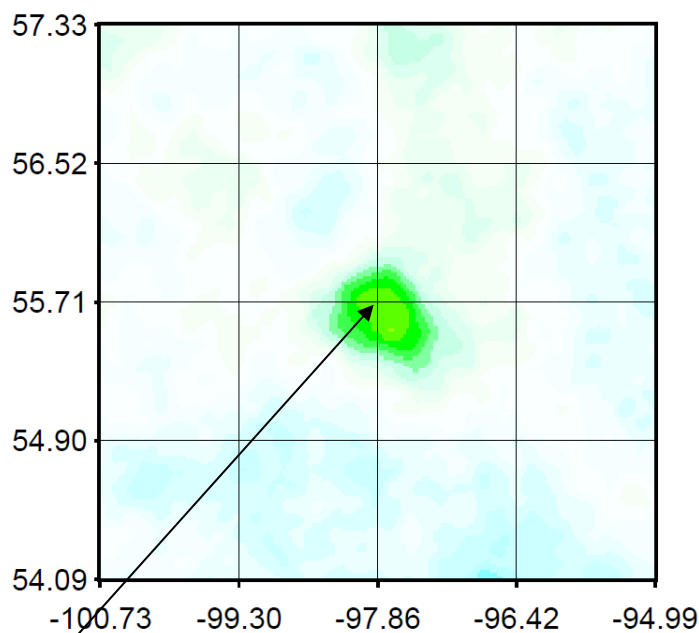
- Satellite data analysis
- Emission estimates from satellite data
- Global SO₂ sources
- Applications for Canada, oil sands
- Validation of satellite data, Pandora spectrometer



Merging satellite data and meteorology

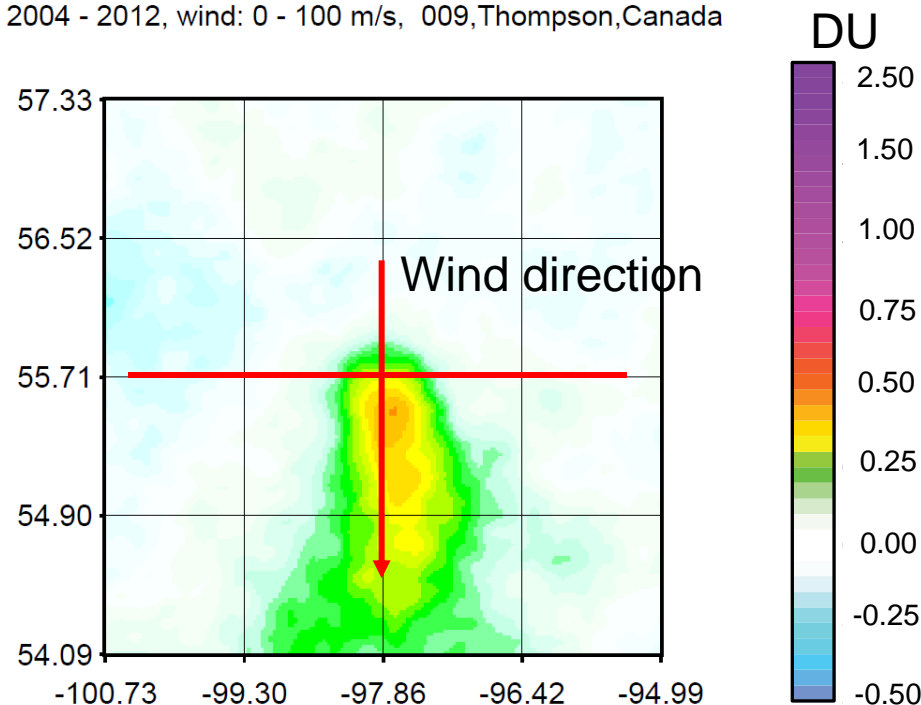
Mean SO_2 from OMI near Thompson, Manitoba (55N, 98 W).

2004 - 2012, wind: 0 - 100 m/s, 009,Thompson,Canada



The same data after rotation of all pixels around the source in a upwind-downwind direction.

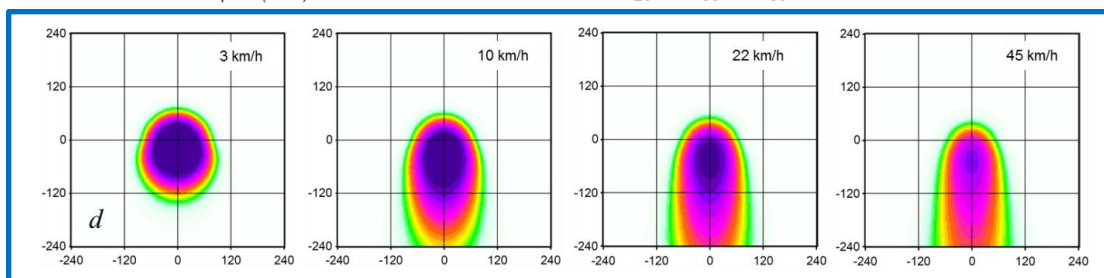
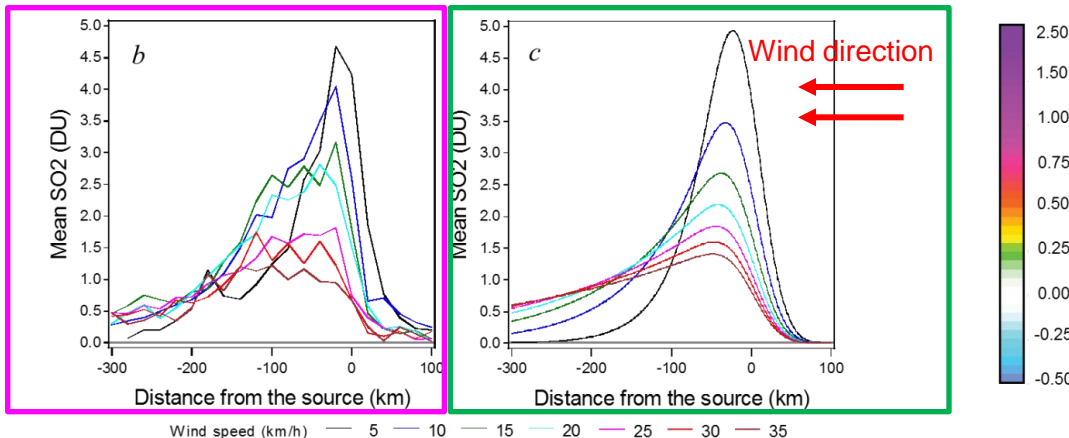
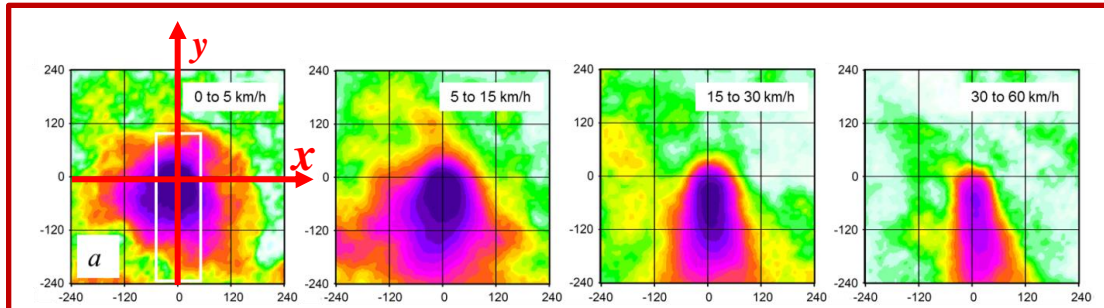
2004 - 2012, wind: 0 - 100 m/s, 009,Thompson,Canada



Downwind decay of pollutants can be studied using a rotation scheme in which the locations of all observation are adjusted so that they have a common wind-direction

The fitting algorithm: $\text{OMI}_{\text{SO}_2} = a f(x,y) g(y,w)$

$f(x, y)$ - Gaussian function, $g(y, w)$ - Exponentially modified Gaussian function;



OMI PCA data:

a. Mean total column SO_2 (in DU) near **Norilsk (69°N, 88°E)** after rotation of all pixels in a upwind-downwind direction for 2005-2013 stratified by the wind speed. The axis show the distance from the source in km.

b. Mean total column SO_2 near Norilsk for different wind speed groups for the area within ± 50 km across the wind direction (the white rectangle in (a)) as a function of the distance from the source (negative for the downwind), after rotation of all pixels in a upwind-downwind direction.

Fitting results:

c. Exponentially modified Gaussian function $g(y, w)$, where y is the distance from the source and w is the wind speed that represents the best fit to Norilsk data on the left.

d. The fitting results for different wind speeds as indicated on the plot.

The fitting algorithm: $\text{OMI}_{\text{SO}_2} = a f(x,y) g(y,w)$

$f(x, y)$ - Gaussian function, $g(y, w)$ - Exponentially modified Gaussian function;

x, y are the coordinates, w is the wind speed

It is assumed that SO_2 concentrations (emitted from a point source) decline with time (t) as $\exp(-\lambda t)$, i.e., with a constant decay rate ($\tau=1/\lambda$)

σ represents the spread due to diffusion

$$f(x, y) = \frac{1}{\sigma_1 \sqrt{2\pi}} \exp\left(-\frac{x^2}{2\sigma_1^2}\right);$$

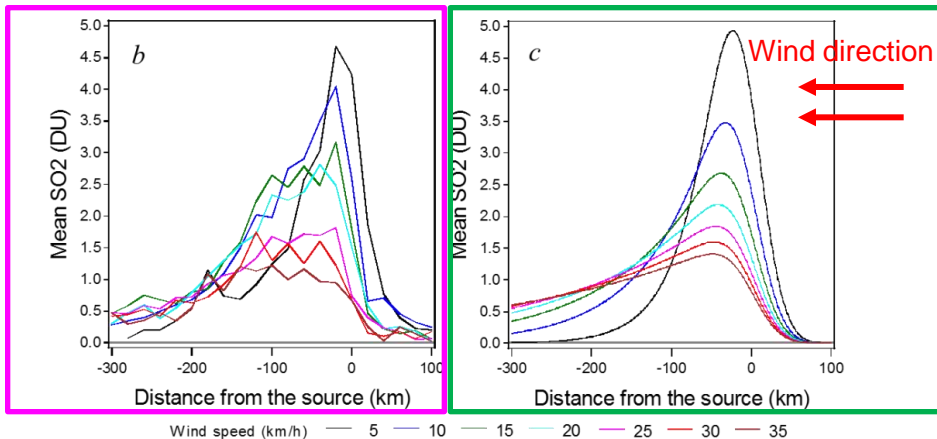
$$g(y, w) = \frac{\lambda_1}{2} \exp\left(\frac{\lambda_1(\lambda_1\sigma^2 + 2y)}{2}\right) \cdot \text{erfc}\left(\frac{\lambda_1\sigma^2 + y}{\sqrt{2}\sigma}\right);$$

$$\sigma_1 = \begin{cases} \sqrt{\sigma^2 - 1.5y}, & y < 0 \\ \sigma, & y \geq 0 \end{cases};$$

$$\lambda_1 = \lambda / w;$$

$$\text{erfc}(x) = \frac{2}{\sqrt{\pi}} \int_x^\infty e^{-t^2} dt$$

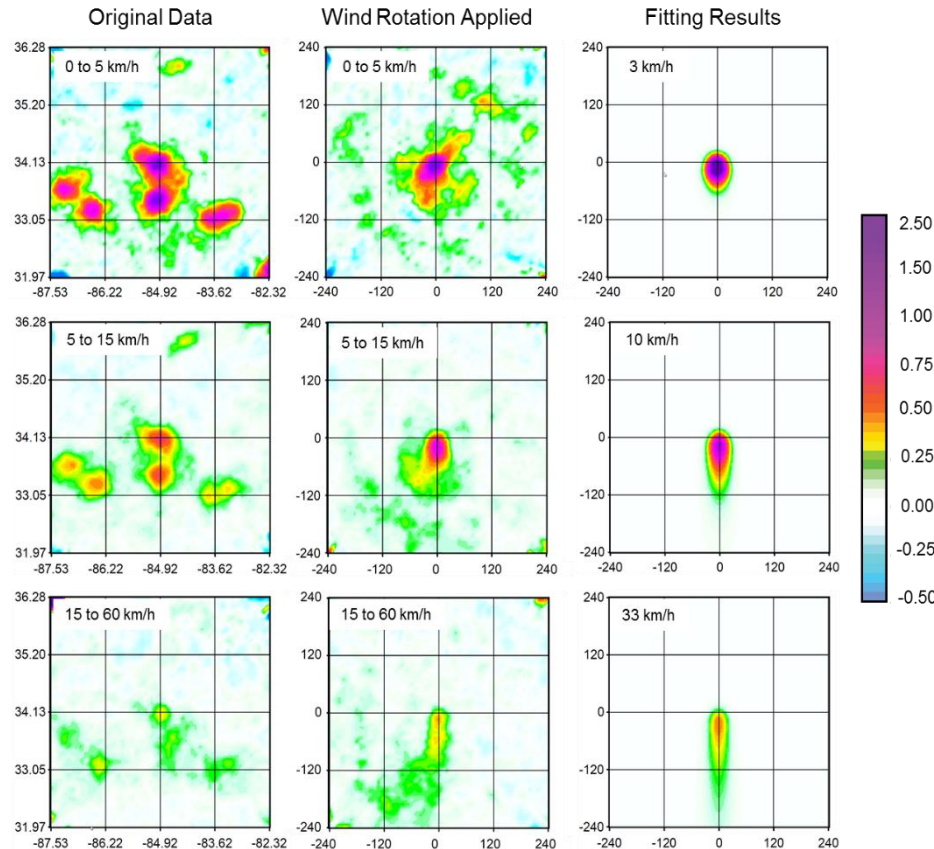
Parameters σ , λ , and a , were estimated from the fit



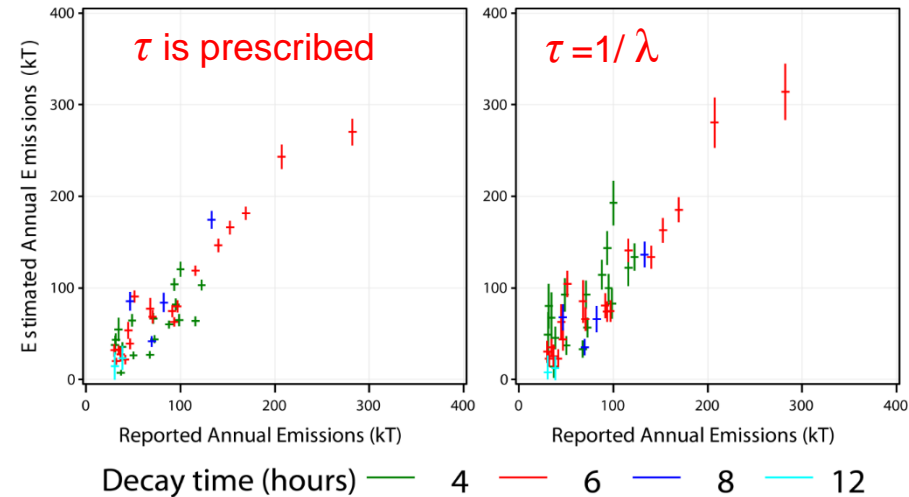
Since $\int_{-\infty}^{\infty} \int_{-\infty}^{\infty} f(x, y) \cdot g(y, w) dx dy = \int_{-\infty}^{\infty} \left(\int_{-\infty}^{\infty} f(x, y) dx \right) \cdot g(y, w) dy = \int_{-\infty}^{\infty} g(y, w) dy = 1$, a is the SO_2 mass

Emissions: $E=a/\tau$, where (1) τ is prescribed or (2) estimated as $1/\lambda$

Estimation of emissions from US major sources using OMI PCA data



The mean column SO_2 values (in DU) for 2005-2007 near the largest US SO_2 source (**Bowen power plant in Georgia**, 170 kt y^{-1} in 2005) located in the center of each plot. The data are stratified by the wind speed. The right column demonstrates the fitting results for different wind speeds as indicated on the plot.



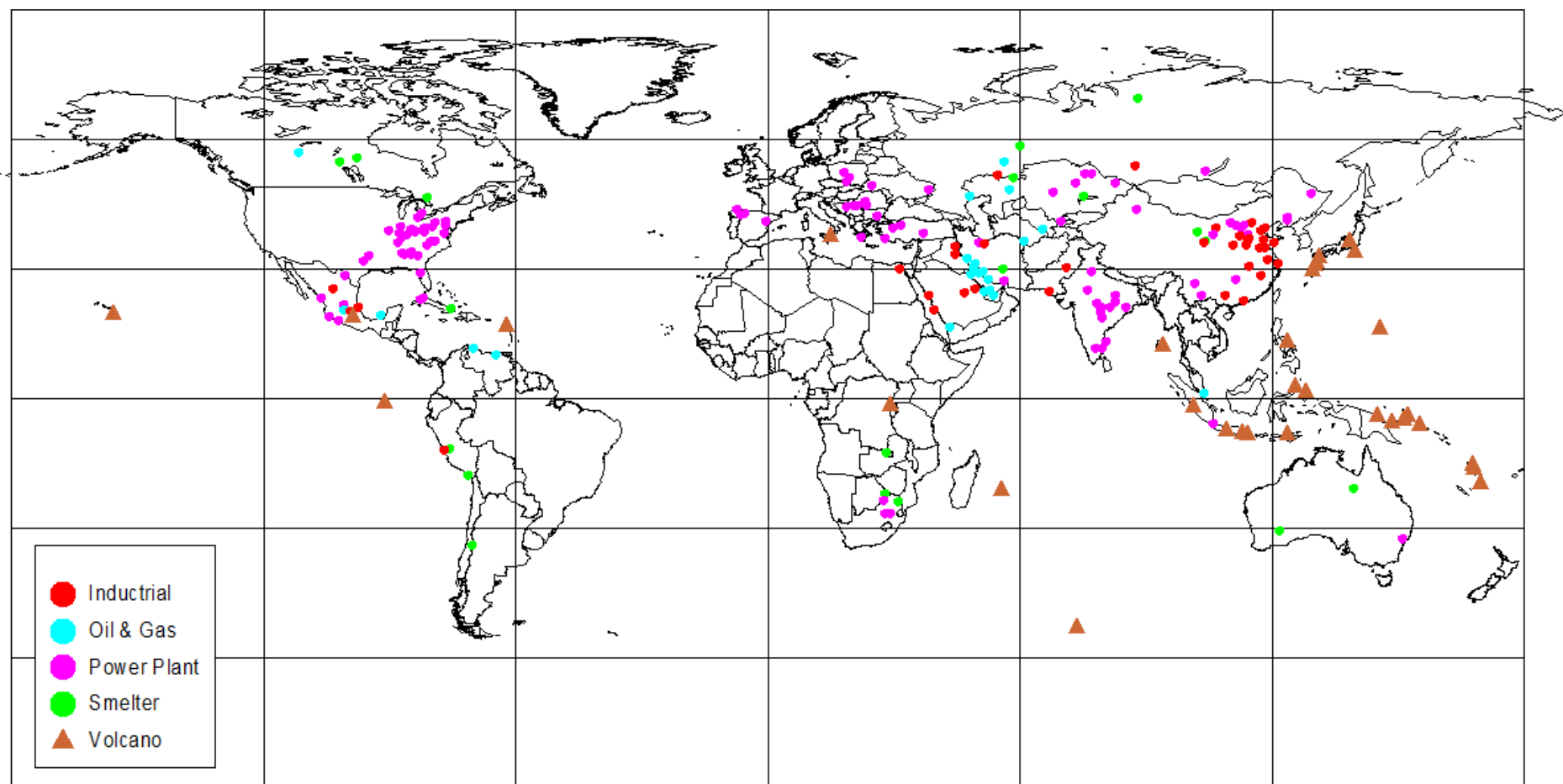
(left) Scatter plots of annual SO_2 emission from the largest US sources in 2005 vs. emissions calculated from OMI data for 2005-2007 assuming a constant the decay time ($\tau=6$ hours) estimated from the best fit of the reported emissions

(right) The same plot but with emissions calculated from OMI data with both parameters estimated from the fit. Emissions are given in kt y^{-1} calculated assuming a constant emission rate.

The error bars represent the one sigma confidence intervals. Different colors indicate estimated decay times ($\tau=1/\lambda$).

OMI SO₂ “catalogue”

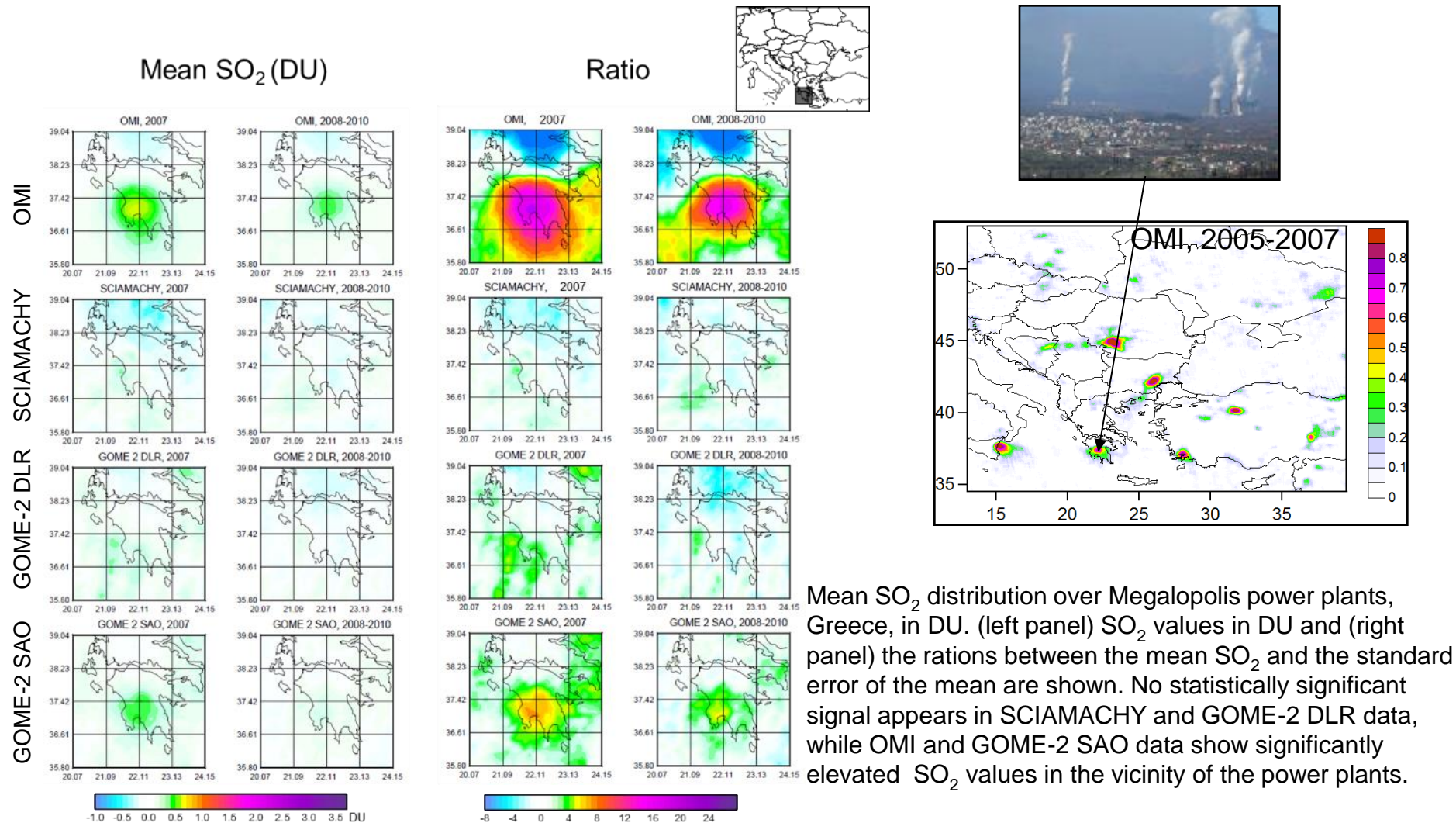
- The detectability levels about 70 kt per year for the old OMI PBL data and ~30 kt per year for OMI PCA data
- >200 sources can be detected
- emissions to be derived, some not currently in inventories



... still work in progress

SCIAMACHY and GOME 2 are less sensitive than OMI.

They can only see ~30 individual sources out of ~200 seen by OMI. Case study: Megalopolis, Greece, one of the largest SO₂ emission sources in Europe (230 kT/y in 2007), in on the border of their sensitivity

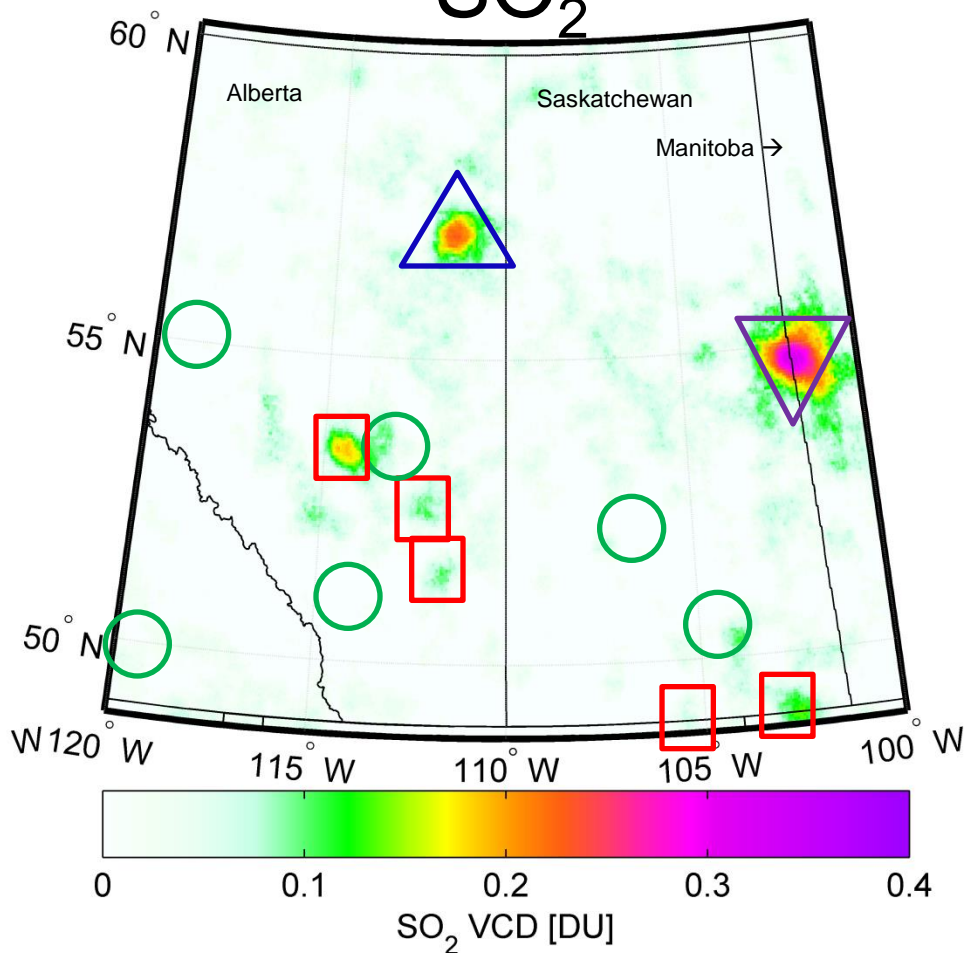
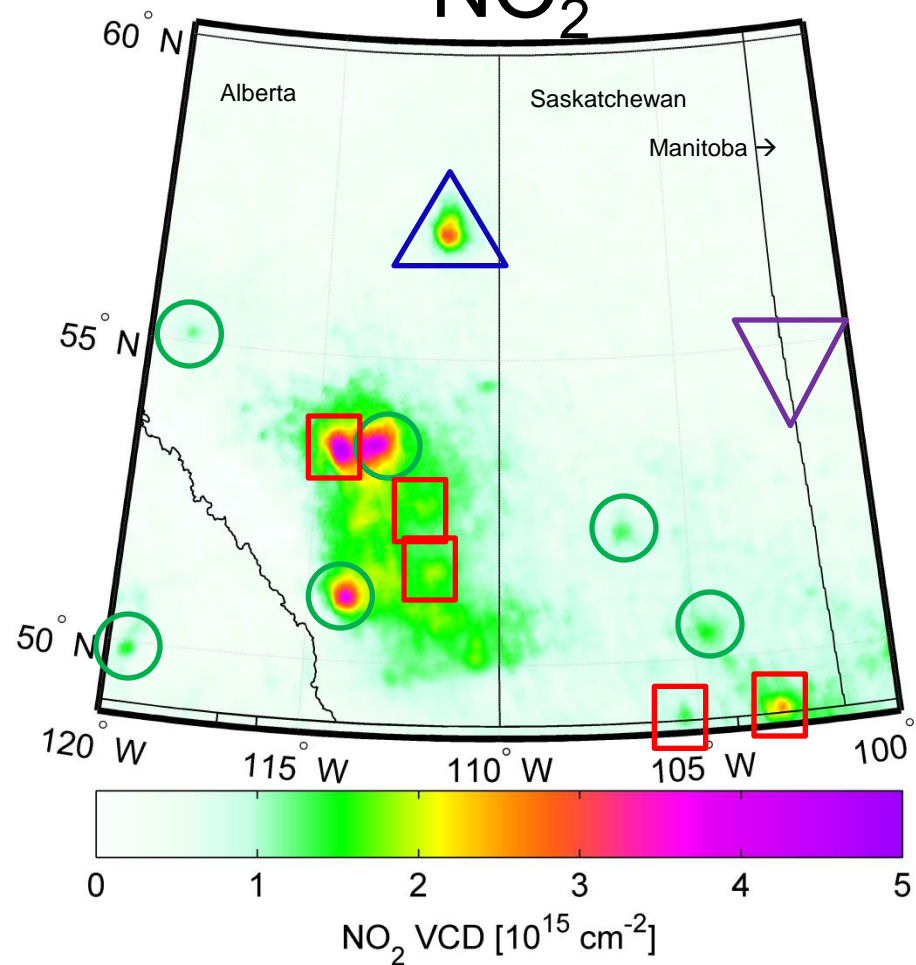


OMI view of Central Canada

NO₂

2005-2007

SO₂



City



Power plant(s)



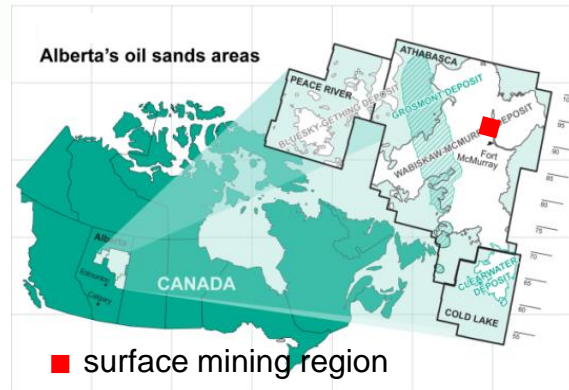
Oil sands



Smelter

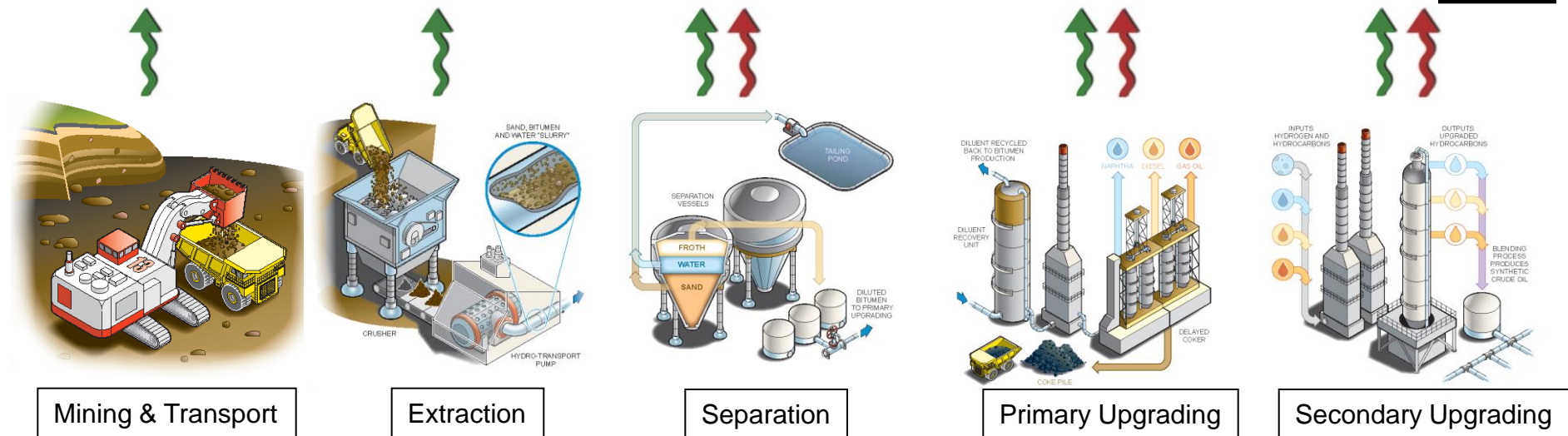
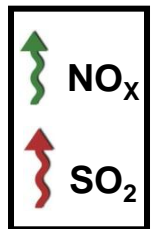


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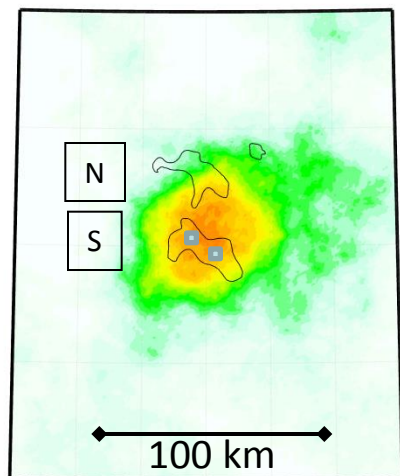
Oil sands monitoring

- Oil sands refers to a type of petroleum deposit in which the oil is very thick and sticky (called “bitumen”) and mixed with sand, water, and clay
- Canada has a proven reserve of ~170 billion barrels with production increase from 0.7 mBPD (2000), 1.8 mBPD (2010), to 3.8 mBPD (2020)
- Surface mining & upgrading processes emit NO_x and SO_2 into the atmosphere

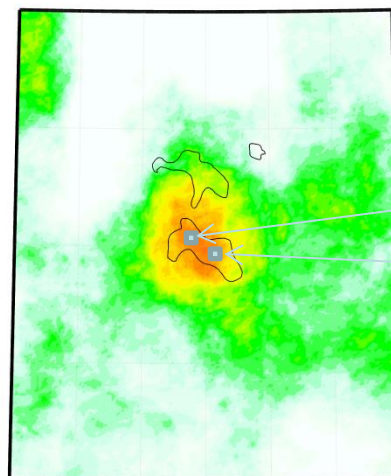




2005-2007



2011-2013



OMI SO₂ over the oil sands

SO₂ emissions due to “upgrading”
(converting bitumen to synthetic
crude)

- Only two significant sources, both in southern, **S**, mining region
- Northern mines pipe bitumen off-site for upgrading

VCD [DU]

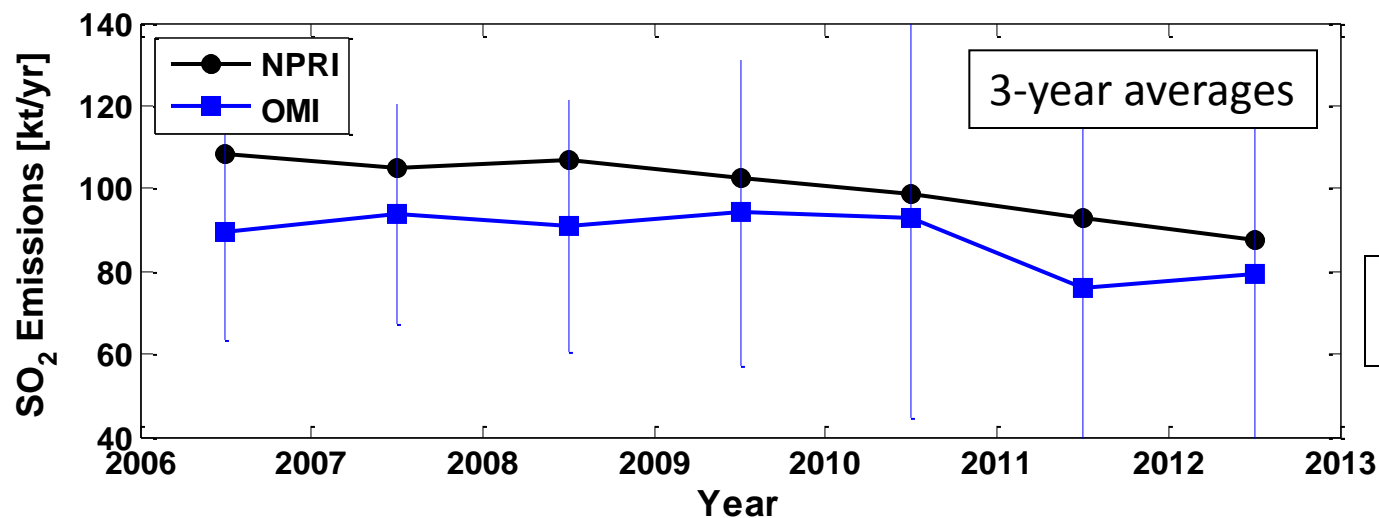
2005 – 2013
average

E(NPRI*) = 100 kt[SO₂]/yr

*NPRI = National Pollutant Release Inventory

E(OMI) = 89±19 kt[SO₂]/yr

τ = 4.3 hours





2005-2007

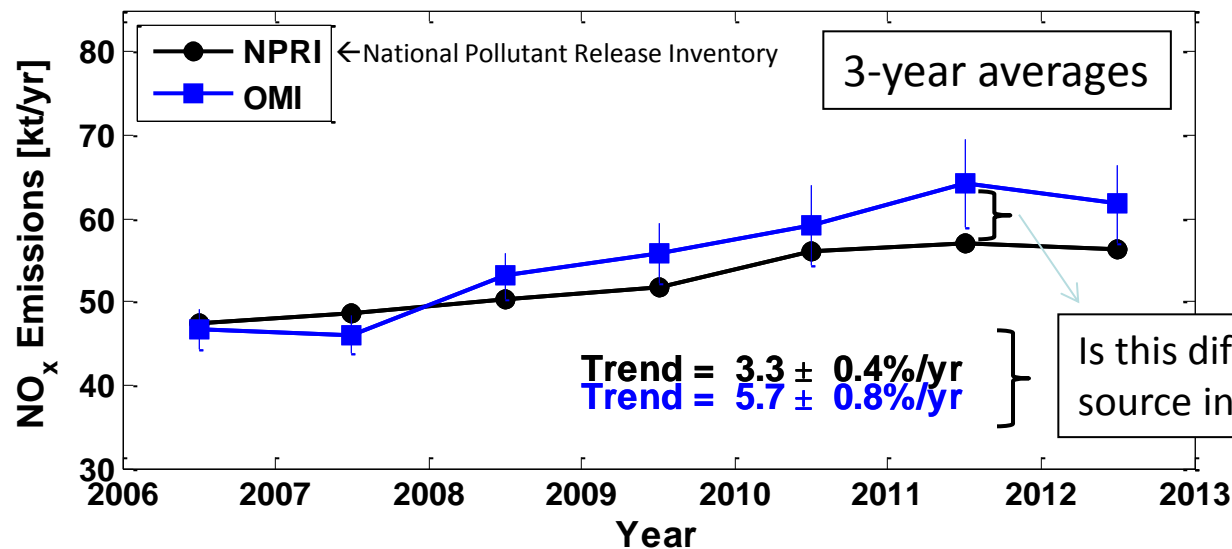
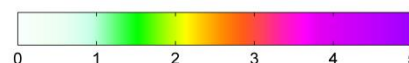
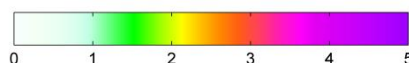
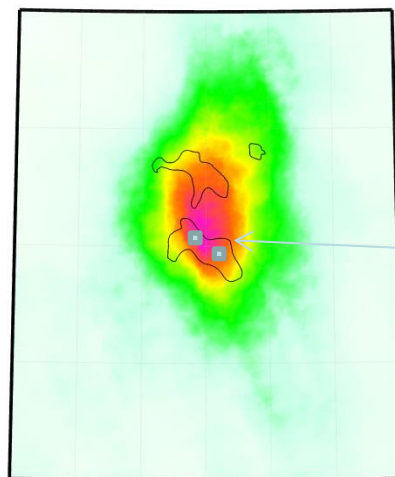
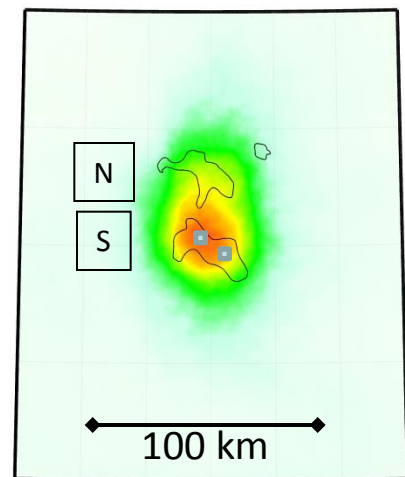
2011-2013

OMI NO₂ over the oil sands

NO₂ emissions due to “upgrading” (~50%) (converting bitumen to synthetic crude) and large vehicles (~50%, more uncertain)

- Two significant point sources **S** (upgraders), both in southern mining region,
- Significant area sources in **N** and **S**
- Change in distribution consistent with expansion into the north

VCD [10^{15} cm^{-2}]



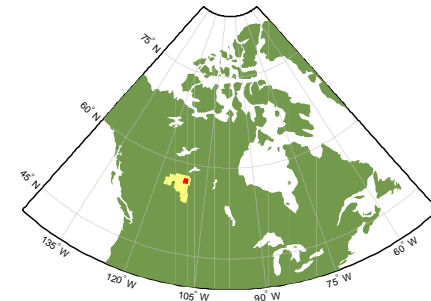
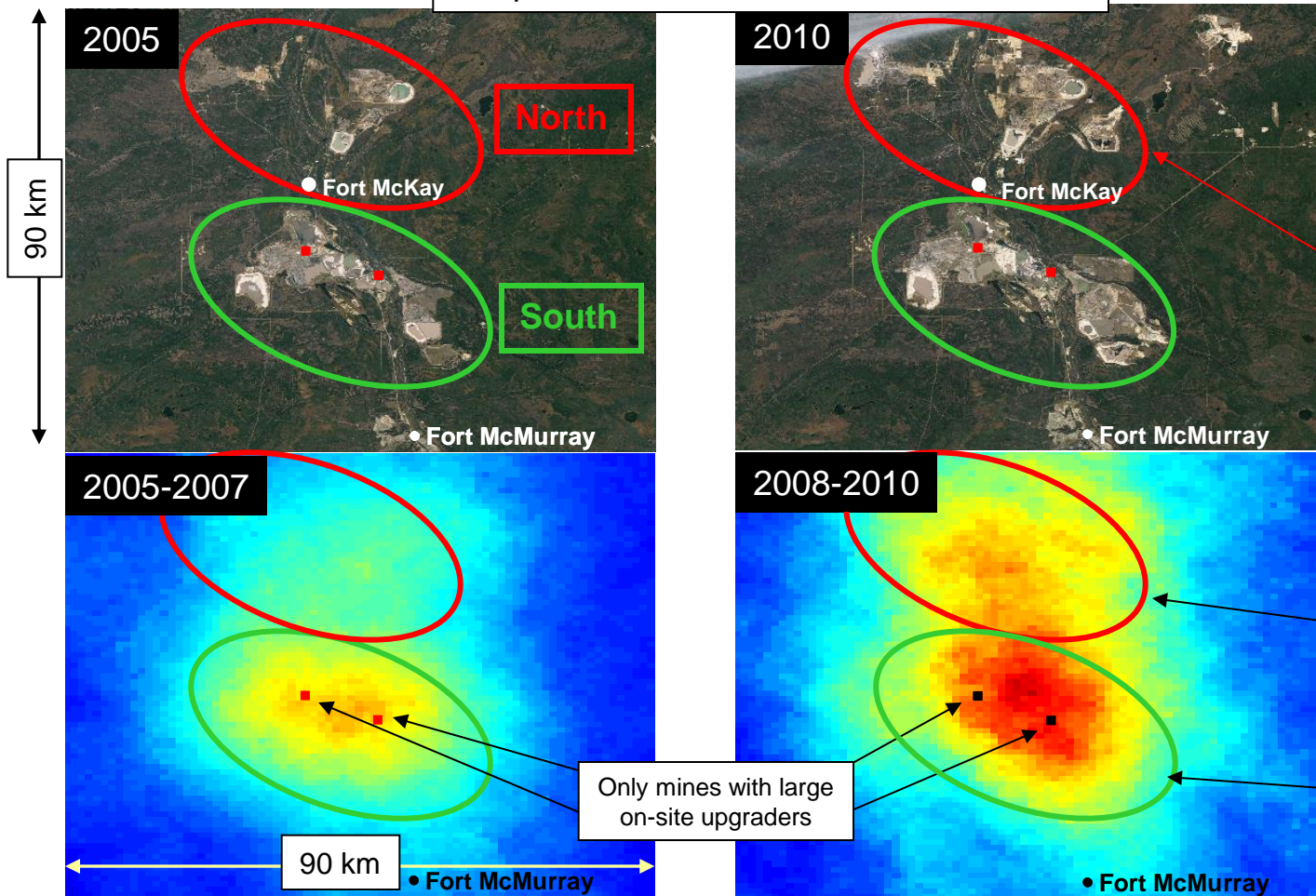
E and τ determined by fitting the downwind decay of NO₂ [similar to Beirle et al., Science, 2011]

NO_x / NO₂ = 1.35 assumed

Is this difference real, from missing source in NPRI (e.g., construction) ?

Estimated NO₂ lifetime is about 2.5 hours

Adapted from McLinden et al., ACP, 2014



Rapid expansion in **North**;
additional mines operational

Landsat Image

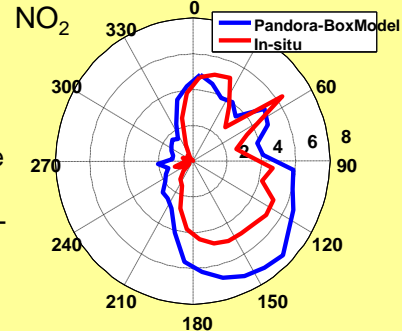
Mean NO_2 from
OMI satellite

NO_2 over **North** likely due to
transportation since no NPRI
(point source) emissions

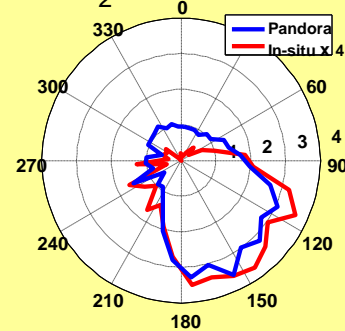
NO_2 over **South** due to some
combination of upgraders
(point sources) and
transportation

Only mines with large
on-site upgraders

Tropospheric

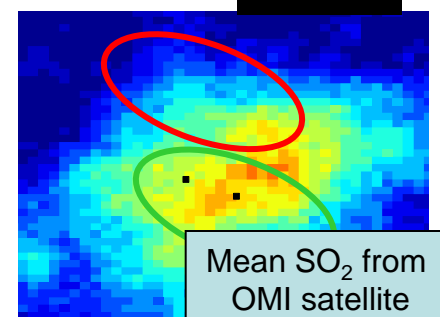


SO_2

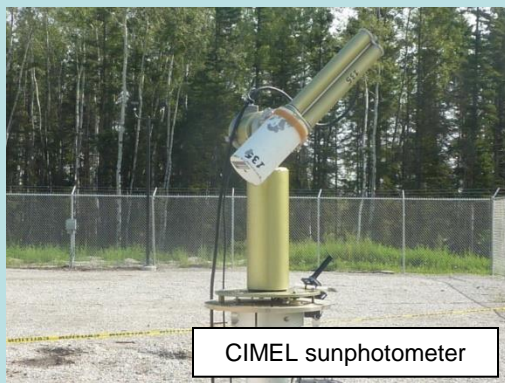


Mean Pandora NO_2 and SO_2 at Fort McKay in 2013 as a function of the wind direction. Both Pandora and in-situ data shows similar patterns: high SO_2 values are associated with south-east winds, while high NO_2 values are observed during south-east and north winds

2005-2010



Mean SO_2 from
OMI satellite



CIMEL sunphotometer

Pandora spectrometer was installed at Fort McKay in August 2013. Pandora operates in DS DOAS mode using spectral range 300 nm-500 nm to retrieve Ozone, SO₂ and NO₂.

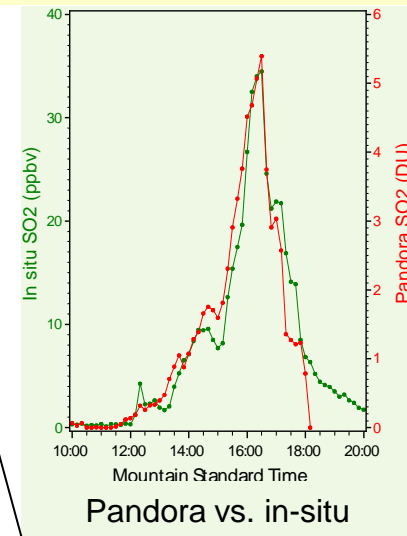
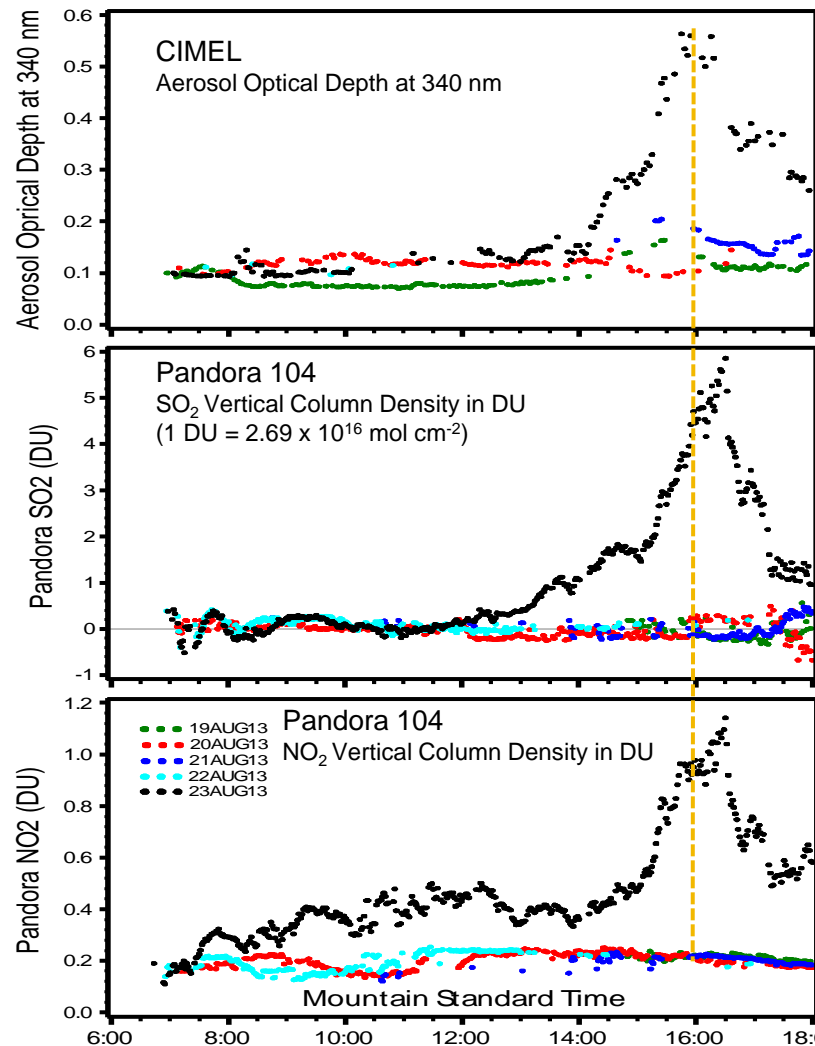


Pandora 104

ment

Remote Sensing Instruments (CIMEL and Pandora), EC site at Fort McKay

Pollution episode at Fort McKay on August 23, 2013



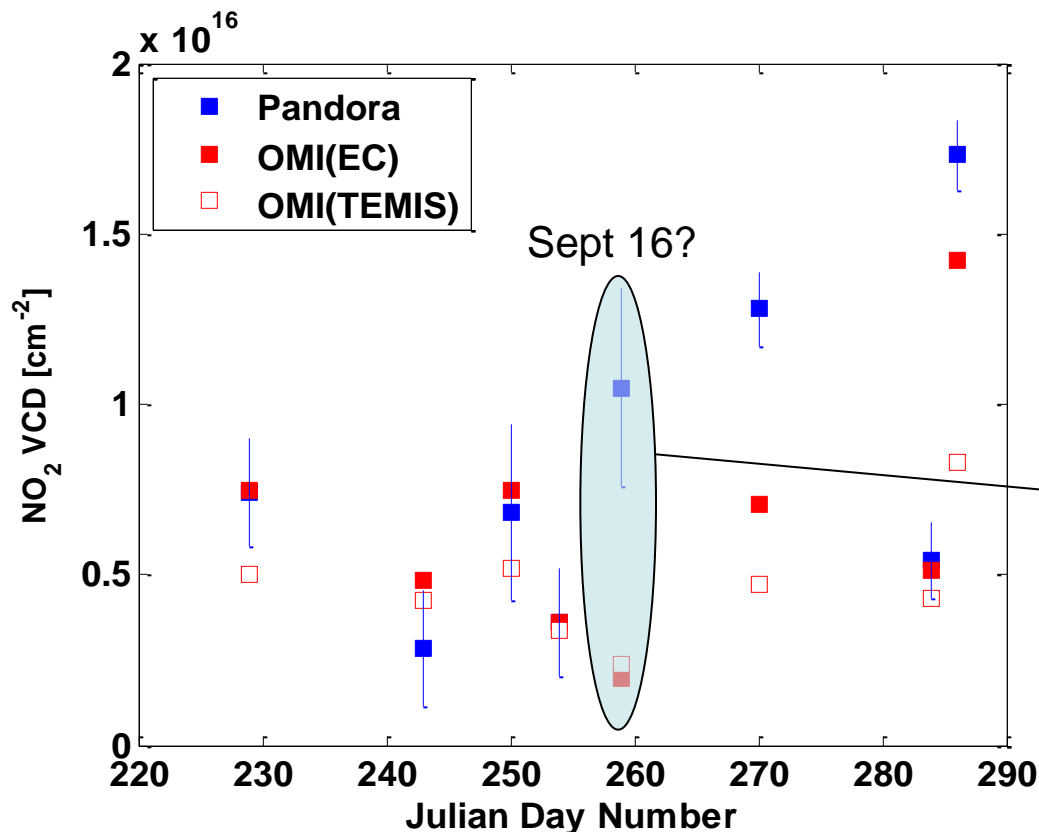
August 23
is in black

Different
colours
represent
different days



Pandora-OMI NO₂ comparison

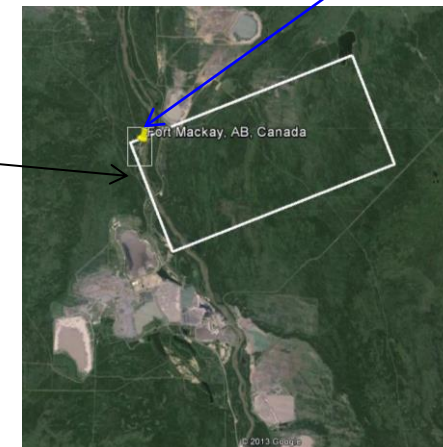
- New EC AMF based on 15 km x 15 km resolution AQ model (McLinden et al., ACP, 2014)
- EC AMF increases NO₂ and SO₂ over the mining area by factors of 2 and 1.4 respectively
- EC AMFs substantially improves agreement as compared to operational algorithms



After filtering and cloud-screening,
only 8 coincidences (in ~60 days)

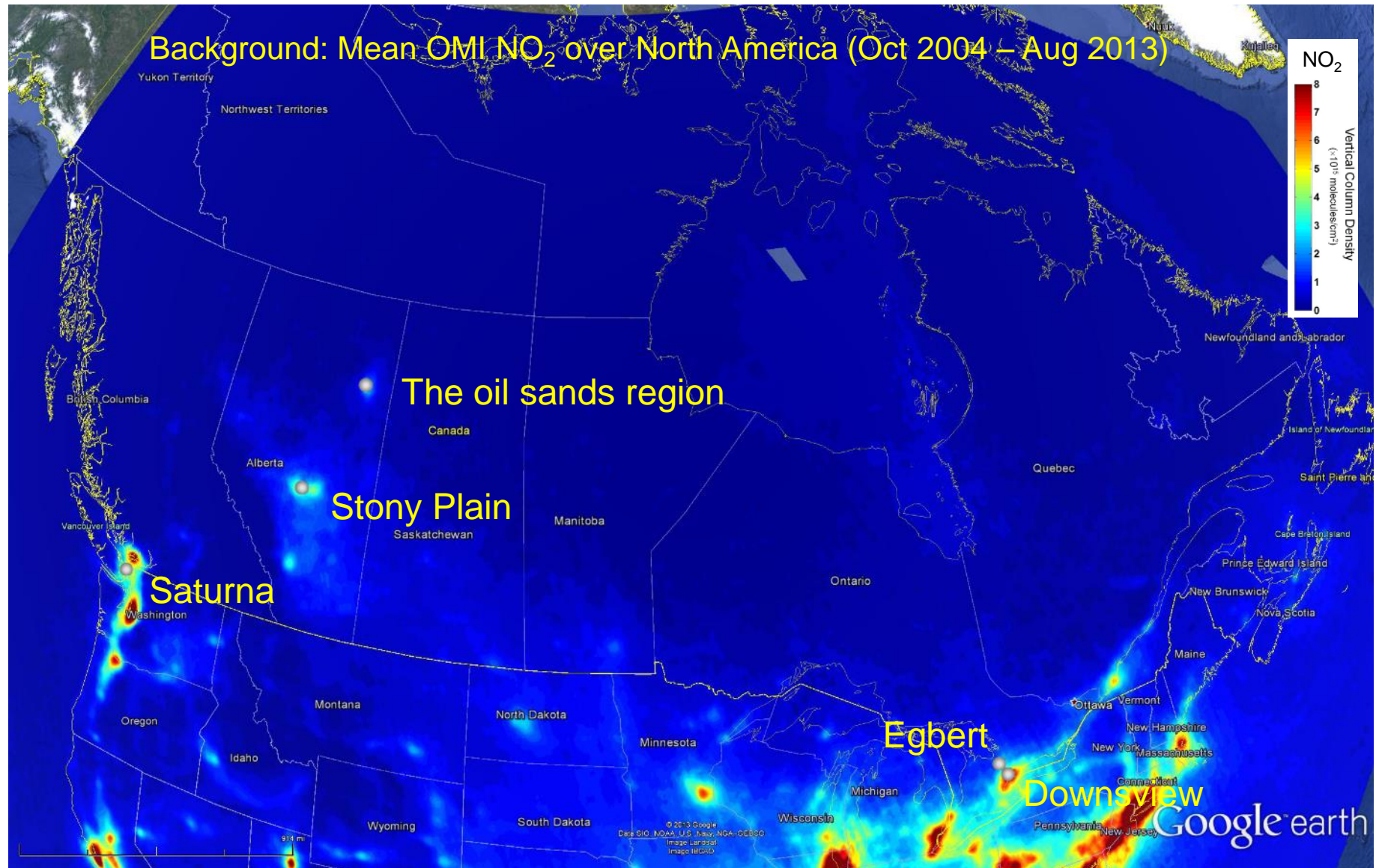
Good quantitative consistency on 5 or
6 days

Pandora location





Possible Canadian ground-based remote-sensing sites for satellite validation





Thanks for your attention!

Fioletov, V. E., C. A. McLinden, N. Krotkov, M. D. Moran, and K. Yang (2011), Estimation of SO₂ emissions using OMI retrievals, *Geophys. Res. Lett.*, 38, L21811, doi:10.1029/2011GL049402.

McLinden, C.A., V. Fioletov, K. F. Boersma, N. Krotkov, C. E. Sioris, J. P. Veefkind, and K. Yang, Air quality over the Canadian oil sands: A first assessment using satellite observations, *Geophys. Res. Lett.*, 39, 4, doi:10.1029/2011GL050273, 2012

Fioletov, V. E., C. A. McLinden, N. Krotkov, K. Yang, D. G. Loyola, P. Valks, N. Theys, M. Van Roozendael, C. R. Nowlan, K. Chance, X. Liu, C. Lee, and R. V. Martin, Application of OMI, SCIAMACHY, and GOME-2 satellite SO₂ retrievals for detection of large emission sources, *J. Geophys. Res.*, 118, doi:10.1002/jgrd.50826, 2013.

McLinden, C. A., Fioletov, V., Boersma, K. F., Kharol, S. K., Krotkov, N., Lamsal, L., Makar, P. A., Martin, R. V., Veefkind, J. P., and Yang, K.: Improved satellite retrievals of NO₂ and SO₂ over the Canadian oil sands and comparisons with surface measurements, *Atmos. Chem. Phys.*, 13, 21609-21664, doi:10.5194/acpd-13-21609-2013, 2014.

Fioletov V.E, C. A. McLinden, N. Krotkov, and C. Li, Lifetimes and emissions of SO₂ from point sources estimated from OMI, *submitted to GRL*, 2014



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