

Field observations of
aerosols and related species
in East Asia
to constrain *model simulations* and
emission inventory

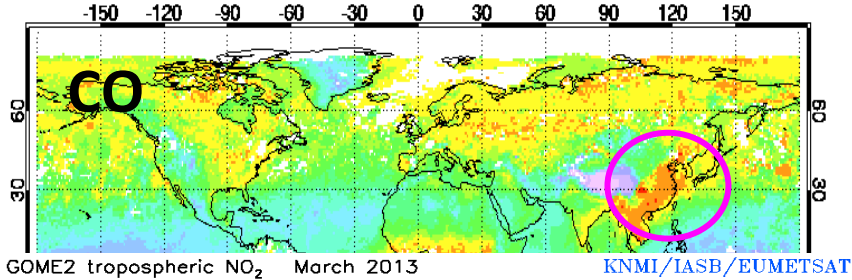


Yugo Kanaya (yugo@jamstec.go.jp)
(RIGC/JAMSTEC)

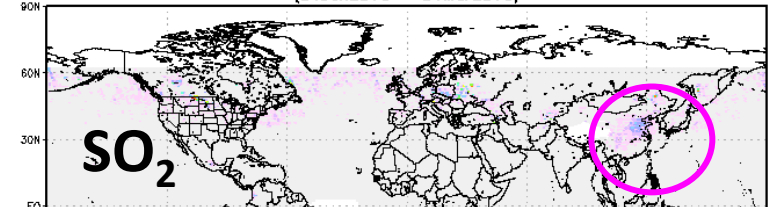
Contributors: F. Taketani, Y. Komazaki, H. Irie, H. Takashima, X.-L. Pan, P. Pochanart, H. Akimoto, H. Tanimoto, S. Inomata, J. Suthawaree, S. Kato, K. Yamaji, Y. Kondo and colleagues, Z. Wang and colleagues, E. Grechko and colleagues

East Asia as hot emission spots

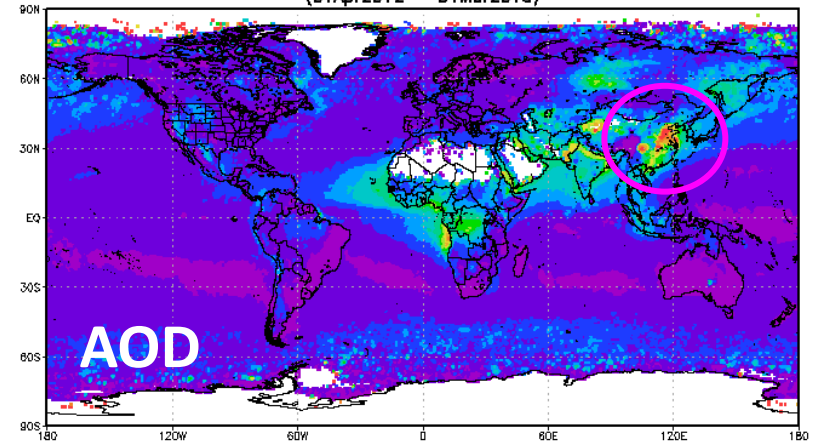
MOPITT CO Column -- 201303



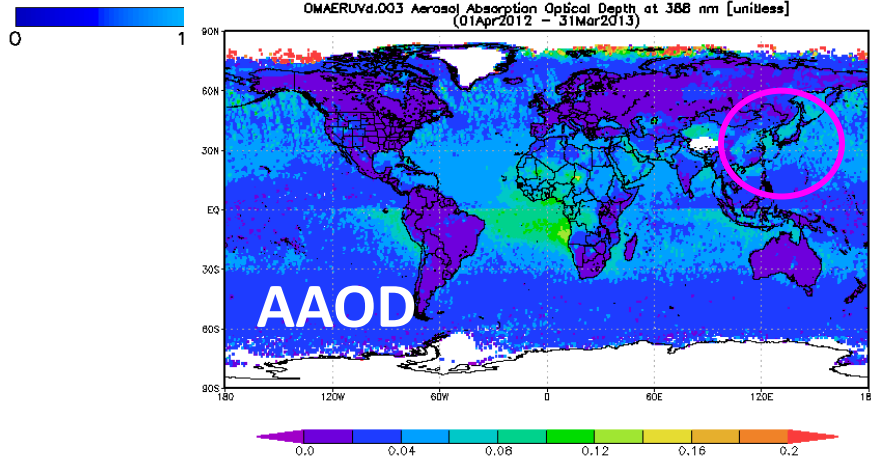
OMSO2e.003 Vertical Column Amount SO₂ (PBL) [DU]
(01Jan2013 - 31Mar2013)



MOD08_D3.051 Aerosol Optical Depth at 550 nm [unitless]
(01Apr2012 - 31Mar2013)



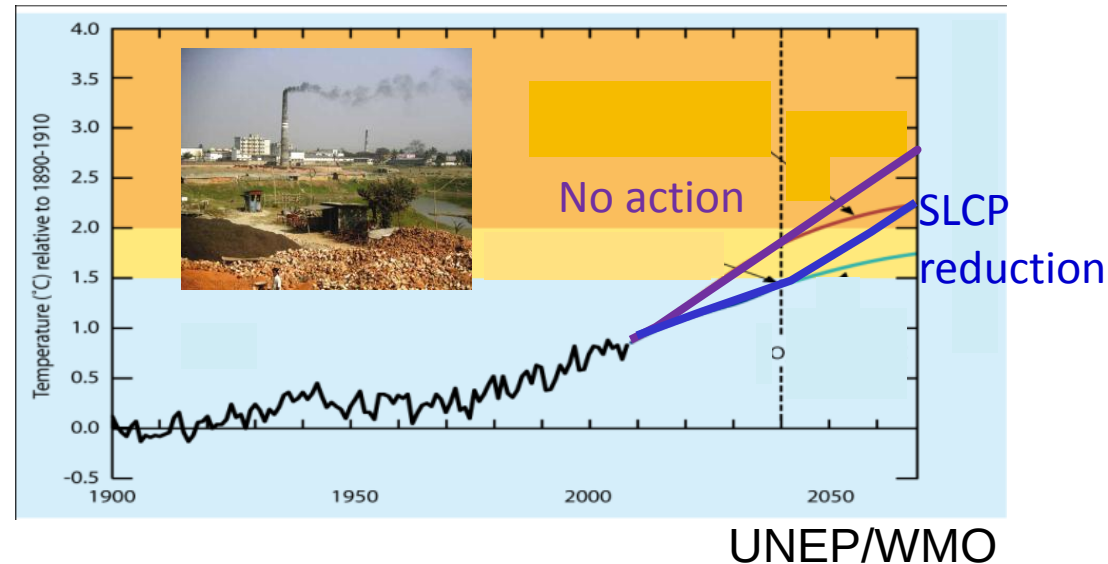
OMAEUVd.003 Aerosol Absorption Optical Depth at 388 nm [unitless]
(01Apr2012 - 31Mar2013)



- still difficult to observe BC and OC from satellite
- demand for field observations

Motivation to study BC + organics in the field

- quantify BC, as SLCP (Short-lived Climate Pollutant) of mitigation target

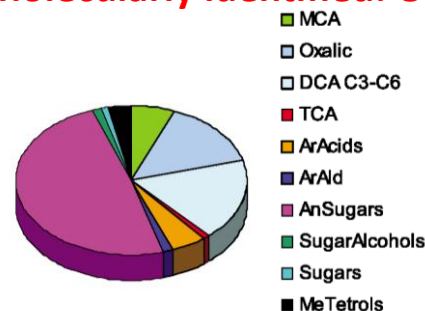


- Major fraction of organics is still unidentified

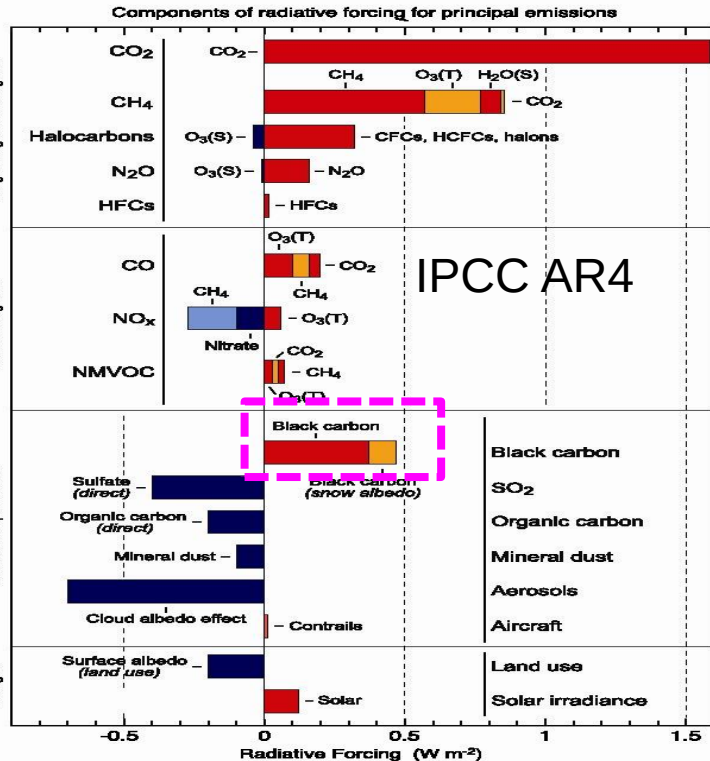
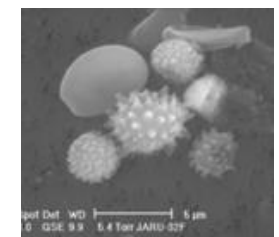
Molecularly identified: Only 10%

90% Unknown: humic-like substance?

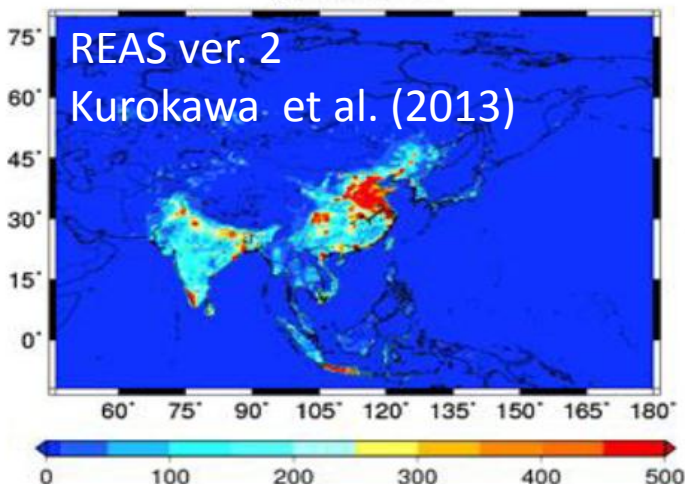
bioaerosols?



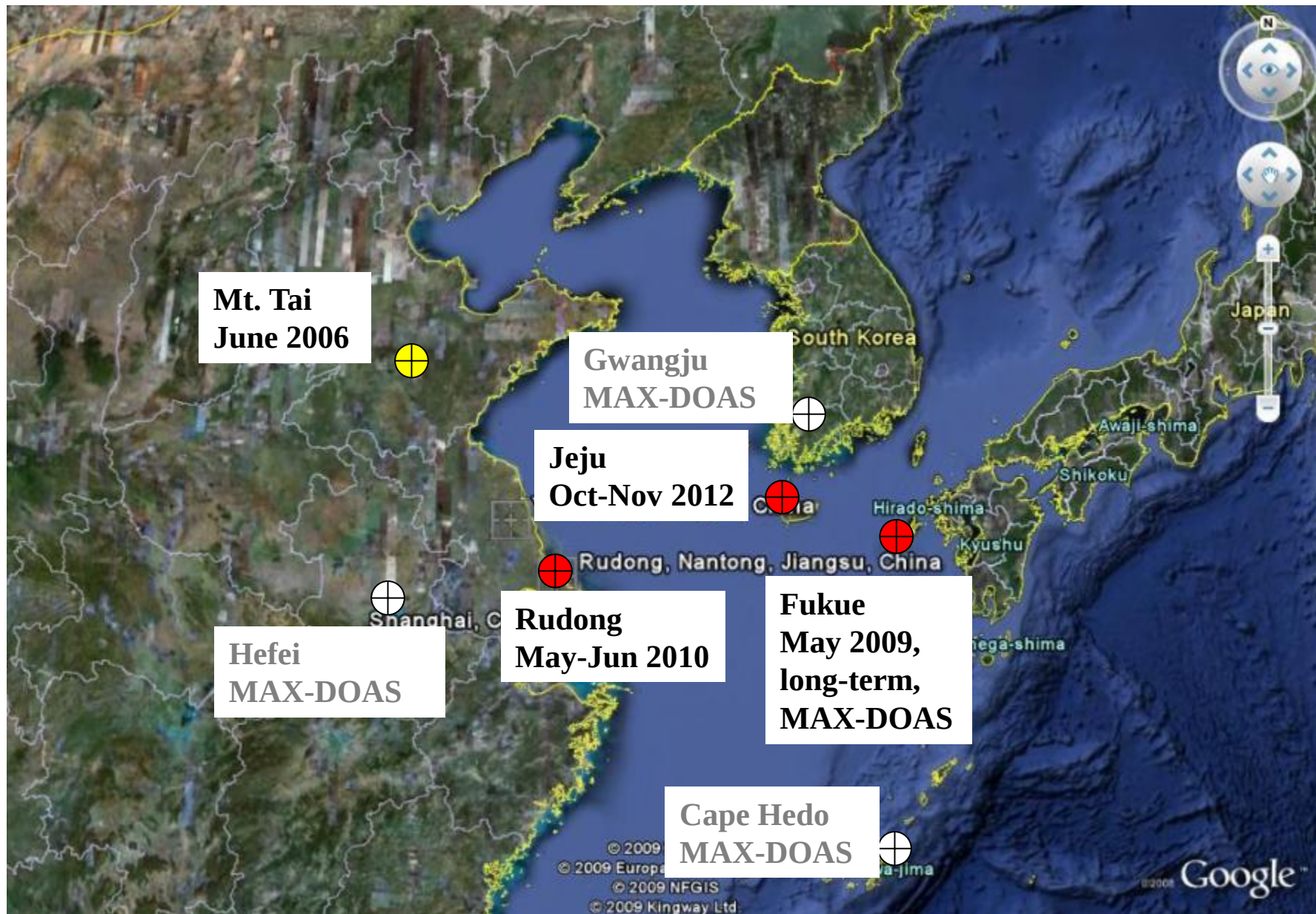
Decesari et al., 2006



(f) BC in 2008



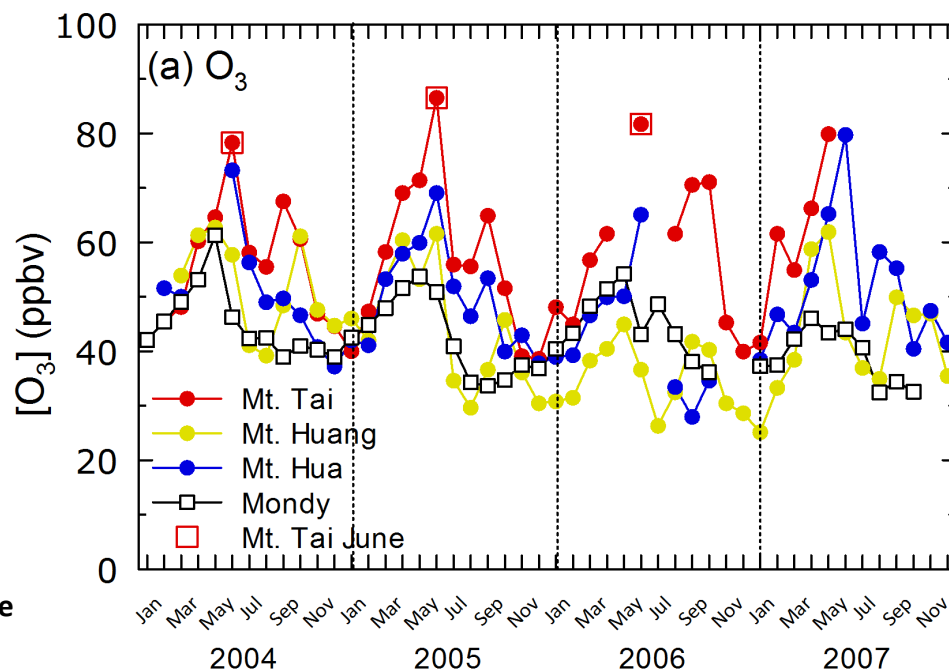
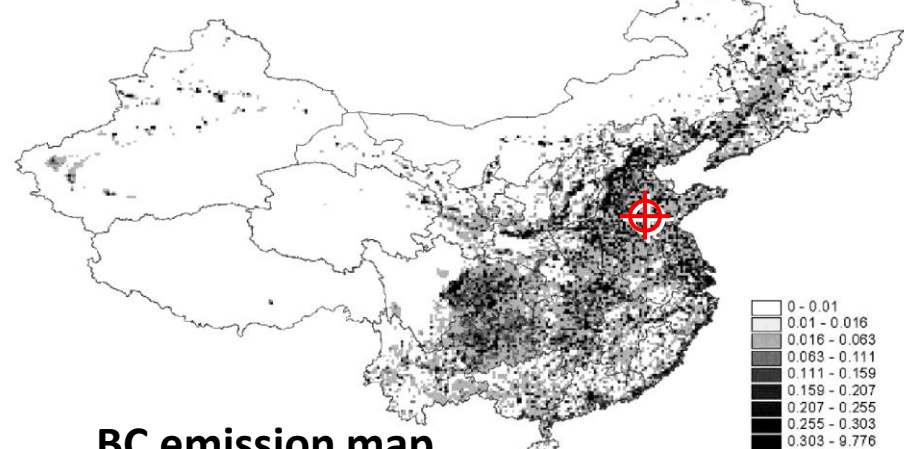
Observational sites



The Mount Tai Experiment 2006

- To measure O_3 , O_3 precursors, aerosol chemical species
- At the top of Mt. Tai, located in the middle of Central East China (CEC), regarded as one of the most significant source regions in the world
- In June, when O_3 and other pollutants show maximum concentrations
- to study O_3 and aerosol chemistry and transport that control regional pollution,
- under close collaboration between observations and model simulations

36.26N, 117.11E 1534 m asl.



Atmospheric
Chemistry
and Physics

ACP special issue: ~20 papers!

The Mount Tai Experiment 2006 (MTX2006): regional ozone photochemistry and aerosol studies in Central East China

Editor(s): F. J. Dentener, S. C. Liu, and L. T. Molina

Overview paper: Kanaya et al., ACP, in press, 2013

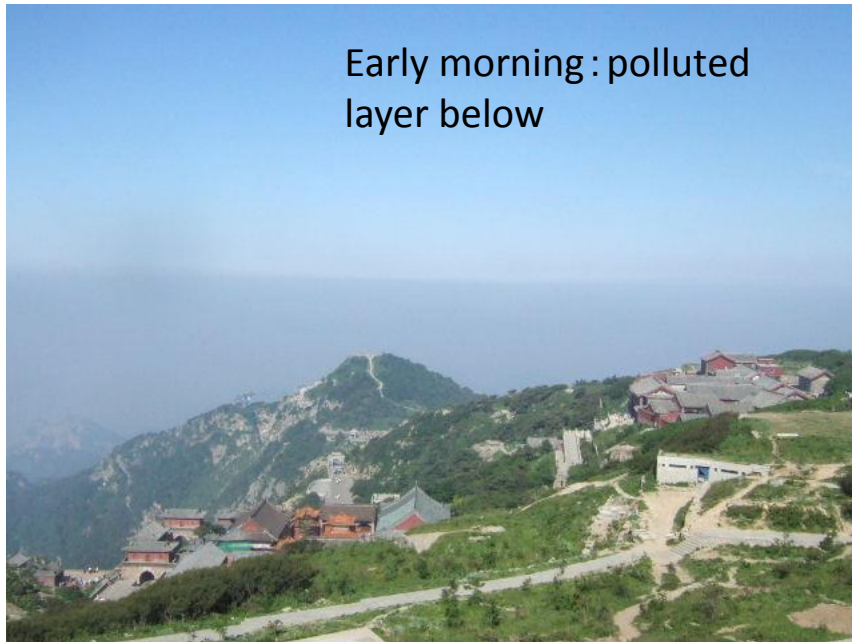
Observed species

Observed species	Method	PI
O ₃ , CO	UV absorption, NDIR	P. Pochanart, JAMSTEC
O ₃	UV absorption	T. Wang, J. Gao, Shandong Univ.
NO/NO _x /NO _y	LED/Molybdenum converter /Chemiluminescence	H. Tanimoto, NIES
NO/NO _x *	Chemiluminescence	T. Wang, J. Gao, Shandong Univ.
NMHCs	Canister/GC-FID/GC-MS	S. Kato, Tokyo Met. Univ.
NMHCs	Canister/GC-MS	Y. Wang, IAP
NMHCs, VOCs	PTR-MS	H. Tanimoto, S. Inomata, NIES
CO ₂	NDIR	Y. Komazaki, JAMSTEC
NO ₂ VCD & AOD from mountain	MAX-DOAS	H. Irie, Y. Kanaya, JAMSTEC
NO ₂ VCD & AOD from foothill	MAX-DOAS	H. Irie, Y. Kanaya, JAMSTEC
Actinic flux & J values	Spectralradiometry	Y. Kanaya, JAMSTEC
T, RH, P, wind	Conventional instruments	Meteorological observatories
Blackcarbon (PM ₁ , PM _{2.5})	Multi-Angle Absorption Photometry	P. Pochanart, JAMSTEC
EC/OC	Thermal-Optical-Transmittance	Y. Komazaki, JAMSTEC
Aerosol absorption	Particle Soot Absorption Photometry	Y. Komazaki, JAMSTEC
Blackcarbon	Aethalometer	T. Wang, J. Gao, Shandong Univ.
Size segregated aerosol chemicals	Cascade impactor/ post analysis	Y. Wang, IAP
Ion analysis of TSP and PM _{2.5}	Sampling/ post analysis	G. Zhuang, Fudan Univ.
Organic gases/aerosols	High volume air sampler, filterpack collections/ post analysis	K. Kawamura, Hokkaido Univ.
PAHs/size segregated aerosols	Sampling/ post analysis	G.Wang, Nanjing/Hokkaido Univ.



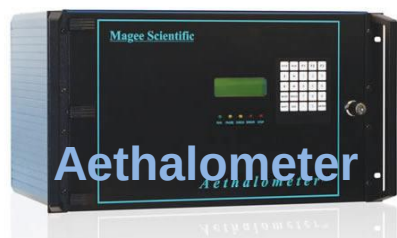
Clean and hazy cases

36.26N, 117.11E 1534 m asl.



Select reliable BC instruments

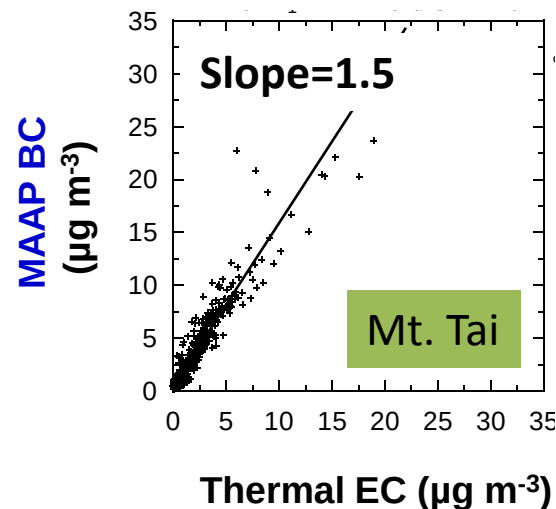
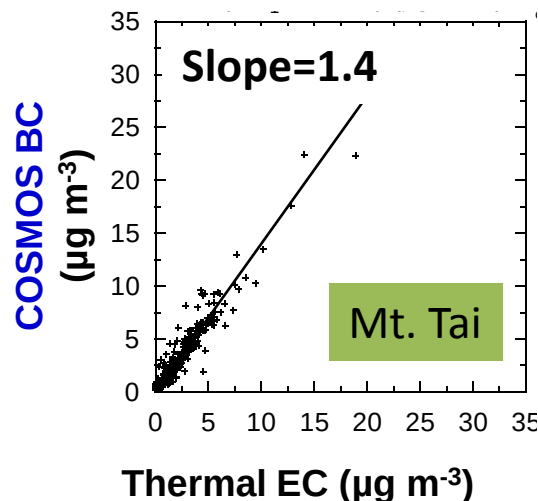
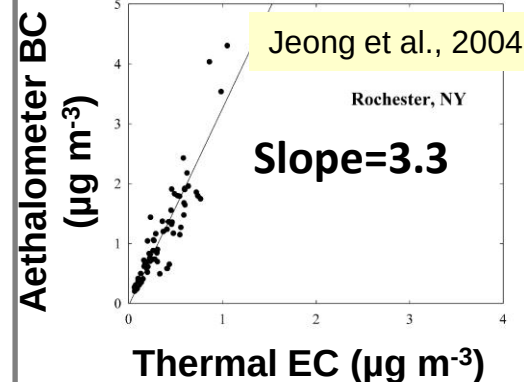
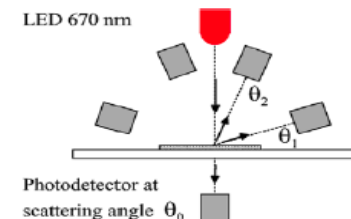
Recommendation: Use COSMOS (heated PSAP) and MAAP, less influenced by scattering particles, and base on ECOC analyzer.



Rochester, NY

Pre-heat to 400C to remove non-refractive scattering particles

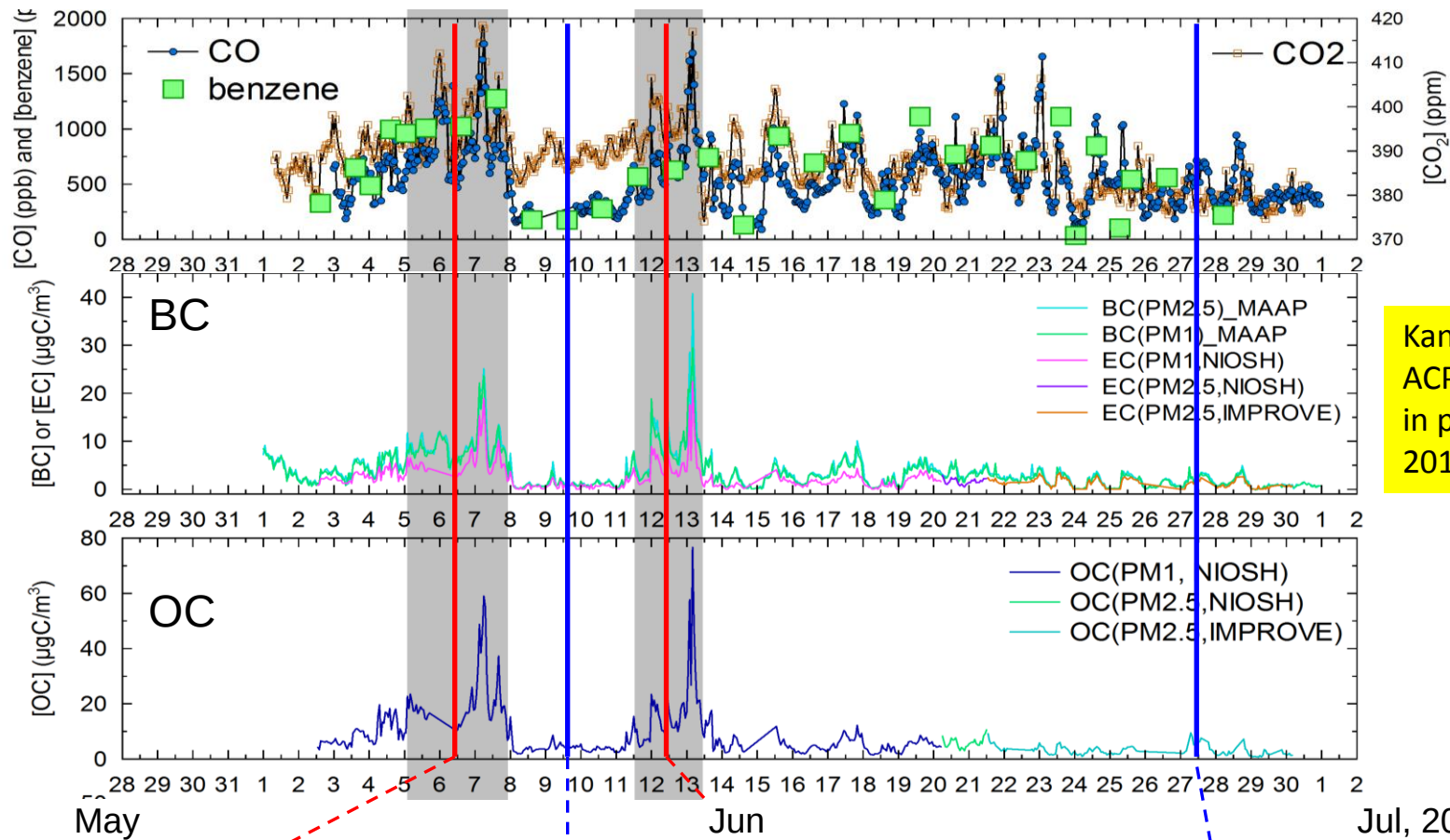
Multi-angle detection to compensate scattering effect



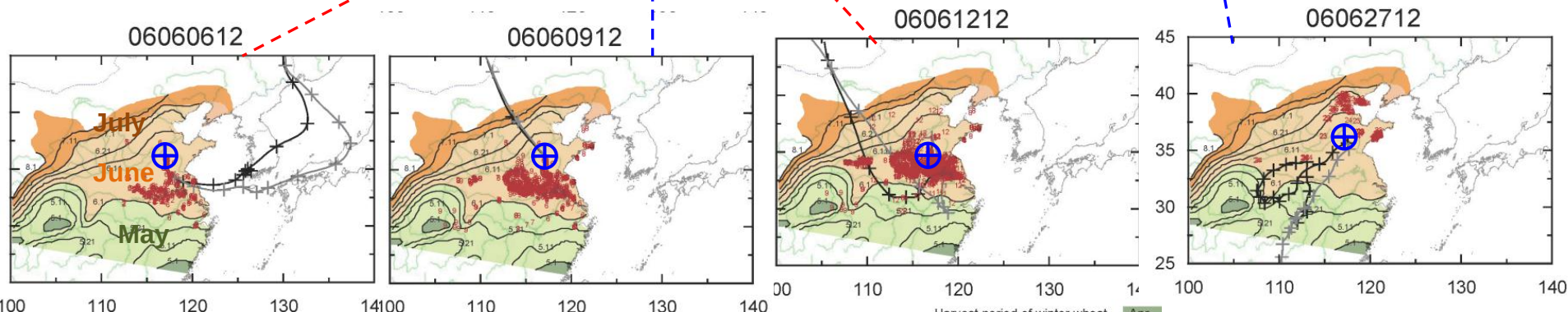
To constrain emission rates better than a factor of 1.5, we need to be able to observe with better accuracy.

Kanaya et al., ACP, 2008

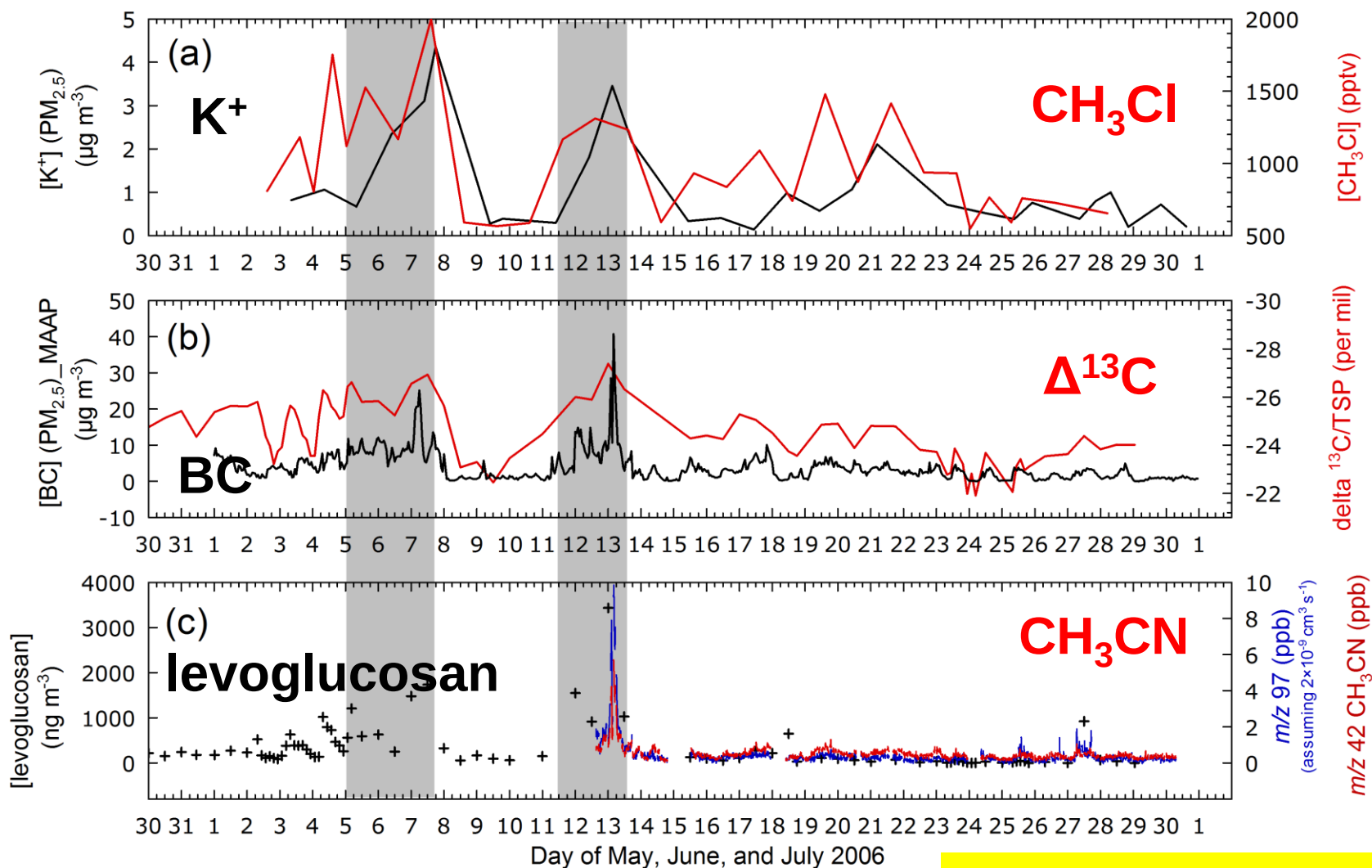
CO, BC, OC influenced by open crop residue burning



Kanaya et al.,
ACP,
in press,
2013



Chemical fingerprints of biomass burning



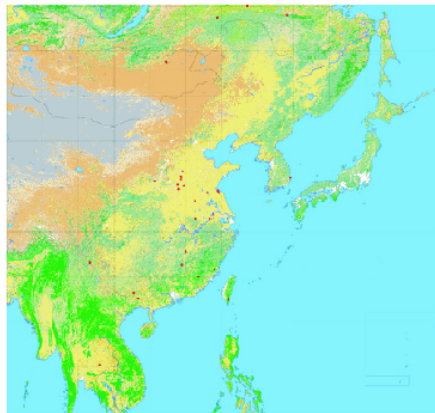
Kanaya et al., in press, ACP 2013

Refinement of OCRB emission using satellite hotspot data

Annual emission: fraction of crop residue burnt in the field [Yan et al. 2006]

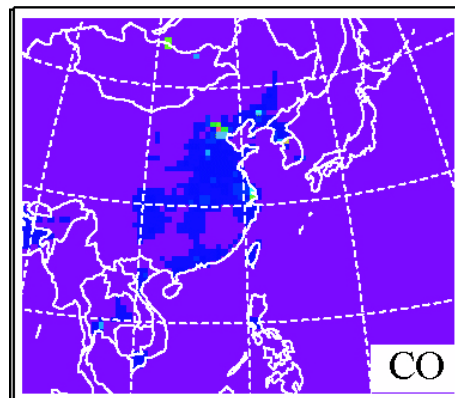
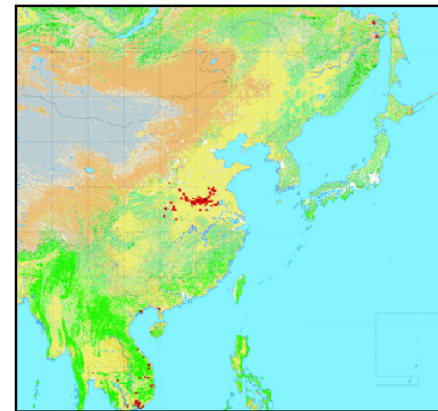
Daily and spatial allocation: based on hotspot distribution (Aqua/Terra MODIS)

28 June 2006



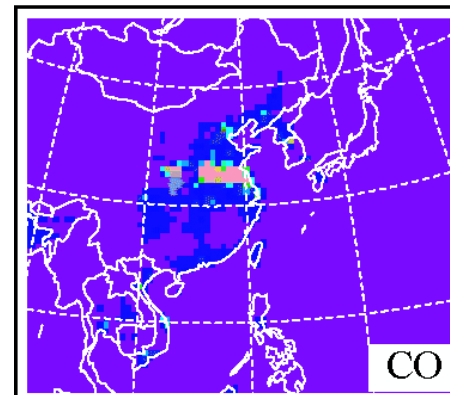
MODIS hotspot

7 June 2006



4000<
mole s⁻¹ grid⁻¹
0

daily CO
emission



Yamaji et al.,
ACP, 2010

Ref: A day with low BB activities

Days with high BB activities

Modeling O₃/aerosol with revised emission inventory

CMAQ/RAMS regional model

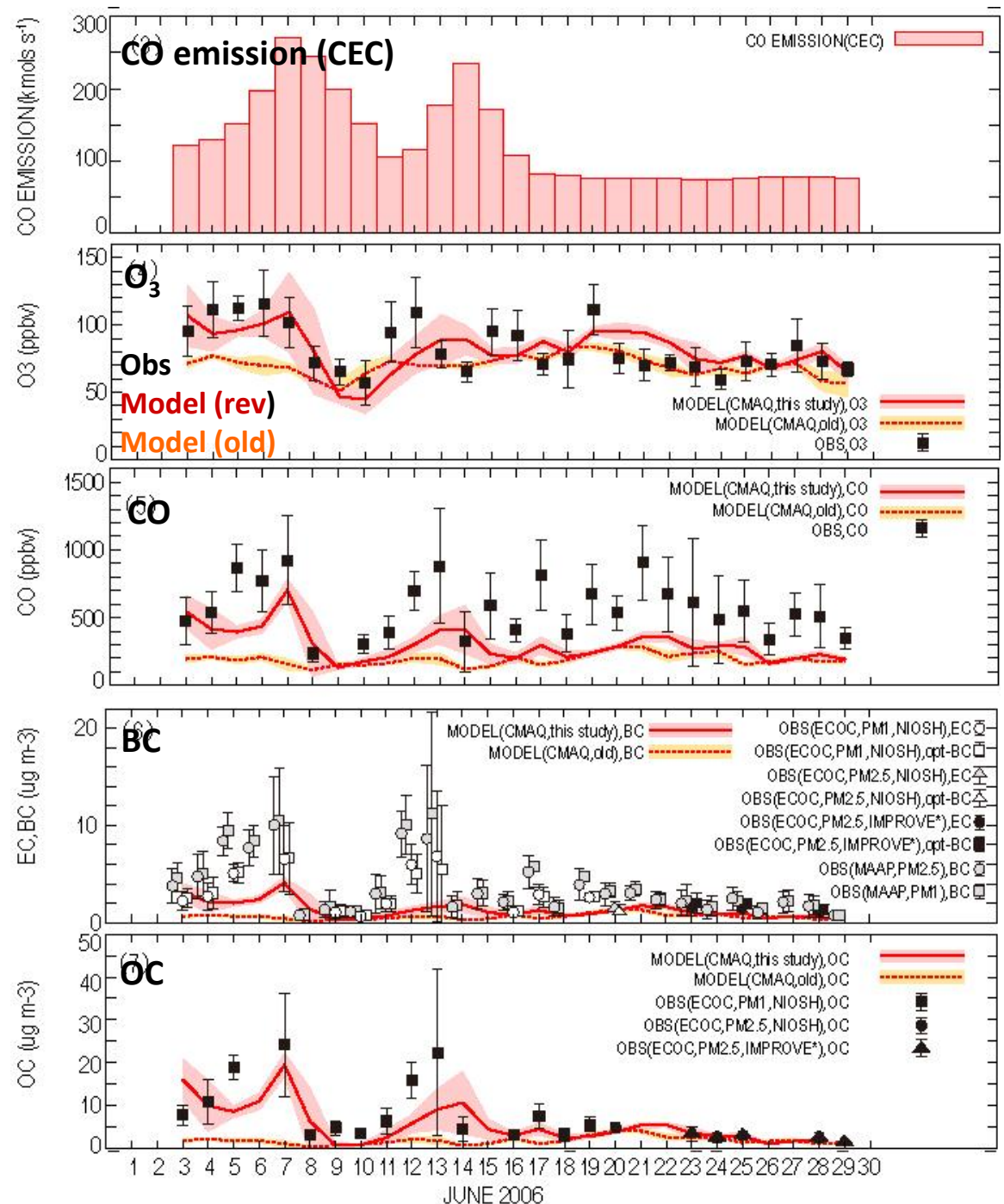
Resolution: 80x80km²

Boundary: Global CTM (CHASER)

Gas chem: SAPRC99

Aerosol: AERO3 (SORGAM, ISSOROPIA, RPM)

OCRB emissions contributed
6% of O₃,
20% of CO,
43% of BC,
and 53% of OC
concentrations over CEC for
the whole month of June



Rudong field campaign 2010

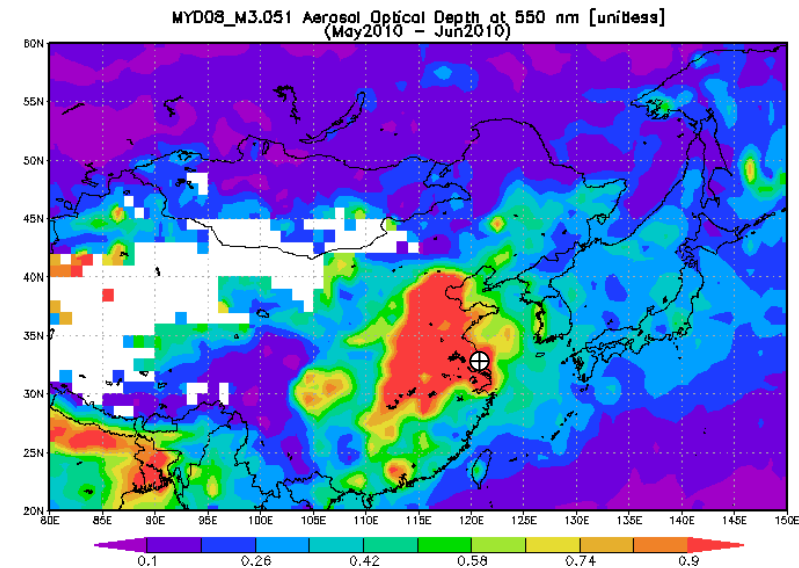
- Observe O_3 /aerosols and their precursors
- in May-June 2010, in seasons with high O_3 /aerosols,
- to **closely observe biomass burning**



100 km north of Shanghai

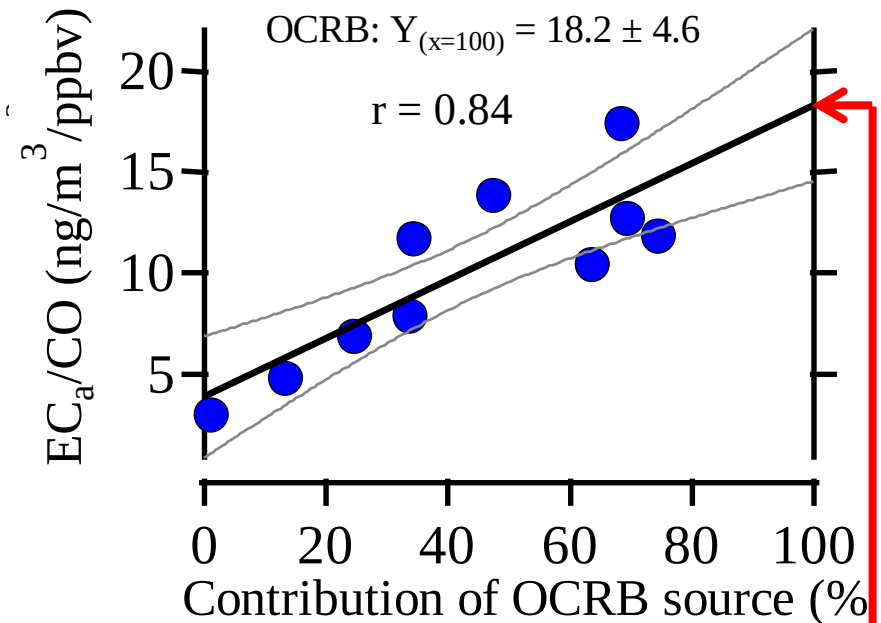
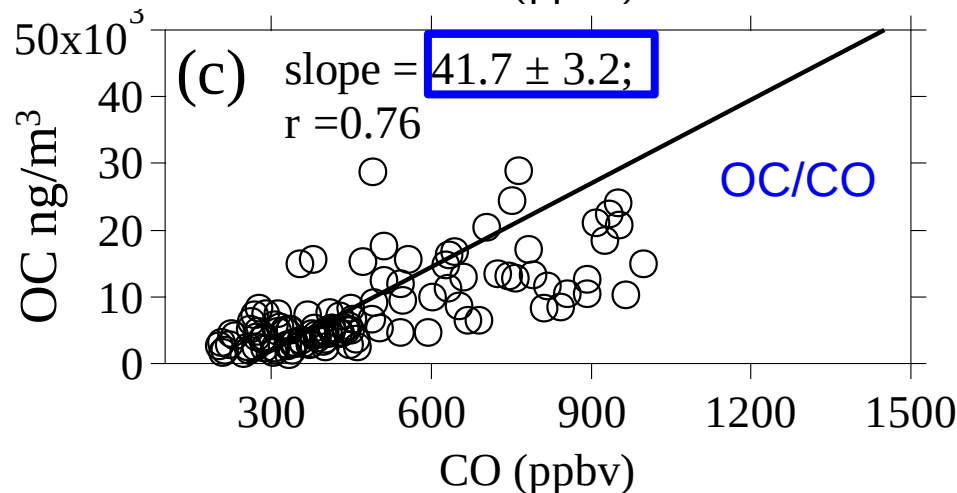
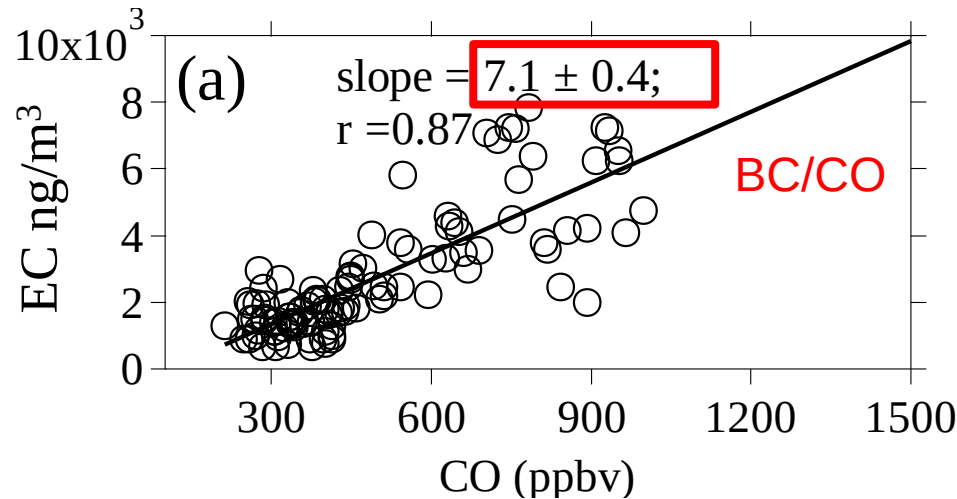


crop residue



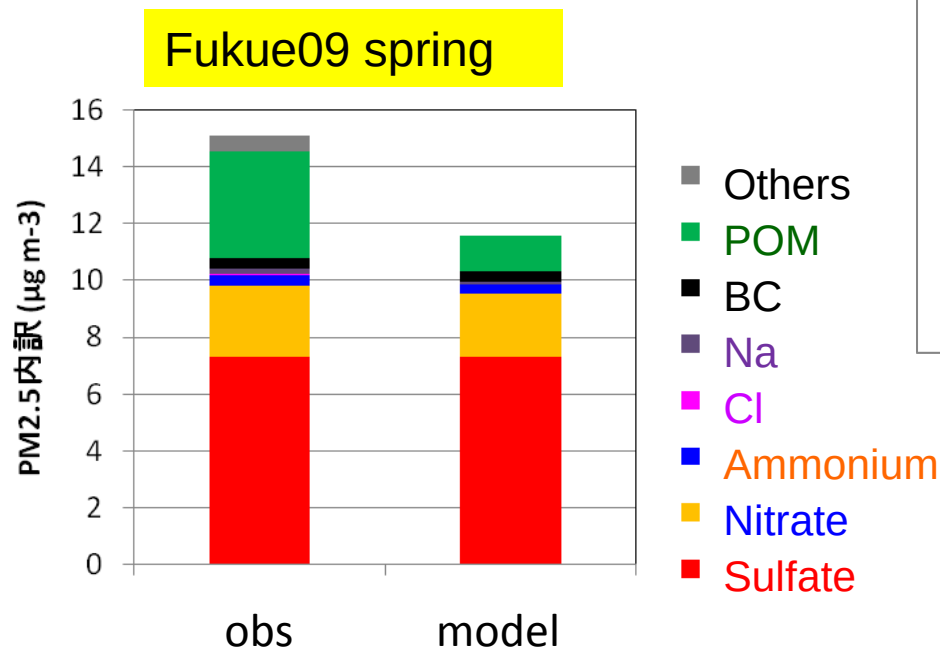
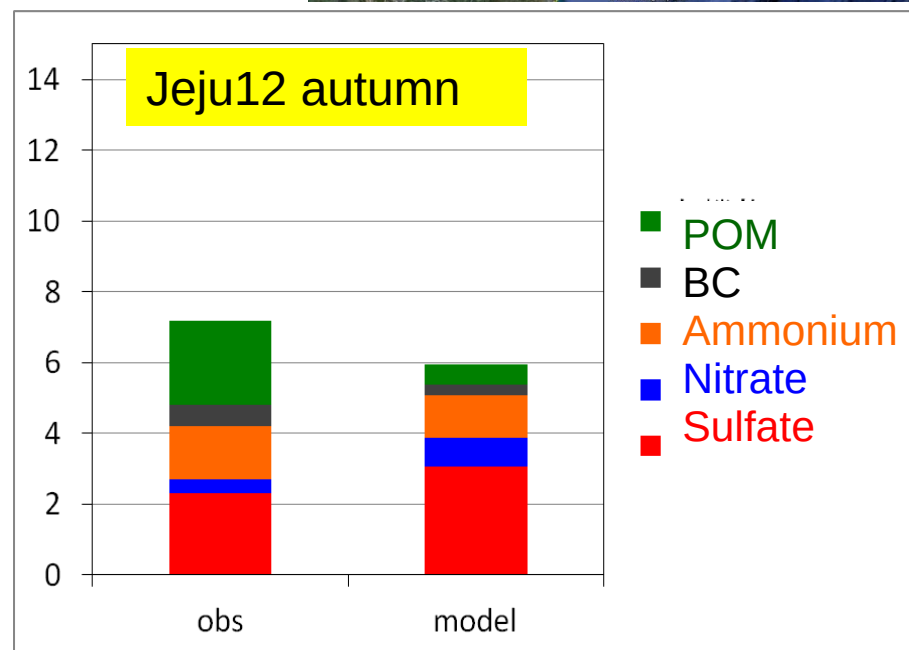
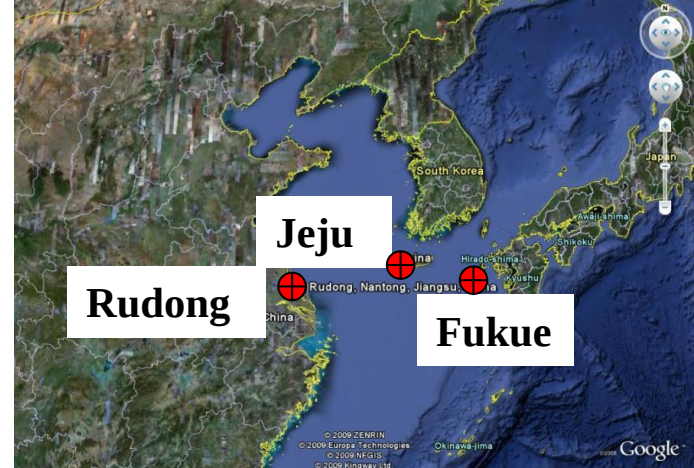
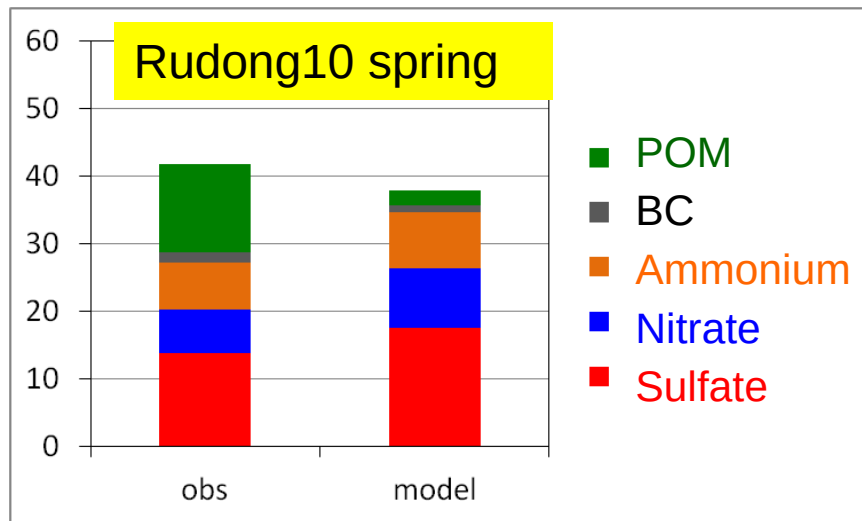
BC/CO emission ratio

$\text{BC/CO} = 7.5 \text{ ng/m}^3/\text{ppbv}$ (Andreae and Merlet, 2001, agricultural waste burning) is often used but the ratio could be larger



Estimated Emission ratio:
BC/CO = 18.2 ng/m³/ppbv
OC/CO = 101 ng/m³/ppbv

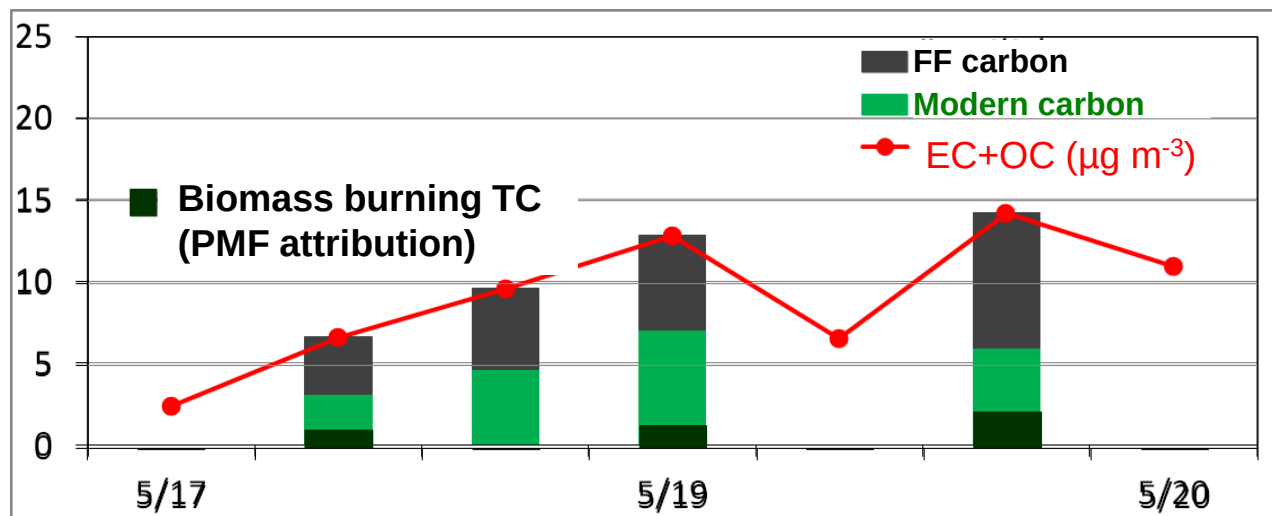
PM_{2.5} composition: obs vs. model ($\mu\text{g m}^{-3}$)



- Common underestimation of **POM** by factors of 4-6
- **SNA** relatively well reproduced

Explore origins of missing carbonaceous aerosols using ^{14}C

(Rudong, May-June, 2010)

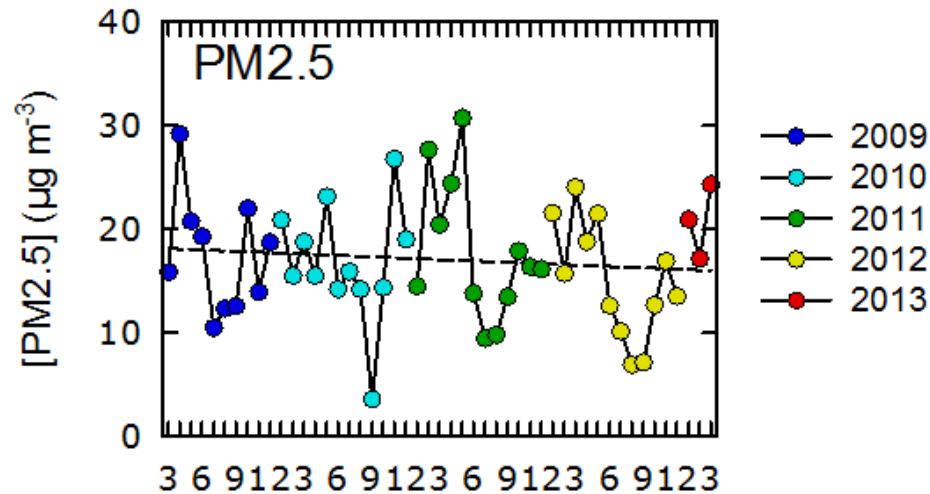


Urban/industrial emission dominant,
But large amount of modern carbon: $3\text{--}7\mu\text{g m}^{-3}$

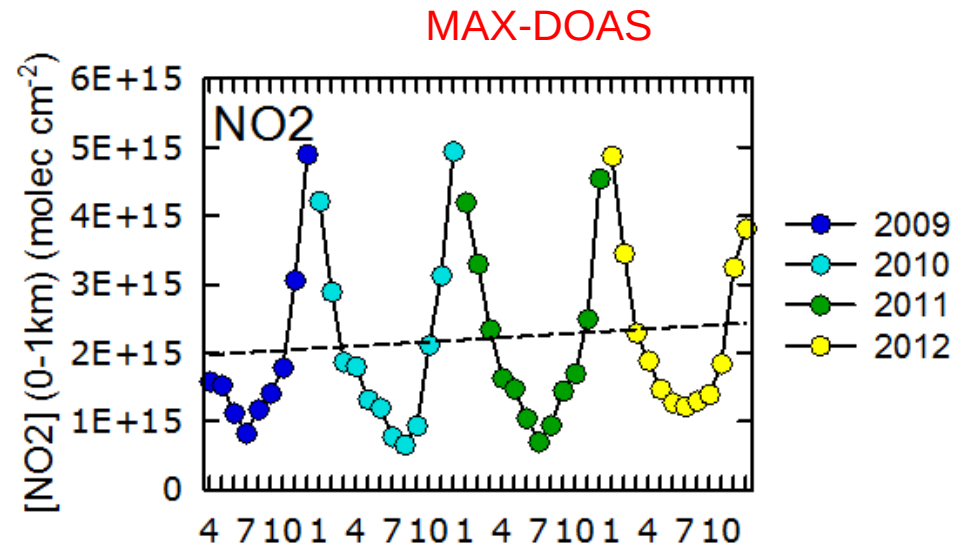
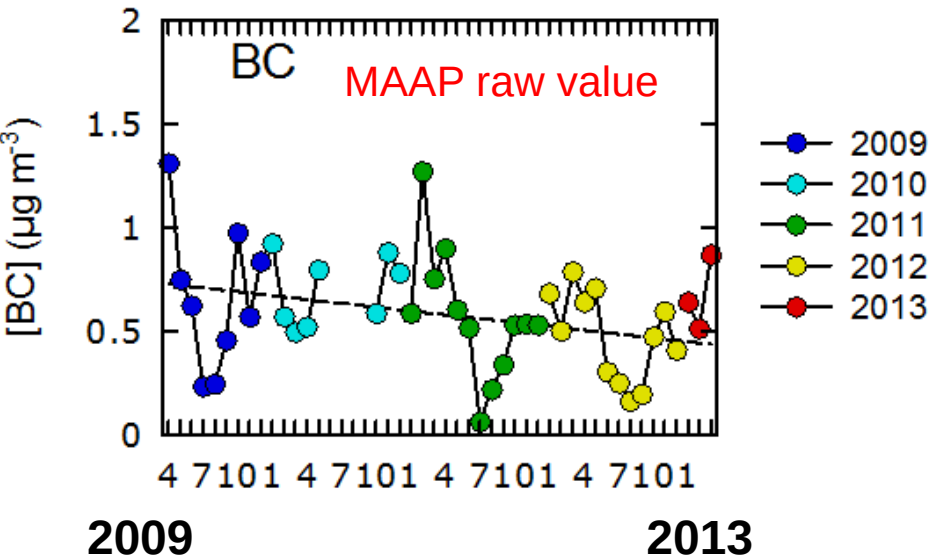
PMF attribution to Biomass burning (TC): only $0\text{--}2\mu\text{g m}^{-3}$

Suggesting more modern carbon from Prim/Sec biogenic OA

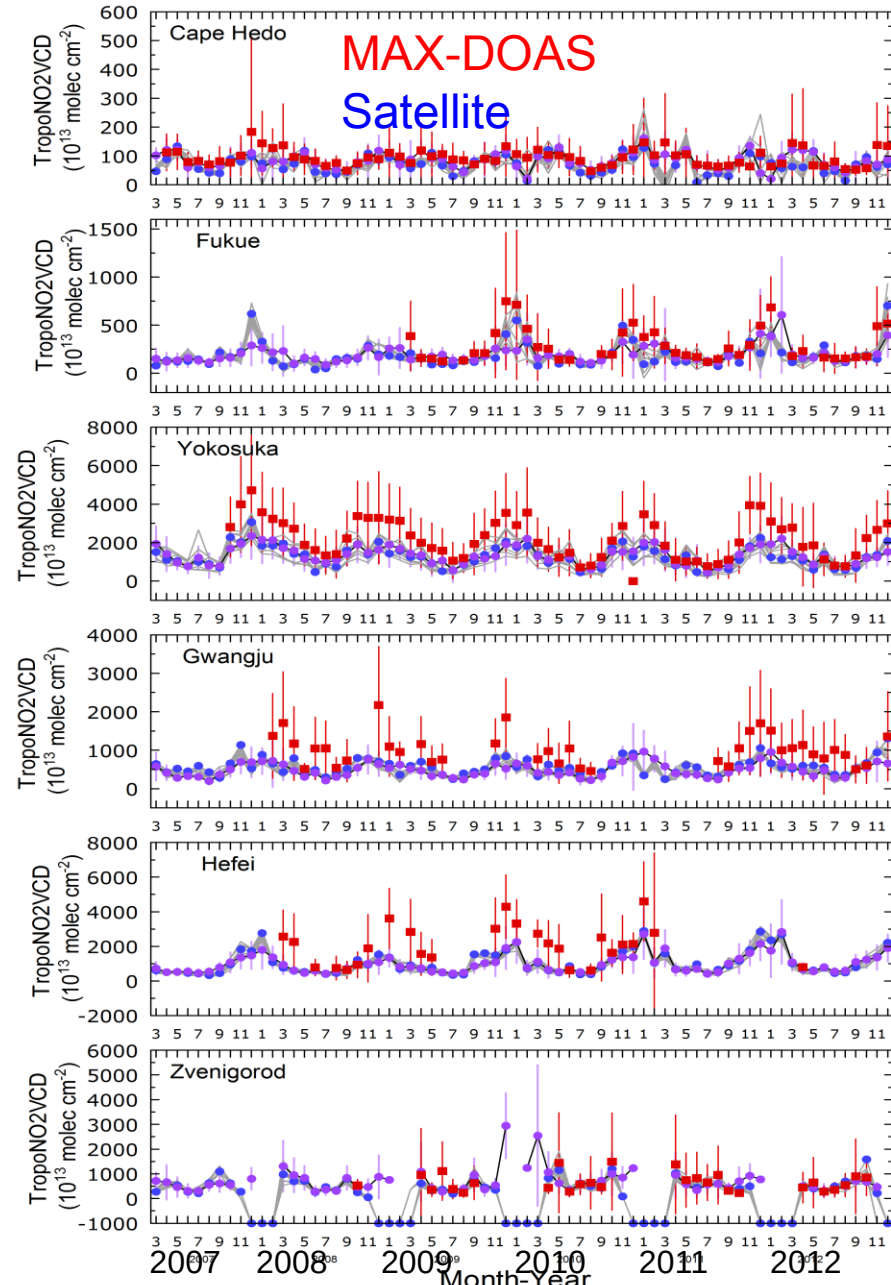
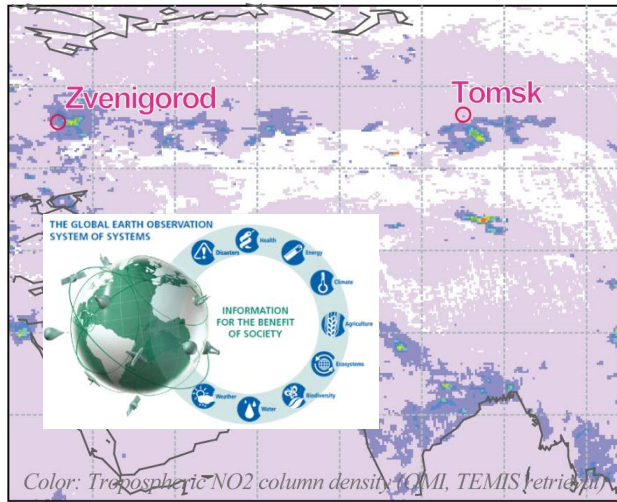
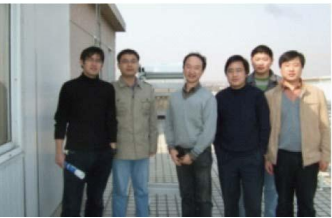
Decreasing BC trend at Fukue? (preliminary)



- Downturn of BC emission started?
- Or coupled to meteorology changes?



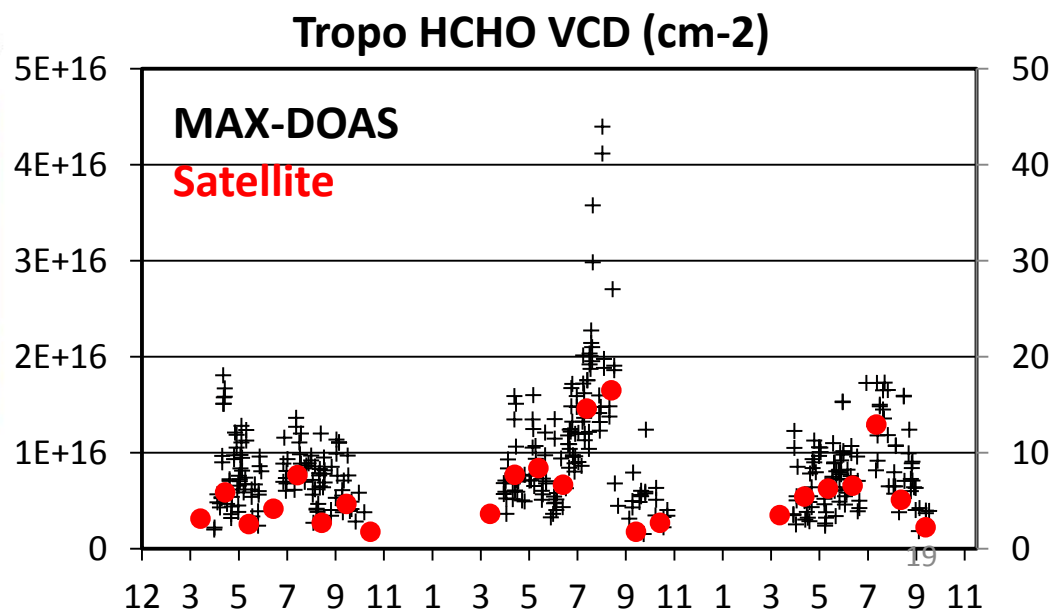
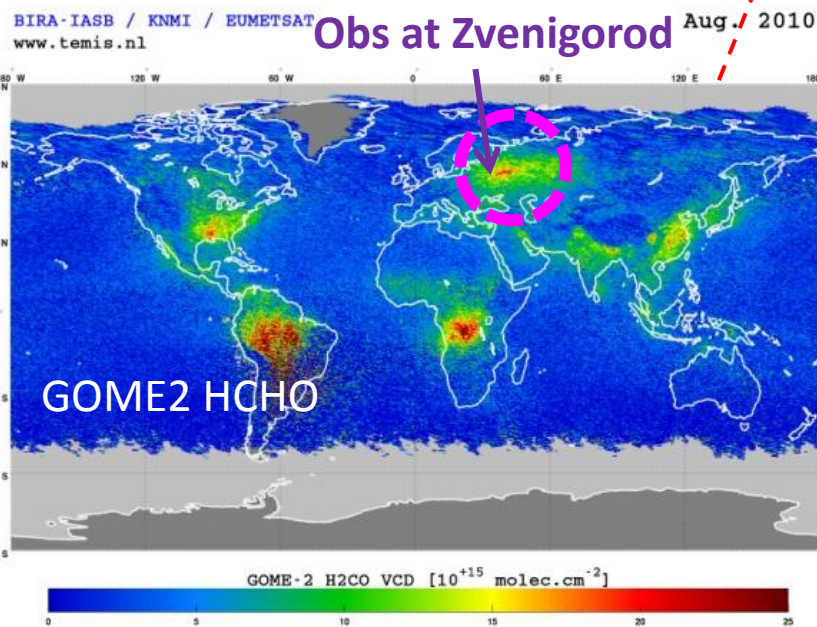
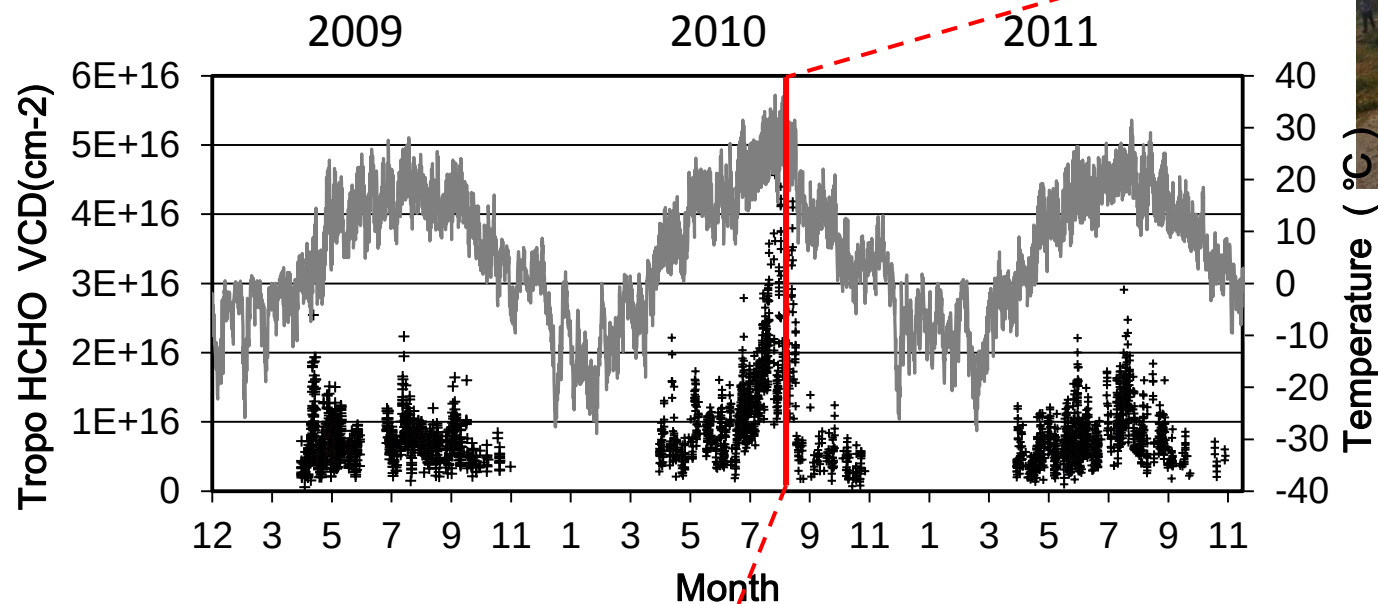
MAX-DOAS network over Asia and Russia



- Wanted: partners for obs in SE/S Asia!
- Climatology of NO₂ and aerosols
- Validation of satellite & model

<http://ebcrpa.jamstec.go.jp/maxdoashp/>

HCHO from Russian Forest Fire 2010

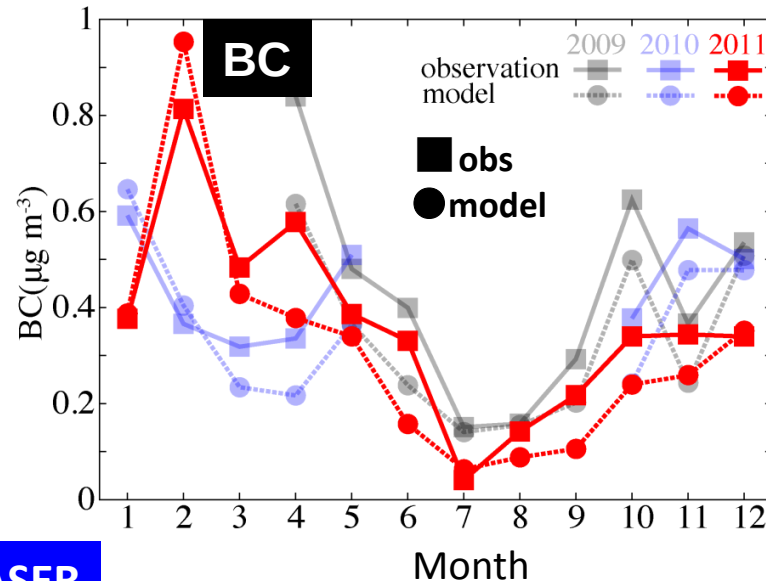
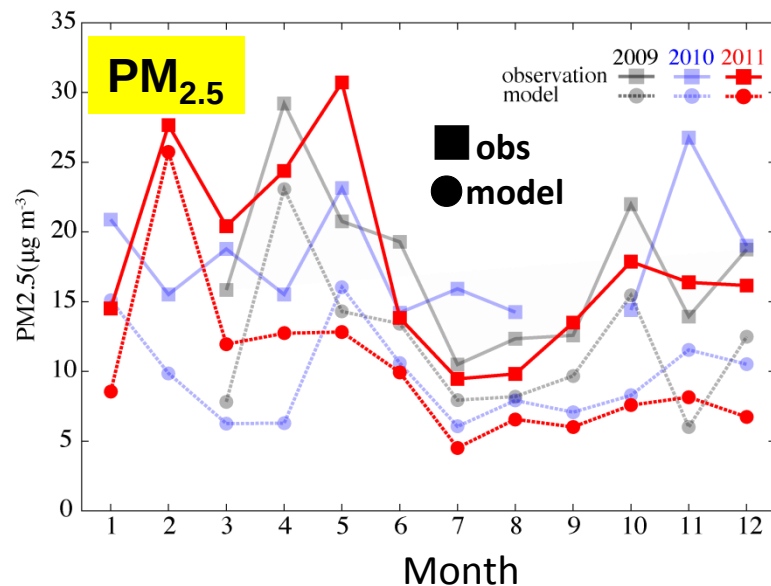


Summary

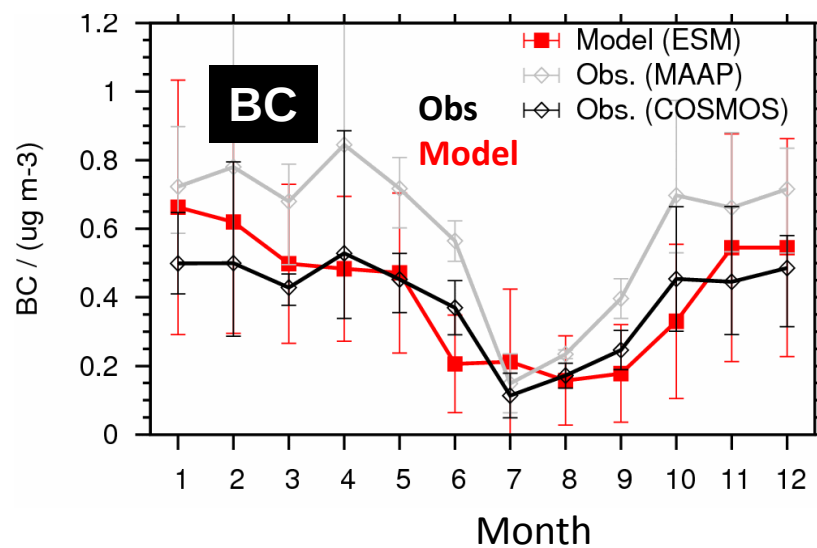
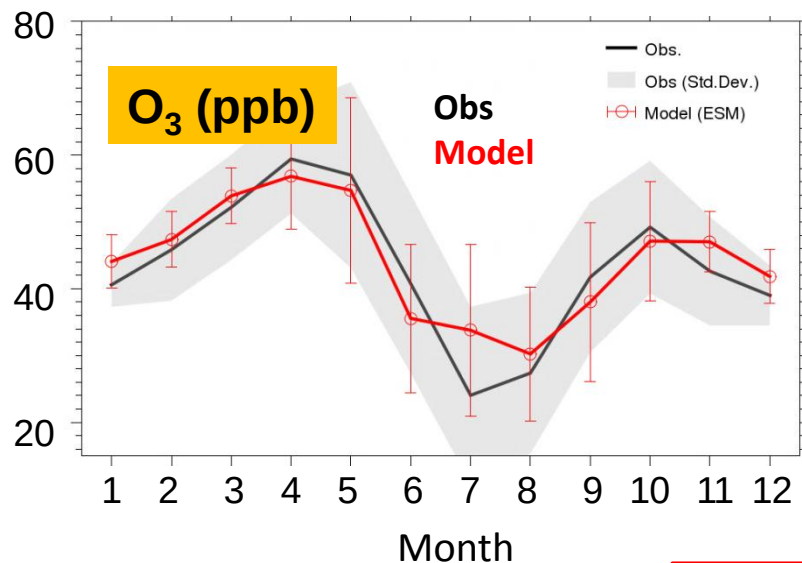
- Regional-scale open crop residue burning (OCRB) in CEC is an importance source of air pollution
- BC emission rates from OCRB could be underestimated by a factor of 2.5
- Organics: underestimated by a factor of 5, at 3 field campaigns;
More natural source suggested by ^{14}C analysis
- Long-term BC reduction is observed at Fukue, needing confirmation
- MAX-DOAS detected HCHO emission from Russian forest fire in 2010:
good agreement with satellite

Comparison of seasonality at Fukue

WRF-CMAQ

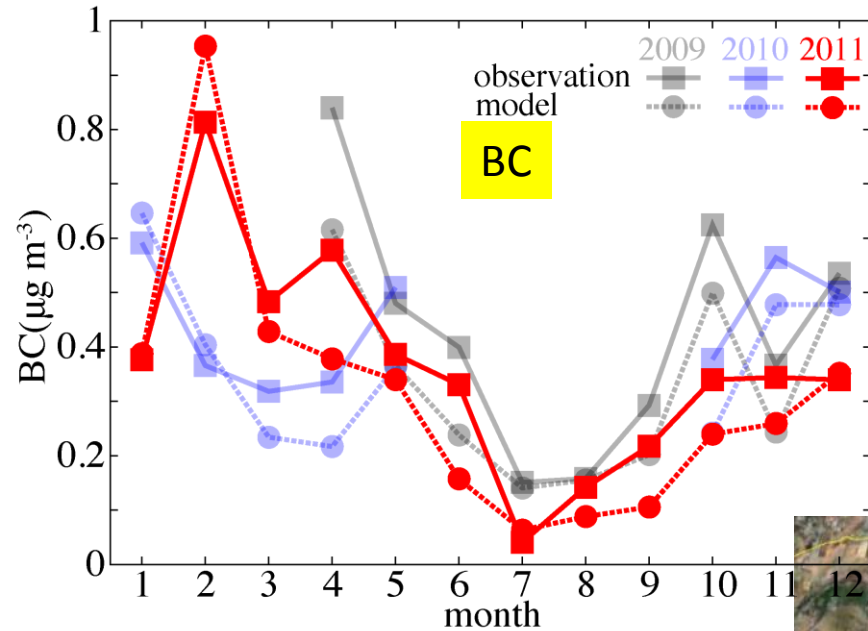


CHASER



Generally good agreement

BC at Fukue, in the downwind, is in agreement with model...



BC emission increase will result in overestimation of BC at Fukue

biomass burning is not a major source
Or the loss rate of BC could also be faster

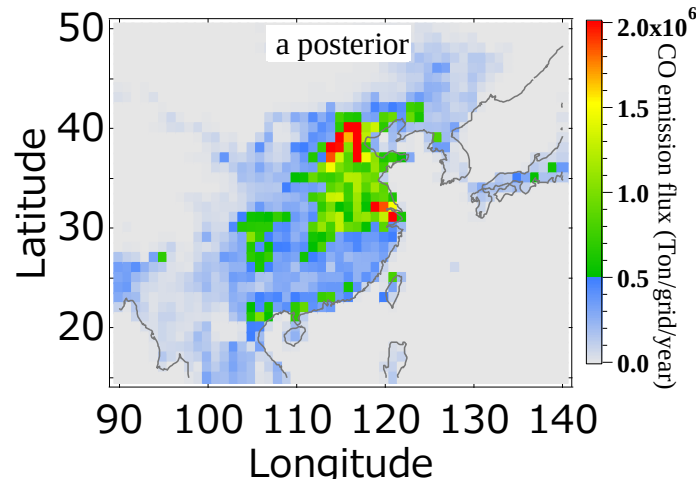
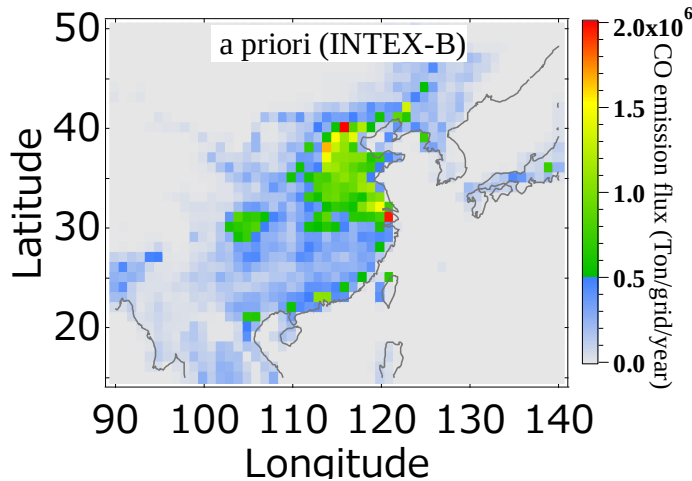
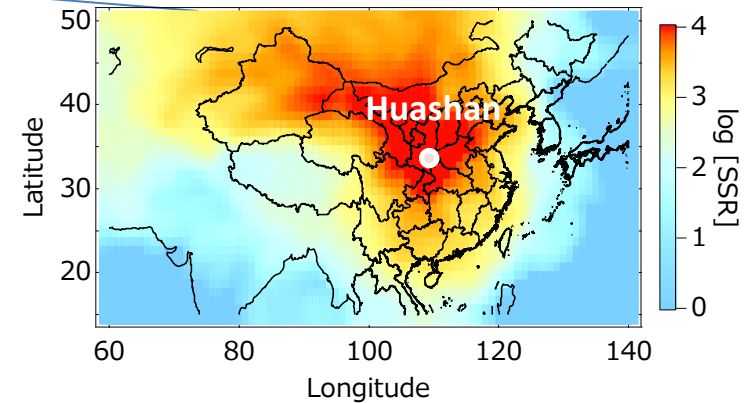
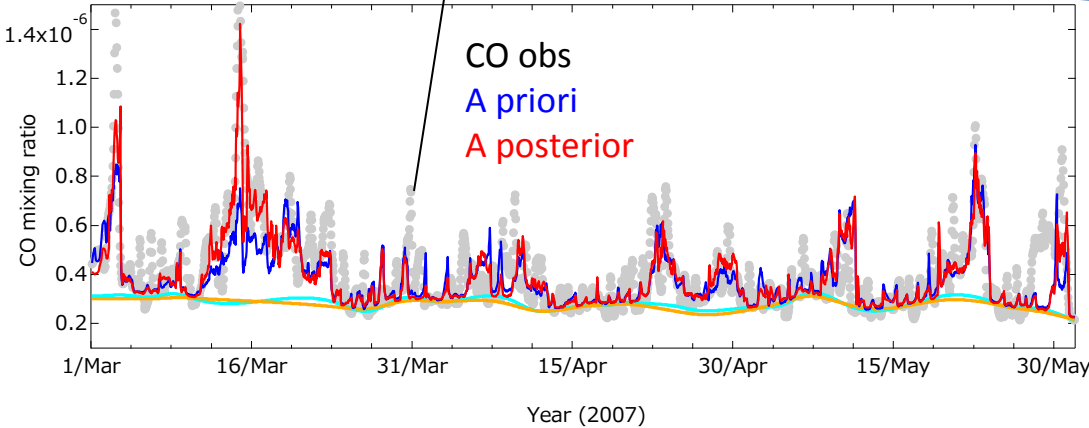
BC in downwind in Japan is well reproduced by the model, suggesting that major BC emission source is different and well constrained, or lifetime of BC is shorter and compensates larger emission rates



Bayesian inversion & WRF_FLEXPART model to optimize CO emission

Pan et al., in preparation

Basic equation: $y_m = M_{m \times n} \cdot x_n + \varepsilon$



- ▶ Need more emission around Beijing, YZRD regions
- ▶ CO emission is likely 30% underestimated
- ▶ BC emission rates, based on BC/CO ratio, also need to be increased further by 30%