



Assessment of cloud related fine mode AOD enhancements based on the AERONET SDA product

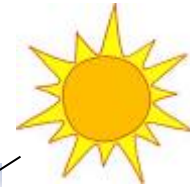
A. Arola, T F. Eck, H. Kokkola, T. Laaksoviita,
A. V. Lindfors, M. Pitkänen and S. Romakkaniemi



"SDA effectively computes the fine mode AOD in mixed cloud-smoke observations". Should one then rather use L1 SDA than L2 to estimate the mean fine mode AOD?



"L2 sun"

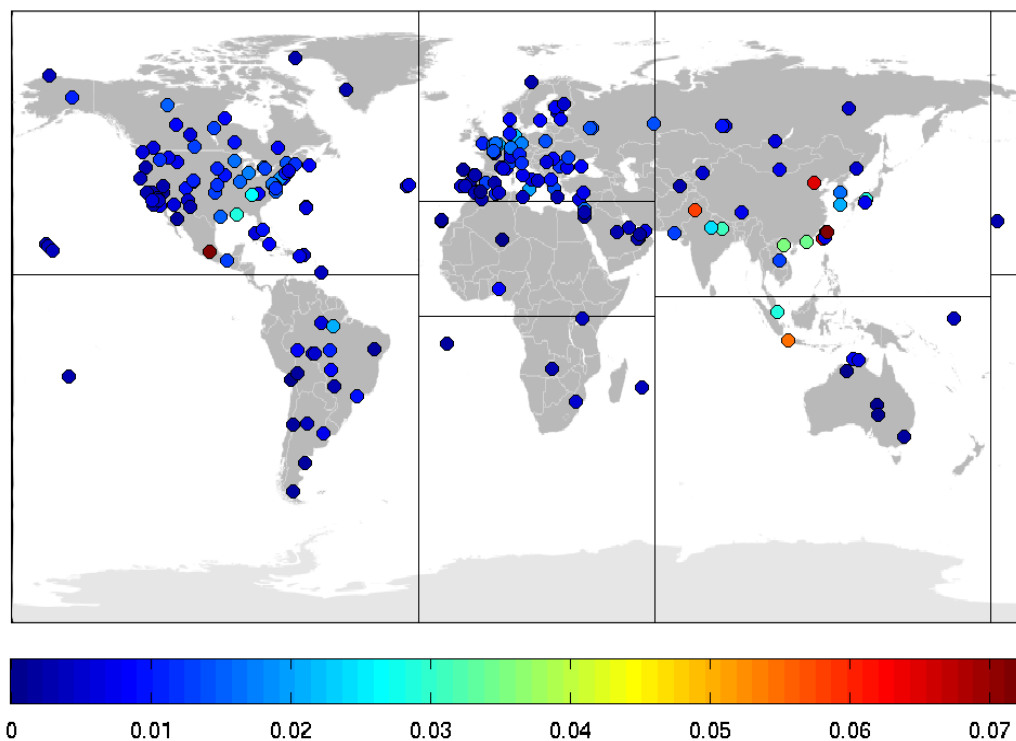


"L1 sun"

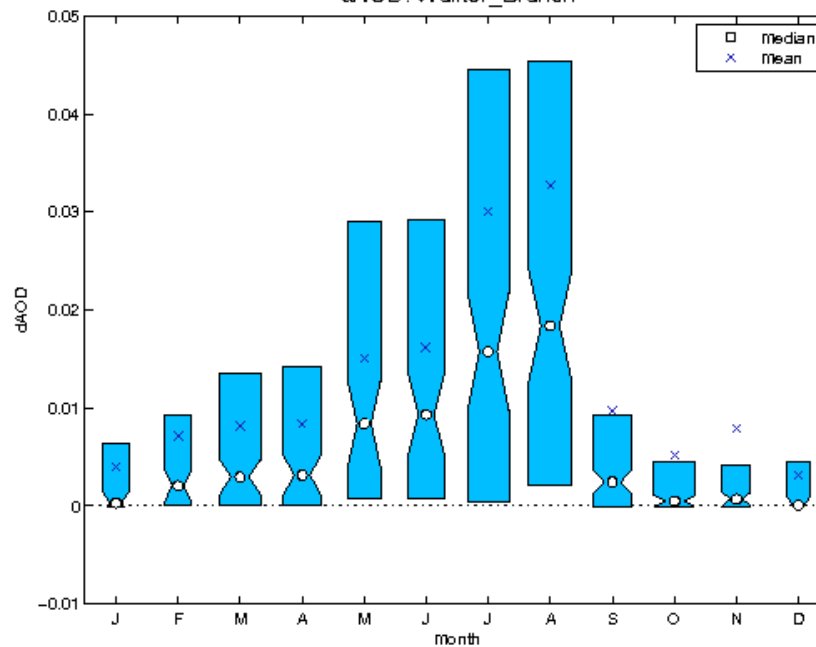


Difference in fine mode AOD between L1 and L2 AERONET data, sampled for the days when both L1 and L2 data were available.

Absolute dAOD, JJA

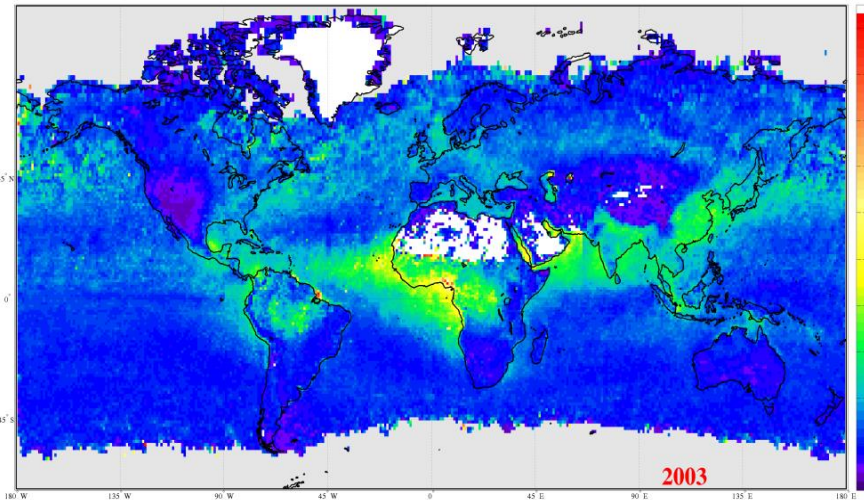


dAOD: Walker_Branch





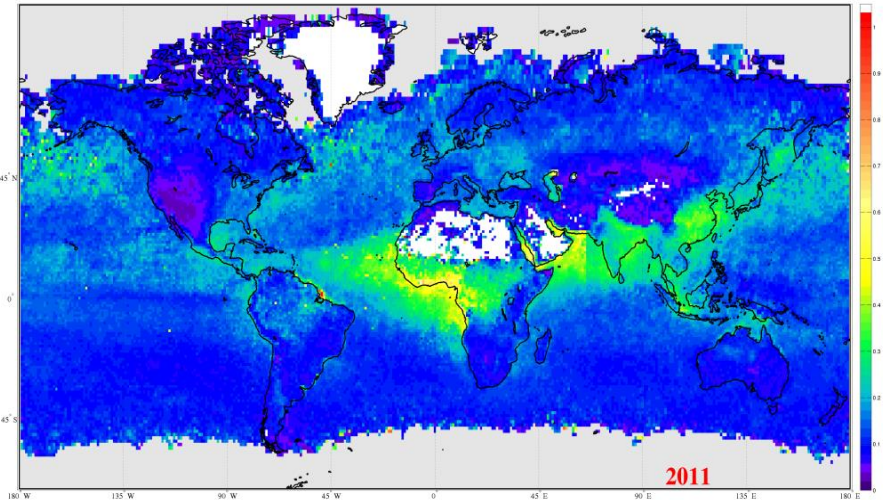
AATSR: 2003-2011



2004

...

2010



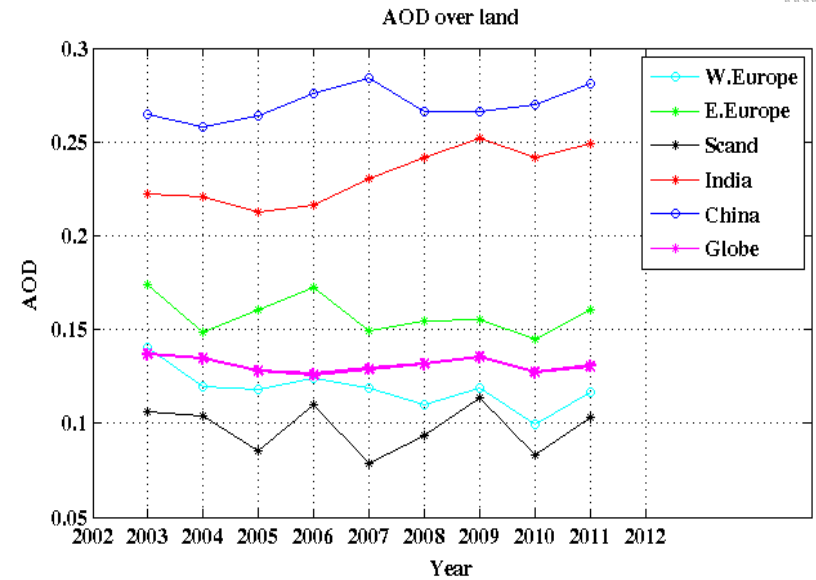
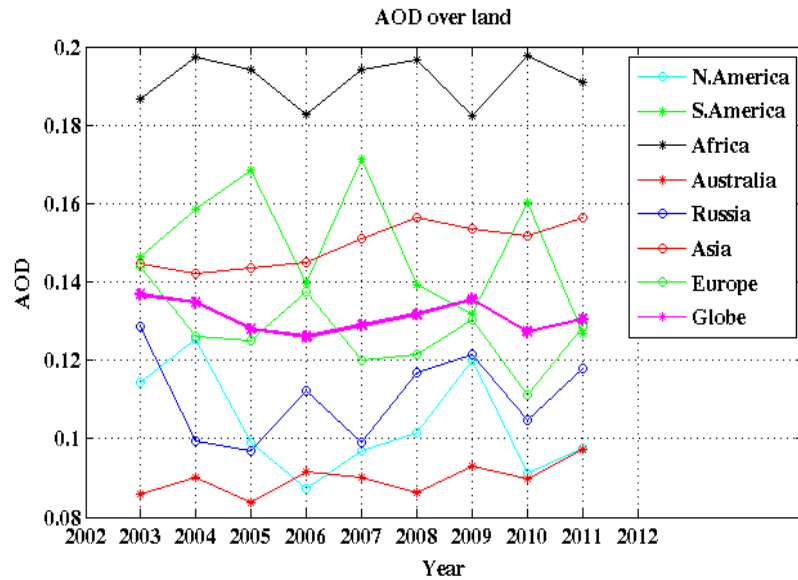
Continents

<

Time series

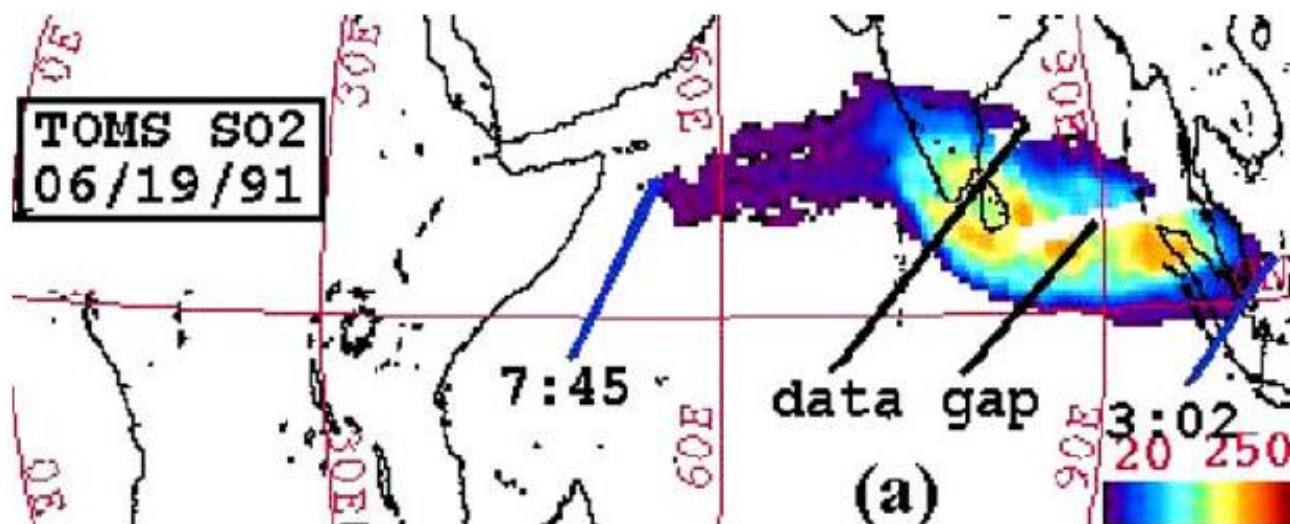
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Regions



Aerosol microphysics simulations of the Mt. Pinatubo eruption with the UKCA composition-climate model

Sandip Dhomse, Graham Mann, Ken Carslaw, et al. (Univ. Leeds, U.K.)



Guo et al.
(GGG,
2004.)

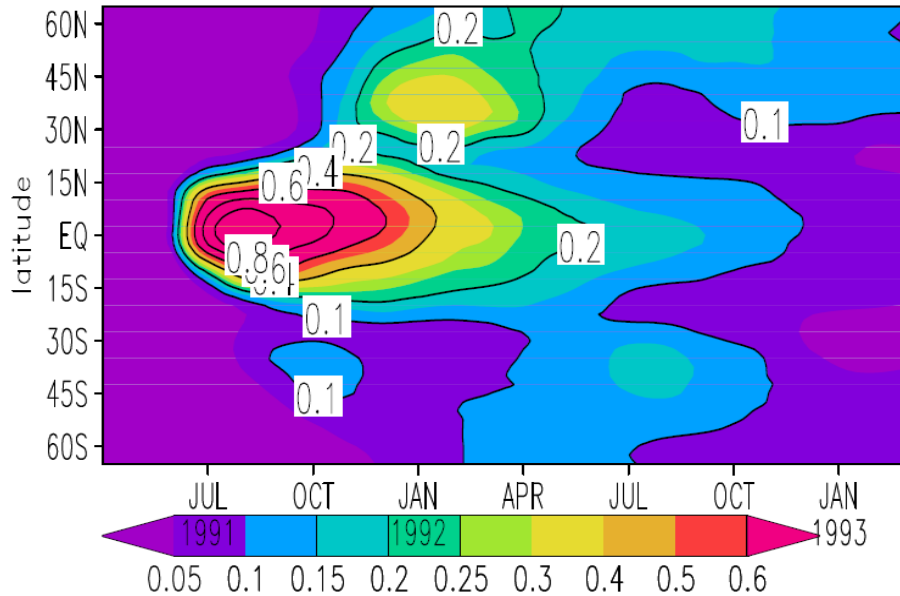
Satellite measurements indicate 14 to 23 Tg of SO₂ (7 to 11.5 TgS) was present in the tropical stratosphere shortly after the eruption.

The stratospheric aerosol loading peaked several months later in the range 19-26 Tg (Lambert et al., 1993). Assuming 59 to 77% sulphuric acid (Grainger et al., 1993) this gives a range of 3.7-6.7 Tg of sulphur.

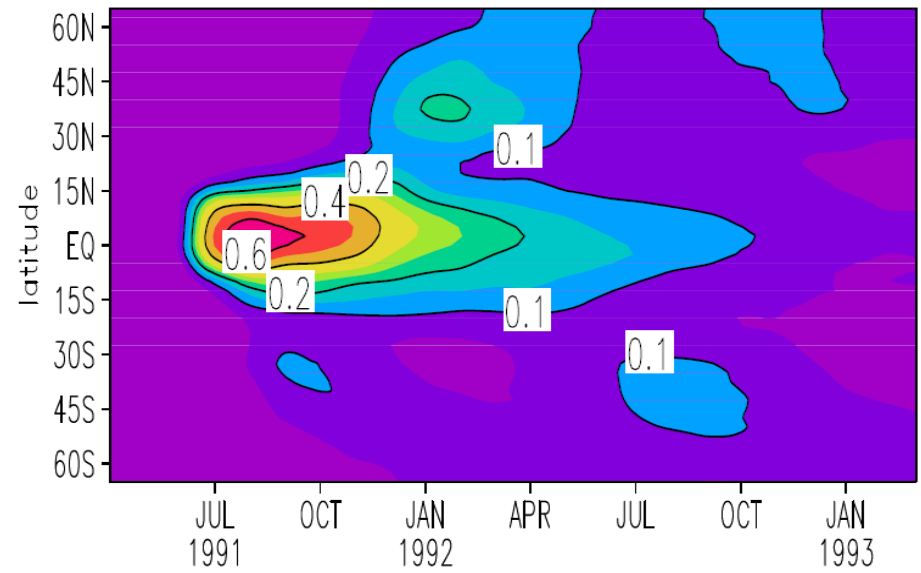
Investigate the eruption's impact on the stratospheric aerosol in UKCA with runs which inject 10 & 20 Tg of SO₂ into the tropical stratosphere

stratospheric aerosol optical properties

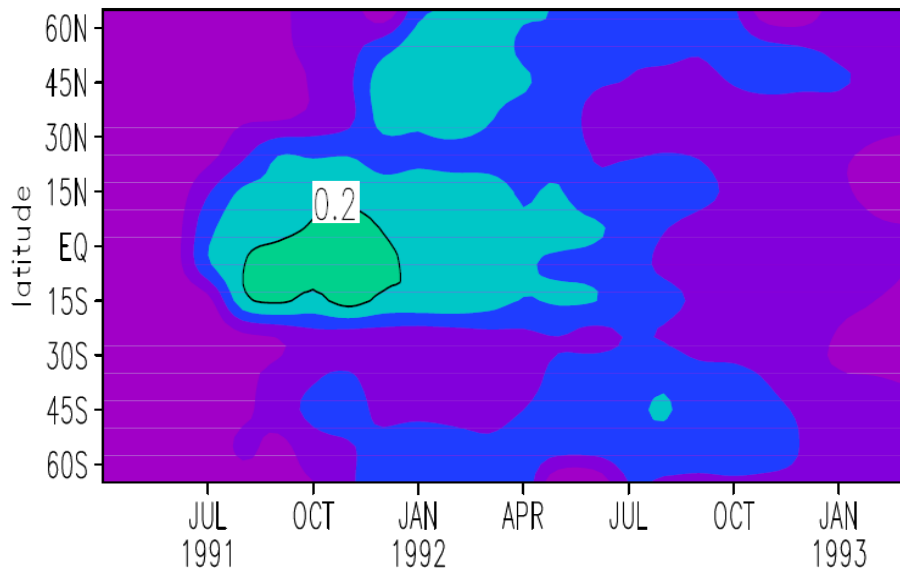
a) UKCA-A_Control20 sAOD (550nm)



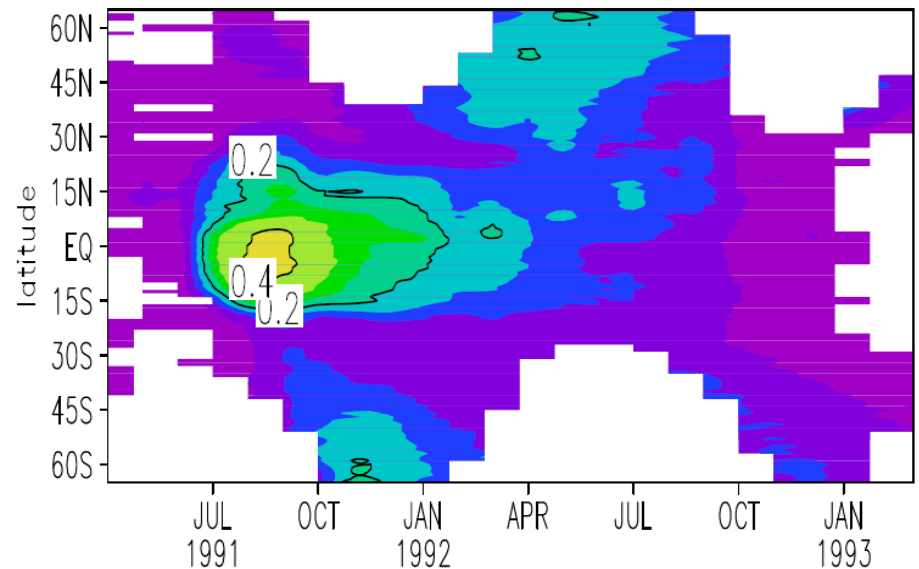
b) UKCA-B_Control10 sAOD (550nm)



c) SAGE AOD (525nm)



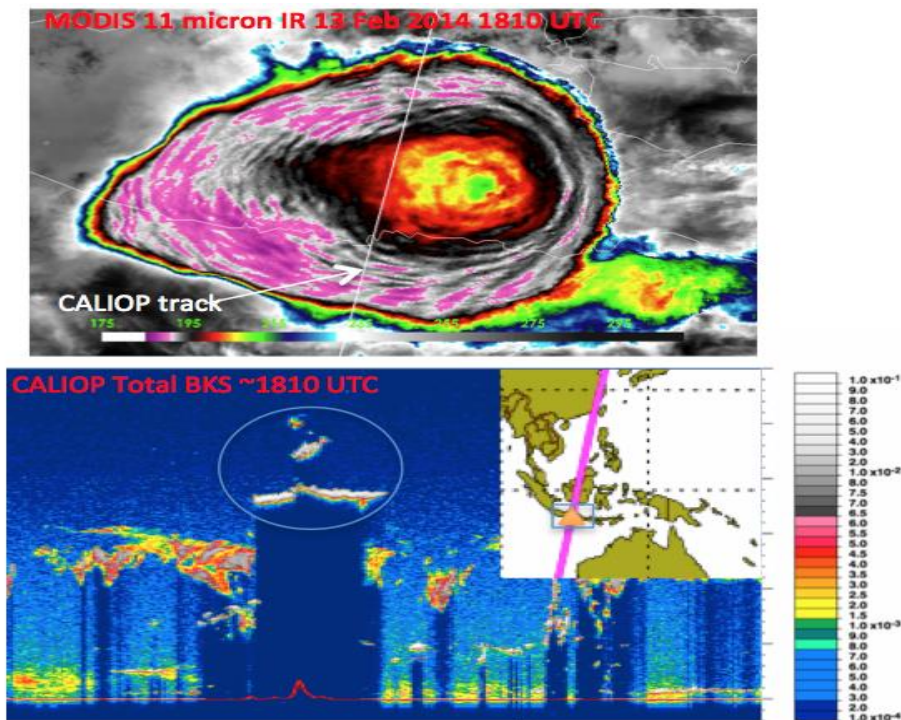
d) AVHRR AOD (600nm)



Klash, 2014: CALIPSO and in-situ balloon measurements of Mt. Kelud volcanic plume; persistence of ash in the lower stratosphere

T. Duncan Fairlie², Jean-Paul Vernier¹, Terry Deshler³, Travis Knepp¹, Murali Natarajan², Katie Foster³, Stan Smith³, Kristopher Bedka¹, Chip Trepte², Larry Thomason², Frank Weinhold⁴

[¹SSAI; ²NASA LaRC; ³U. Wyoming; ⁴ETH, Zurich]



The eruption of Mt. Kelud : 14 Feb 2014:

MODIS(Aqua) Brightness Temperature (11 micron), and CALIPSO total attenuated backscatter curtain showing main volcanic plume ~18-19 km altitude, extending as high as 26 km.

KLASH deployment:

10-day balloon field experiment in Darwin (Australia) May, 2014.

Rapid Response, with critical support from NASA HQ (Considine, Kaye), CALIPSO (Trepte), SAGE (Thomason), Australian BOM (Atkinson), CASA.

Objectives: CALIPSO validation, ash confirmation, size, volatility, RF.



- Flew 4 dual backscatter (COBALD) sondes under medium balloons
- Flew combined optical particle counter (OPC) with COBALD under large balloon

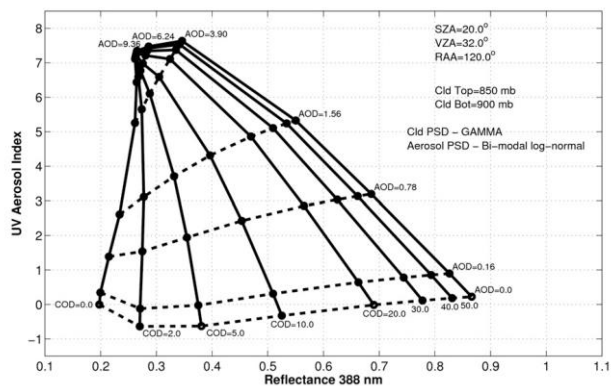


Retrieval, Inter-comparison, and Validation of Above-cloud Aerosol Optical Depth from A-train Sensors

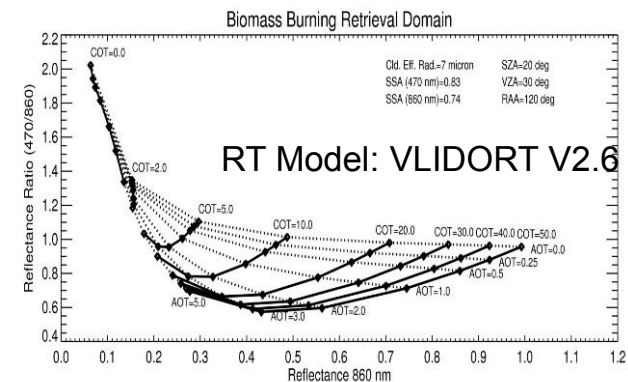
HIREN JETHVA, O. TORRES, P. K. BHARTIA, L. A. REMER, J. REDEMANN, S. E. DUNAGAN, J. LIVINGSTON, Y. SHINOZUKA, M. KACENELENOBOGEN, M. SEGAL-ROSENHEIMER, ROB SPURR

The presence of absorbing biomass burning and dust aerosols above clouds pose greater potentials of exerting positive radiative effects (warming) whose magnitude directly depends on the aerosol loading above cloud, optical properties of clouds and aerosols, and cloud fraction

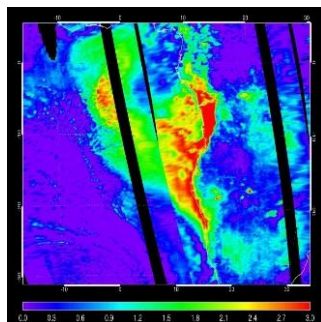
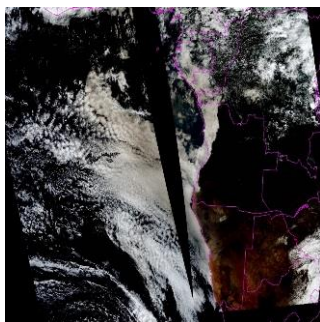
Near-UV Retrieval Domain



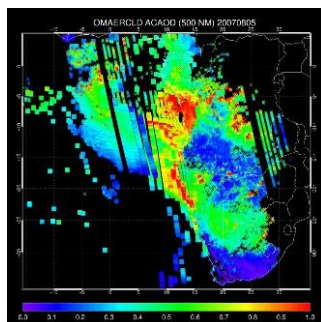
VIS/SWIR Retrieval Domain



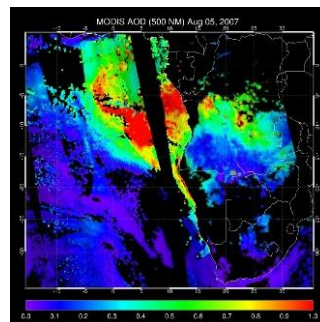
Aqua/MODIS RGB
Aura/OMI
UV Aerosol Index



OMI AOD

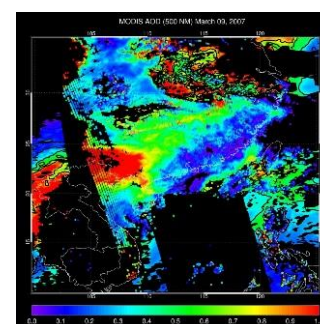
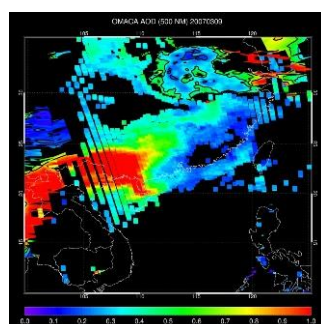
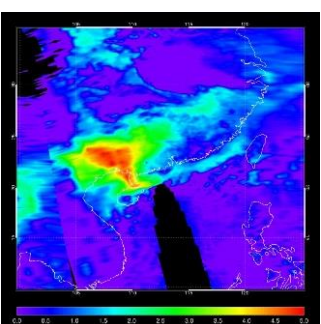
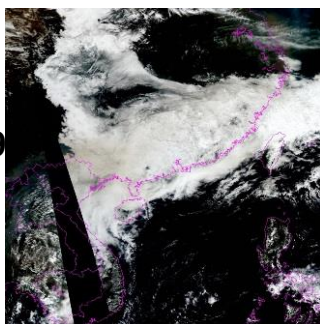


MODIS AOD



Aug 05
2007

Mar 09
2007



Input Parameters

TOA reflectance
OMI: 354 and 388 nm
MODIS: 470 and 660/860 nm
(ocean/land)

A priori:

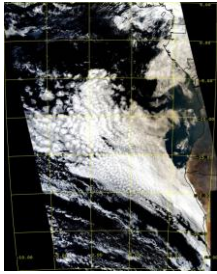
Aerosol Model (size distribution, real/imaginary part of refractive indices, aerosol vertical profile, cloud droplet distribution (modified-Gamma), Cloud top/bottom pressure

Output Parameters

Above-cloud AOD
(at 388 for OMI and at 660/860 nm for MODIS)
aerosol-corrected COD

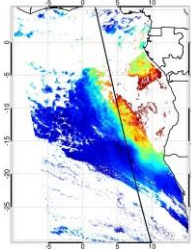
A-train Inter-comparison of Above-cloud AOD

MODIS RGB



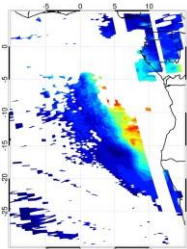
MODIS ACAOD

Jethva et al. (2013)

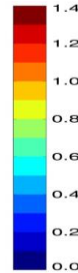
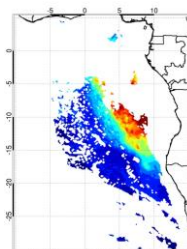


OMI

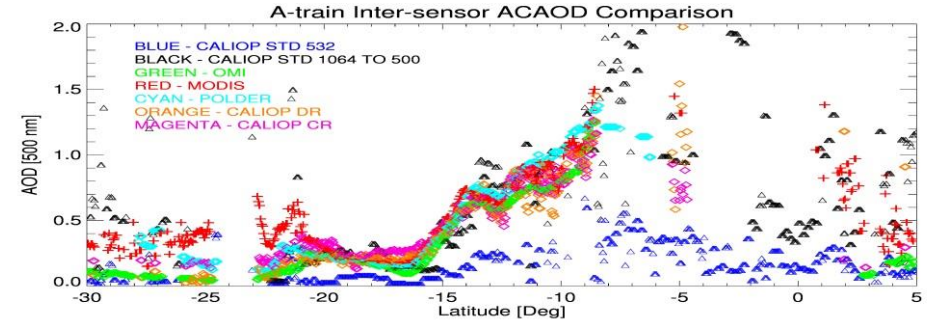
Torres et al. (2012) Waquet et al. (2009, 2013)



POLDER



Aug 02, 2007

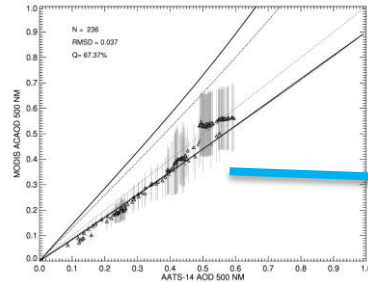
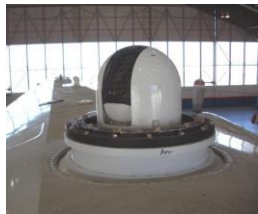


Ames Airborne Tracking Sunphotometer (AATS)
Spectrometer for Sky-Scanning, Sun-Tracking Atmospheric Research (4STAR)

Validating Above-cloud AOD

AATS-14

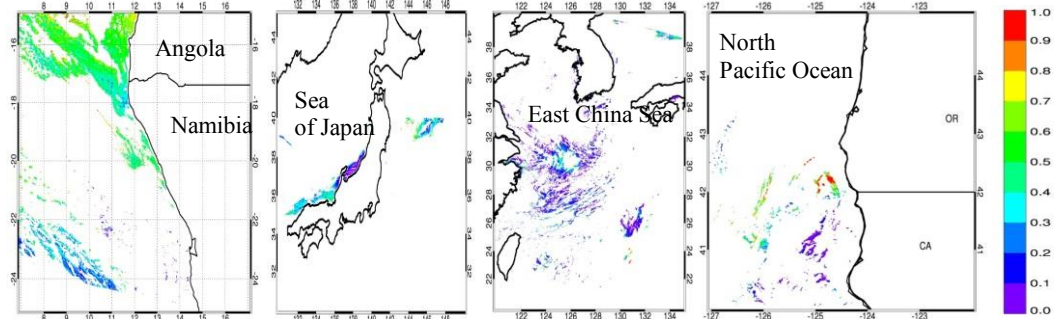
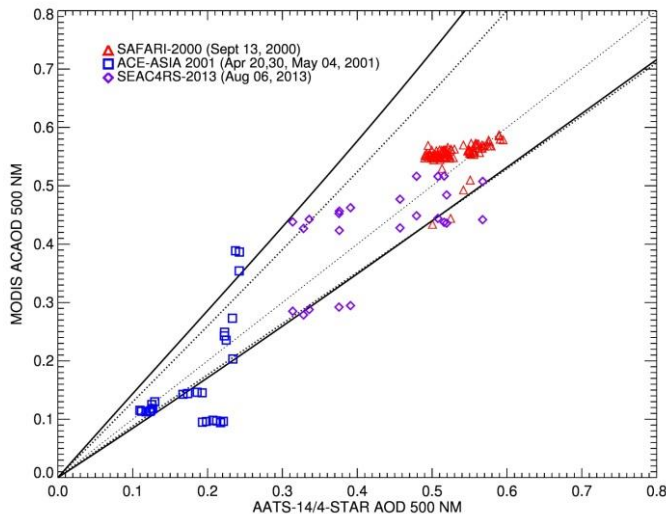
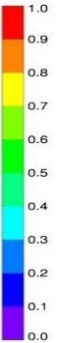
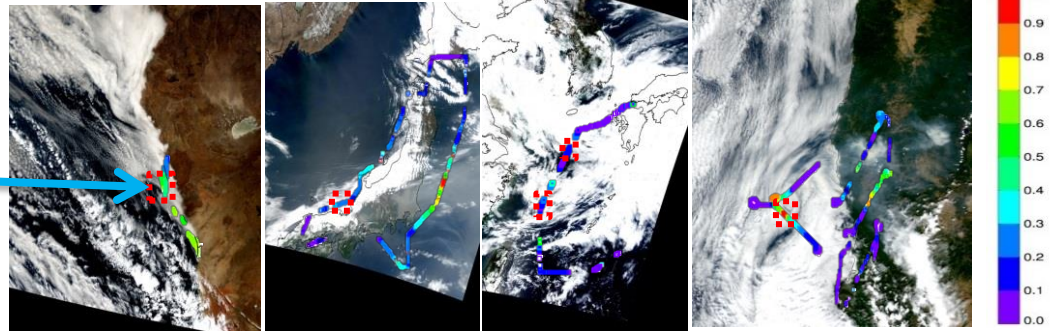
4STAR



SAFARI-2000

ACE-ASIA 2001

SEAC4RS-2013



- Most satellite-airborne matchups falling within the predicted uncertainties in the above-cloud AOD retrieval (-10% to 50%)
- The co-retrieved COD was found to be either equivalent to the MODIS operational cloud product for lower AODs (ACE-ASIA) or higher by 30-35% for more absorbing aerosol event of SAFARI-2000

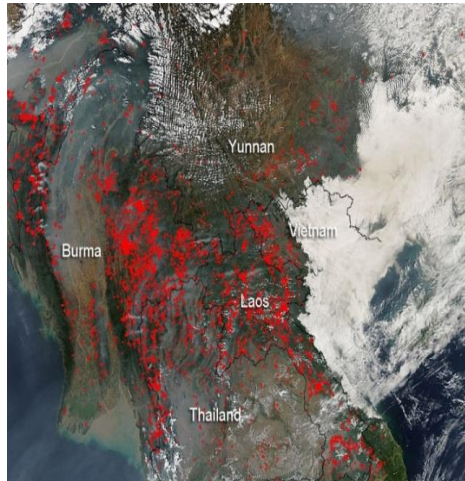
Impacts of Aerosol Induced by Wildfire over Indochina Peninsula on East Asian Climate

Yiquan Jiang¹, Xiaohong Liu^{1*}, Yun Qian² and Kai Zhang²

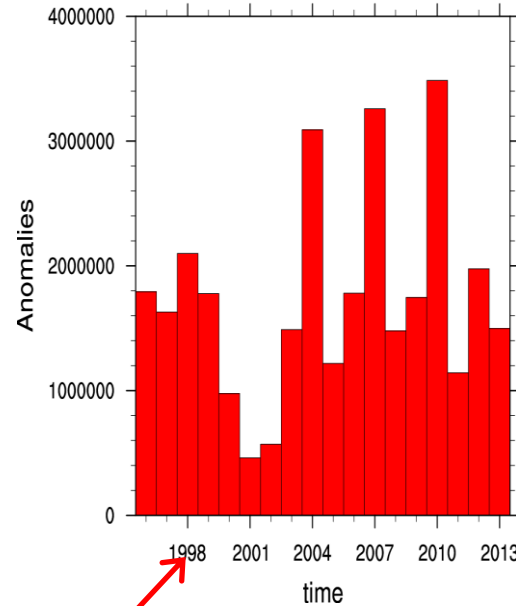
1 Department of Atmospheric Science, University of Wyoming

2 Pacific Northwest National Laboratory, Richland, Washington, USA

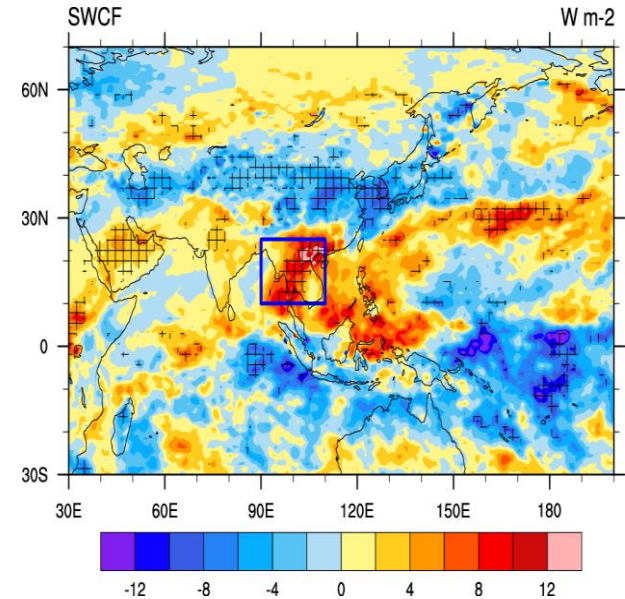
How fire aerosols from Indochina affects climate?



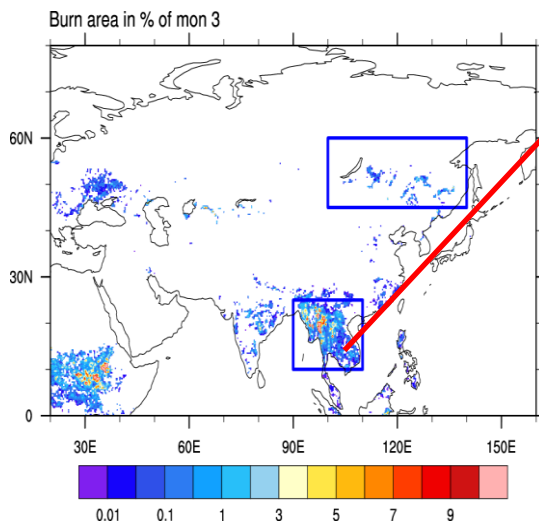
March South Asia Burnarea (1.748e+06)



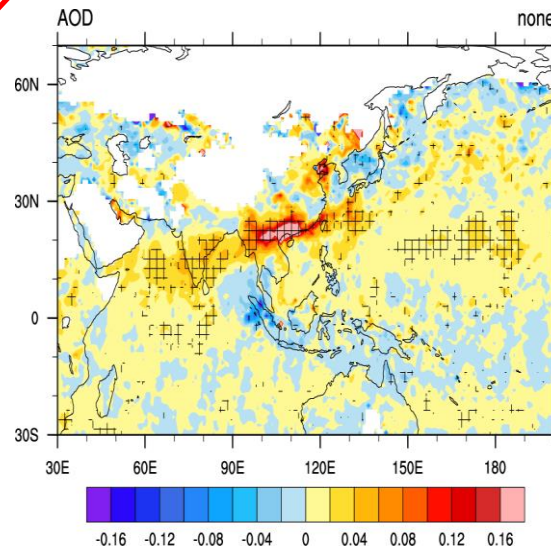
Regressed SWCF



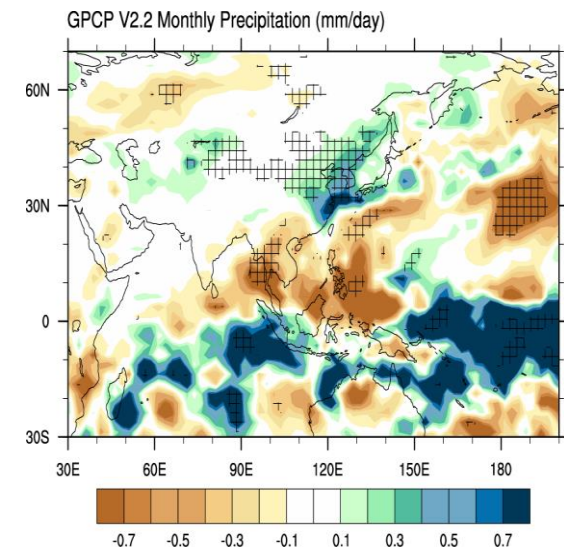
GFED Burn area of March



Regressed AOD



Regressed Precipitation



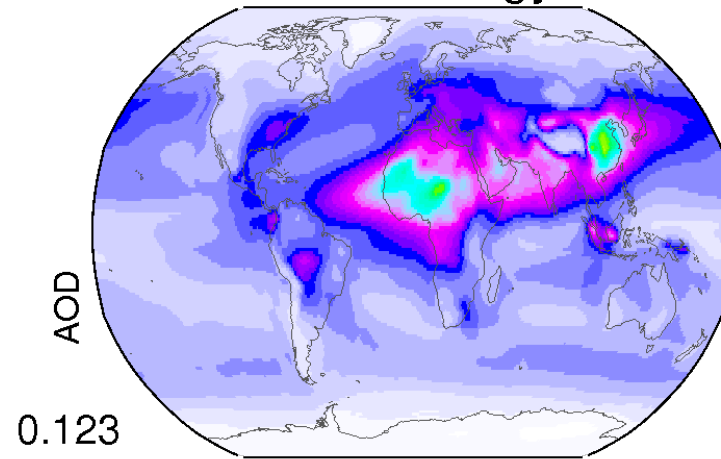
The MPI-M Aerosol Climatology

Stefan Kinne

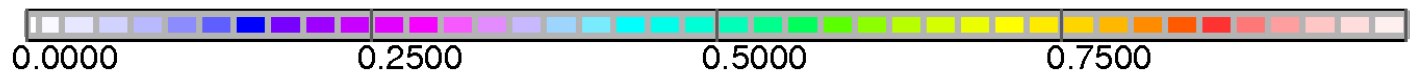
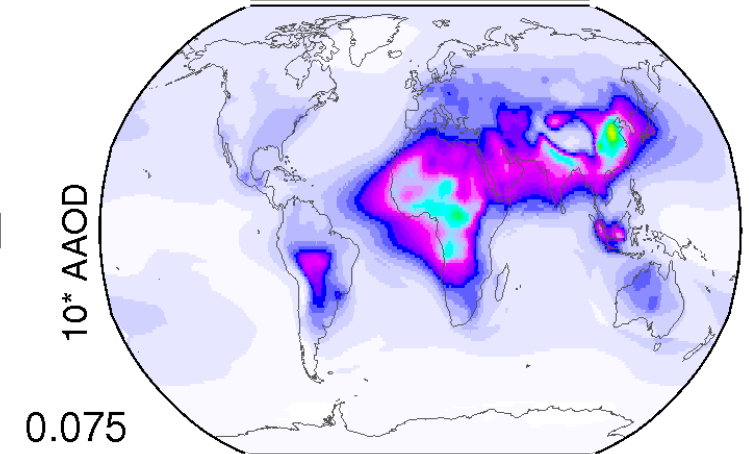
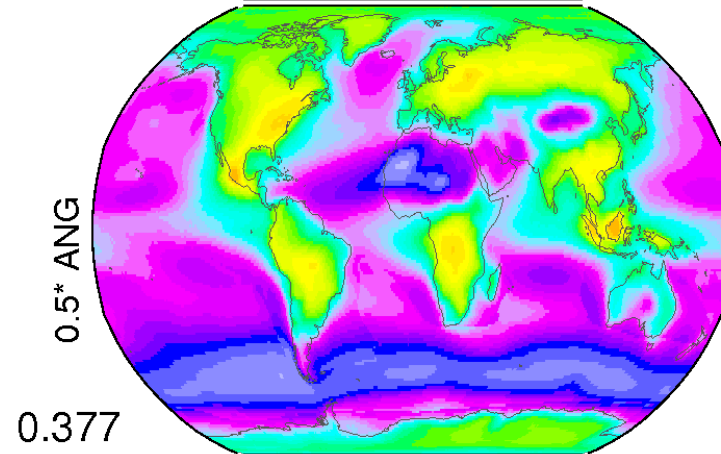
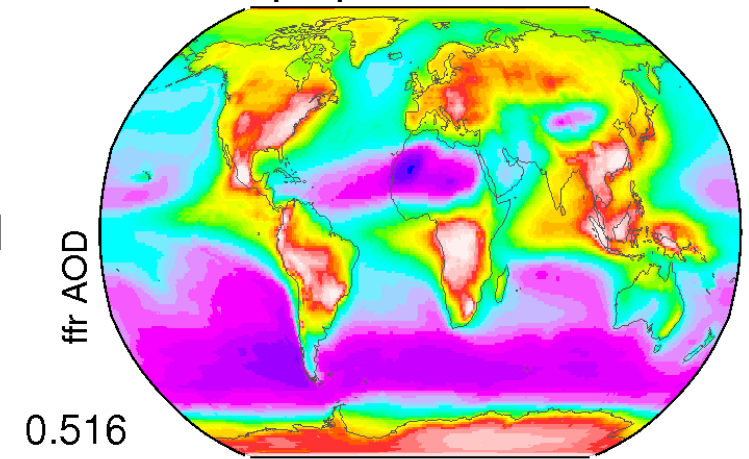
AeroCom



MAC-v2 climatology



aerosol properties at 550nm



annual global maps →

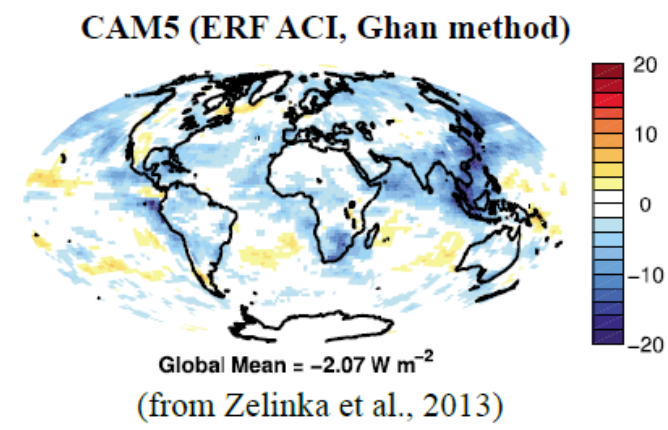
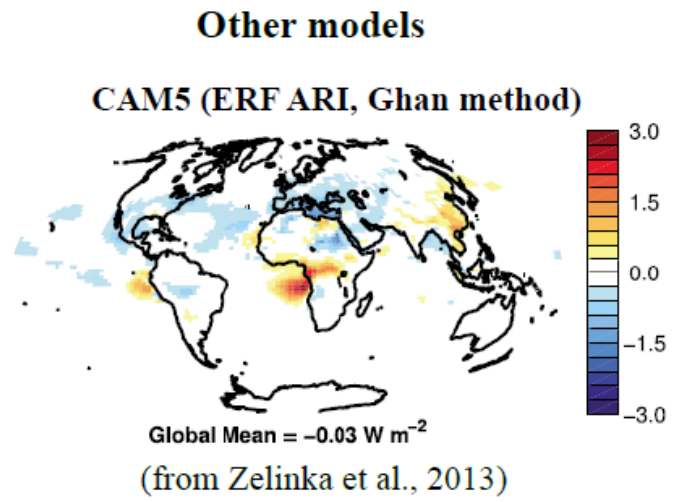
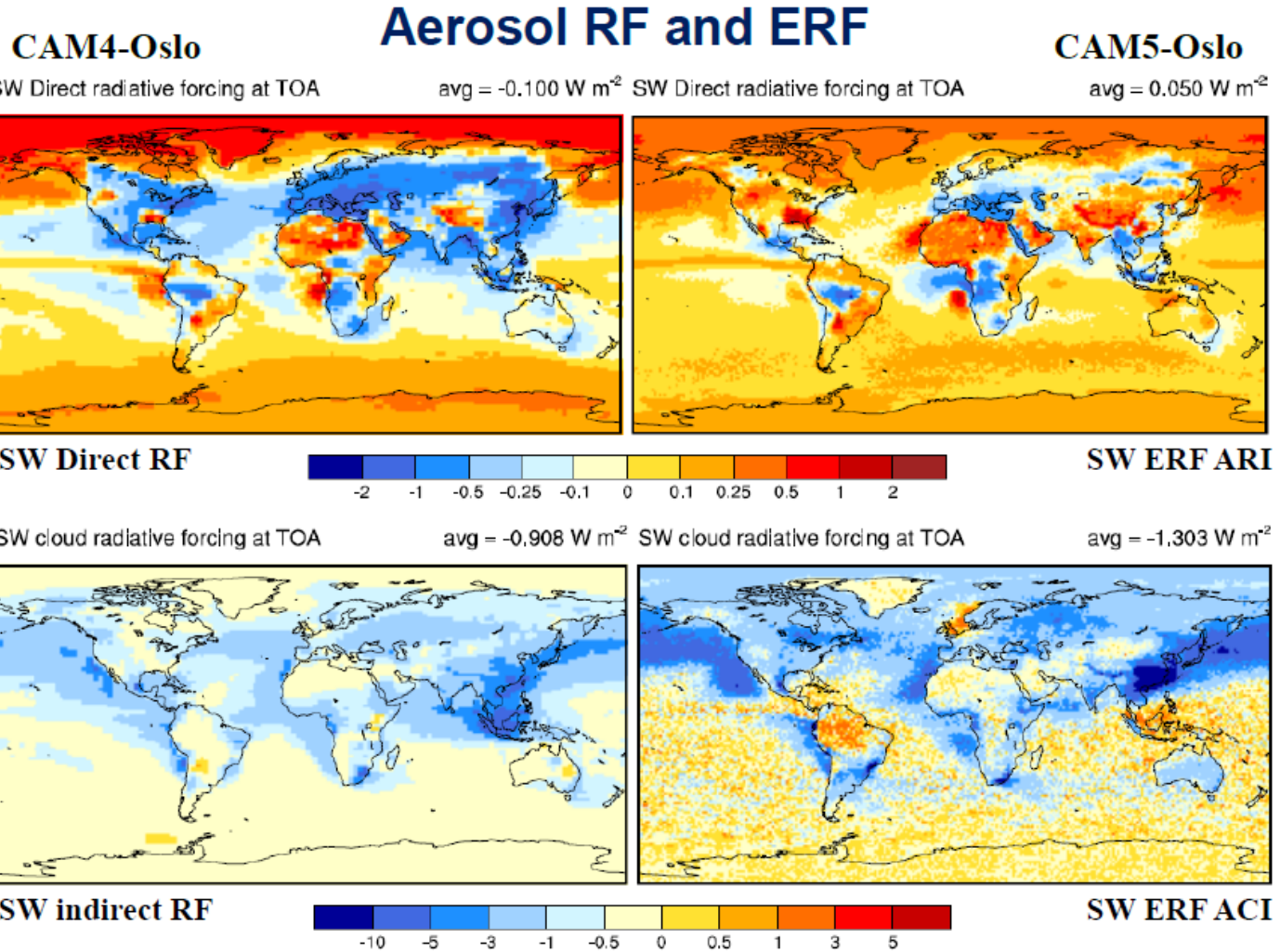
applications

- a general reference
- shortcut, when opt/rad properties are needed
- with rad. transfer - a tool for sensitivity studies
- obs. connection helps identify model biases
- CCN estimates are a path to indirect effects

always think about simplifications, if they work
... not to get lost in complexity space

Preliminary estimates of Aerosol Effective Radiative Forcing in CAM5-Oslo

A. Kirkevåg, A. Grini, T. Iversen, D. Olivié, M. Schulz and Ø. Seland



CAM5-Oslo is a version of CAM5 where schemes for aerosol chemistry, physics and interaction with clouds originally developed for CAM4-Oslo/NorESM1 will exist as options alongside with the modal aerosol modules (e.g. MAM3). Note: the aerosol coupling with ice nuclei is still as in CAM5 MAM3, and the treatment of wet-scavenging and aerosol activation is not yet consistent → only preliminary results!

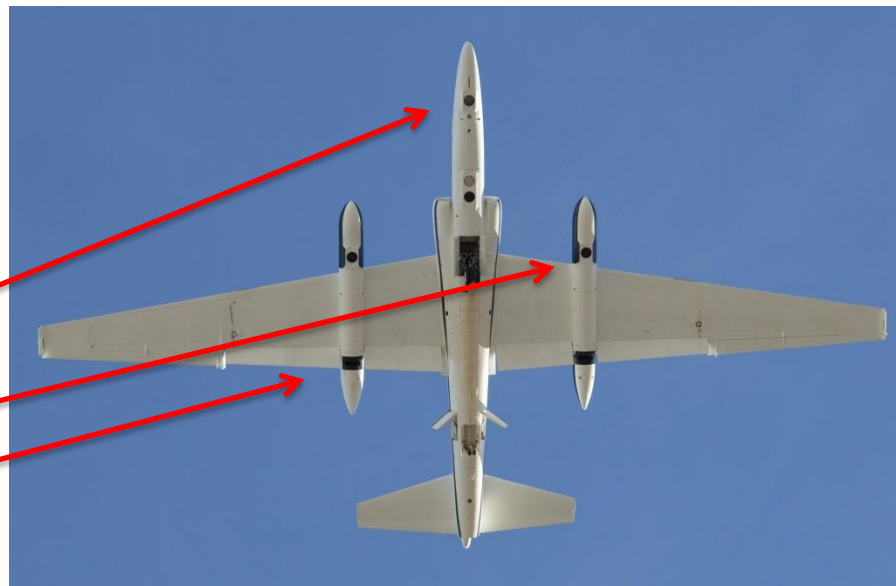
Polarimeter Definition EXperiment (PODEX) Level 1 comparisons

Kirk Knobelspiesse, Jens Redemann

NASA Ames Research Center

Airborne polarimeters relevant to NASA
ACE, PACE missions:

AirMSPI, NASA/JPL
PACS, UMBC
RSP, NASA GISS



PODEX goal: develop and inter-compare polarimeter aerosol / cloud retrievals

	Type	Approximate polarimetric accuracy @reflectance=0.2	# view angles	ER-2 Nadir ground resolution	355	380	410	445	470	550	555	660	670	766	865	870	935	960	1593	1880	2263	total # obs. per pixel
AirMSPI	Photoelastic modulation, imager	1%: Step & Stare mode; 0.5%: sweep mode	varies, 1 to 31	7m+9m smear	hatched	hatched		hatched	blue		hatched	blue		blue	blue		hatched					up to 420
PACS	Philips prisms + linear polarizers, imager	?	varies, max ~65	37m + smear?					blue	blue	blue		blue	blue		blue						up to 1170
RSP	Wollaston Prisms, not an imager	0.075%	~152	277m+277m smear			blue		blue		blue		blue		blue		blue	blue	blue	blue	blue	~4100

 Reflectance
  Reflectance+polarization

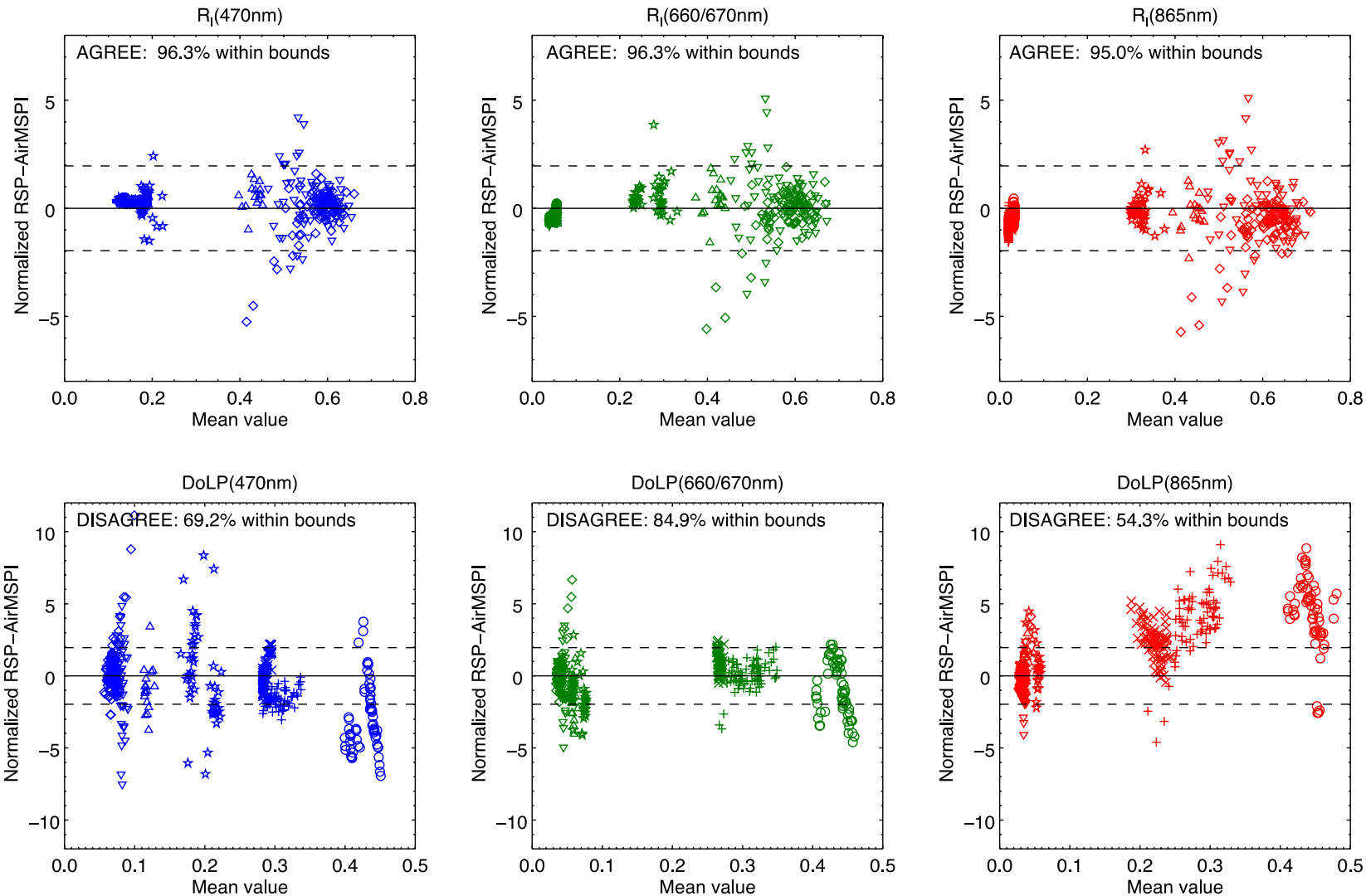
Level 1 intercomparisons (AirMSPI & RSP only, PACS data not available)

more details: earthscience.arc.nasa.gov/sgg/ACEPWG/

32

Comparison normalized by uncertainty

New!!



Reflectance compares well, but...

Degree of Linear Polarization (DoLP) differences are greater than uncertainty estimates

Evaluation of observed and modelled aerosol lifetimes

- using radioactive tracers of opportunity and an ensemble of 19 global models



Contact:
nik@nilu.no

N. I. Kristiansen¹, A. Stohl¹, T. Christoudias², D. Kunkel³, B. Croft⁴, J. Pierce⁴, R. Martin⁴, T. Bergman⁵, H. Kokkola⁵, Y.H. Lee⁶, D. Shindell¹⁶, G. Pitari⁷, G. Di Genova⁷, H. Zhang⁸, S. Zhao⁸, O. A. Søvde⁹, H. Wang¹⁰, K. Zhang¹⁰, X. Liu¹¹, N. Evangelou¹², Y. Balanski¹², K. Tsigaridis¹³, S. Bauer¹³, H. Klein¹⁴, S. Leadbetter¹⁵, D. J. L. Olivie¹⁴, M. Schulz¹⁴

1: NILU-Norwegian Institute for Air Research, Kjeller, Norway; 2: Cyprus Institute; 3: Institute for Atmospheric Physics, Johannes Gutenberg-University Mainz, Germany; 4: Department of Physics and Atmospheric Science Dalhousie University; 5: Finnish Meteorological Institute; 6: NASA Goddard Institute for Space studies, New York; 7: University of L'Aquila, Italy; 8: Chinese academy of meteorological science; 9: Center for International Climate and Environmental Research - Oslo (CICERO), Oslo, Norway; 10: Pacific Northwest National Laboratory (PNNL), Richland, WA, USA; 11: University of Wyoming; 12: Laboratoire des Sciences du Climat et de l'Environnement, CEA-CNRS-UVSQ, Gif-sur-Yvette, France; 13: NASA Goddard Institute for Space Studies and Columbia Earth Institute, New York, NY, USA; 14: Norwegian meteorological institute; 15: Met Office, Exeter, UK; 16: Earth and Ocean Sciences, Nicholas School of the Environment, Duke University, Durham, NC 27708

Aim

Evaluate measured and modelled accumulation-mode aerosol lifetimes.

Measurements

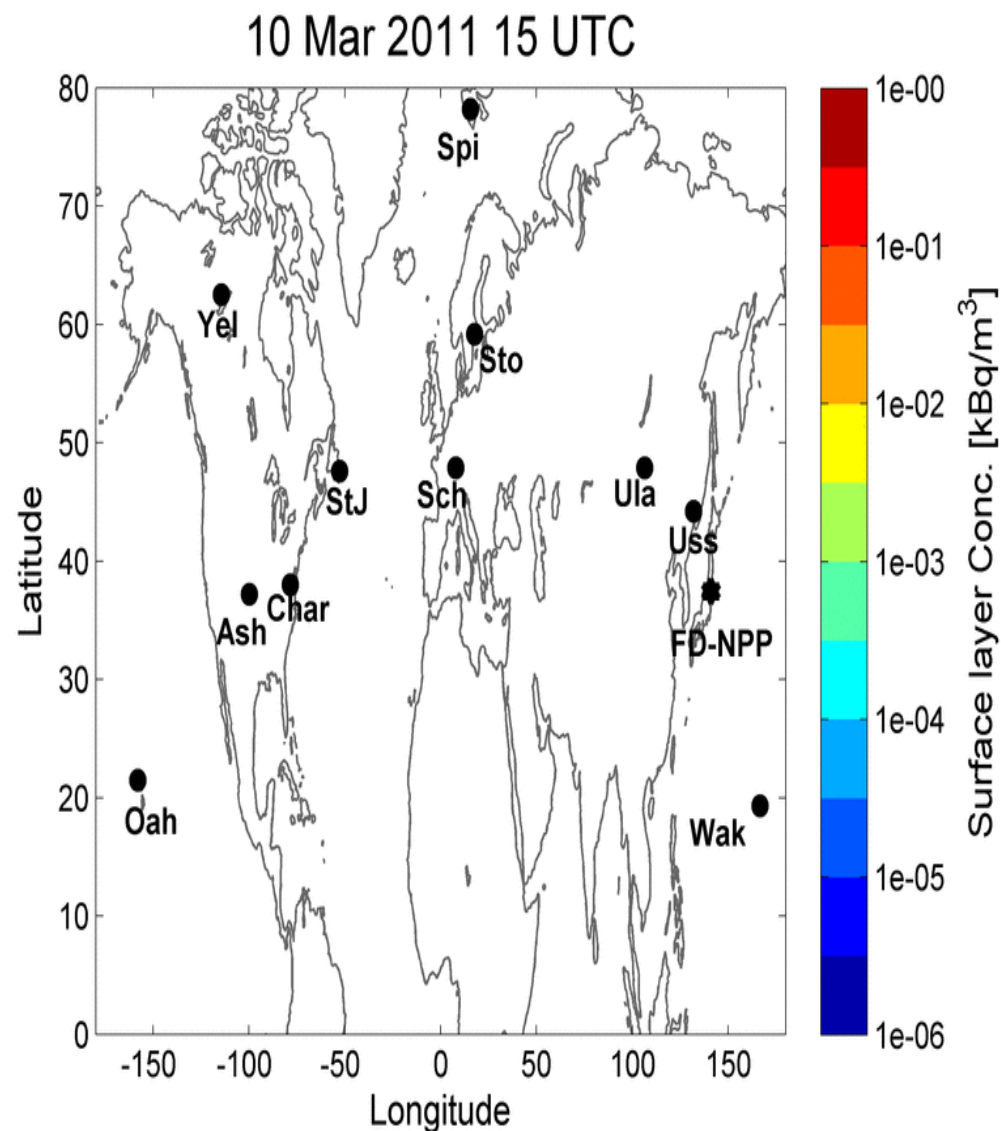
CTBTO station data of radioactive isotopes (aerosol-bound cesium, passive tracer xenon) released during the Fukushima accident of March 2011.

Models

19 global models simulated the transport of the radioactive isotopes using identical emissions.

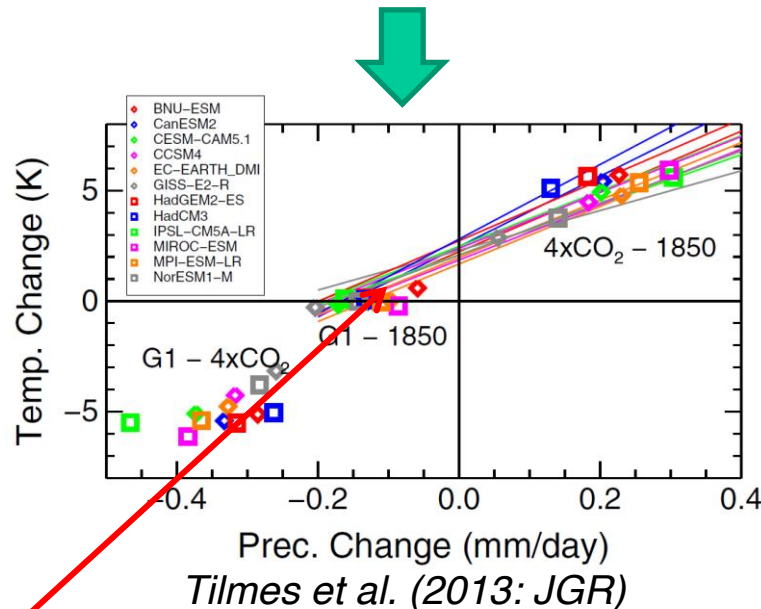
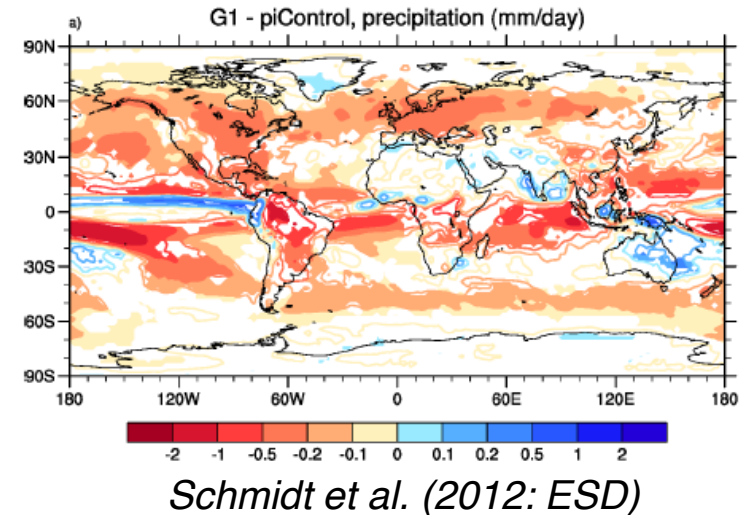
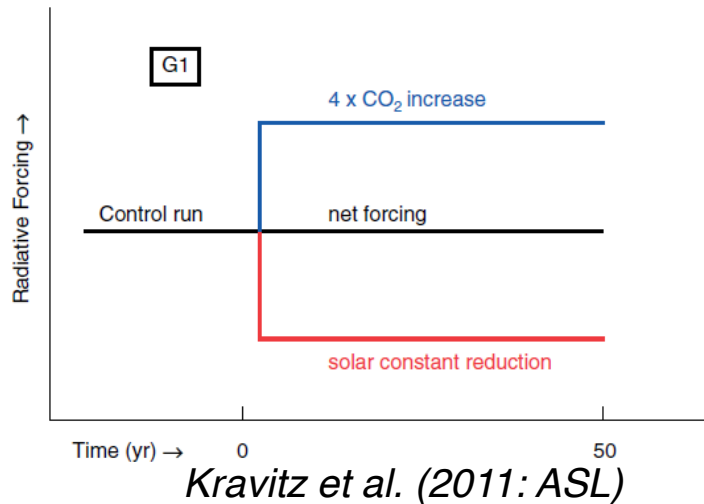
Key question

To what extent can the models reproduce the observed loss of aerosol mass with time?



Climate Engineering and the Hydrological Cycle

Jón Egill Kristjánsson (Univ. Oslo) Helene Muri (Univ. Oslo), Hauke Schmidt (MPI-M)



GHG&AP

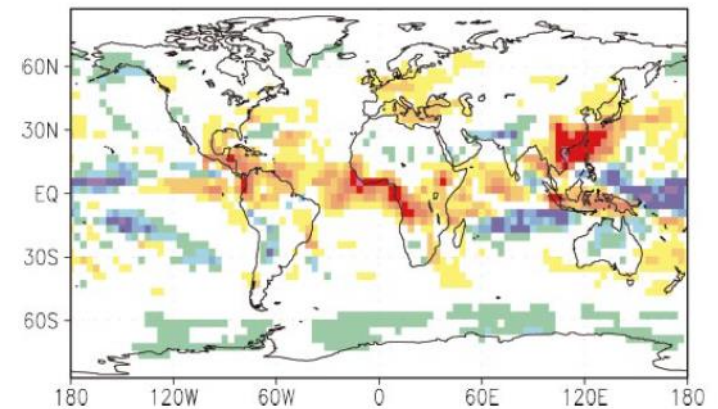
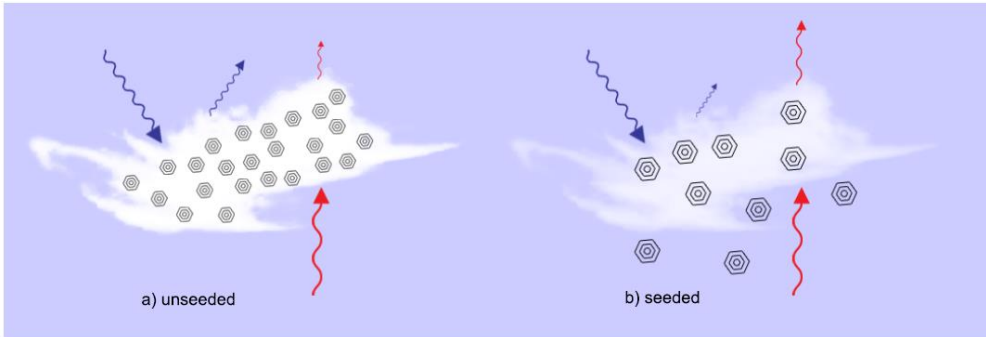


FIG. 5. Annual mean precipitation differences (mm day⁻¹) between PD and PI

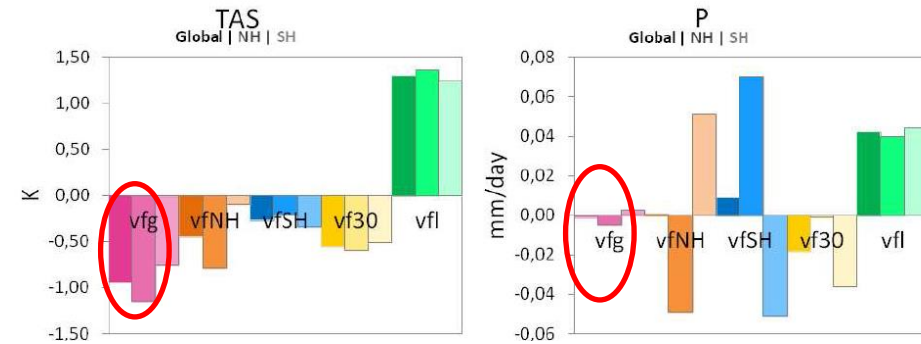
Feichter et al. (2004: J.Climate)

A ~5% reduction in P for $\Delta T=0$

Cirrus Cloud Thinning

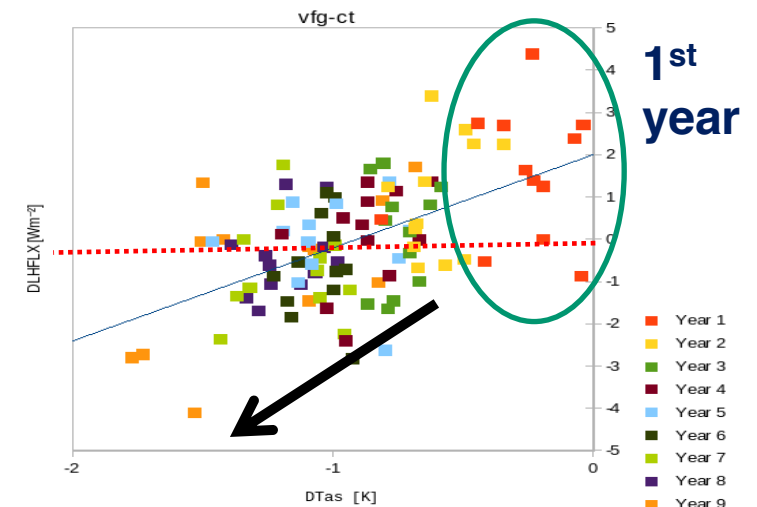


Storelvmo et al. (2013: GRL)



Muri et al. (2014: JGR)

- **CCT => Cooling, but no reduction in precipitation. Why?**
- **Fast feedbacks** dominated by the nature of the forcing: Enhanced Latent Heat Flux in 1st year
- **Cirrus Cloud Thinning cools the troposphere => Latent Heat Flux enhanced**

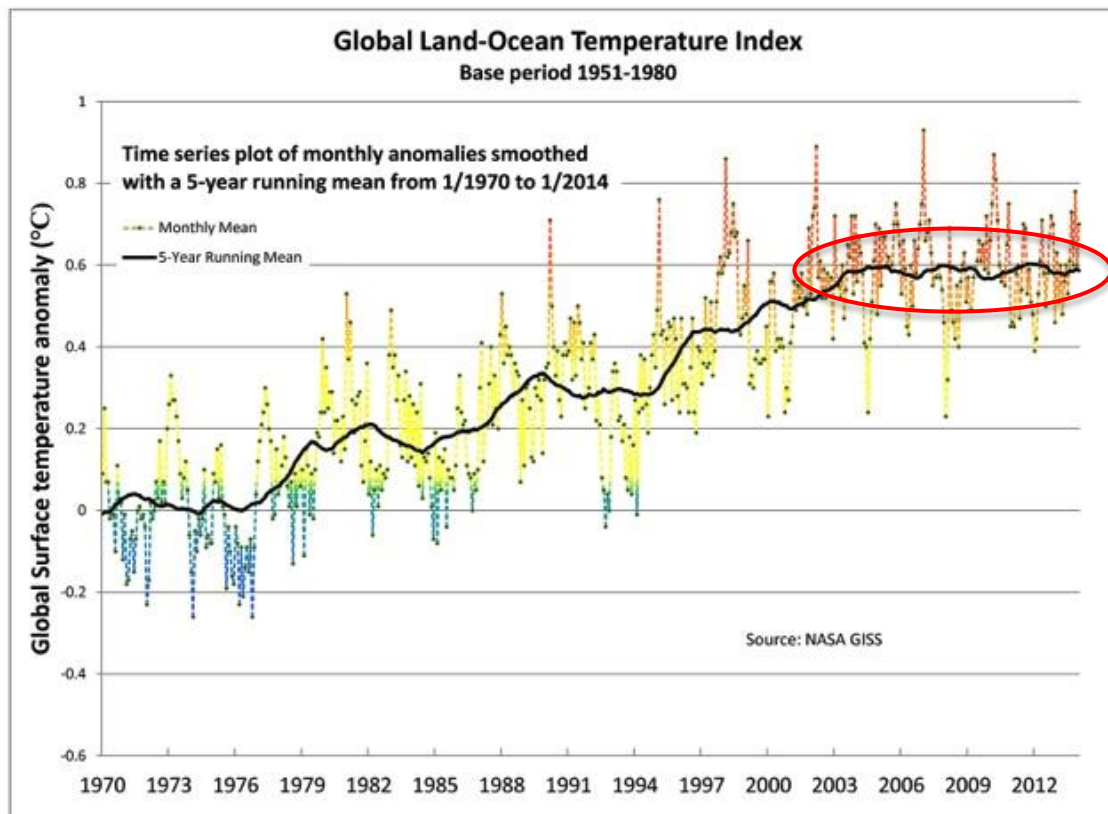


$$LH + SH = F_{SW+LW,surf}^{\downarrow} - F_{SW+LW,surf}^{\uparrow} - (F_{SW,TOA}^{\downarrow} - F_{SW+LW,TOA}^{\uparrow})$$

- Different CE techniques have **very different influences** on the hydrological cycle
- **Cirrus Cloud Thinning: Avoids suppression of the hydrological cycle**

Aerosol climate impact and its regional modulations in the 2000ies

Thomas Kühn, A.-I. Partanen, A. Laakso, Z. Lu, T. Bergman, S. Mikkonen, H. Kokkola, H. Korhonen, P. Räisänen, D. Streets, S. Romakkaniemi, A. Laaksonen

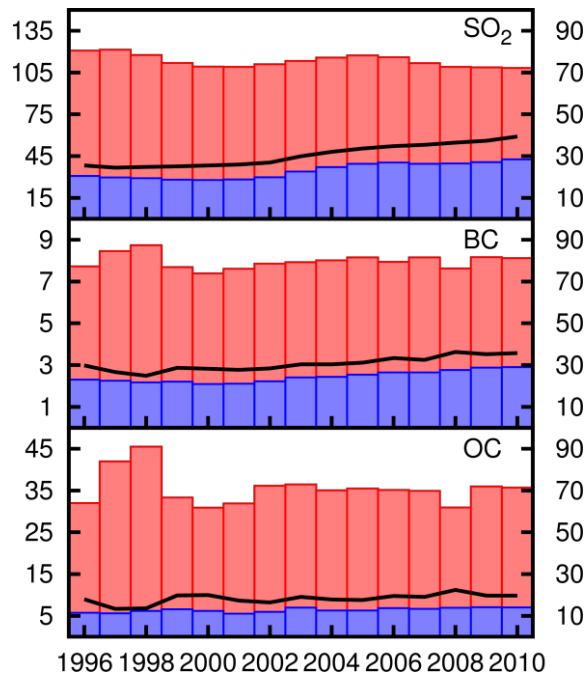


global warming "hiatus"

- ocean heat content
- solar variability
- ?asian aerosols?
- ...

Aerosol climate impact and its regional modulations in the 2000ies

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Aerosol climate impact and its regional modulations in the 2000ies

Thomas Kühn^{1,2}, A.-I. Partanen³, A. Laakso³, Z. Lu⁴, T. Bergman³, S. Mikkonen¹, H. Kokkola³, H. Korhonen³, P. Räisänen³, D. Streets⁴, S. Romakkaniemi¹, A. Laaksonen^{1,4}
 1: University of Eastern Finland, Kuopio, Finland; 2: Finnish Meteorological Institute, Helsinki, Finland;
 3: Finnish Meteorological Institute, Kuopio, Finland; 4: Decision and Information Sciences Division, Argonne, USA

Introduction

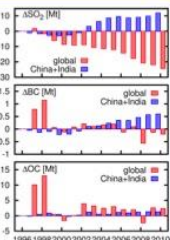
Increased anthropogenic aerosol emissions in South and East Asia have been suggested as one possible cause for the hiatus in global warming during the last 15 years. While European and North American aerosol emissions have continuously decreased since the 1980s, emissions in China and India have started increasing at the same time and, although total global aerosol emissions have decreased, aerosol effects on the global energy budget are expected to enhance towards the equator.

Setup

Our aerosol emission inventories are composed of the inventories by Lu et al. for China and India and the AeroComII-ACCMIP inventories [Riahi et al. 2011]. All other anthropogenic influences are kept fixed between simulations.

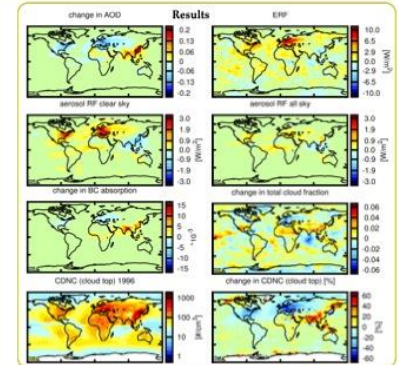
The graph to the right shows yearly changes in anthropogenic aerosol emissions since 1996 both globally and in China and India alone.

We used the aerosol-climate model ECHAM5-HAM2 to simulate the contribution of anthropogenic aerosols to the global energy budget between 1996 and 2010.



References

Bergman, T., A. Partanen, A. Laakso, Z. Lu, T. Bergman, S. Mikkonen, H. Kokkola, H. Korhonen, P. Räisänen, D. Streets, S. Romakkaniemi, and A. Laaksonen (2014), Climate impact of aerosol emissions since 1980: Cooling, air quality, and climate change, *Atmos. Chem. Phys.*, 14, 2121–2139.
 Lu, Z., D. Streets, and G. He, (2011), Global aerosol emission inventories based on a high-resolution grid, *Atmos. Chem. Phys.*, 11, 1011–1027.
 Partanen, A.-I., A. Laakso, Z. Lu, T. Bergman, S. Mikkonen, H. Kokkola, H. Korhonen, P. Räisänen, and A. Laaksonen (2014), A review of contemporary high-resolution aerosol emission data, *Atmos. Chem. Phys.*, 14, 1011–1027.



Conclusions

The re-distribution of anthropogenic aerosol emissions appears to have a net warming effect (RF=+0.09 W/m², RRF=+0.06 W/m², ERF=+0.42 W/m²), making it unlikely to be the cause of the hiatus in global warming.



Planned co-ordinated experiments for the new SPARC initiative Stratospheric Sulphur and it's Role in Climate (SSiRC)

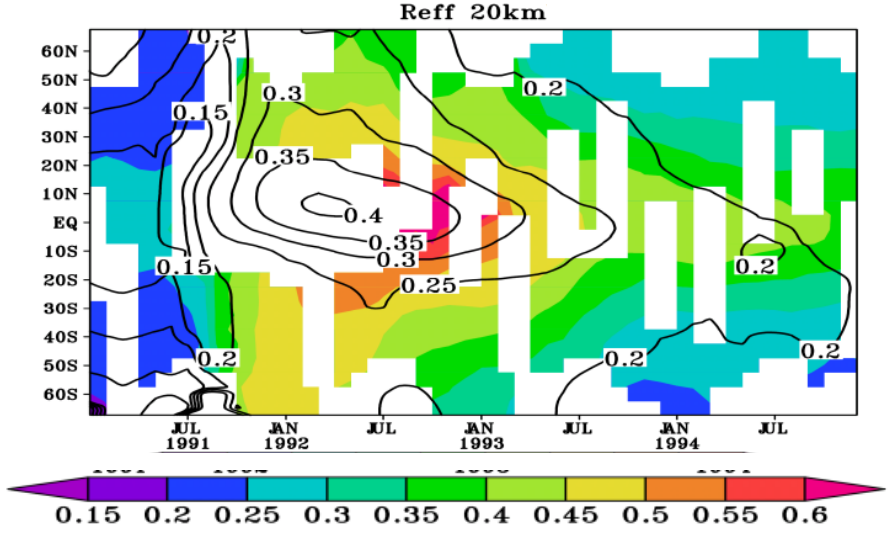
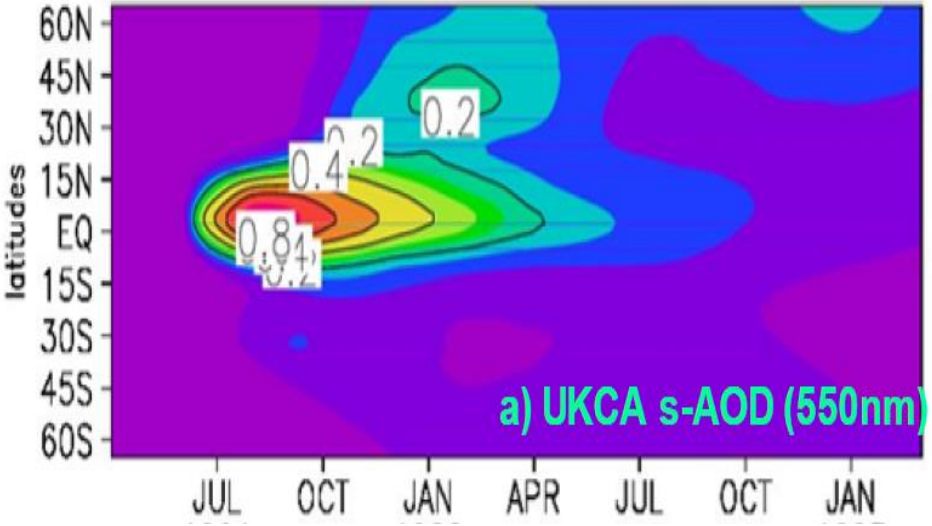
Claudia Timmreck, Graham Mann, Matt Toohey, Rene Hommel, Lindsay Lee, Valentina Aquila, Jason English, Mian Chin, Christoph Bruhl, Ryan Neely.

- New SPARC activity “Stratospheric Sulfur and its Role in Climate” (SSiRC) initiated to better understand changes in stratospheric aerosol and its precursor gaseous sulphur species
- One element of SSiRC is an intercomparison of A-GCMs which have interactive stratospheric aerosol modules
- Three co-ordinated experiments planned to intercompare background stratospheric aerosol, the perturbation through the Pinatubo period and the transient record between 1998 and 2013.
- Pinatubo experiment involves each model running perturbed physics ensemble with emulators used to quantify uncertainty in a range of key stratospheric aerosol properties and associated radiative forcings.

PinatubO Emulation in Multiple models (POEMS)

Quantify & attribute uncertainty via Gaussian emulation

Graham Mann, Ken Carslaw, Lindsay Lee, Sandip Dhomse et al. (Univ. Leeds, UK)

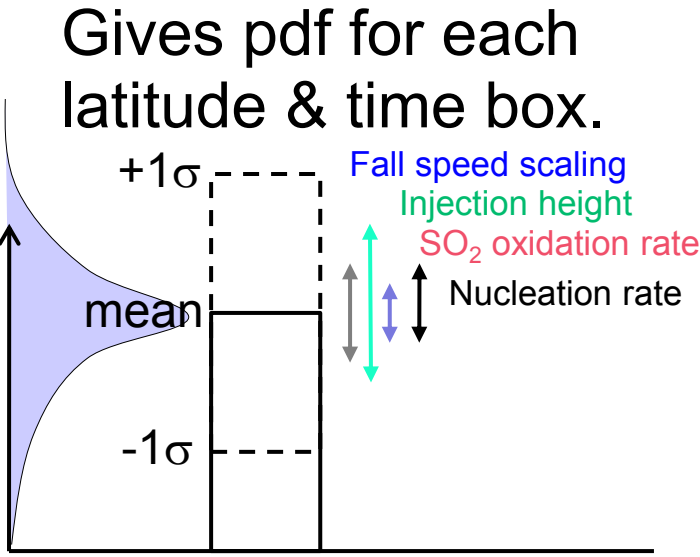


New statistical approach to quantify the magnitude & causes of uncertainty in Pinatubo radiative forcing predicted by stratospheric aerosol predicting GCMs

1. Perturbed Physics Ensemble of Pinatubo simulations with CCM

2 Use Gaussian emulators conditioned on CCM Pinatubo PPE.

3. Run full Monte Carlo of simulations with fast emulator for full variance-based sensitivity analysis.



- title: "The aerosols in the CNRM global and regional climate models"
- synthesis: evaluation and use of a prognostic aerosol scheme, derived from the GEMS/MACC scheme of the ECMWF IFS
 - ✓ main primary aerosols and sulfate; 12 added prognostic fields
 - ✓ a number of adaptations, for instance new dust emission scheme and modulation of biomass burning emissions ($\times 2$)

Simulations performed, evaluated and analysed

- ❑ with the global climate model
 - ✓ nudged and free, SST imposed, 1.4 deg (hor.), 2004, transient 1993-2012
 - ✓ AOD evaluation against satellite, MAC-v1 and AERONET monthly data

- ❑ with the regional climate model
 - ✓ coupled ocean/atmosphere, 50 km (hor.), summer 2012, transient 1980-2012
 - ✓ AOD evaluation against satellite, and AERONET monthly data over a large Mediterranean region
 - ✓ various analyses performed, e.g., analysis of the direct radiative forcing using prognostic aerosols and a climatology of these aerosols

How much brown carbon is emitted?

ECHAM6
spin-up
(3 months)

Monthly
AERONET
BrC fractions
in the
atmosphere

ITERATION TOOL

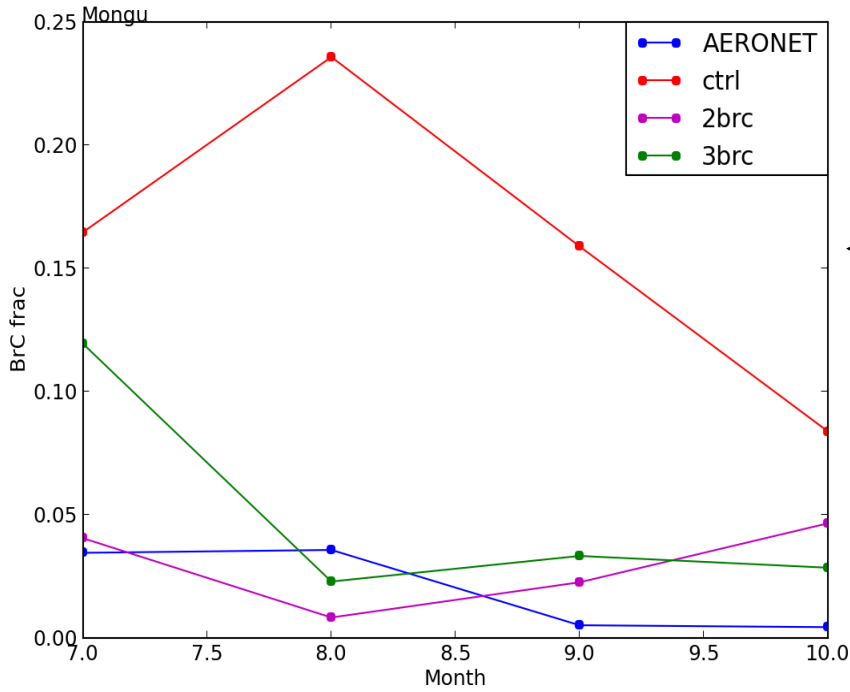
Kalman filter setup:

- number of emission sources (2/3)
- initial guess for BrC emission fraction (0.66)
- state noise std (0.15)
- perturbation size (0.01)
- std for initial guess (0.33)
- measurement noise std (0.005)

- model runs for 15 months (3 month spin-up) x number of emission sources

Monthly
BrC emission
fractions for
Biomass burning
Biogenic emissions
(Fossil fuel)

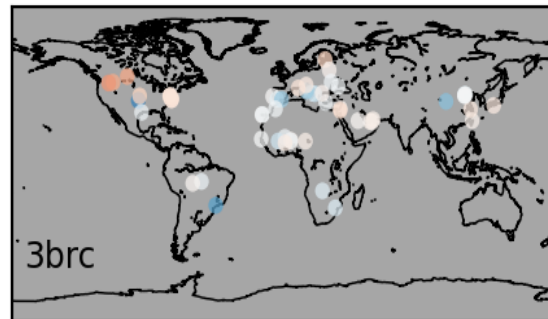
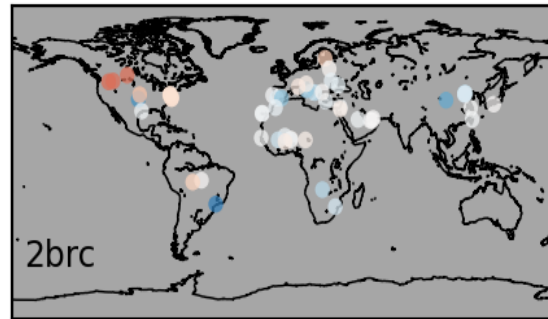
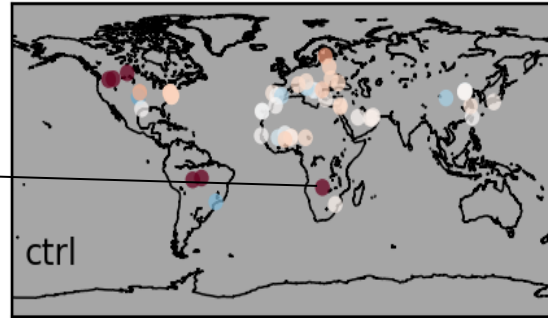
Preliminary results



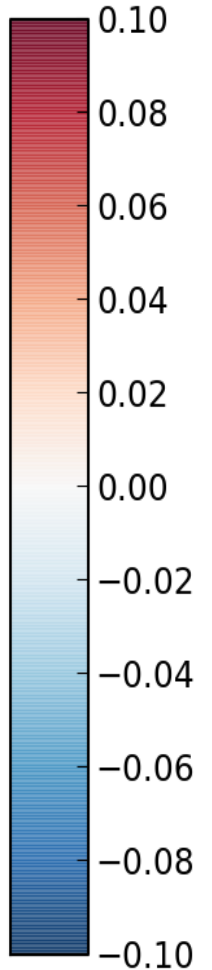
BrC emission fractions

- ctrl: Fossil=0
Biomass=0.66
Biogenic=0.66
- 2brc: Fossil=0
Biomass=optimized
Biogenic=optimized
- 3brc: Fossil=optimized
Biomass=optimized
Biogenic=optimized

model - AERONET BrC fraction
200608



Absolute
difference



Comparison of C5 & C6 MODIS dark target algorithm & validation

Munchak, Levy, Mattoo & Petrenko

- MODIS C5 aerosol products extensively used by modeling community, it is well understood and characterized.
- C6 recently available for MODIS-Aqua (L2 & L3), will soon be operational for MODIS-Terra
- This poster shows major changes to algorithm and details each change's effect on AOD

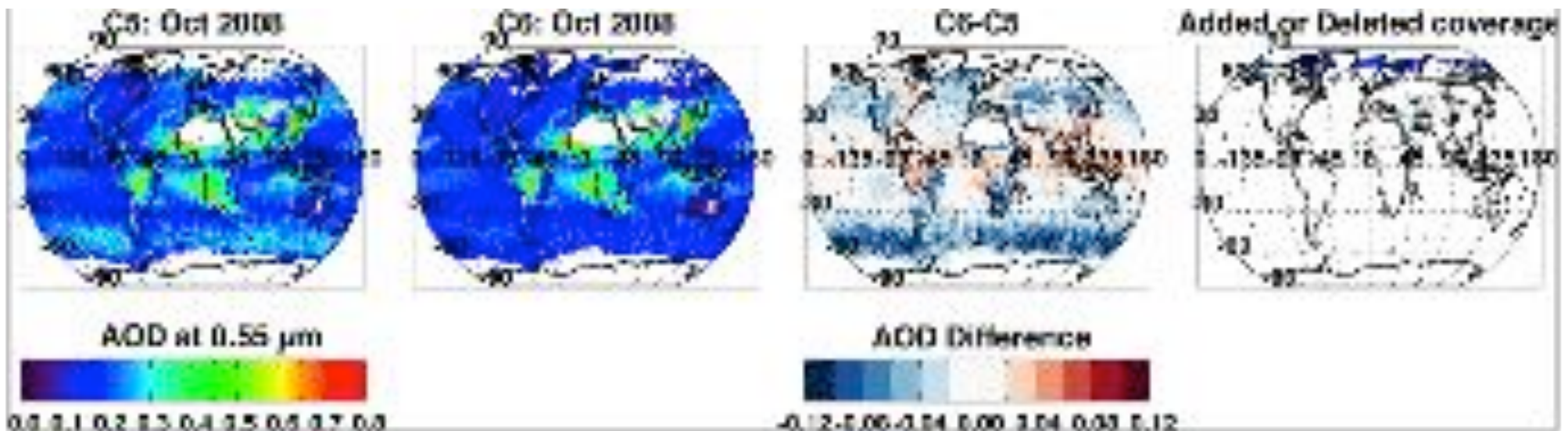
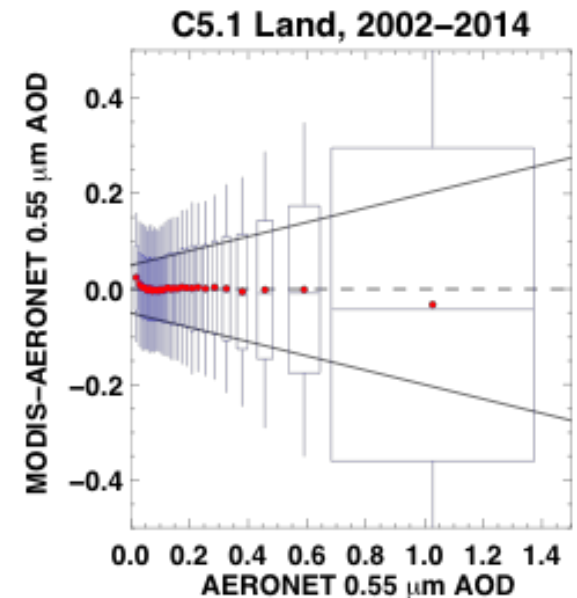
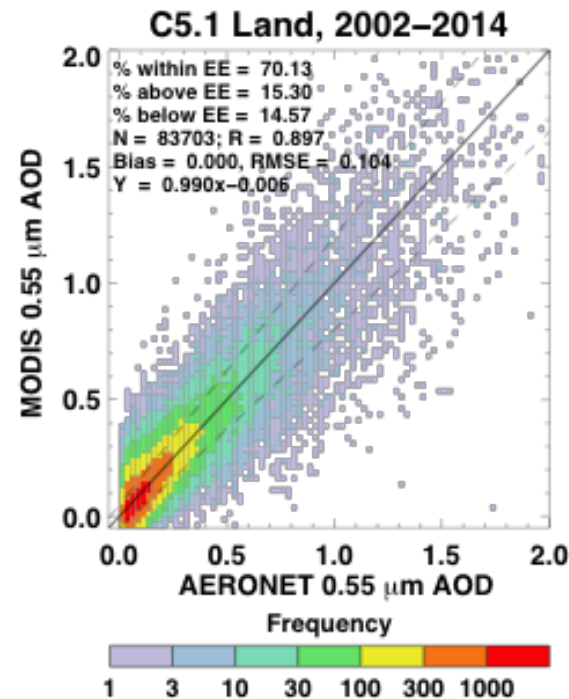
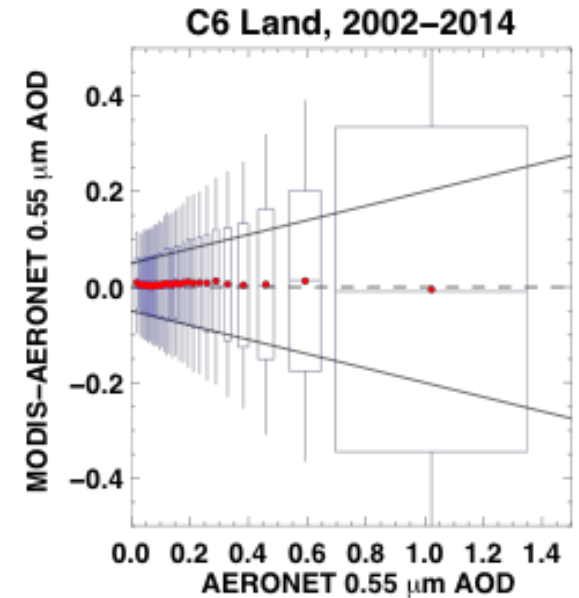
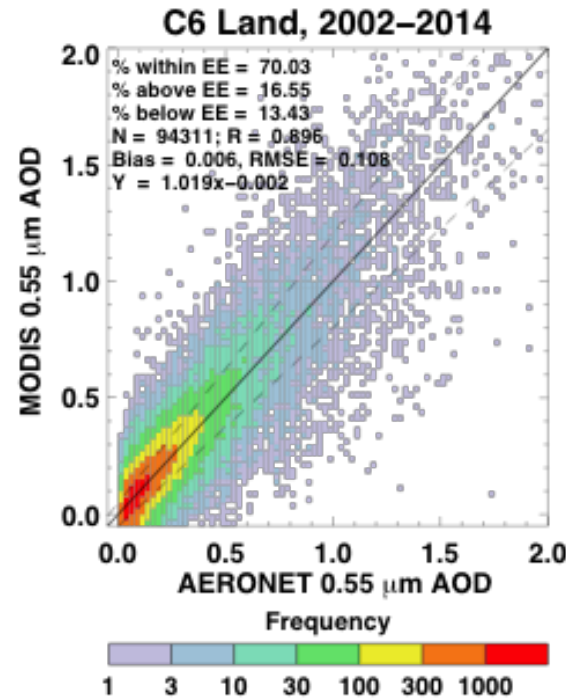


Figure from Levy et al., 2013

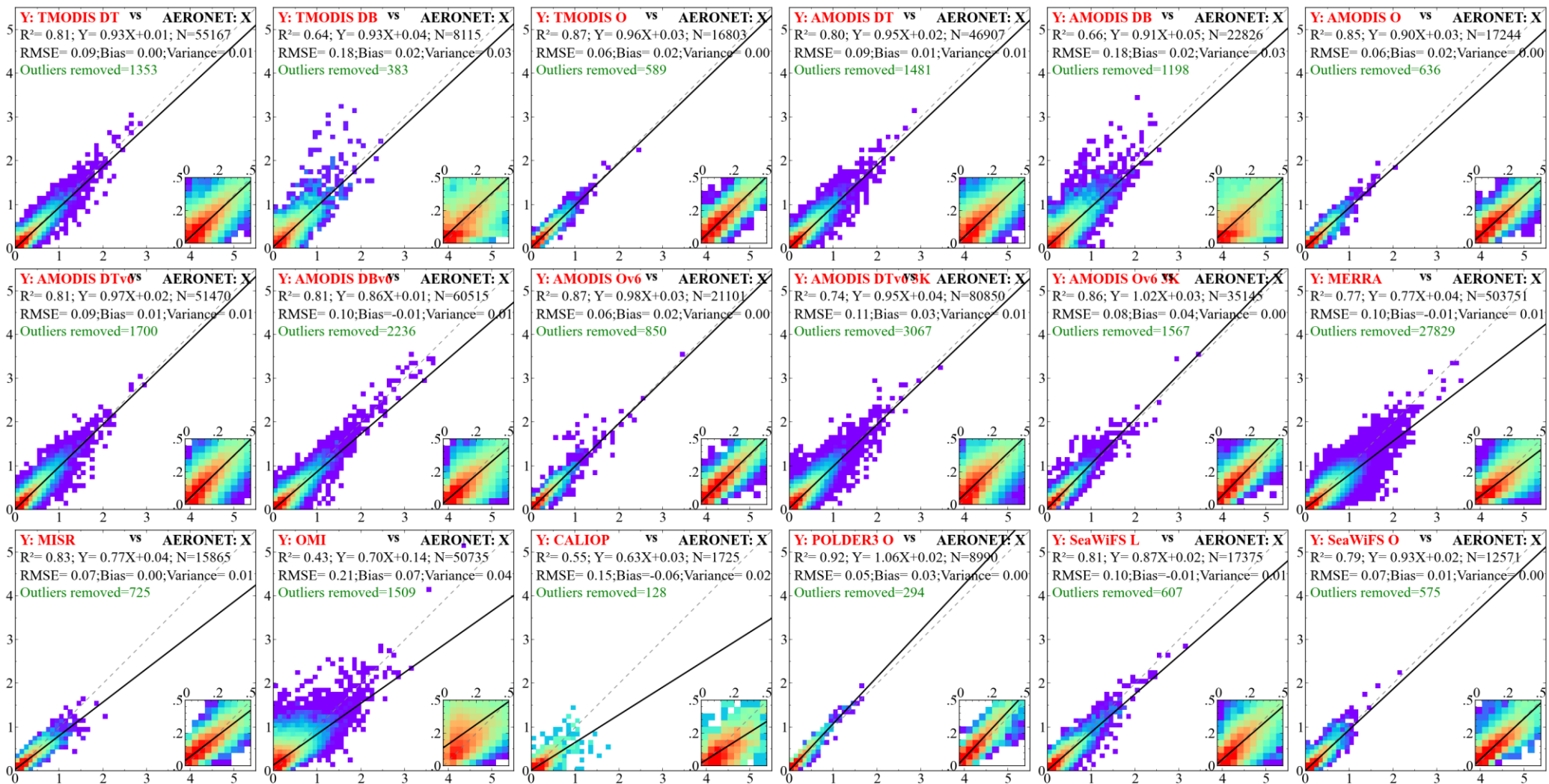
- Poster shows global validation with AERONET for both C5 & C6.
- Regional validation is shown for C6
- Curious to hear from modelers about how MODIS is currently used, and what steps are needed to transition better from C5 to C6.



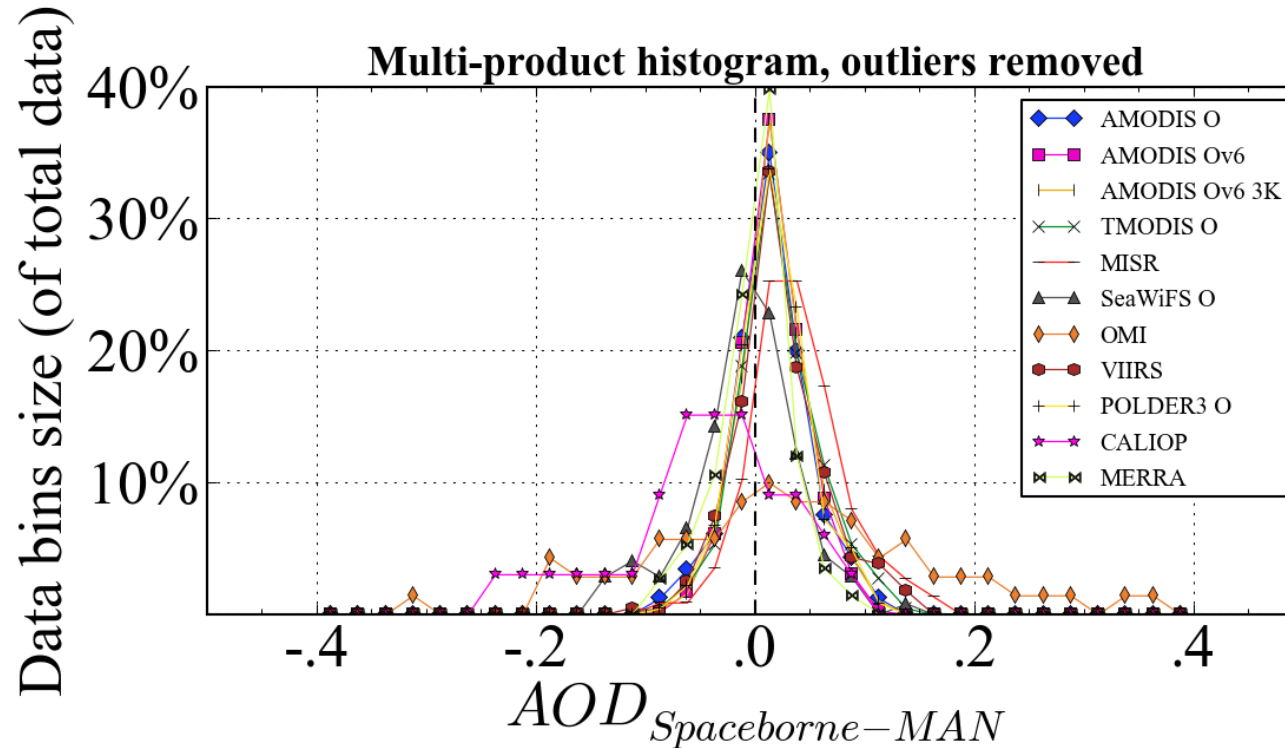
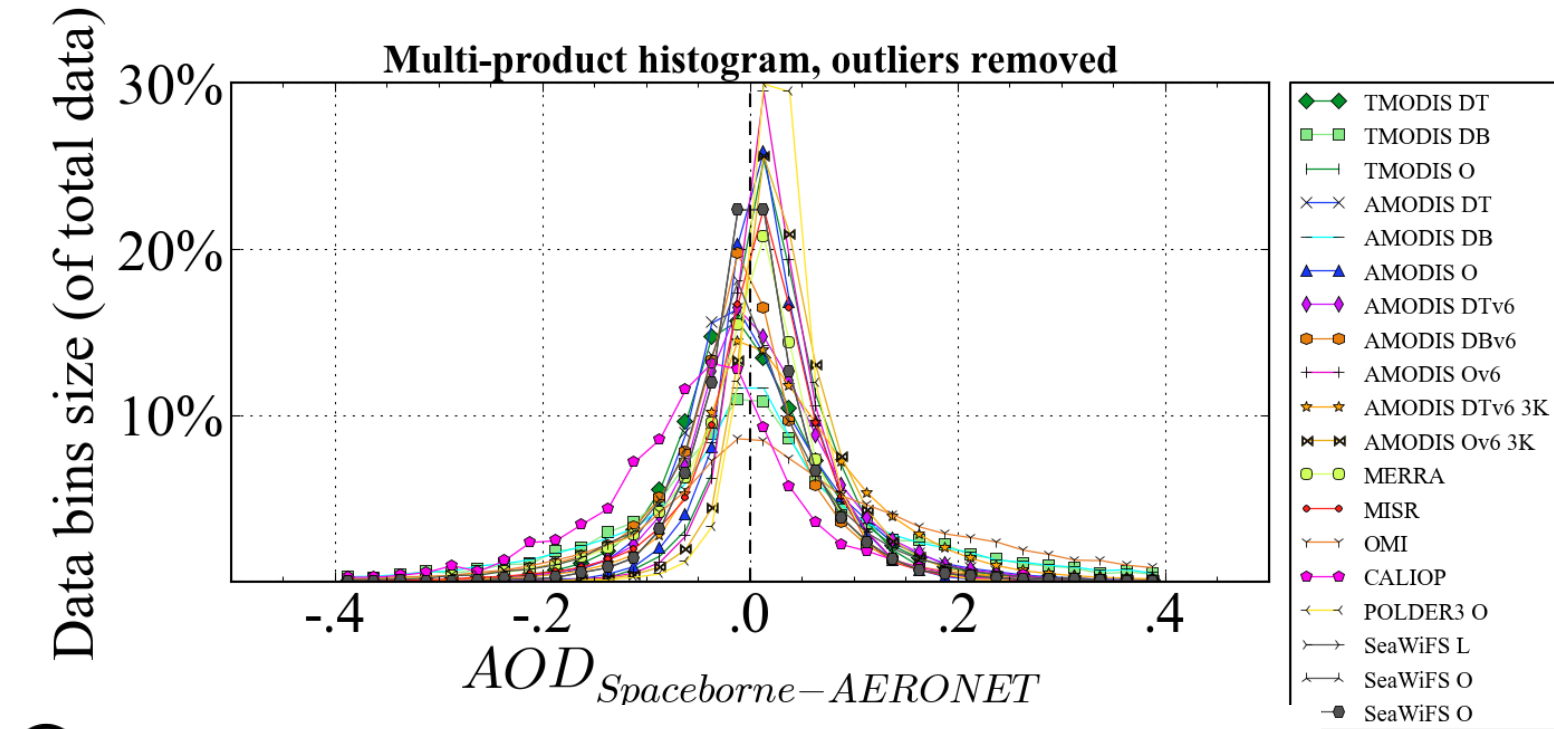
Joint Accuracy Assessment of Aerosol Retrievals from Multiple Satellite Sensors and GEOS-5 model

Maksym Petrenko, Alexander Smirnov, Charles Ichoku, Arlindo da Silva

NASA Goddard Space Flight Center, code 613, Greenbelt, MD 20771, USA.



Error Distri of Multiple Products Relative to AERONET and MAN

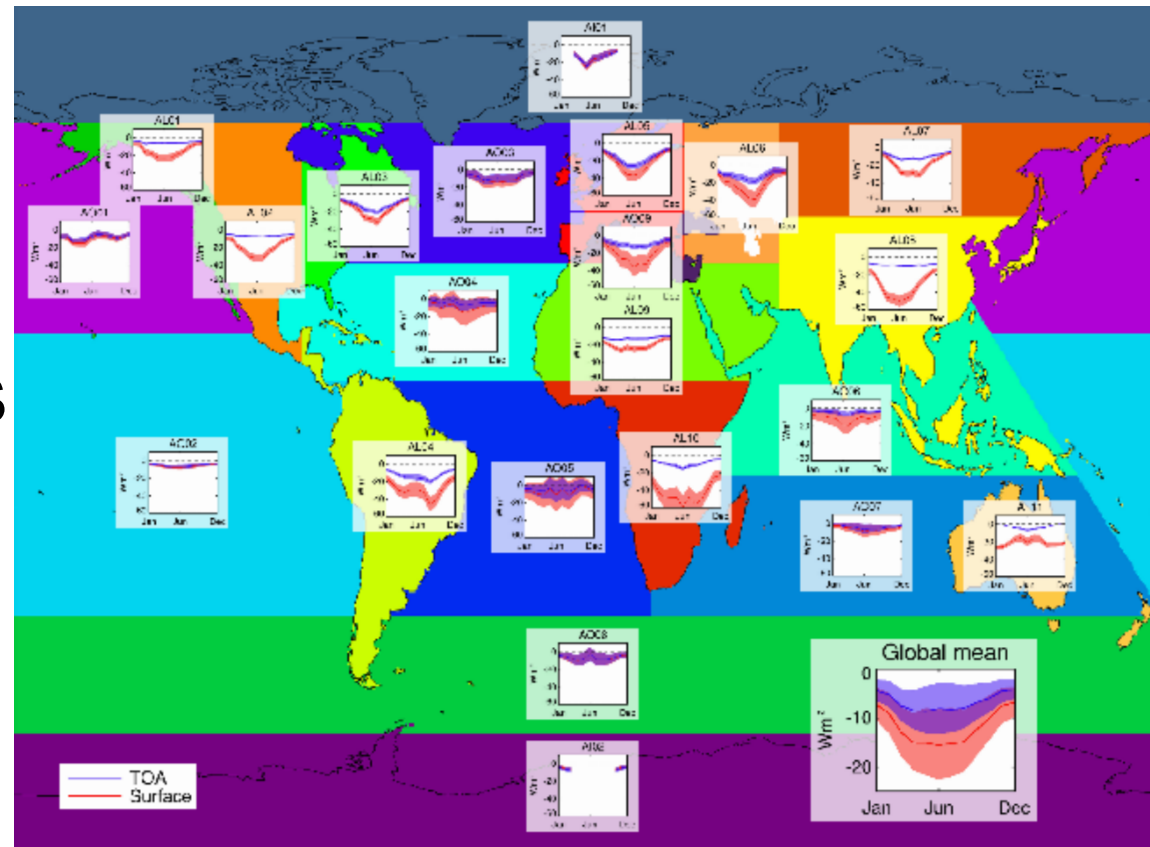


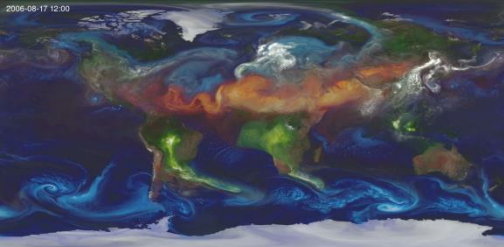
ORAC: The Optimal Retrieval of Aerosol and Cloud

- ORAC is a generalised optimal estimation scheme to retrieval cloud, aerosol, and surface properties from visible and/or infrared satellite imagers.
 - Currently supports (A)ATSR, AVHRR, MODIS, and SEVIRI
 - Additional sensors can be implemented as desired
- The algorithm has been used to produce various aerosol and cloud datasets
 - Work under way to harmonize these into a single retrieval code

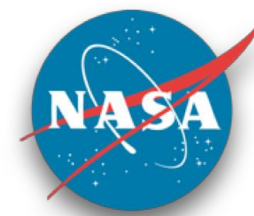
ORAC: The Optimal Retrieval of Aerosol and Cloud

- Validation of aerosol and cloud products produced within the ESA Climate Change Initiative presented
- GlobAEROSOL product was used to estimate the aerosol direct effect over land and sea regions globally





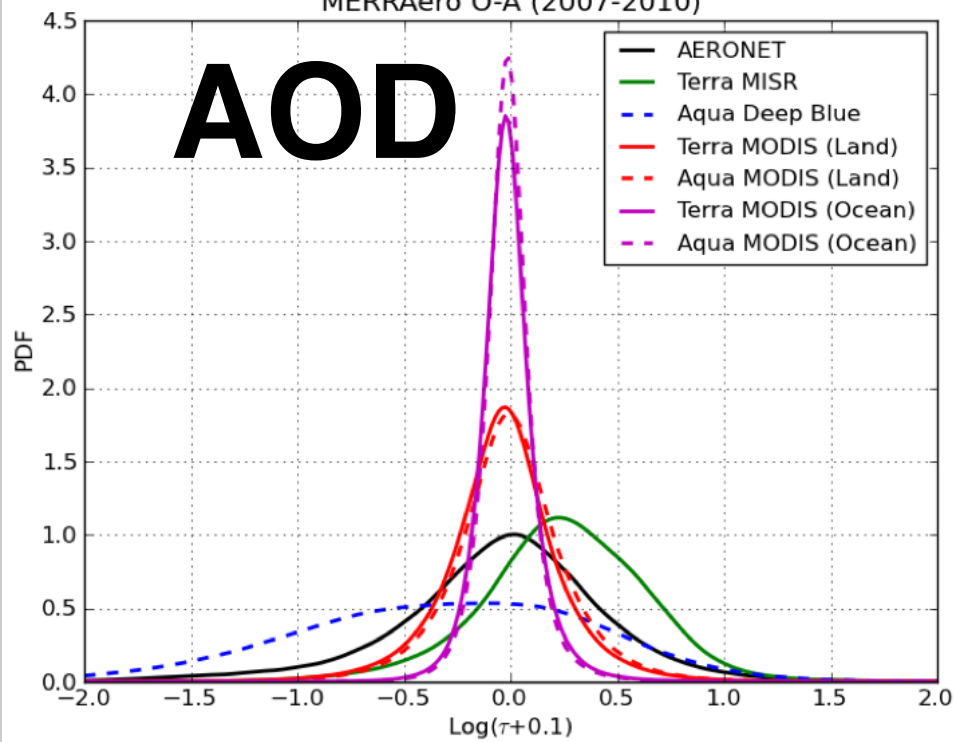
C. A. Randles, V. Buchard, P. R. Colarco, A. da Silva, A. Darmenov, E. Nowottnick, V. Aquila², and R. Govindaraju



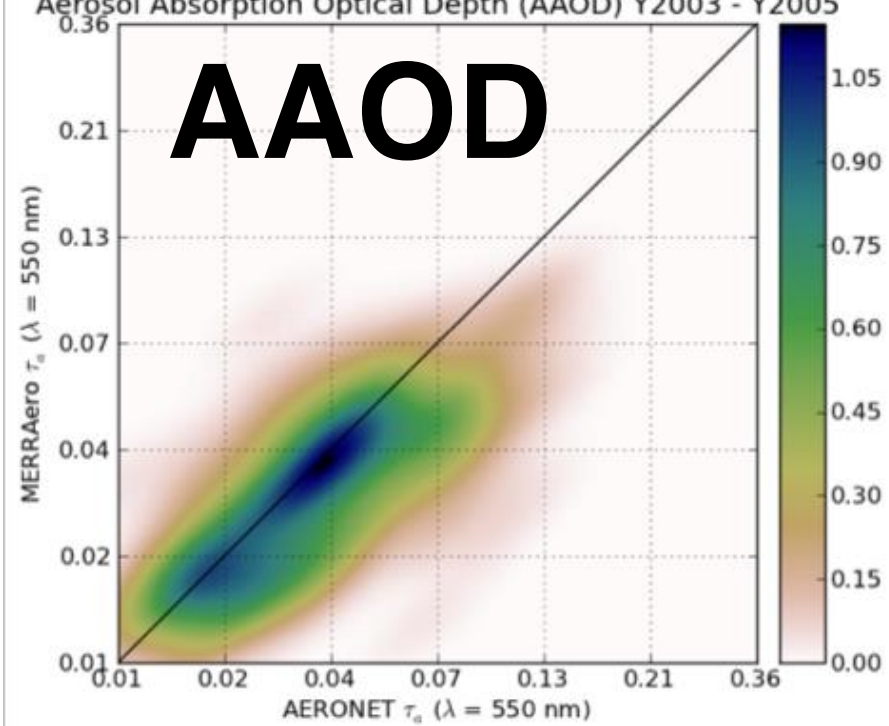
The MERRAero Aerosol Reanalysis: Evaluation and Climate Study Applications

Feature	Description
Model	GEOS-5 Earth Modeling System (w/ GOCART) Constrained by MERRA Meteorology (Replay) Land sees obs. precipitation (like MERRA <i>Land</i>) Driven by QFED daily Biomass Emissions
Aerosol Data Assimilation	Local Displacement Ensembles (LDE) MODIS reflectances AERONET Calibrated AOD's (550 nm Neural Net) Stringent cloud screening
Period	mid 2002-present (Aqua + Terra) 2000-mid 2002 (Terra only)
Resolution	Horizontal: nominally 50 km Vertical: 72 layers, top ~85 km
Aerosol Species	Dust, sea-salt, sulfates, organic & black carbon

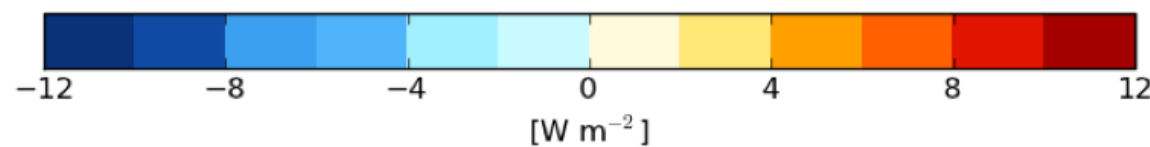
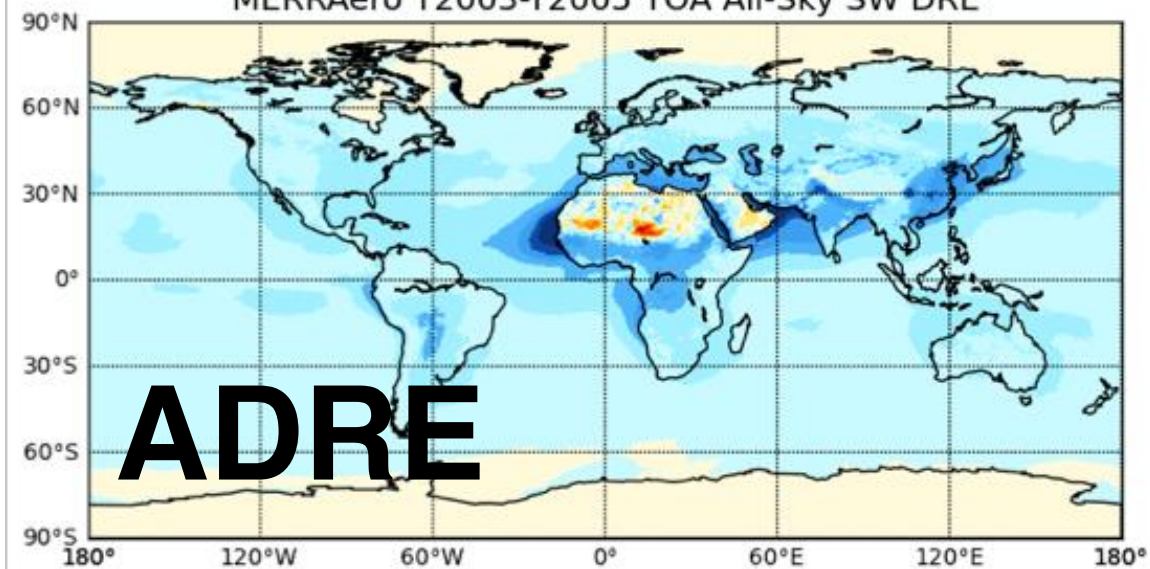
MERRAero O-A (2007-2010)



Aerosol Absorption Optical Depth (AAOD) Y2003 - Y2005



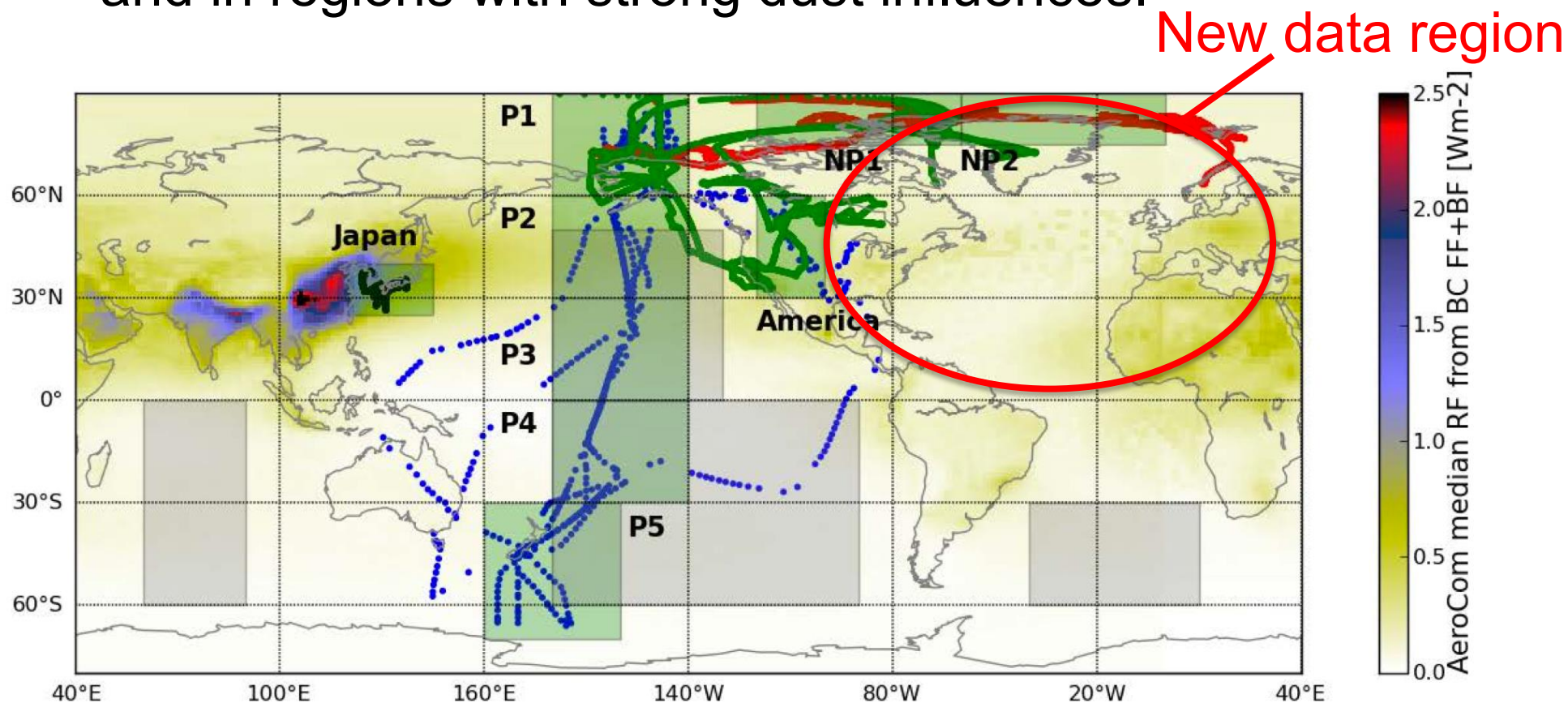
MERRAero Y2003-Y2005 TOA All-Sky SW DRE



AeroCom/BC Measurement Comparison

J. P. Schwarz and B. Weinzierl

Samset et al., 2014, Bond et al., 2013, and Koch et al., 2009, highlight the need for continued evaluation of BC MMR close to source regions (and at altitude), and in regions with strong dust influences.

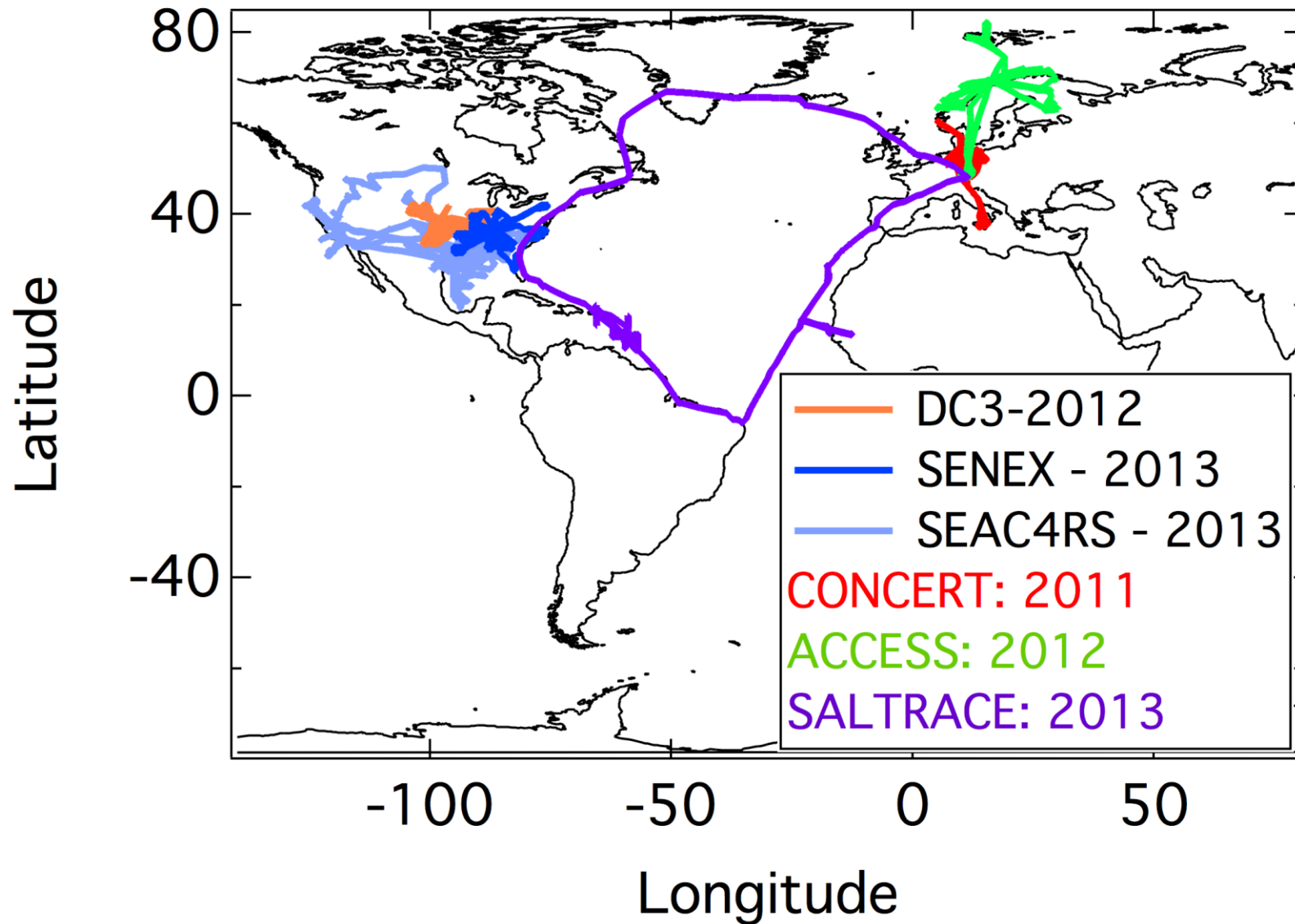


Samet et al., ACPD, 2014

AeroCom/BC Measurement Comparison

J. P. Schwarz and B. Weinzierl

Multiple SP2 data sets obtained in
DLR/NOAA/NASA Campaigns.



Estimating Anthropogenic Aerosol Indirect Effects Through Cirrus Clouds using CAM5.1 with Different Ice Nucleation Parameterizations

Xiangjun Shi¹, Xiaohong Liu¹, Kai Zhang²

1. University of Wyoming
2. Pacific Northwest National Laboratory

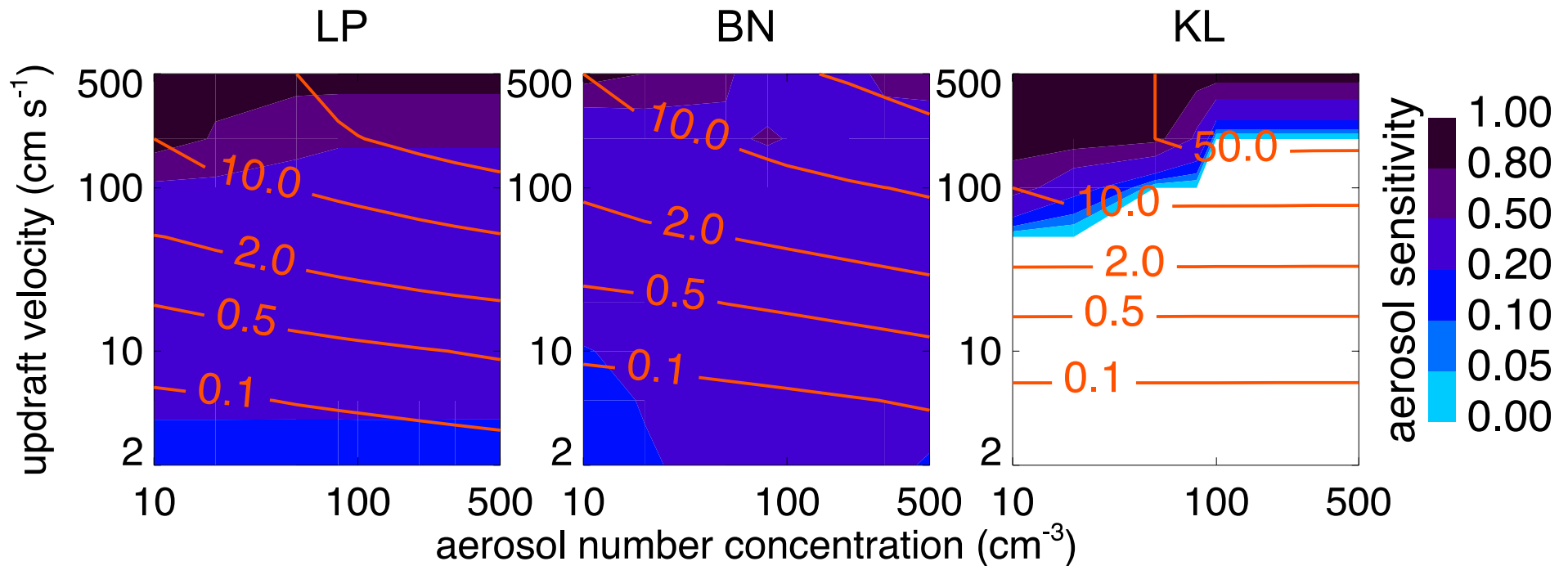
Anthropogenic Aerosol Indirect Effects through cirrus clouds on climate.

Names	CF	LWCF	SWCF	IWP	LWP	CDNI	CDNC	PRECC	PRECL	PRECT
LP _{PD}	-27.84	25.86	-53.70	19.19	45.32	1.50	1.40	2.07	0.89	2.96
BN _{PD}	-27.81	25.46	-53.27	18.77	45.09	1.45	1.39	2.08	0.89	2.97
KL _{PD}	-28.15	25.06	-53.21	18.52	45.20	1.50	1.41	2.08	0.89	2.97
LP _{PI}	-27.97	25.50	-53.47	18.99	45.26	1.41	1.40	2.07	0.89	2.96
BN _{PI}	-27.91	25.13	-53.04	18.58	45.01	1.37	1.40	2.08	0.89	2.97
KL _{PI}	-28.02	25.01	-53.03	18.53	45.12	1.47	1.40	2.09	0.89	2.98
Δ LP	0.13	0.36	-0.23	0.20	0.06	0.09	0	0	0	0
Δ BN	0.10	0.33	-0.23	0.19	0.08	0.08	-0.01	0	0	0
Δ KL	-0.13	0.05	-0.18	-0.01	0.08	0.03	0.01	-0.01	0	0

[?] Liu and Penner (2005) ;Barahona and Nenes (2009) ;Kärcher et al. (2006)

Aerosol Sensitivity Parameter (η_α)

$$\eta_\alpha = d(\ln Ni) / d(\ln Na).$$

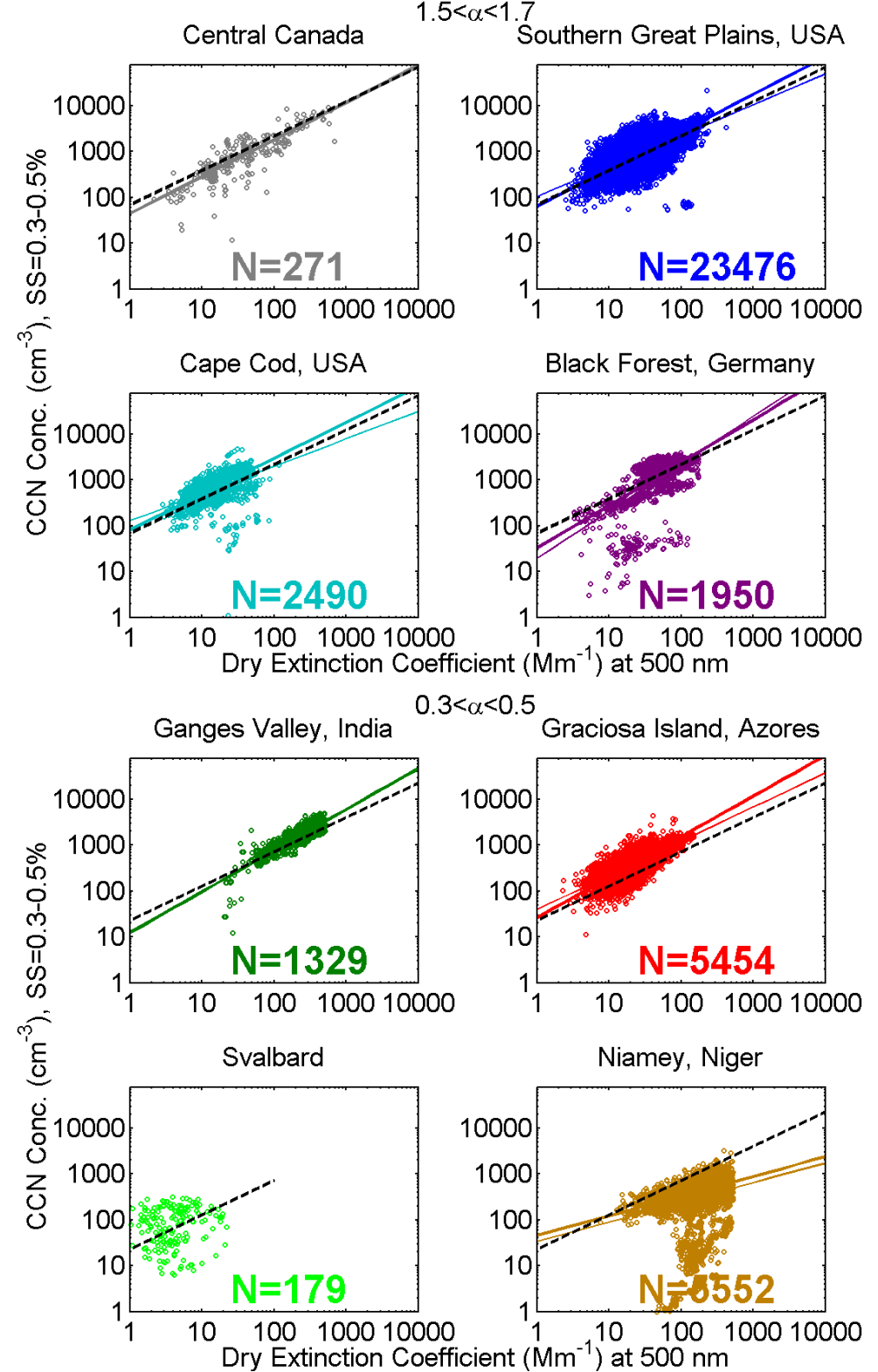


Ice crystals number concentration contoured as a function of vertical velocity and sulfate aerosol number concentration. Colors indicate the aerosol sensitivity parameter η_α . Results from pure homogeneous freezing experiments using ice nucleation parameterizations.

The relationship between CCN concentration and aerosol extinction in situ observations for dried particles

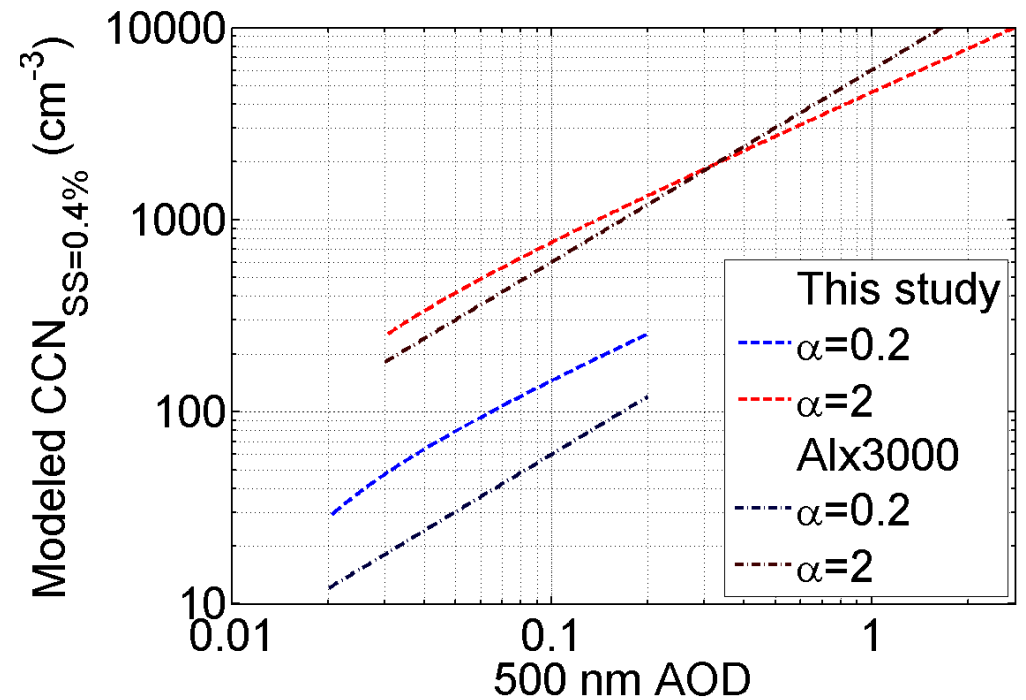
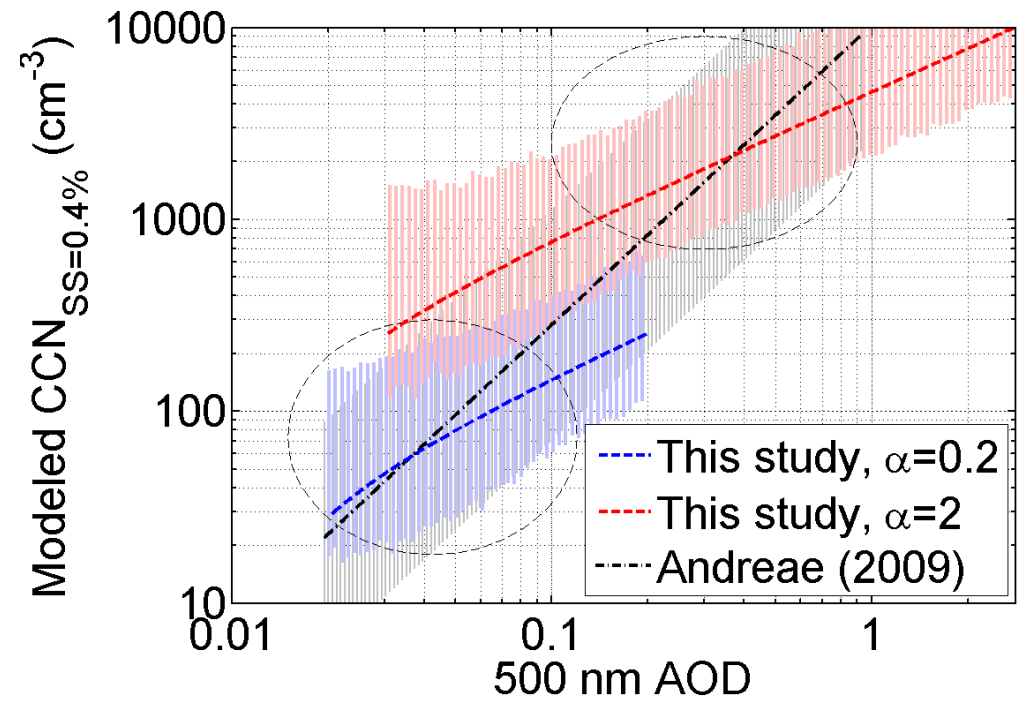
$$CCN_{SS \sim 0.4\%} (\text{cm}^{-3}) = 10^{0.4\alpha + 1.2} \sigma^{0.75}$$

σ : ext (Mm^{-1}), α =Angstrom Exp.



The relationship between CCN concentration and aerosol extinction

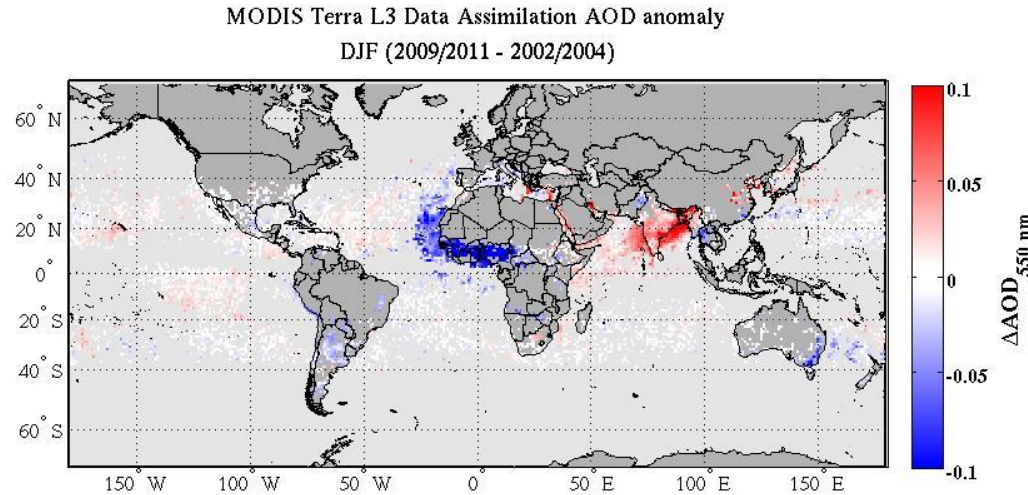
implications on satellite-based CCN estimates



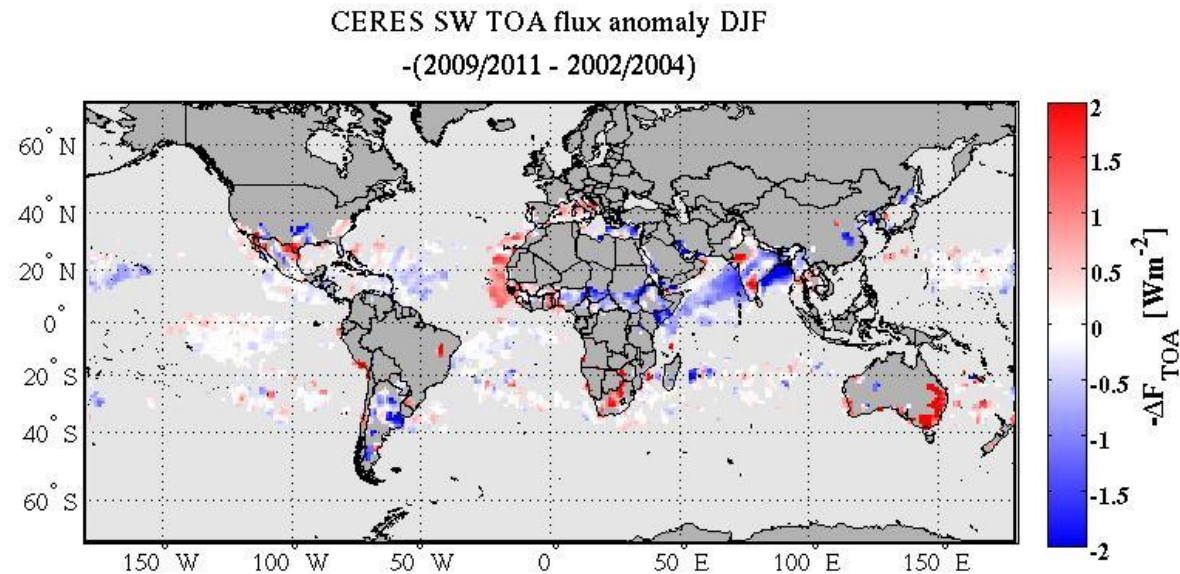
Sundström A.-M. et al.

DECADAL CHANGES IN CERES CLEAR-SKY SHORTWAVE TOA FLUXES: WHAT CAN WE SAY ABOUT AEROSOL CONTRIBUTION?

AOD anomaly
winter (DJF)
(2009/2011
2002/2004)

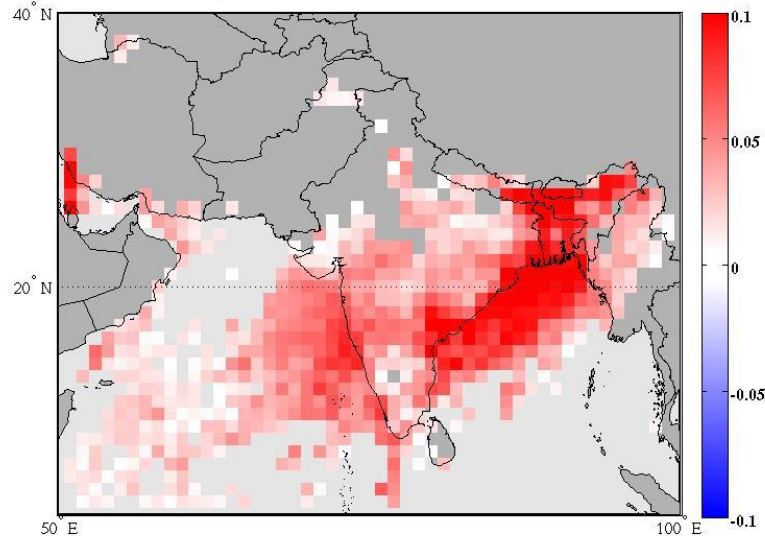


CERES clear-sky SW
TOA flux anomaly
winter (DJF)
(2009/2011
2002/2004)



MODIS AOD anomaly

MODIS L3 data assimilation AOD anomaly DJF (2009/2011 - 2002/2004)
for pixels where MODIS & MISR $\Delta AOD > 0$



Simulations

”Dark” surface:

□ AOD=+0.05

□ SW Flux_{TOA} = 0.2...3 Wm⁻²
(abs.... scatt. type)

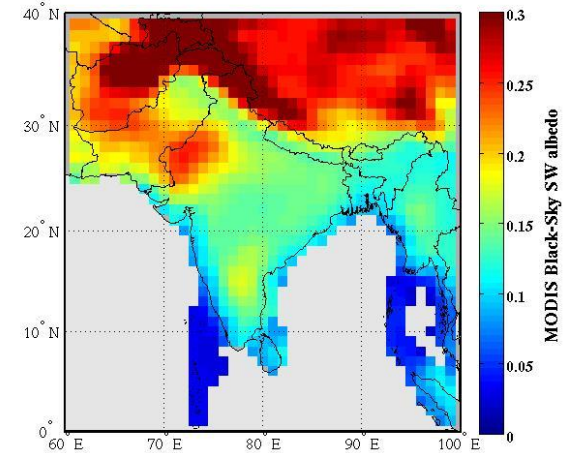
Bright surface:

AOD=+0.05

□ SW Flux_{TOA} = 0.6...3 Wm⁻²
(abs.... scatt. type)

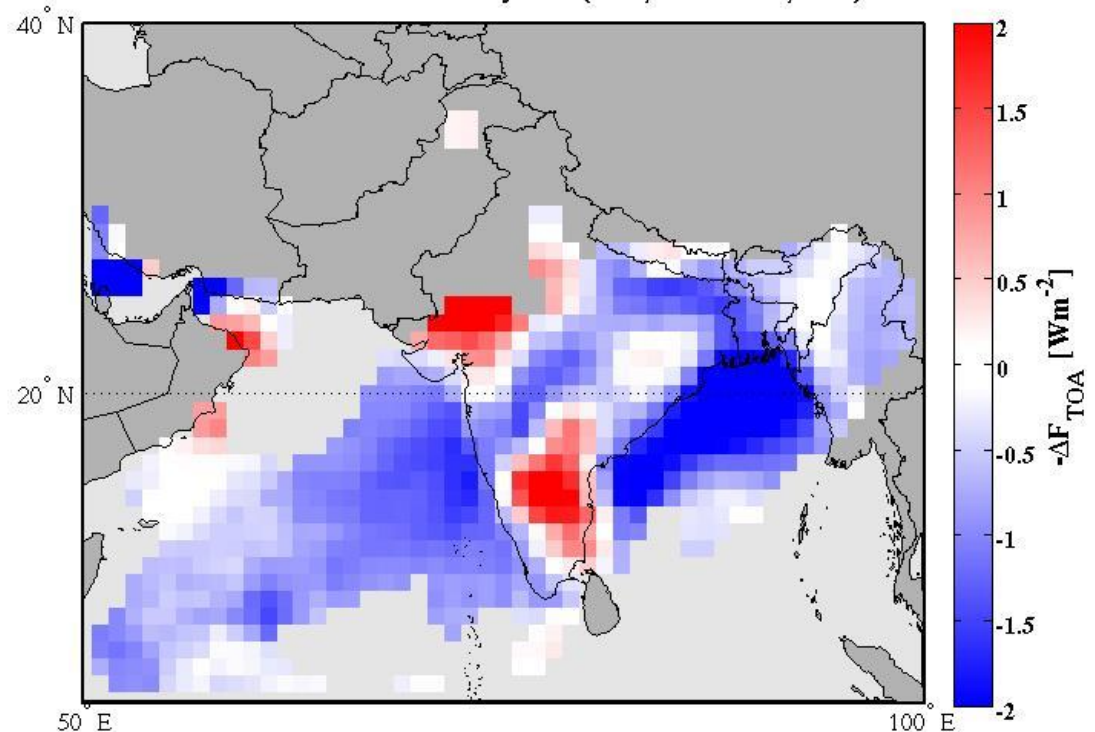
+ precipitable water & surface anomalies

MODIS Black-Sky Albedo



SW Clear-Sky TOA flux anomaly

CERES SW TOA flux anomaly DJF (2009/2011 - 2002/2004)

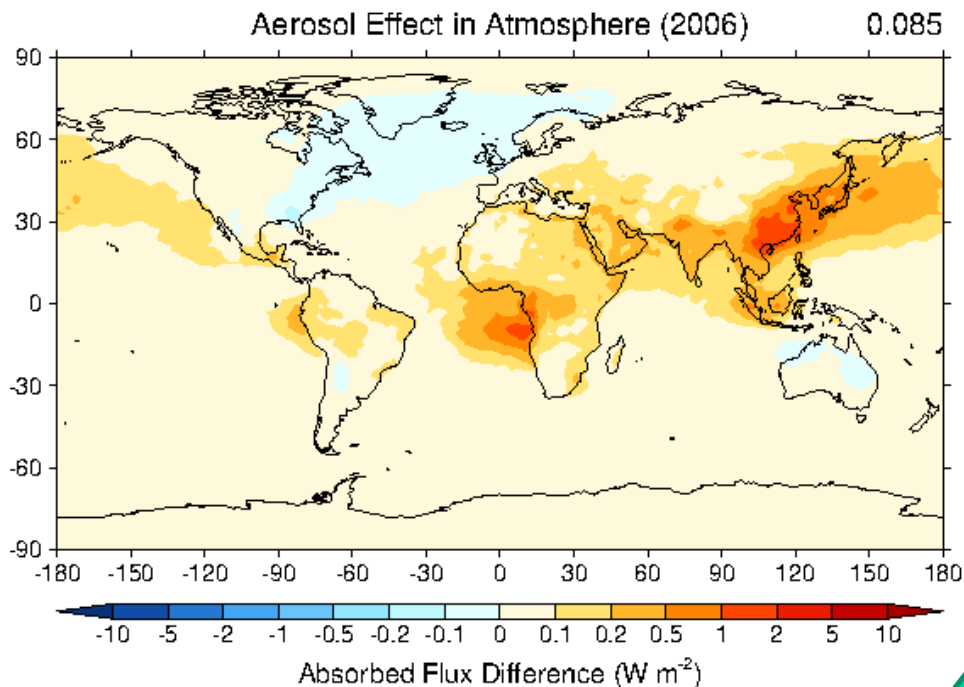


Clear-sky and all-sky direct radiative forcing estimates based on TM5 and a doubling-adding radiative transfer model using observed clouds

Michiel van Weele, Ana Ruiz-Garces, Twan van Noije, Jan-Willem Meijerink, Ping Wang, Piet Stammes

Reduce modelled uncertainties in direct aerosol effect/forcing = $(\Delta B)^2 * (\Delta M)^2 * (\Delta E)^2$

B = Burden Determined by emissions and residence times
M = Mass extinction coefficient Determined by load and water uptake
E = Radiative Efficiency Determined by clouds, radiative transfer



Approach

- (1) Chemical transport model TM5
Aerocom Phase 2 simulations
- (2) Radiative transfer model DAK
Direct radiative effect / rad. forcing
- (3) Observed clouds FRESCO/SCIAMACHY
2006

Results

- (i) Cloud impact on atm. absorption
- (ii) All sky radiative efficiency per AOD

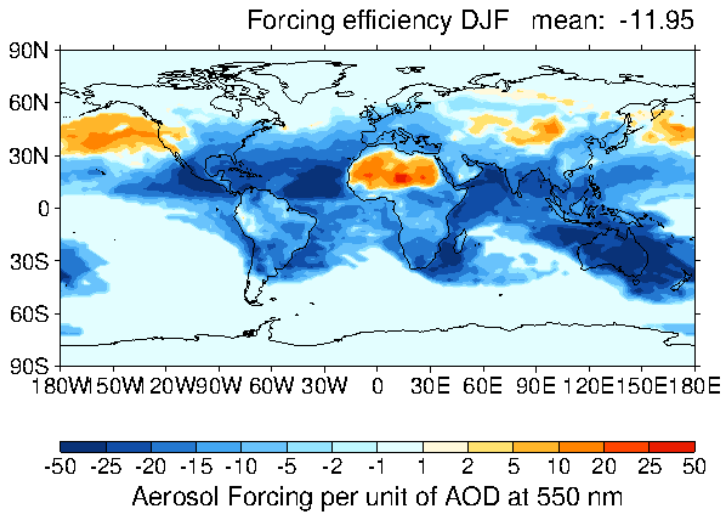
Positive (and slightly negative) effects of clouds on atm. absorption

Clear-sky and all-sky direct radiative forcing estimates based on TM5 and a doubling-adding radiative transfer model using observed clouds

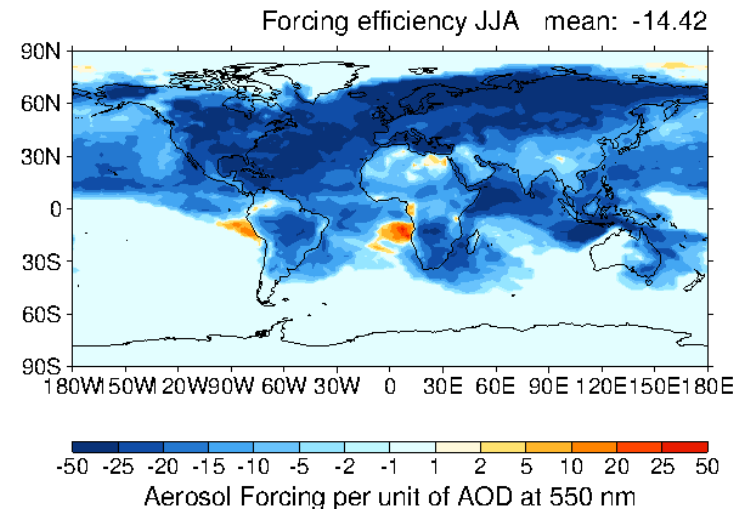
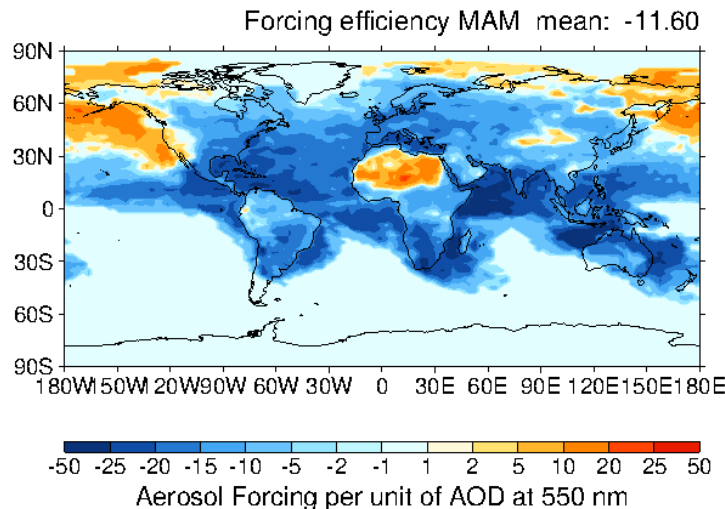
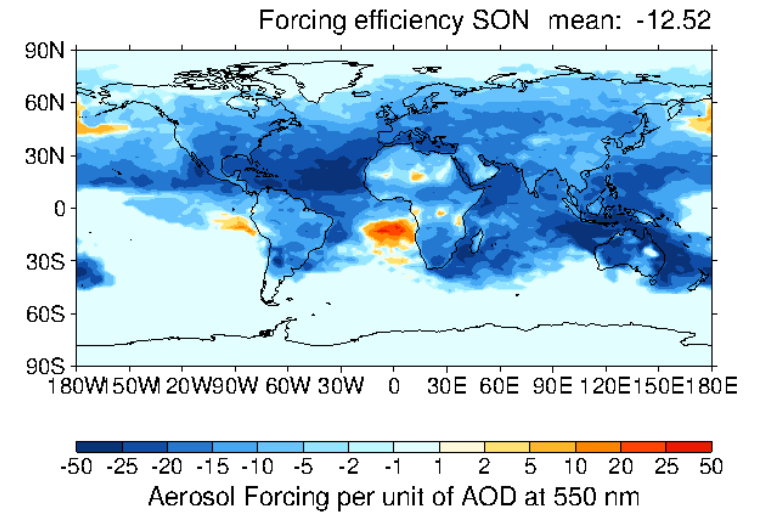
Michiel van Weele, Ana Ruiz-Garces, Twan van Noije, Jan-Willem Meijerink, Ping Wang, Piet Stammes

KNMI The Netherlands

All sky aerosol forcing efficiency per season



Annual mean
direct radiative
forcing (at TOA):
 $F \approx -12.6 \tau$
Clear sky:
 $F \approx -24.5 \tau$



Top-down Estimates of SO₂ Degassing Volcano Emissions Using In Situ SO₂ Measurements and the WRF-STILT Model, a Case Study at the Turrialba Volcano, Costa Rica

Xin Xi¹ (xin.xi@nasa.gov), Matthew S. Johnson¹, Matthew Fladeland¹, David Pieri², Jorge Andres Diaz³, Seongeun Jeong⁴, Geoff Bland⁵

¹NASA Ames Research Center; ²NASA Jet Propulsion Laboratory. ³University of Costa Rica, Costa Rica. ⁴Environmental Energy Technologies Division, ⁴Lawrence Berkeley National Laboratory ⁵Wallops Flight Facility, NASA Goddard Space Flight Center



Through a case study at the Turrialba Volcano, Costa Rica (which is assigned an extraeruptive rate in AeroCom), our **research goals** are

- 1) to develop an inverse estimate of volcanic degassing rates by applying a high-resolution receptor-oriented analysis on in situ SO₂ measurements.
- 2) to examine the impact of top-down SO₂ emission fluxes on regional-scale atmospheric compositions.

Main findings:

WRF-STILT model is able to accurately connect measurement locations and the volcanic SO₂ source.

The top-down estimate of SO₂ degassing flux from the Turrialba Volcano is higher than the AEROCOM extraeruptive (posteruptive) rate by a factor of 10⁴ (100).

Sensitivity model tests using GEOS-Chem show the top-down SO₂ flux leads to large increases in the SO₂ and SO₄²⁻ concentrations near the source and in downwind regions, which implies that using the AEROCOM inventory underestimates the natural SO₂ contribution from the Turrialba Volcano.

**Improvement of cloud microphysics in the
aerosol-climate model
BCC_AGCM2.0.1_CUACE/Aero, evaluation
against observations, and updated aerosol
indirect effect**

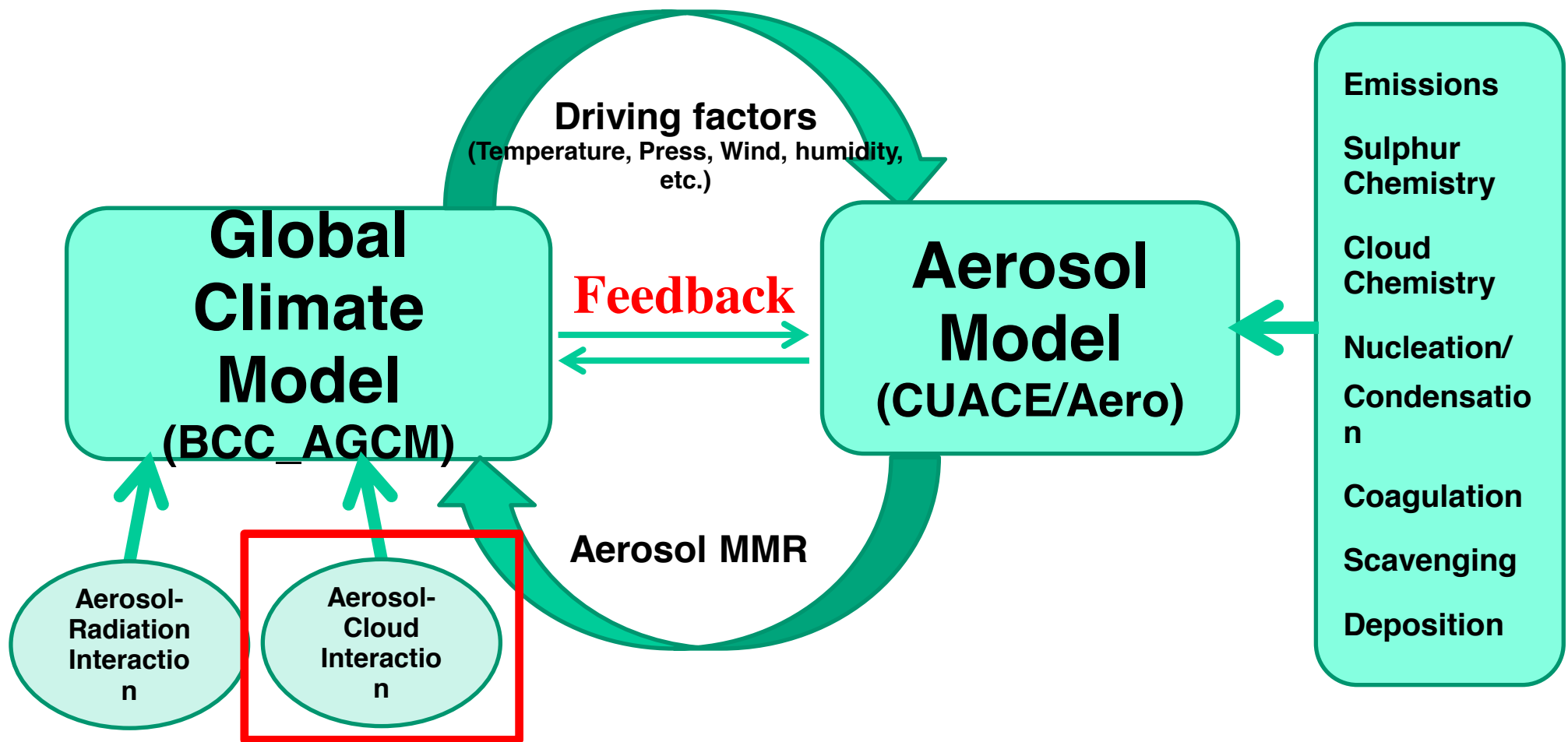
Hua Zhang

(huazhang@cma.gov.cn)

National Climate Center, CMA, Beijing, China

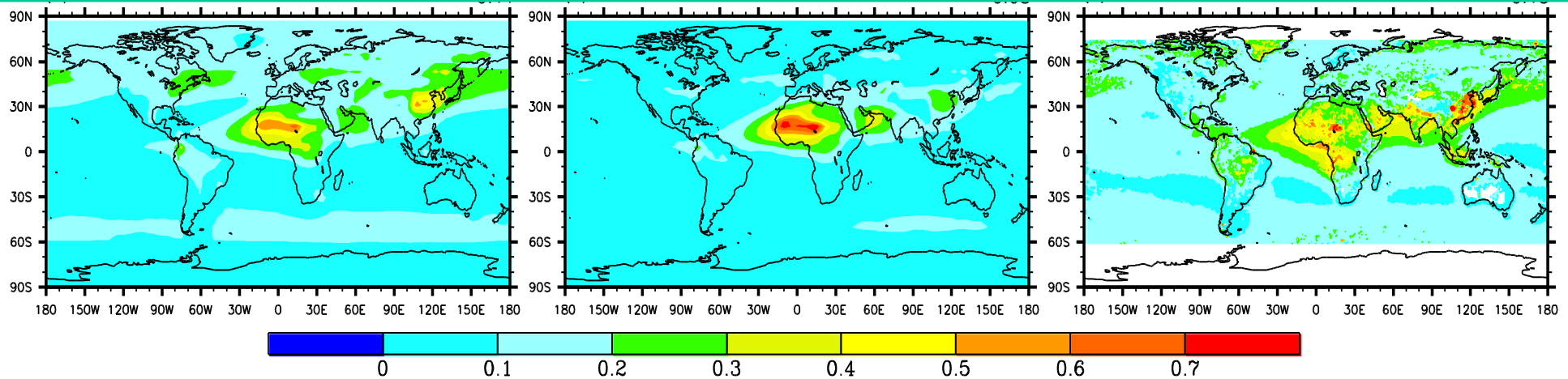
Zhili Wang and Peng Lu

Aerosol-Climate Model (BCC_AGCM2.0.1_CUACE/Aero)



We implemented **two-moment cloud microphysical scheme Of Morrison and Gettelman (2008)** into this model instead of the original one-moment bulk cloud microphysical scheme , and evaluated the new model.

Global distributions of simulated and observed annual mean AOD at 550 nm. (a) New Model, (b) Old Model and (c) MODIS&MISR.

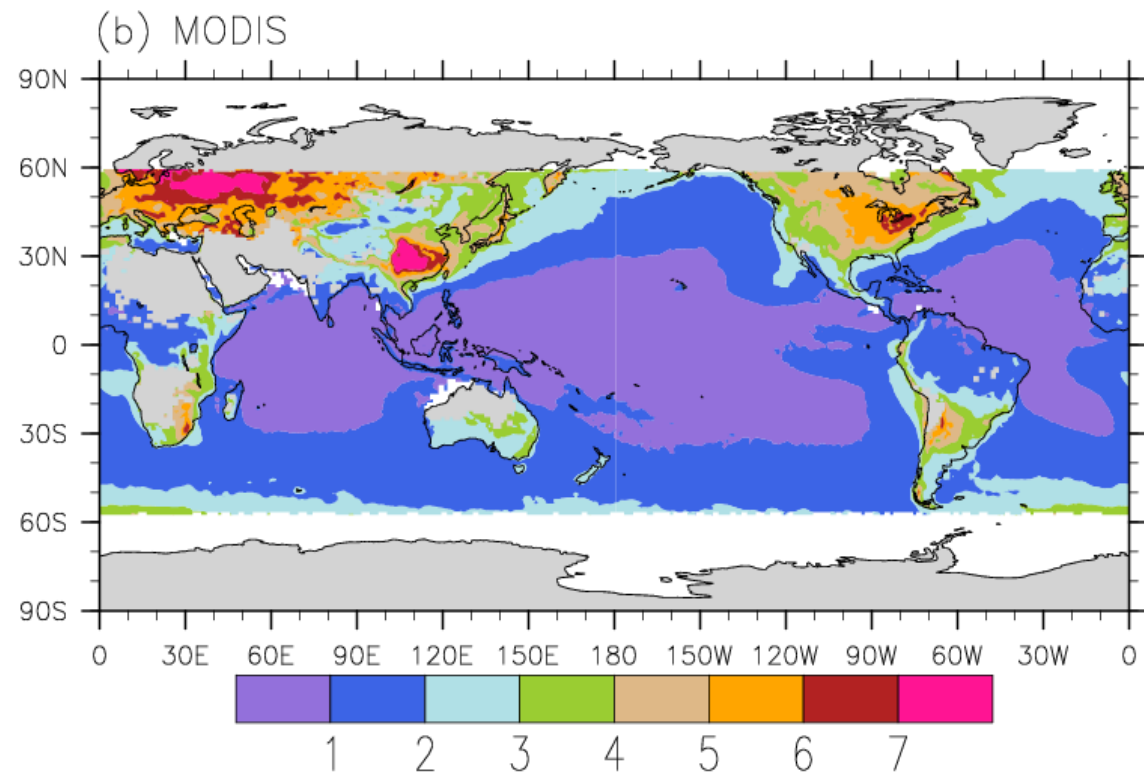
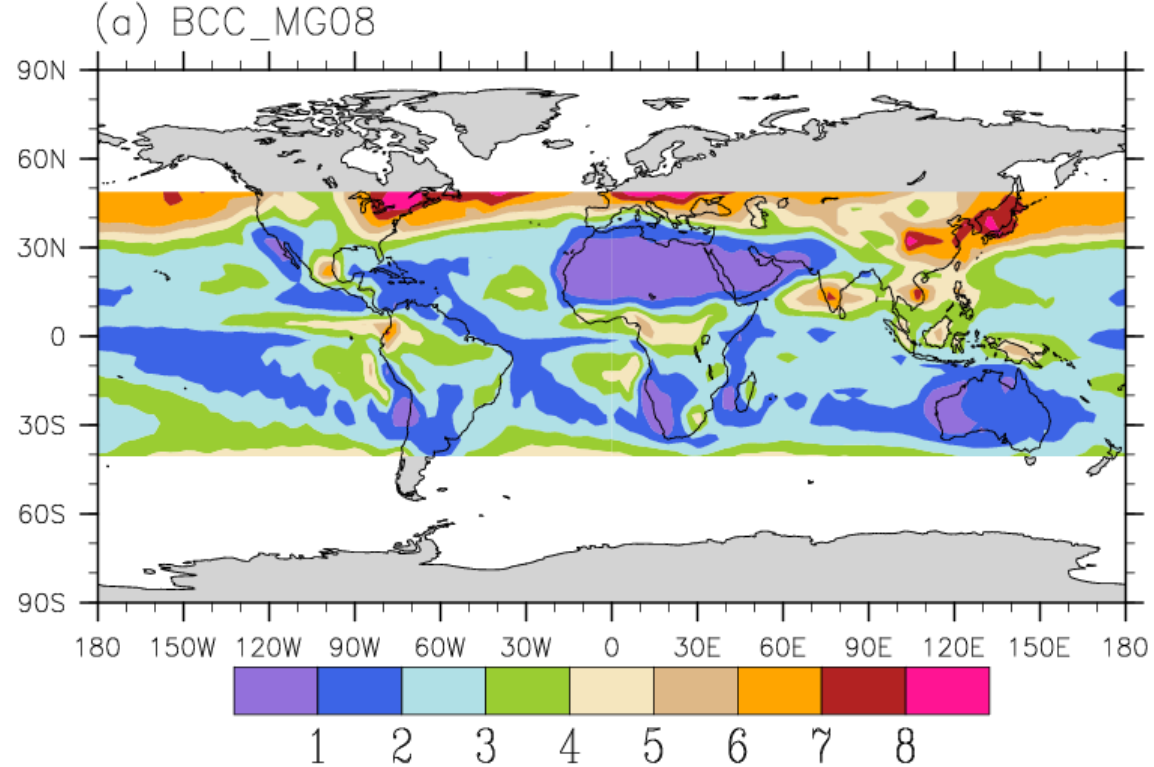


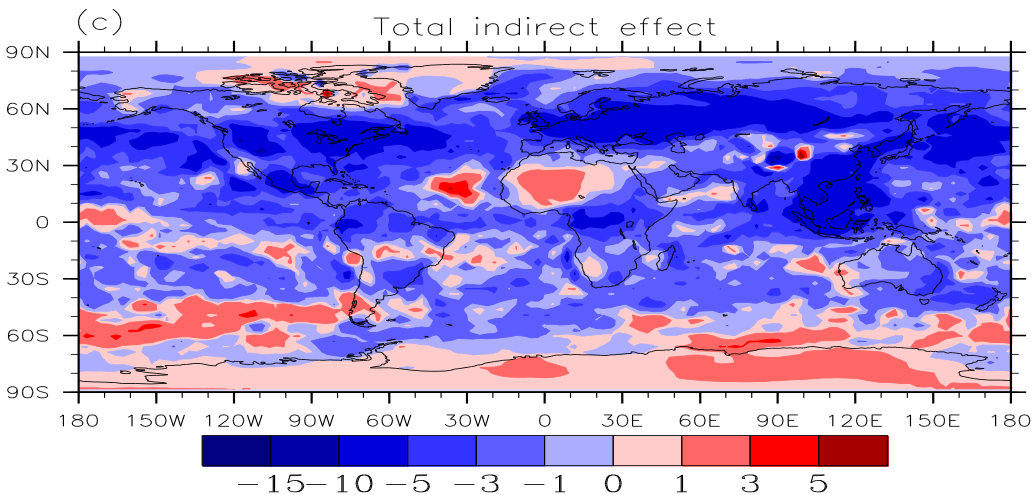
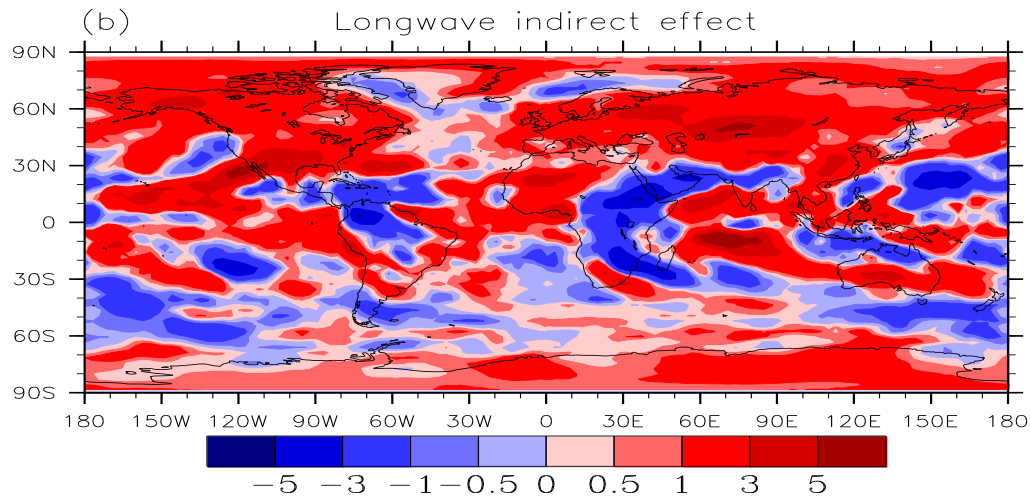
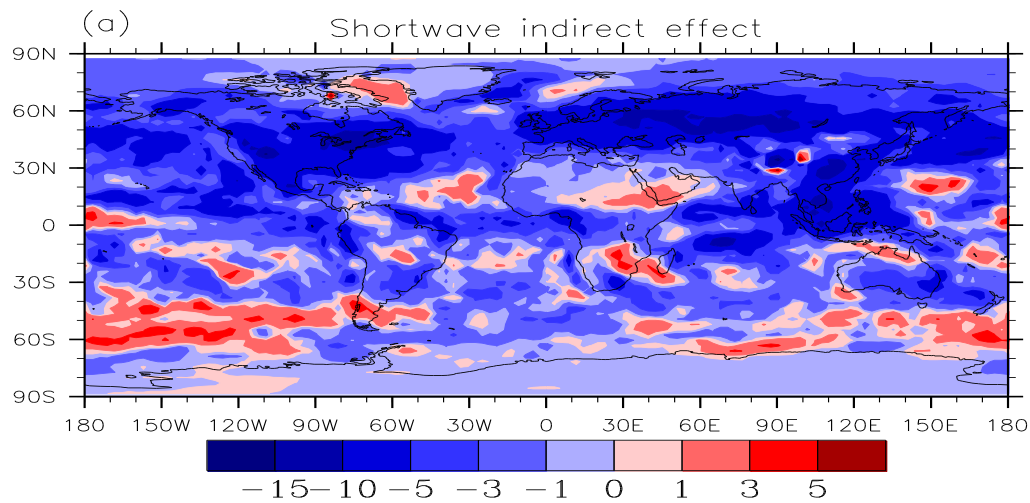
Global Budgets for Aerosols and Cloud Water (CW)^a

	BCC_MG08				BCC_RK98			
	Source	Burden	Sink	Lifetime	Source	Burden	Sink	Lifetime
SU	70.2	0.97	20.2 (D), 52.9 (W)	5.0	–	0.69	16.7 (D), 55.4 (W)	3.6
BC	7.7	0.084	4.9 (D), 2.6 (W)	4.0	–	0.069	4.7 (D), 2.8 (W)	3.3
OC	66.1	0.79	39.1 (D), 26.6 (W)	4.4	–	0.68	38.0 (D), 27.3 (W)	3.8
DU	3846.4	20.7	3049.0 (D), 799.2 (W)	1.96	5535.6	21.3	4080.3 (D), 1459.1 (W)	1.4
SS	33320.3	7.2	31171.4 (D), 2154.7 (W)	0.079	34737.4	7.4	31688.5 (D), 3046.2 (W)	0.078
CW	6.3×10^7	4.2×10^7	4.0×10^7	243.3	25.7×10^7	7.1×10^7	8.7×10^7	100.8

The cloud water lifetime in new model is significantly longer than that in old model, resulting in longer lifetimes and larger burdens of aerosols in new model.

**Annual mean distributions
of column cloud droplet
number concentration
(unit: 10^{10} m^{-2}). (a) Model,
(b) MODIS.**





**Anthropogenic
Aerosol Indirect effect.**

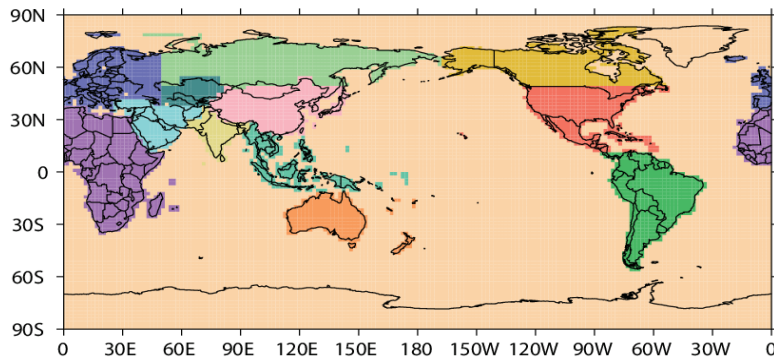
**Global mean value:
-1.9 Wm⁻²**

Investigating the Vertical Distribution and Source Attribution of Black Carbon over the Pacific Ocean

Jiachen Zhang, Junfeng Liu, George A. Ban-Weiss, and Shu Tao

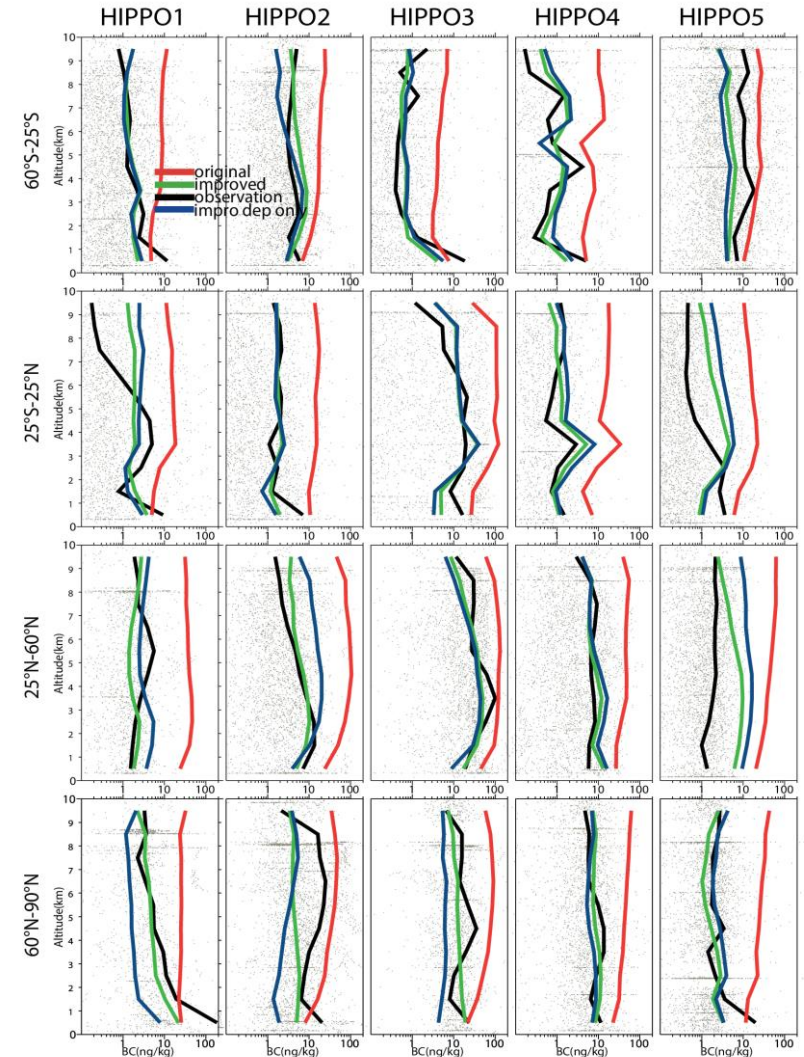
① Improve model's performance

- Implement physically-based dry& wet deposition schemes to MOZART-4
- Optimize aging rates according to different source regions



Thirteen defined source regions

	CA	SU	EU	MA	EA	ME	NA	SE	IN	AF	SA	AU	RR	Mean bias (Improved)	Mean bias (Original)
HIPPO1 Jan	200	120	60	120	4	48	60	4	4	4	60	4	4	3.4	26.4
HIPPO2 Nov	200	200	120	60	4	4	4	4	4	4	200	4	4	1.7	13.2
HIPPO3 Apr	200	200	200	200	24	60	4	24	38	48	4	4	200	1.4	6.6
HIPPO4 Jun	48	4	120	4	4	200	4	8	4	4	60	4	4	1.0	10.6
HIPPO5 Aug	60	4	12	4	4	4	4	4	4	60	4	27	4	2.2	18.7

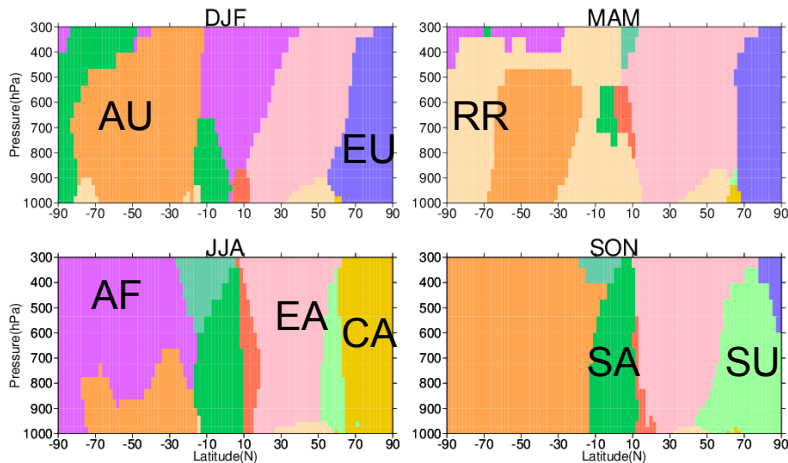


The vertical profiles of BC simulated by the improved model is closer to the HIPPO observations than the original model (Total biases on average are reduced by a factor of 5).

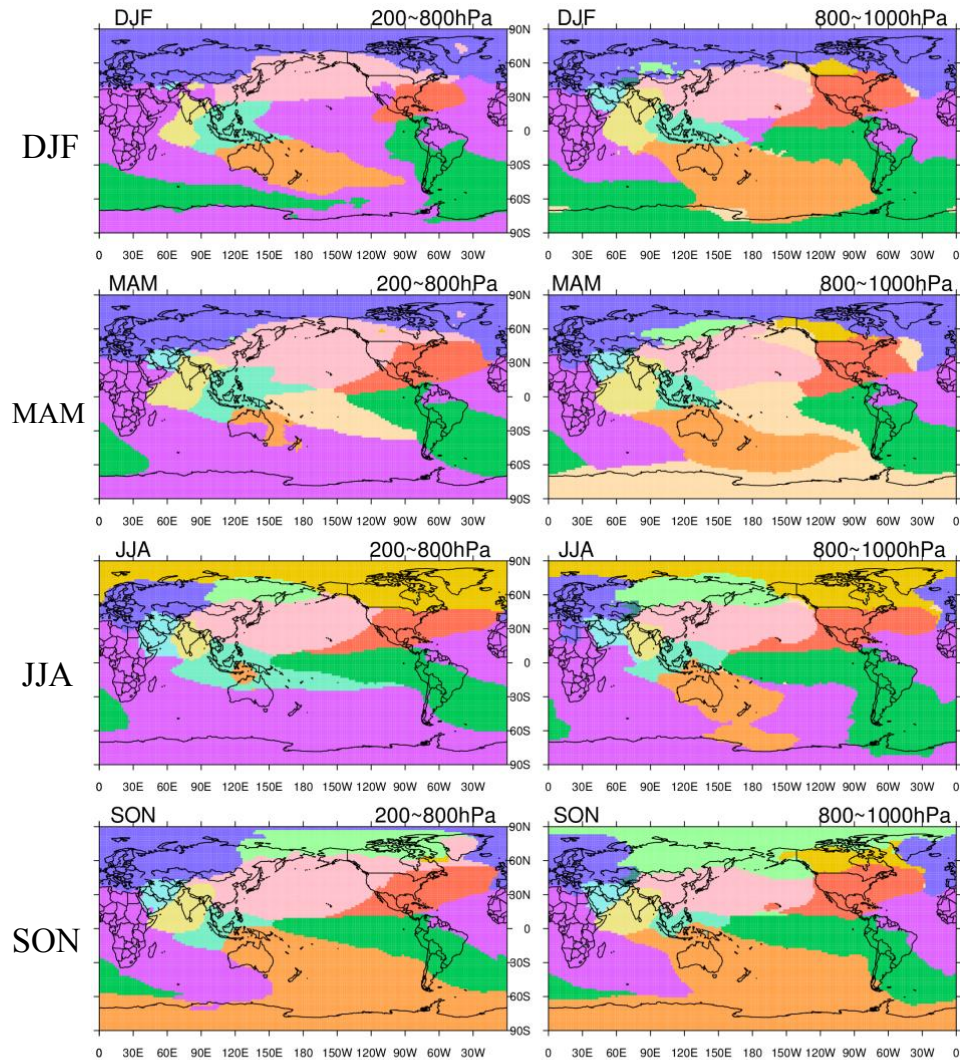
Vertical Distribution and **Source Attribution** of Black Carbon

② investigate regional contributions in different altitudes

The climate response of Black carbon (BC) depends on its altitude. [Ban-weiss et al., 2011; Samset et al., 2013]



The dominant regional contributors to zonal mean BC concentrations over the central Pacific (130°W-150°E)



The dominant regional contributors to BC burdens in the free troposphere (left column) and boundary layer (right column) in different seasons.

- BC in the boundary layer is dominated by local sources.
- BC in mid-upper troposphere over the Pacific ocean is influenced mostly by BC sources from **East Asia, Africa, South America and Australia.**