

First global overview on the representation of water uptake by ten Global Climate Models using a new in-situ benchmark hygroscopicity dataset

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HYGROSCOPICITY:

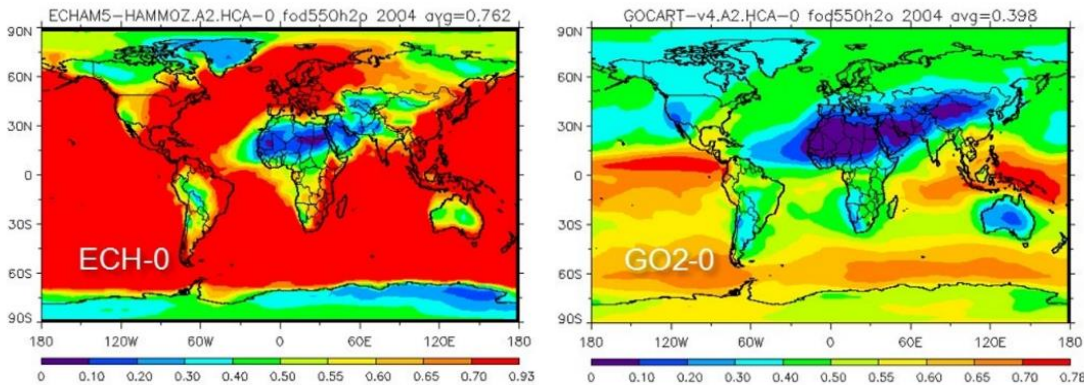
Since aerosol particles can take up water, they can change in size and chemical composition depending on the ambient relative humidity (RH)

SCATTERING ENHANCEMENT FACTOR

$$f(RH, \lambda) = \frac{\sigma_{sp}(RH, \lambda)}{\sigma_{sp}(RH_{dry}, \lambda)}$$

Interestingly, most models are doing well in reproducing the total aerosol optical depth (AOD), but a closer look into the individual components reveals discrepancies between them

Fraction of aerosol optical depth due to water:



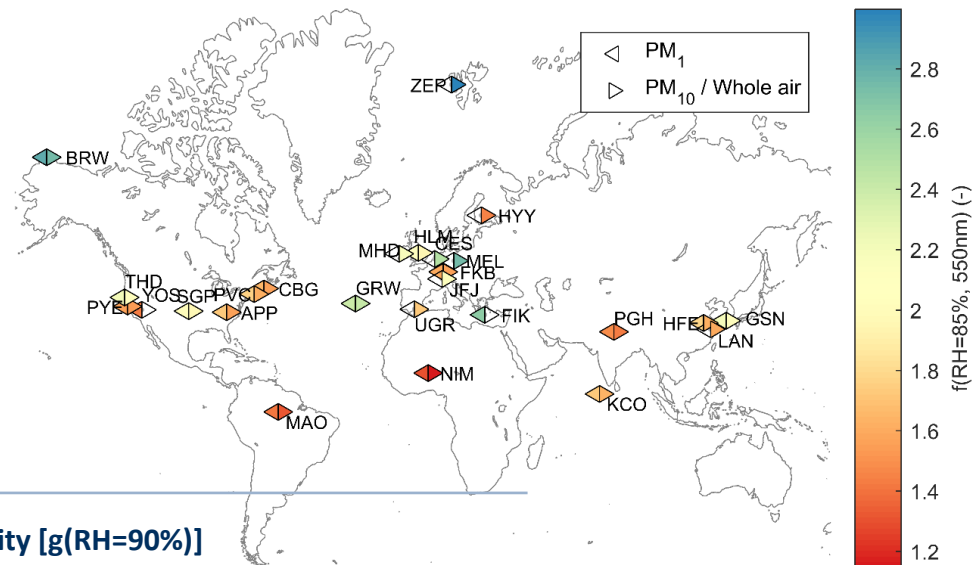
Figures from Mian Chin (NASA Goddard)

How well do Earth System Models represent aerosol optical hygroscopic growth?



Hygroscopicity benchmark dataset

Burgos et al., 2019:
 $f(RH)$ measurements from in-situ sites around the globe used to create a benchmark dataset



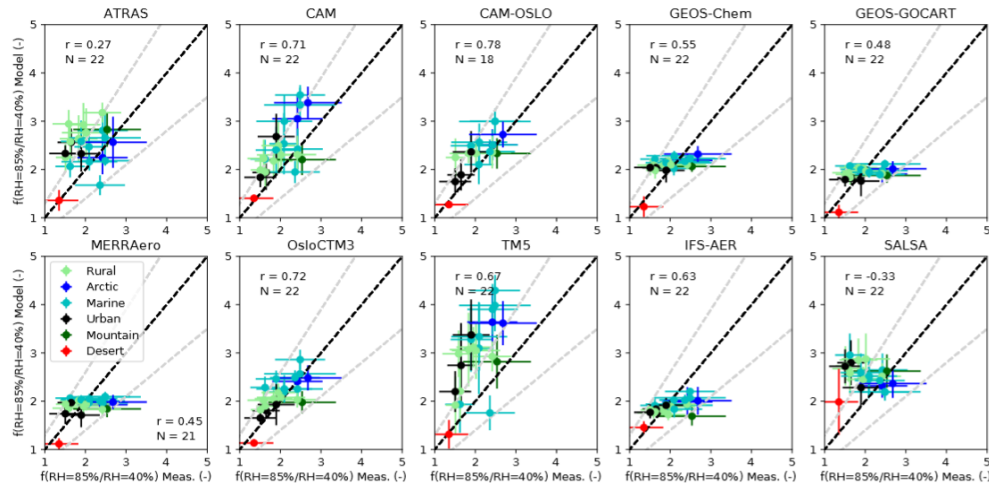
MODEL	Chemical composition	Mixing State	Hygroscopicity [g(RH=90%)]					
			parameterization	ss	so4	bc	oa	dd
ATRAS	bc,so4,oa,ss,dd + no3/nh4	I	κ -Köhler Theory	2.25	1.87	1.0	1.24	1.0
CAM	bc,so4,oa,ss,dd	I	κ -Köhler Theory	2.25	1.77	1.0	1.24	1.2
CAM-Oslo	bc,so4,oa,ss,dd	I,E	κ -Köhler Theory	2.28	1.77	1.0	1.31	1.2
GEOS-Chem	bc,so4,oa,ss,dd + no3/nh4	E	Modified GADS	2.38	1.64	1.4	1.64	1.0
GEOS-GOCART	bc,so4,oa,ss,dd	E	Modified GADS	1.9-2.1	1.8	1.4	1.6	1.0
MERRAero	bc,so4,oa,ss,dd	E	Modified GADS	1.9-2.1	1.8	1.4	1.64	1.0
OsloCMT3	bc,so4,oa,ss,dd + no3/nh4	I	Own development	2.3-2.4	1.72	1.0	1.46	1.0
TM5	bc,so4,oa,ss,dd + no3/nh4	I, E	Own development	-	-	1.0	1.0	1.0
IFS-AER	bc,so4,oa,ss,dd + no3/nh4	E	Own development	2.36	1.73	1.0	1.64	1.0
SALSA	bc,so4,oa,ss,dd	E	Own development	2.4	1.9	1.0	1.5	1.0

Model data: INSITU project - AeroCom Phase III

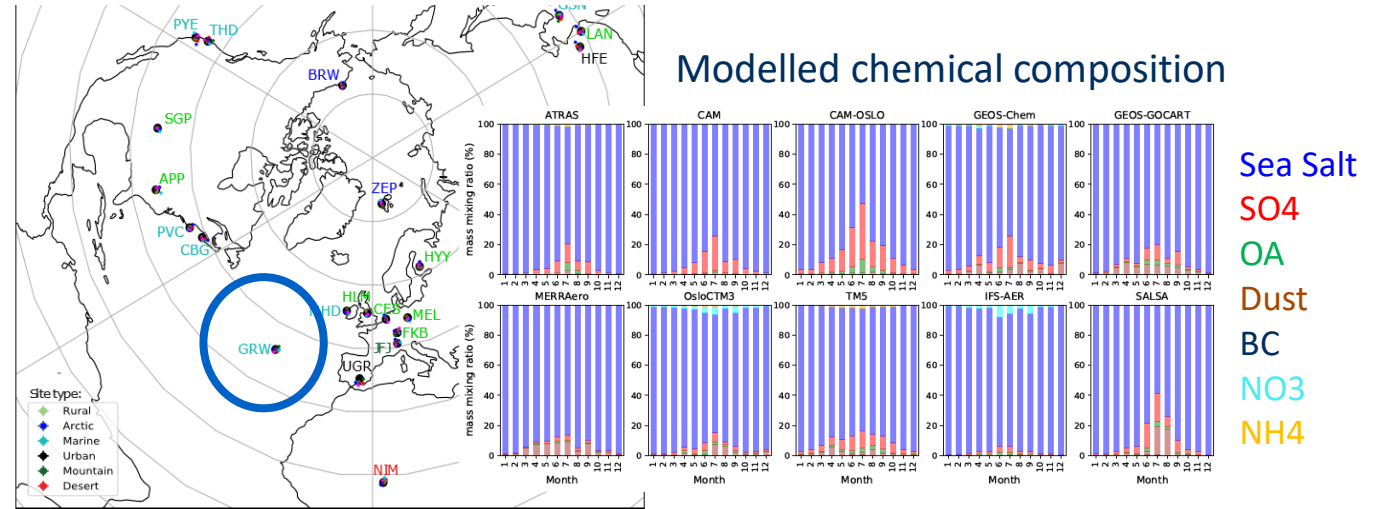
Main characteristics implemented by each model

Main Results:

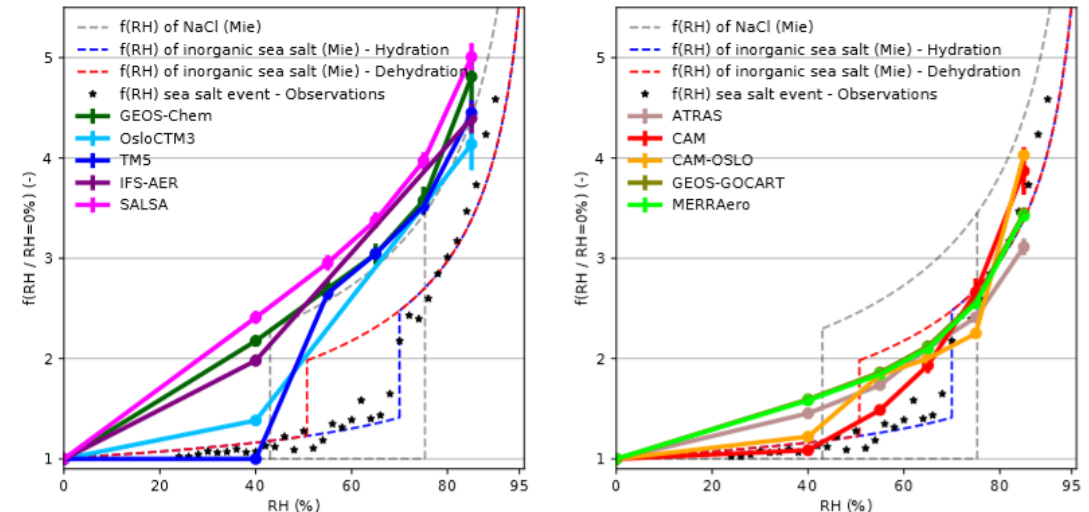
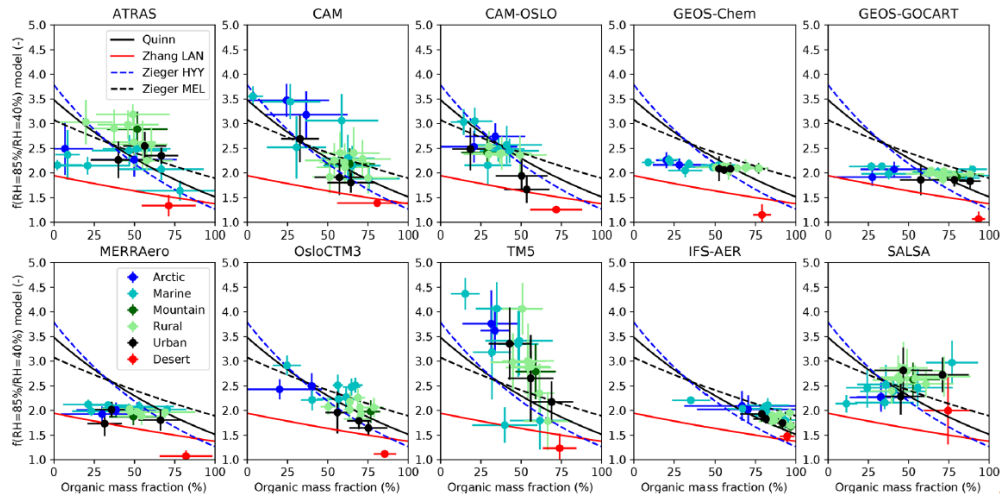
(1) Modeled vs Measured $f(RH)$



(3) $f(RH)$ of sea salt dominated aerosol at Graciosa



(2) $f(RH)$ vs. organic mass fraction



- (Some) models reproduce the range in measured $f(RH)$
- (Some) models: similar $f(RH)$ -OMF relationship as parameterizations
- Diversity of behaviors: Good correlation / Inverse correlation

- (Some) models treat sea salt as NaCl
- (Some) models: simulate inorganic sea salt (wrong kappa for NaCl)
- TM5: correctly models salt to be fully solid (=NaCl) as expected, while other models assume growth at $RH < 40\%$

Conclusions :

- GEOS-family models assign **too much hygroscopicity to all species** -> narrow $f(\text{RH})$ range
- Models that **best** reproduce the observed relationship between $f(\text{RH})$ and OMF are those that assume **lower hygroscopicity for organics** – this allows these models to simulate a wider range of $f(\text{RH})$.
- Different **assumptions** about the **hygroscopicity of sea salt** explain some model variation at a marine location -> some models assume sea salt can be represented by NaCl, while others do not

Recommendations:

1. Update the **hygroscopic parameterization** of some components (e.g. **sea salt**) and parameterizations based on OPAC
2. **Reproducing observational-based parameterizations** of $f(\text{RH})$ using chemical mass fractions
3. Compare models and measurements at **similar conditions**:
 - > Models: calculate σ at the same variable RH conditions as the measurements
 - > Measurements: control $\text{RH}_{\text{dry}} < 40\%$ + maintaining a narrower RH_{dry} distribution

Proposed AeroCom activities:

1. Experiment with **common/improved hygroscopicity scheme** for all ESMs
2. **$f(\text{RH})$ for individual aerosol components** from models (see sea spray comparison)
3. Take into account other variables (e.g. **size distribution**)
4. Study the influence on aerosol (e.g. AOD, radiative forcing, lifetime) and cloud properties **with improved hygroscopicity scheme**
5. Is it possible to model **backscattering**? Is it available already? (We could calculate forcing efficiency at dry and wet conditions)