Climate sensitivity to marine organic aerosol emissions in ACME v0 Susannah M. Burrows, S. Elliott, R. Easter, X. Liu, P.-L. Ma, P. Cameron-Smith, H. Wang, B. Singh, K. Zhang, S. Ghan, P. Rasch

Objective and approach

Zonal mean seasonal changes in aerosol and cloud properties

Accumulation mode number Accumulation mode number concentration emissions (surface model layer)

Boundary-layer CCN concentrations [S=0.1%]



Top panels: zonal seasonal means of model

Submicron organic aerosol mass at selected stations

2000 - 1000 - 0 -	west of Portugal
3000 - 2000 - 1000 -	west of Namibia
300 - 200 -	La Reunion Island

1 2 3 4 5 6 7 8 9 10 11 12

MOA: marine organic aerosol

POA: Primary organic

Marine organic matter in sea spray aerosol:

- Organic matter from ocean phytoplankton enters sea spray aerosol through bubble bursting mechanism, and is a previously missing aerosol source.
- Natrual background aerosols are an important controlling factor for aerosol indirect effects.
- The influence of ocean biogeochemistry on sea spray chemistry is represented in the OCEANFILMS model (Burrows et al., ACP, 2014; Elliott et al., GRL, 2014), which has been implemented in ACME.
- Sensitivity experiments were conducted to determine the model's simulated aerosol chemistry and climate response.

Simulation configurations:

10-y atmosphere-only simulations.

One control experiment and 4 sensitivity cases: "int, add"; "int, replace"; "ext, add"; "ext, replace"

- "int": marine organics internally mixed with sea salt aerosol
- "ext": marine organics externally mixed with sea salt aerosol
- "add": organics increase number and mass of emitted sea spray
- "replace": number and mass of emitted sea spray remain constant

This poster focusses on results from "int, add", which observations suggest is the most realistic case.

Simulated sea spray chemistry and change in number:













North Pacific Ocean 1

North Pacific Ocean 2

New Caledonia

dominates organic mass in submicron aerosol primarily at remote locations where continental sources are small.

Geographic distribution of seasonal changes in aerosol and cloud properties



Shaded, outlined regions are significant at p<0.1% level by two-sided Welch's unequal variances t-test.

Shaded areas surrounded by dark contour indicate regions of statistically significant differences from control simulation a p<0.1% level, by two-sided Welch's unequal variances t-test, treating each year of a 10-y simulation as a sample (N=10).

Simulated and observed seasonal cycles

OMF in Longhurst biogeographical provinces

of sea spray chemistry varies between ocean biomes. seasonal



Comparison with observed seasonal cycles



Summary and Impact

Summary:

- Representing marine organic matter in sea spray, a previously missing aerosol source, improves representation of the natural background aerosol, an important factor controlling the magnitude of aerosol indirect effects.
- Marine organic matter has important seasonal impacts on remote marine aerosol concentrations, particularly in the Southern Ocean, where

summertime aerosol number is **doubled** in some regions by marine organics.

• In the Northern Hemisphere, summertime marine aerosol number and composition are impacted, but marine sources are small relative to natural and anthropogenic continental sources, and climate impacts are mostly not statistically significant.

Previous work inferred that marine organic aerosol contributes to determining seasonal and large-scale spatial patterns of cloud drop number concentration over the Southern Ocean (McCoy, Burrows, et al., Sci. Adv., 2015); magnitude of simulated reponses is consistent with this.

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climatemodeling.science.energy.gov/acme ENERGY Burrows, S.M., Ogunro, O., Frossard, A.A., Russell, L.M., Rasch, P.J. and Elliott, S.M., 2014. A physically based framework for modeling the organic fractionation of sea spray aerosol from bubble film Langmuir equilibria. Atmos. Chem. Phys, 14(24), pp.13-601. Elliott, S., Burrows, S.M., Deal, C., Liu, X., Long, M., Ogunro, O., Russell, L.M. and Wingenter, O., 2014. Prospects for simulating macromolecular surfactant chemistry at the ocean-atmosphere boundary. Environmental Research Letters, 9(6), p.064012.