

Aerosol nucleation and growth, SOA
formation, upcoming NorESM simulations at
University of Helsinki

Risto Makkonen
NorESM WS, Stockholm, 24.10.2014

Evaluation of aerosol number concentrations in NorESM with improved nucleation parameterization

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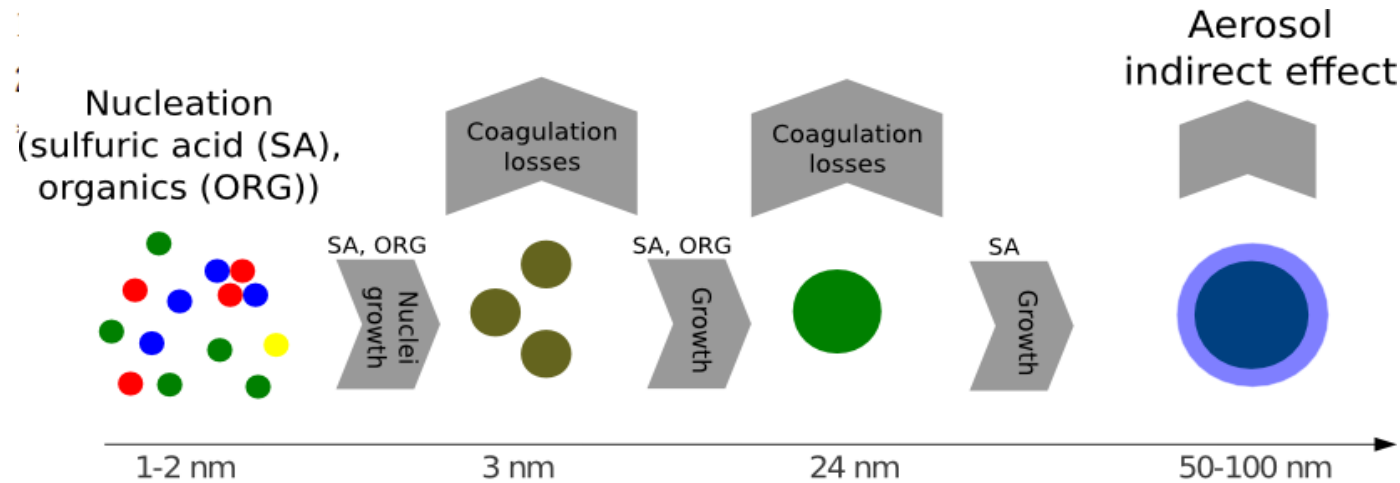


Table 1. Simulated sensitivity experiments. The simulated climate is identical in all experiments except ActNuc_BC24_Online, where aerosol–climate interactions are switched on.

Short name	Nucleation	BSOA	BC _{FF} radii	Additional information
NorESM1-M	On*	On	11.8 nm	As in Kirkevåg et al. (2013)
NoNuc_BC12	Off	On	11.8 nm	
NoNuc_BC24	Off	On	23.6 nm	
ActNuc_BC12	Eq. (1)	On	11.8 nm	
ActNuc_BC24	Eq. (1)	On	23.6 nm	
OrgNuc_BC12	Eq. (4)	On	11.8 nm	
ActNuc_BC12_NoSOA	Eq. (1)	Off	11.8 nm	
ActNuc_BC24_NoSOA	Eq. (1)	Off	23.6 nm	
ActNuc_BC24_Online	Eq. (1)	On	23.6 nm	Aerosols affect model meteorology
ActNuc_BC24_Nuc10	Eq. (1)	On	23.6 nm	SO ₄ nucleation-mode diameter set to 10 nm

*Original NorESM1-M assumed that after condensation, all excess sulfuric acid nucleates as new particles.

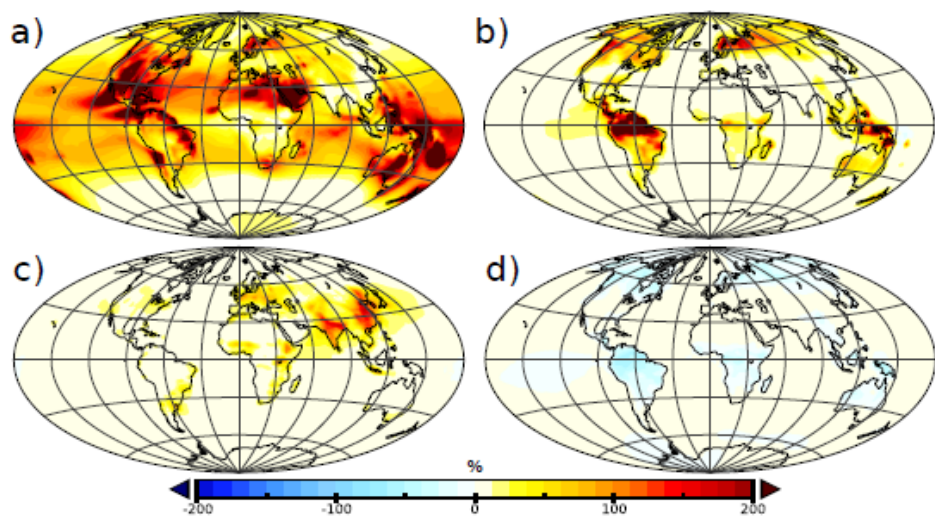
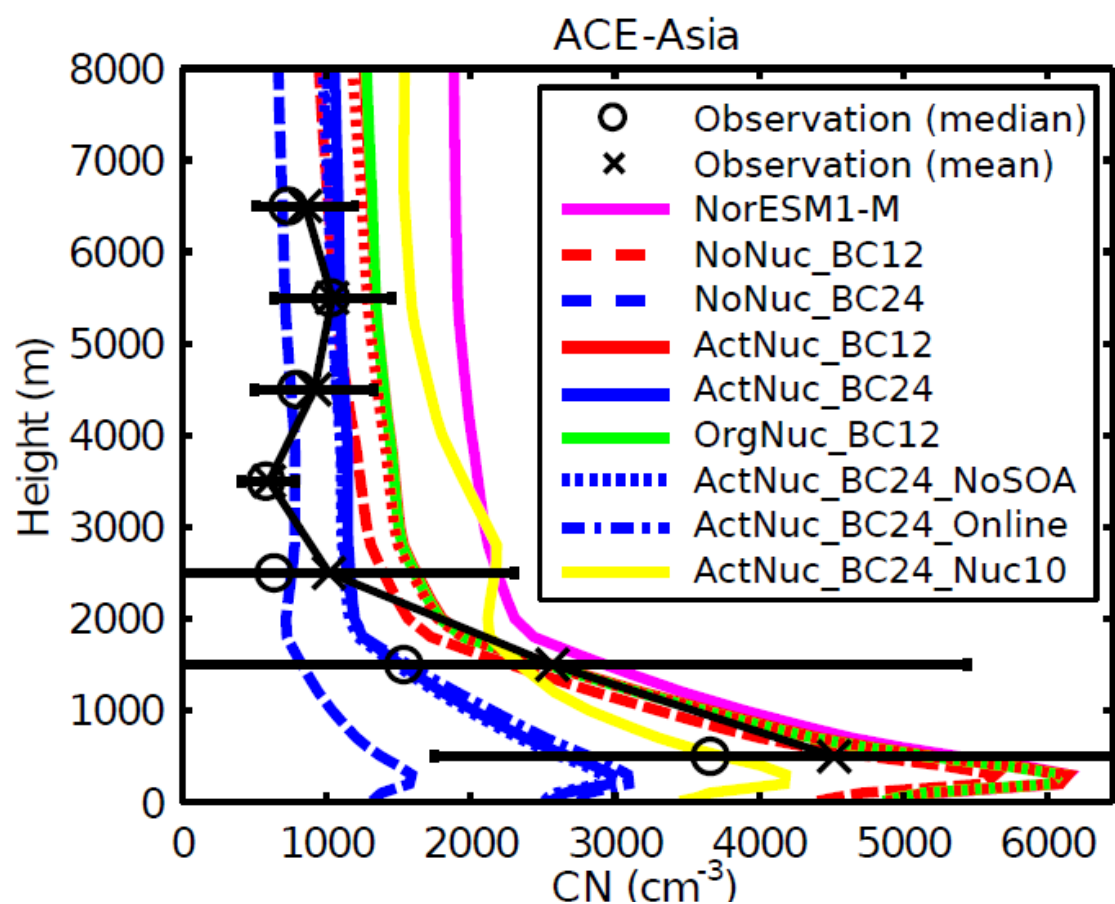
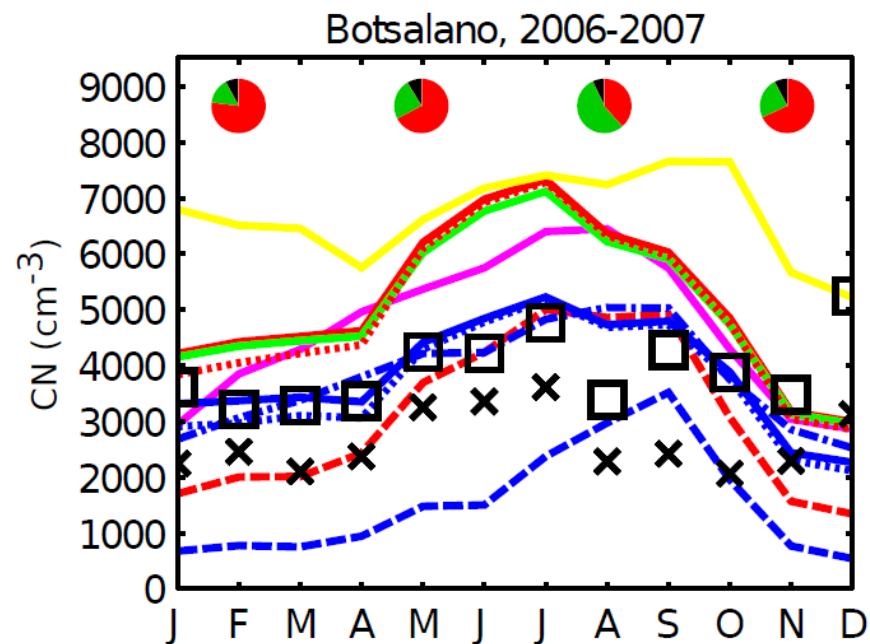


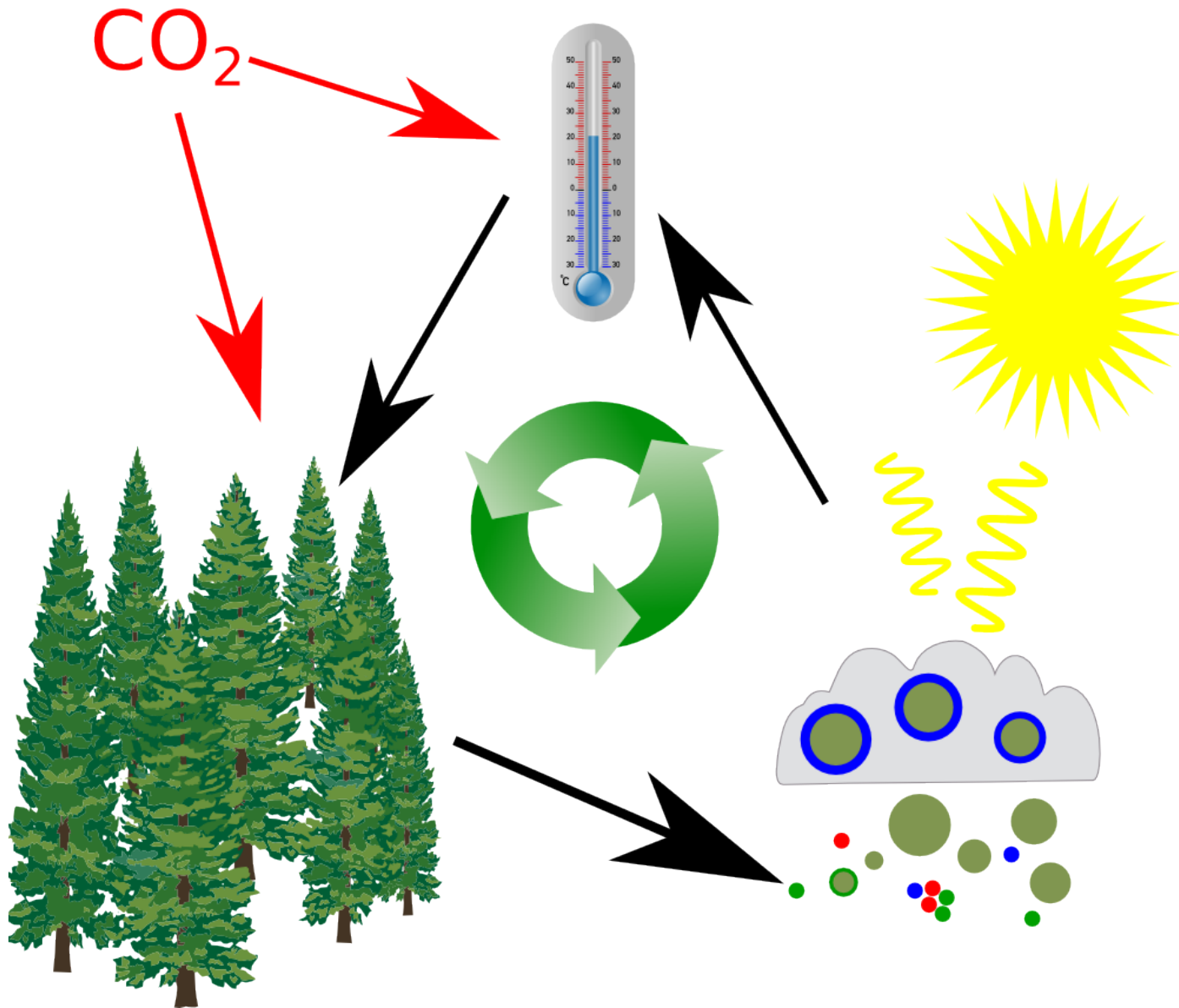
Fig. 2. Surface-level aerosol number concentration difference fields (%) from sensitivity simulations, due to (a) nucleation (ActNuc_BC12-NoNuc_BC12), (b) SOA formation (ActNuc_BC12-ActNuc_BC12_NoSOA), (c) BC emission size (ActNuc_BC12-ActNuc_BC24) and (d) nucleation parameterization (ActNuc_BC12-OrgNuc_BC12). Differences are calculated against ActNuc_BC12 simulation (Fig. 1).

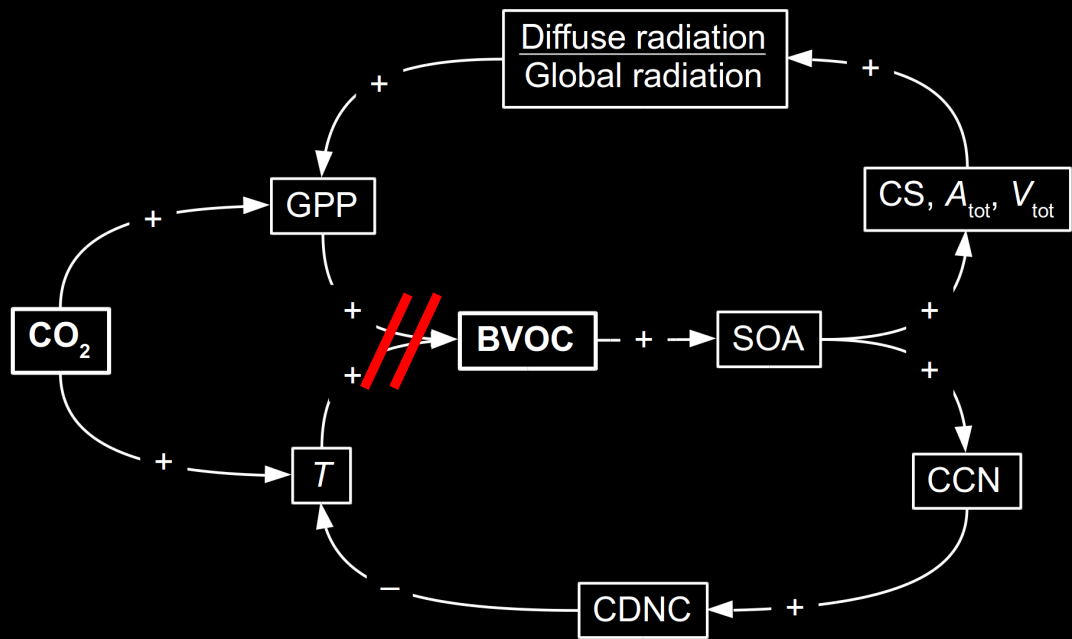
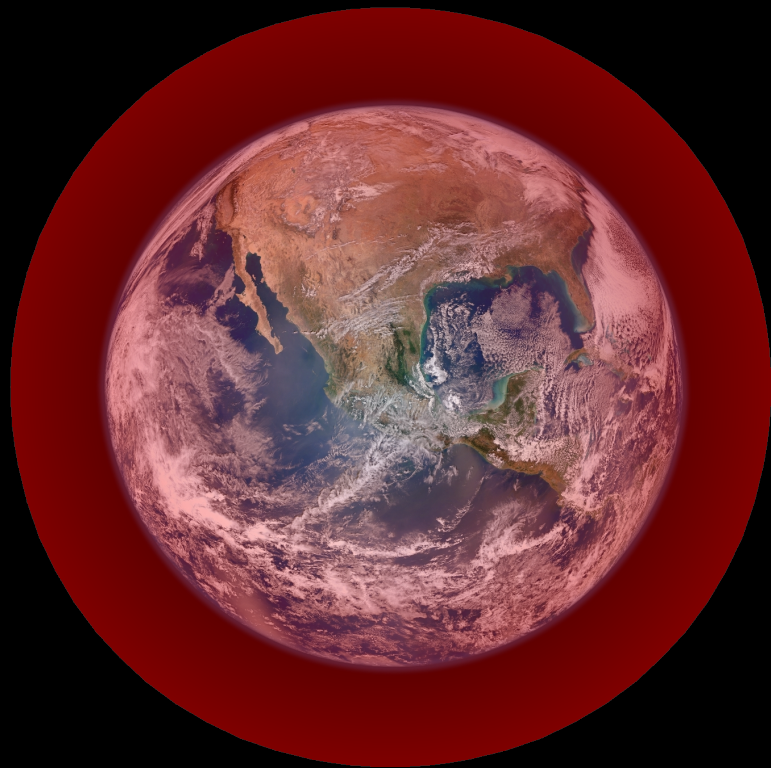
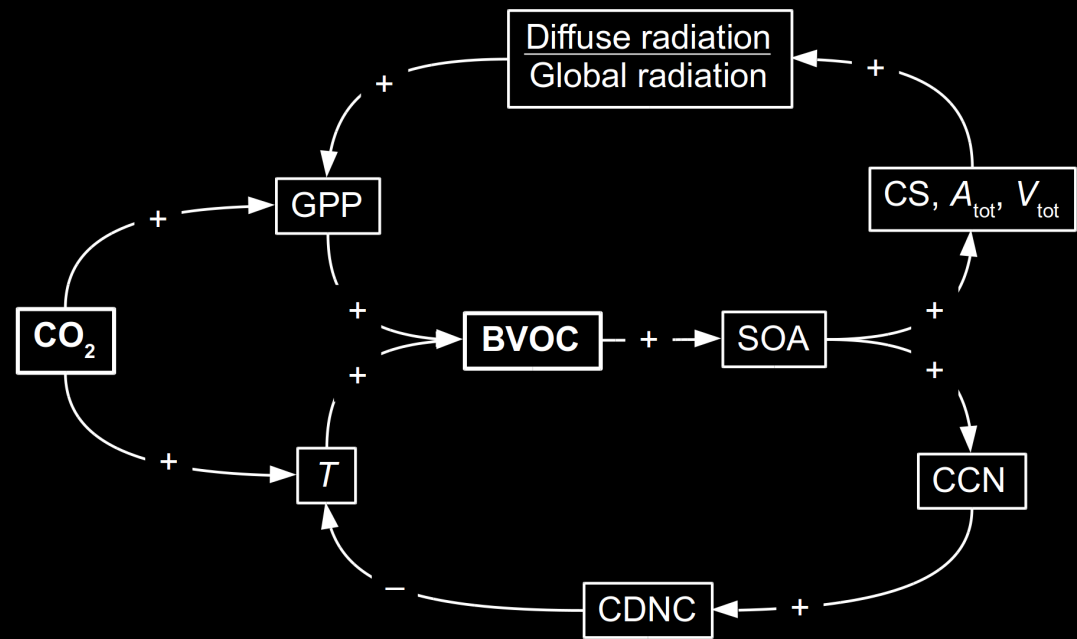
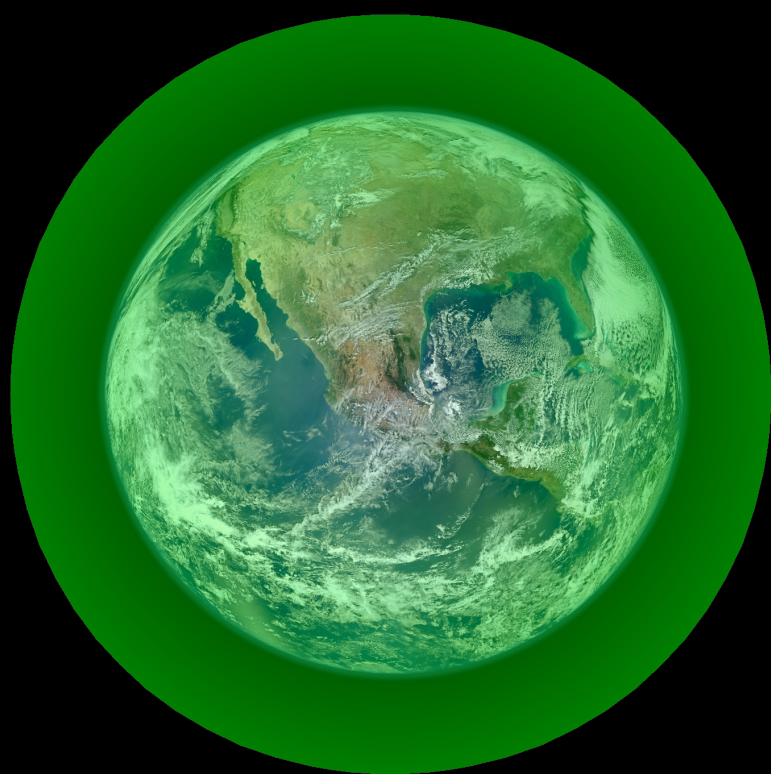
	All BIAS (%)	R^2
NorESM1-M	33	0.38
NoNuc_BC12	-27	0.30
NoNuc_BC24	-64	0.35
ActNuc_BC12	20	0.40
ActNuc_BC24	-5	0.41
OrgNuc_BC12	20	0.39
ActNuc_BC12_NoSOA	2	0.38
ActNuc_BC24_NoSOA	-25	0.42
ActNuc_BC24_Online	-8	0.43
ActNuc_BC24_Nuc10	81	0.38

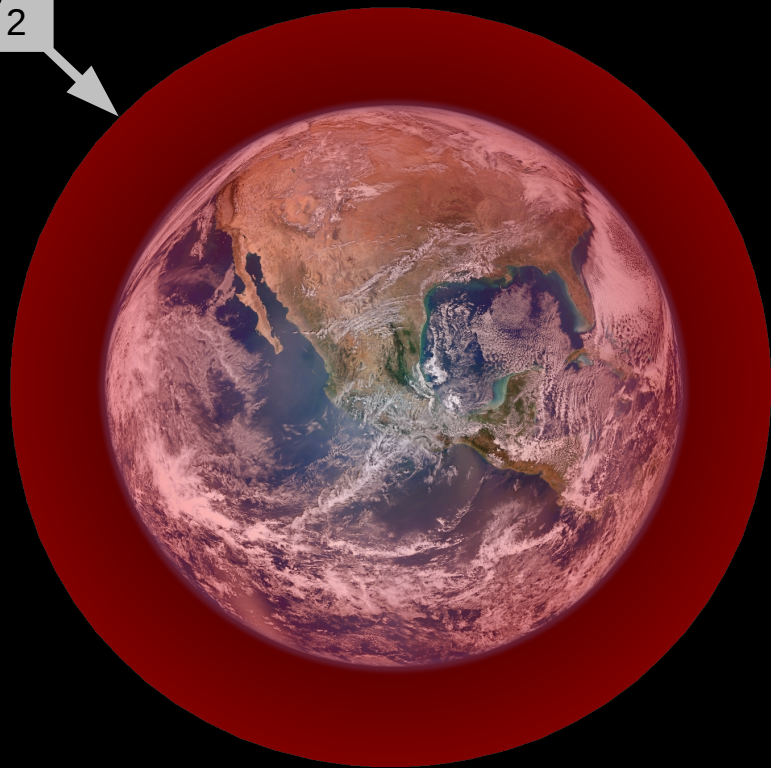
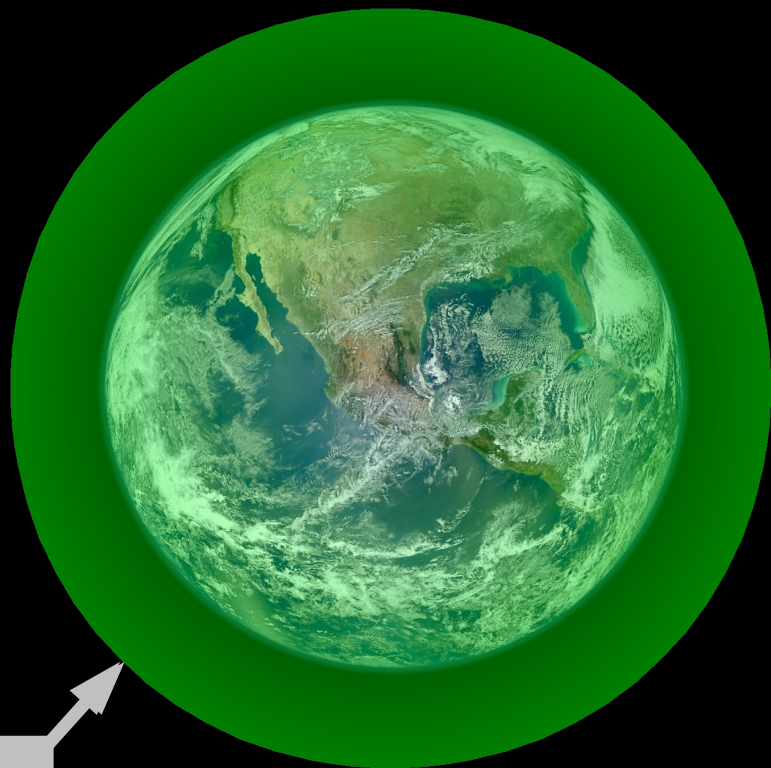
Aerosol-Climate Feedbacks in Earth System Simulations

- CRAICC WP8: synthesis
- BACCHUS task 4.4: Biosphere-atmosphere-cloud-climate interactions and feedbacks
- CSC Grand Challenge ACFESS: 6 million CPU hours

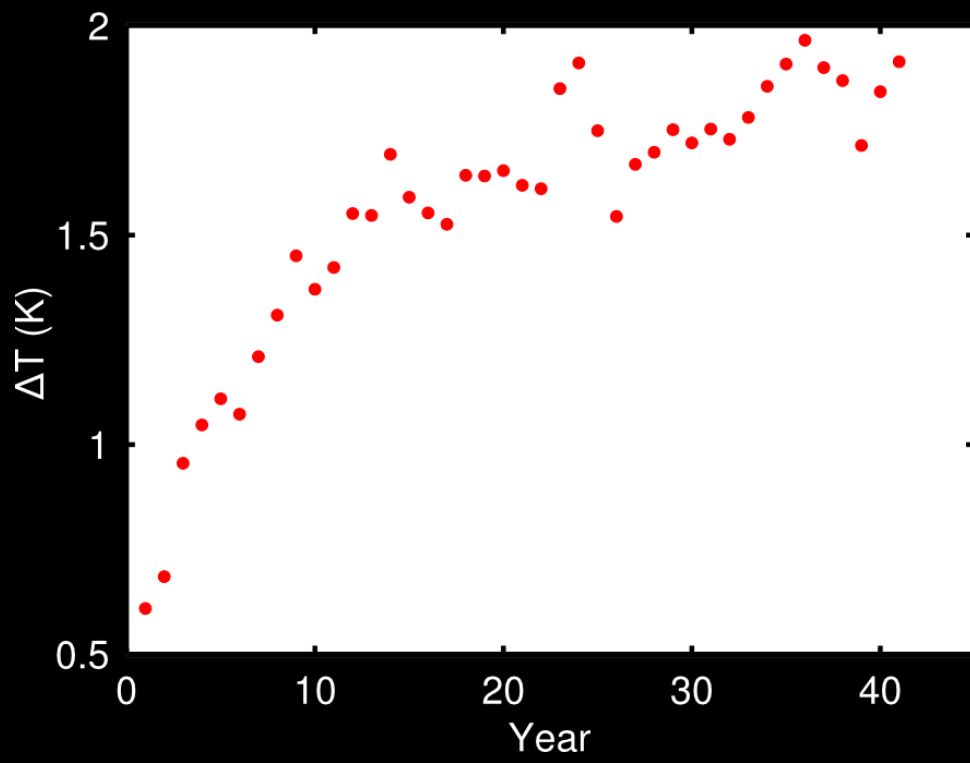
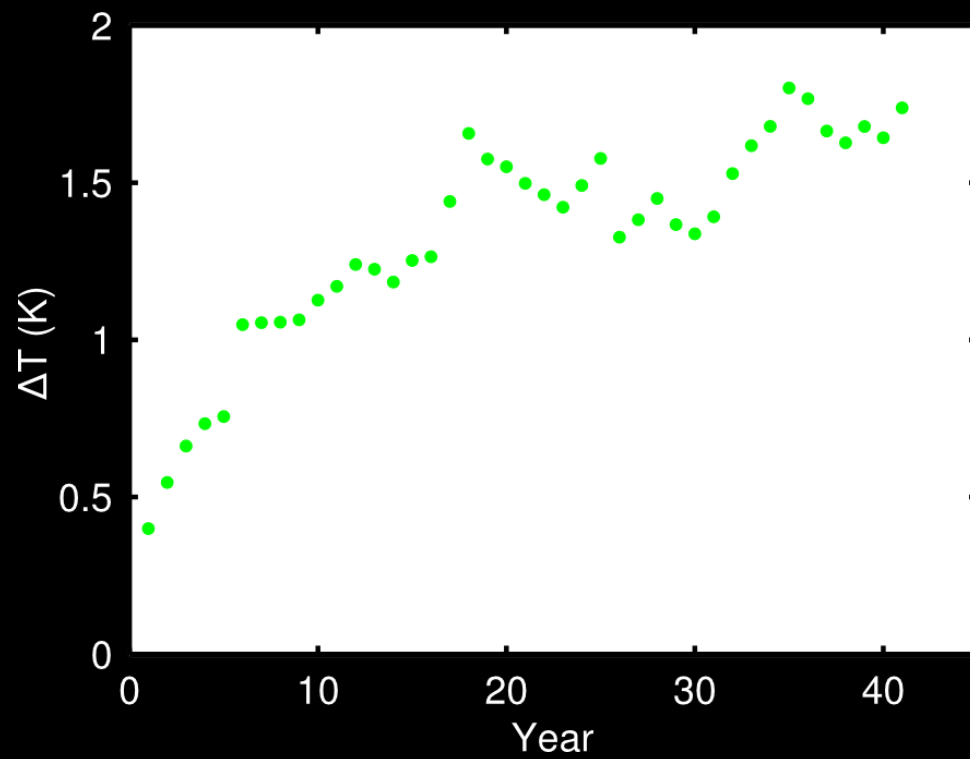
BVOC-aerosol-climate feedback

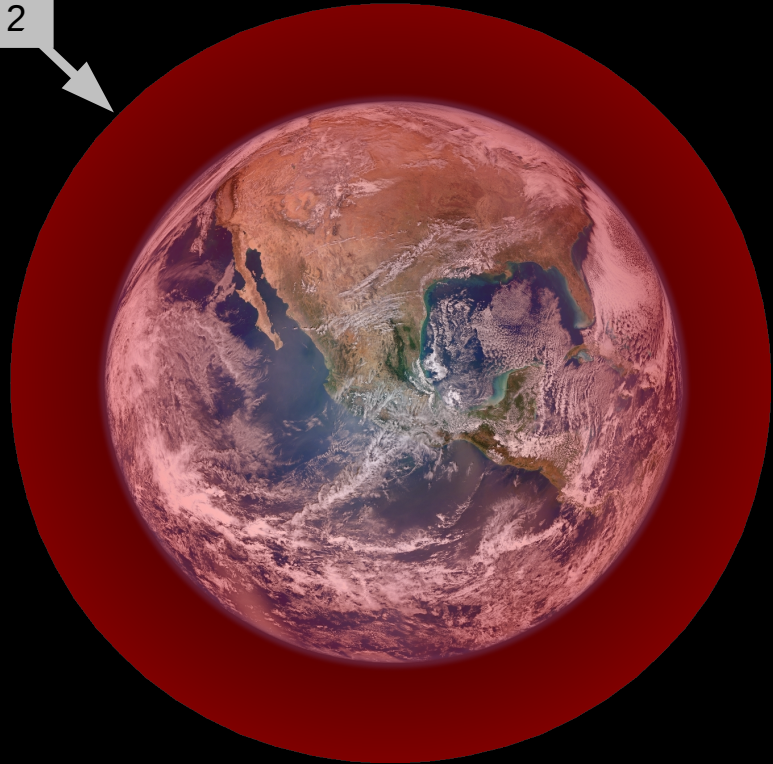
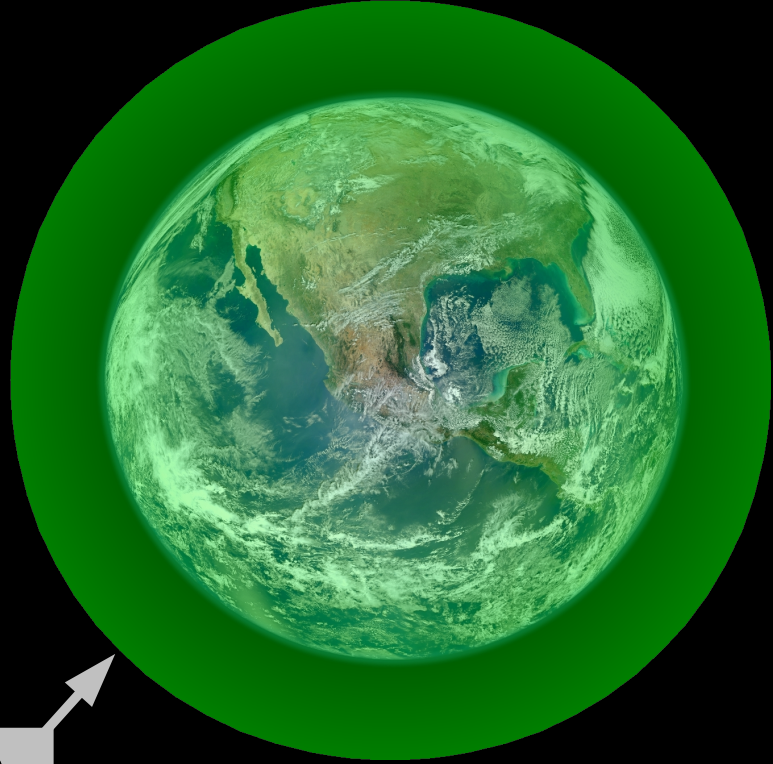






2xCO₂

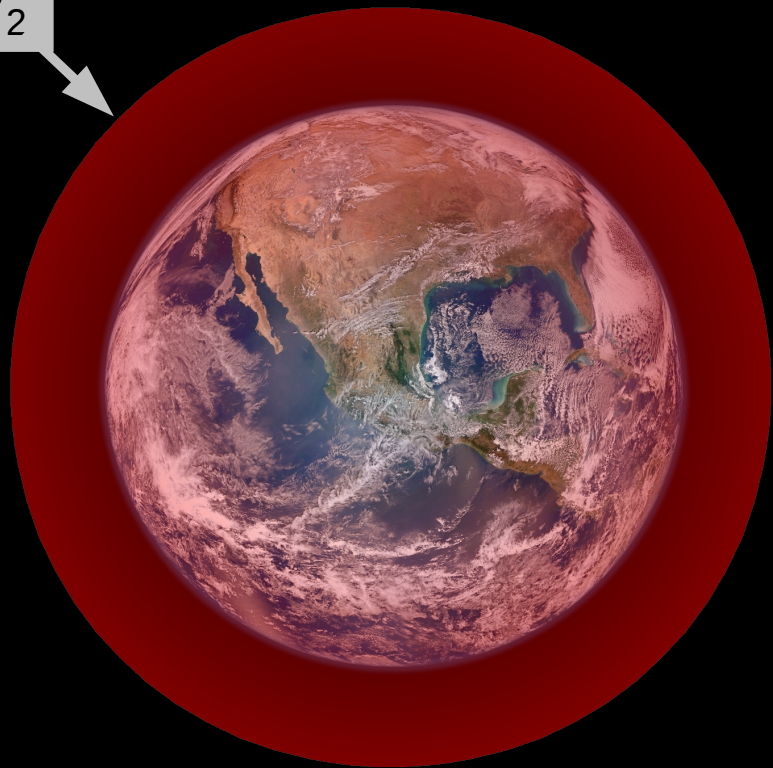
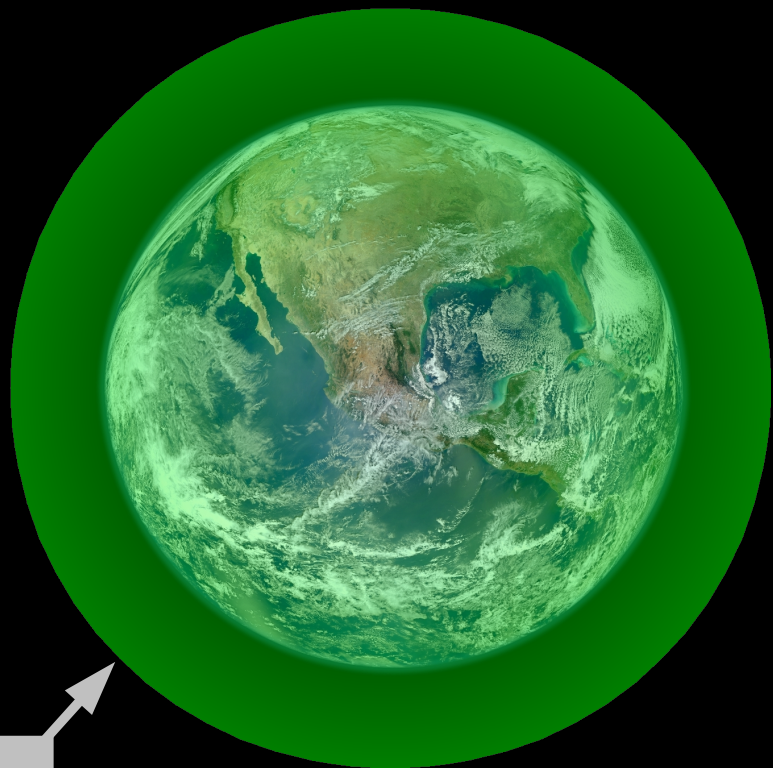




2xCO₂

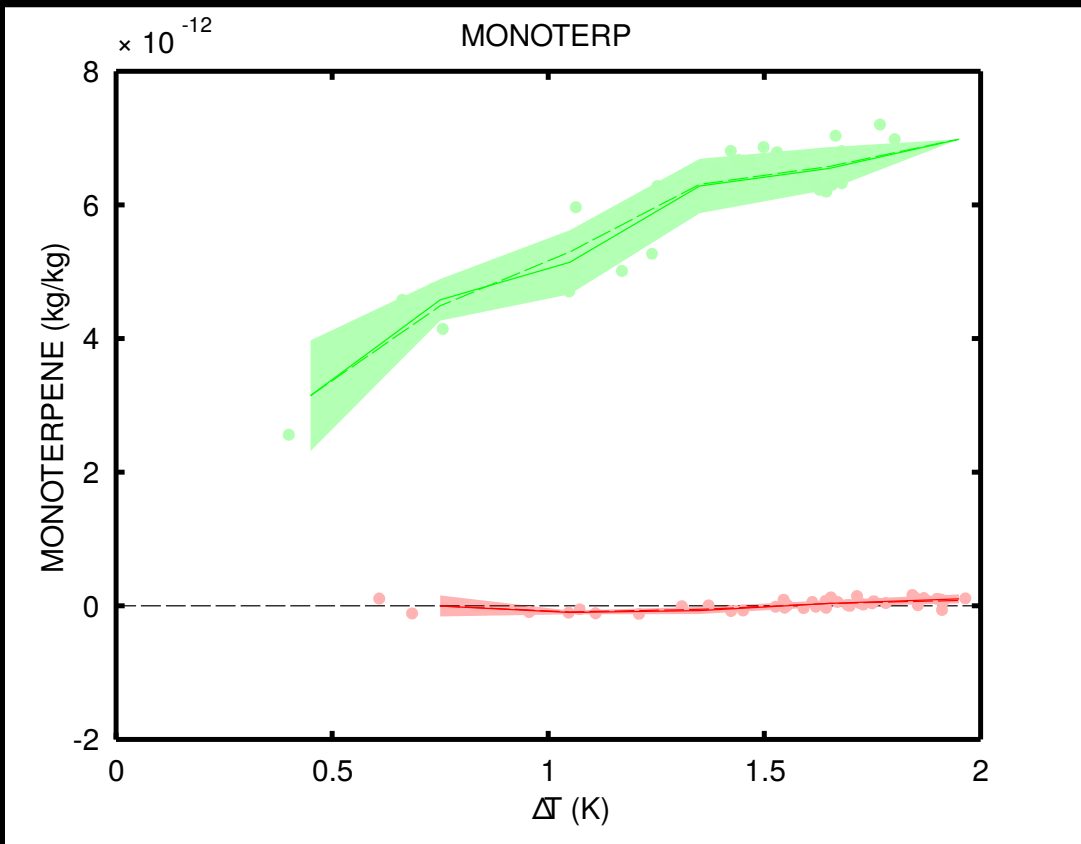


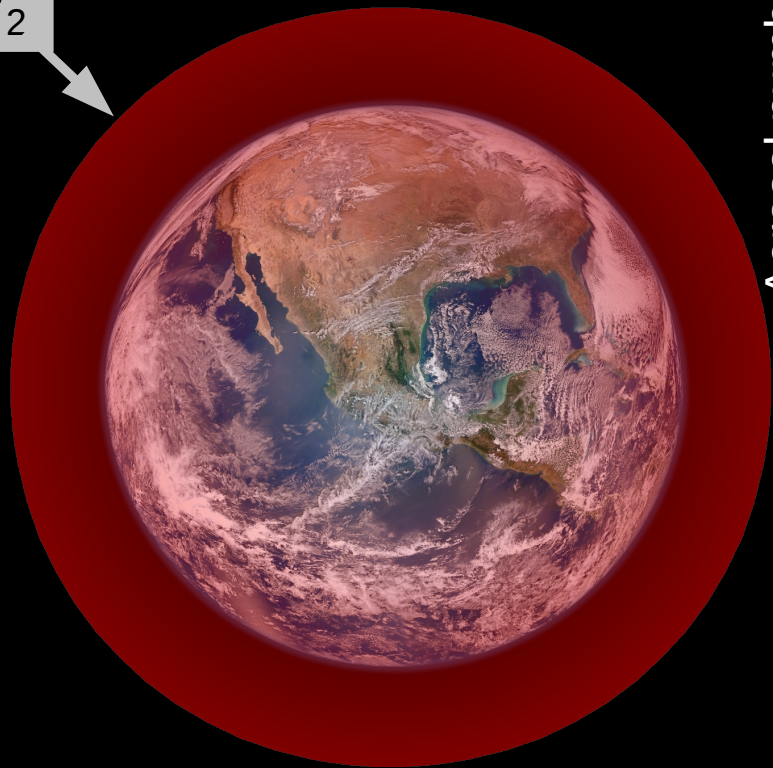
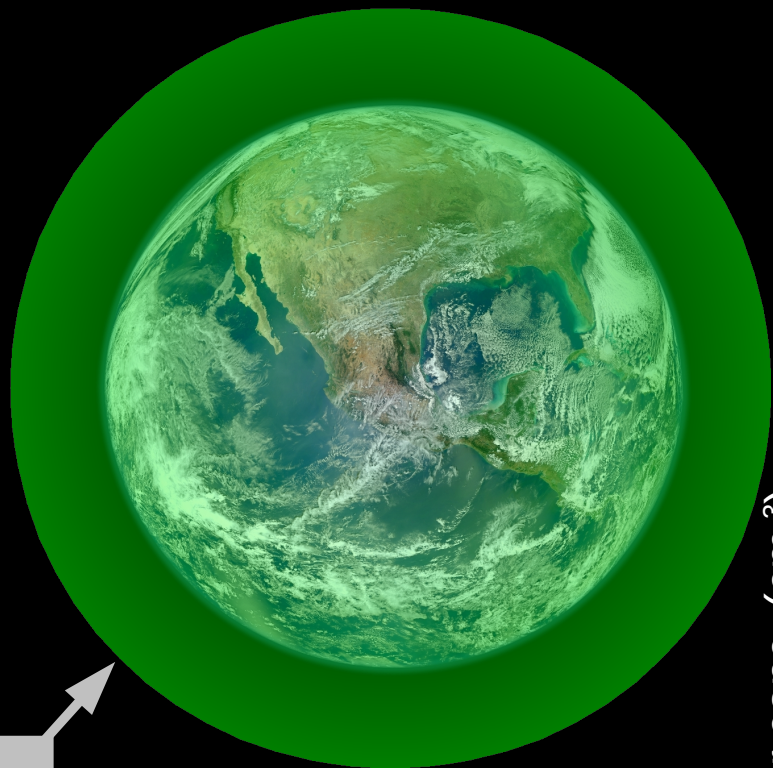
Total precipitation



2xCO₂

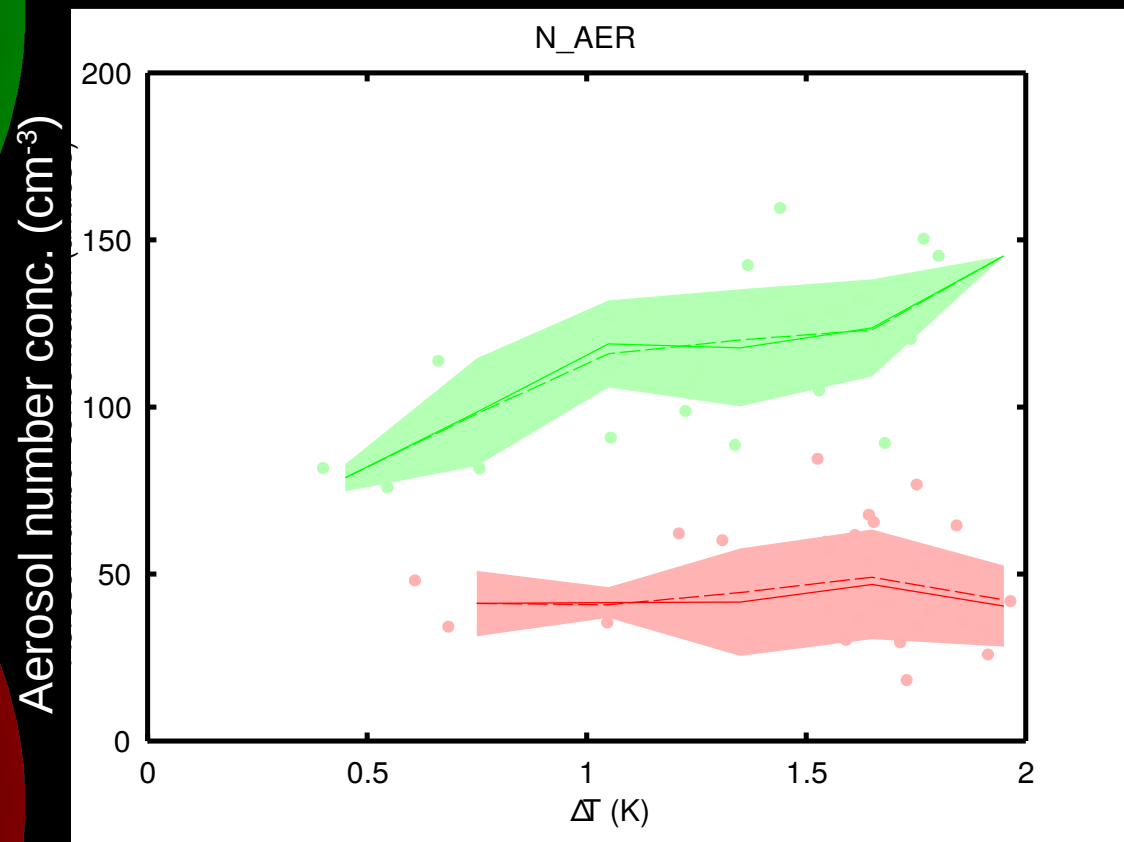
Monoterpene concentration

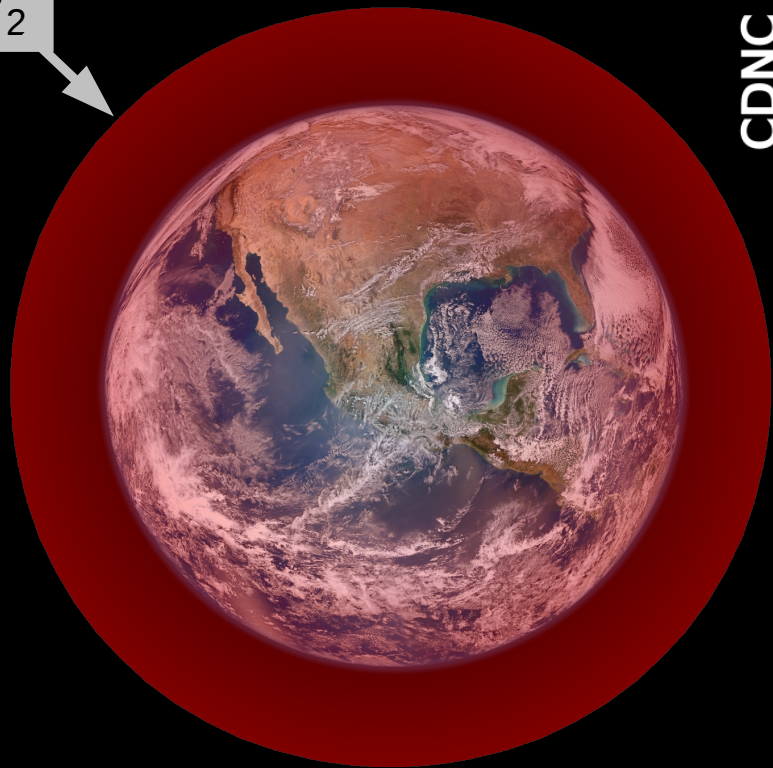
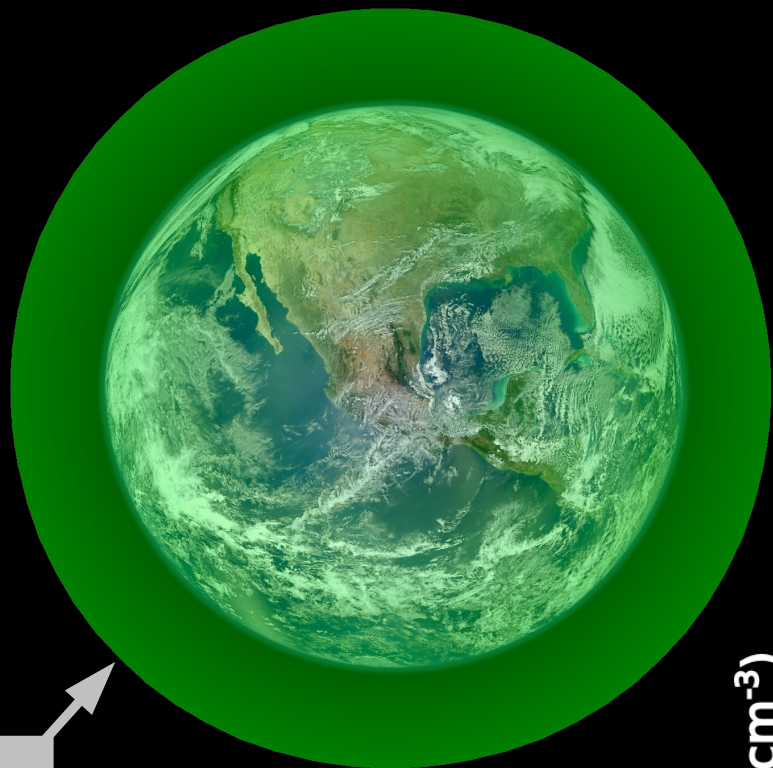




2xCO₂

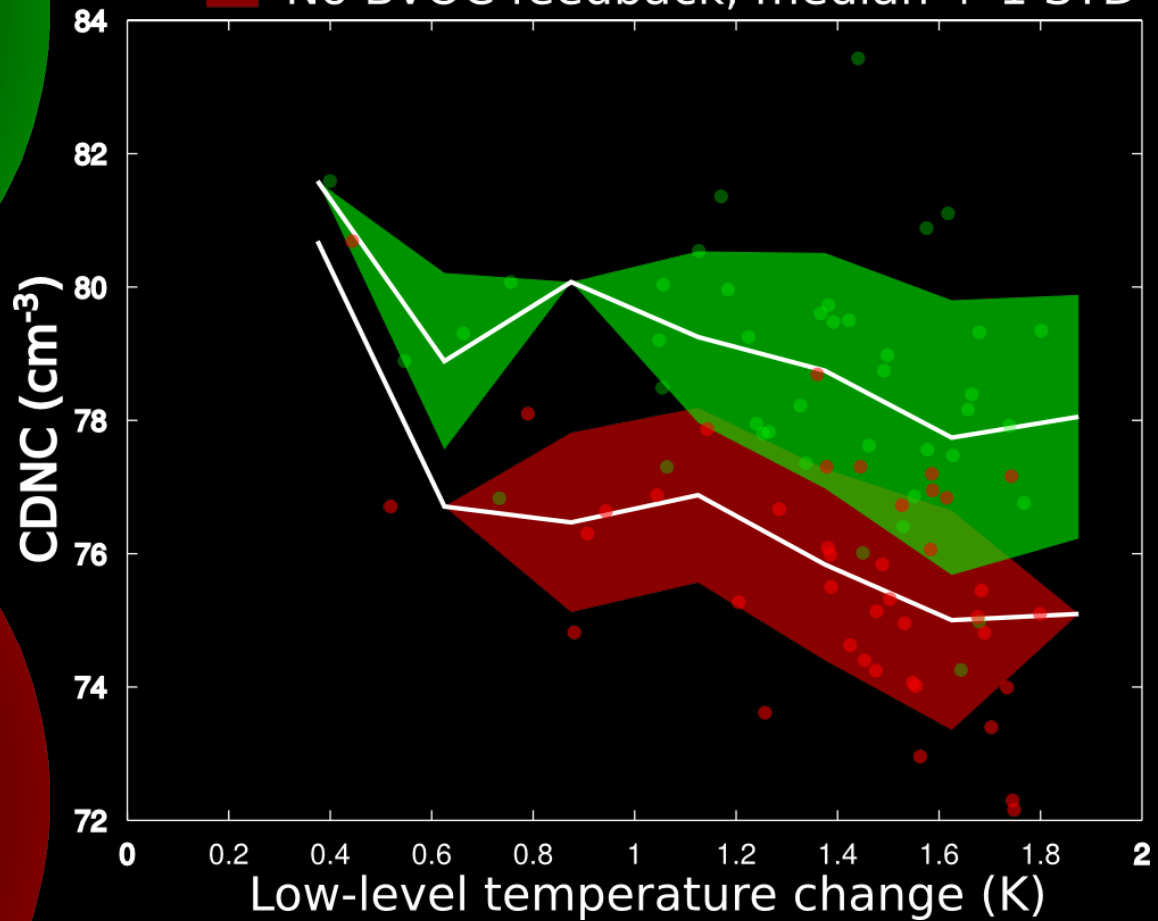
Aerosol number concentration

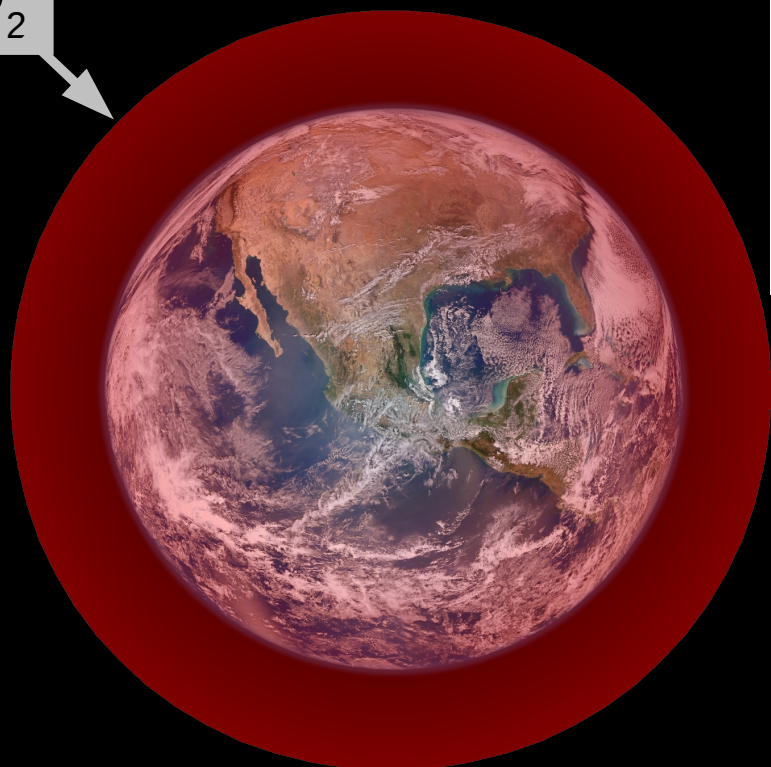
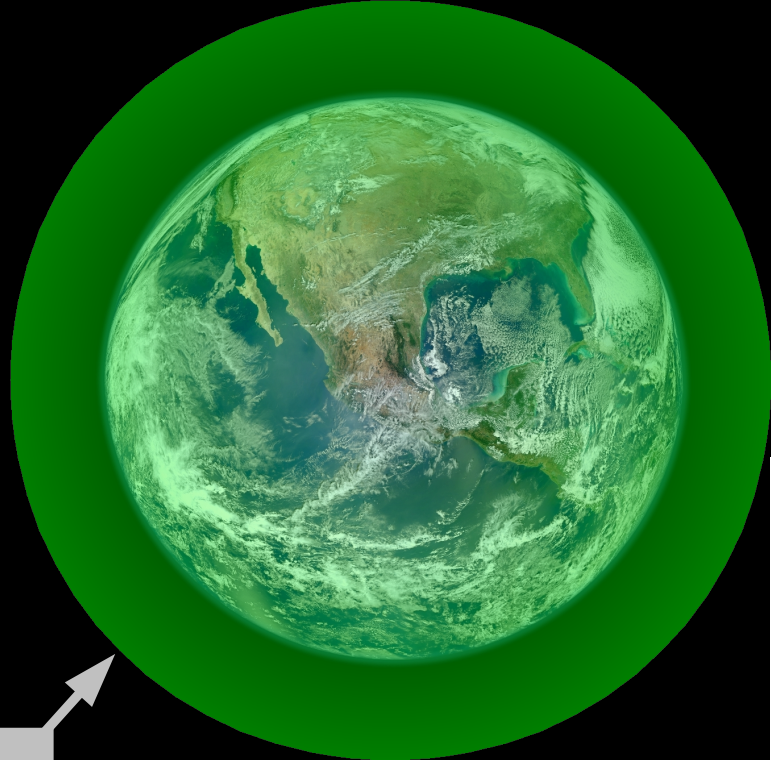




2xCO₂

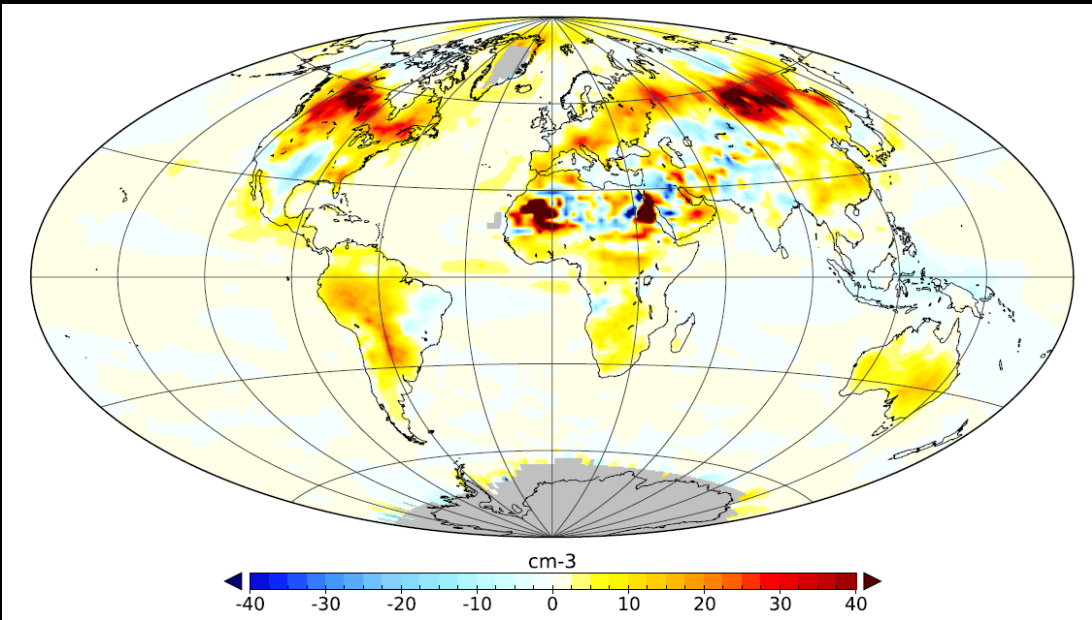
■ BVOC feedback, median + 1 STD
■ No BVOC feedback, median + 1 STD

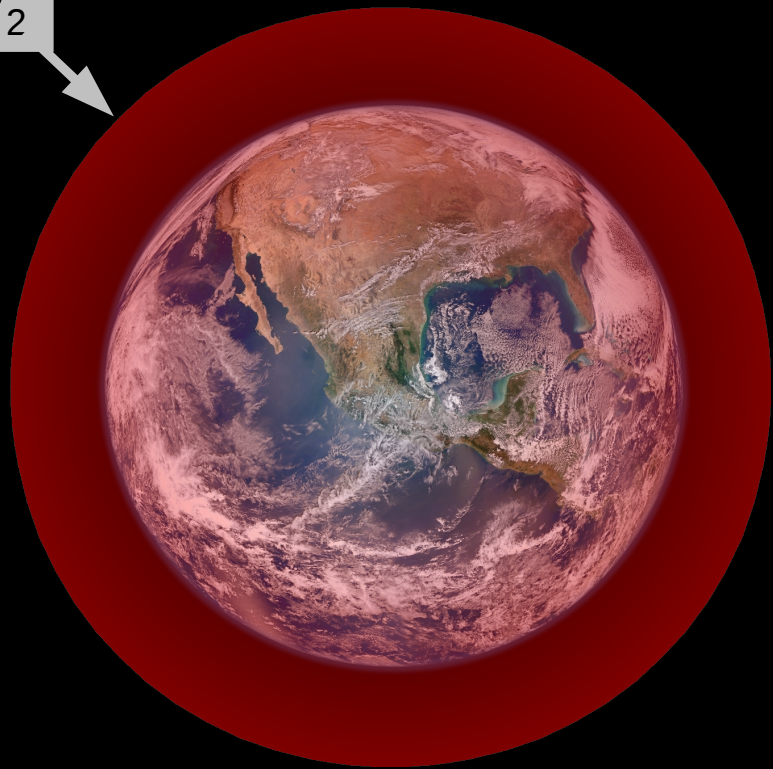
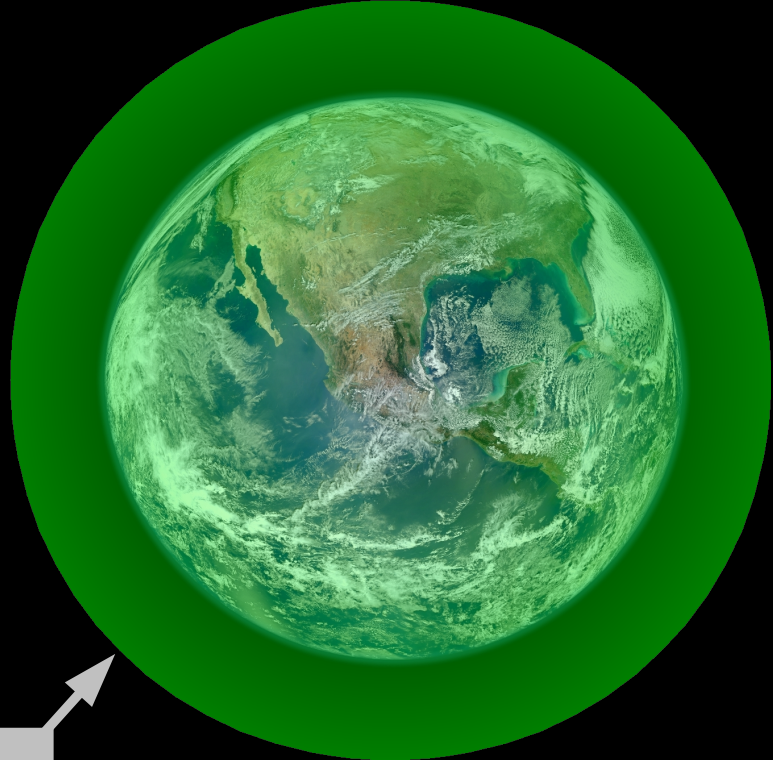




2xCO₂

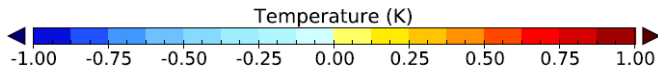
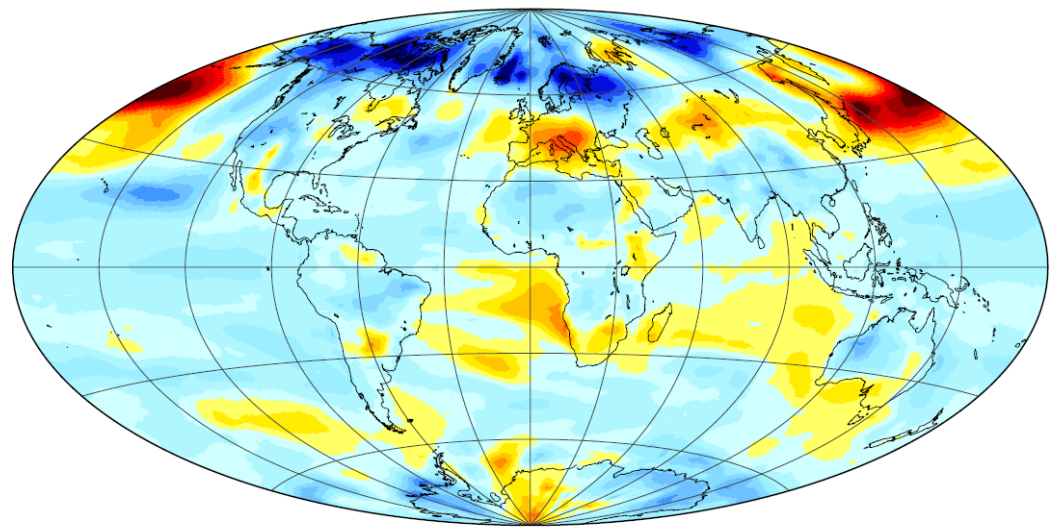
CDNC: **FEEDBACK** – **NO FEEDBACK**

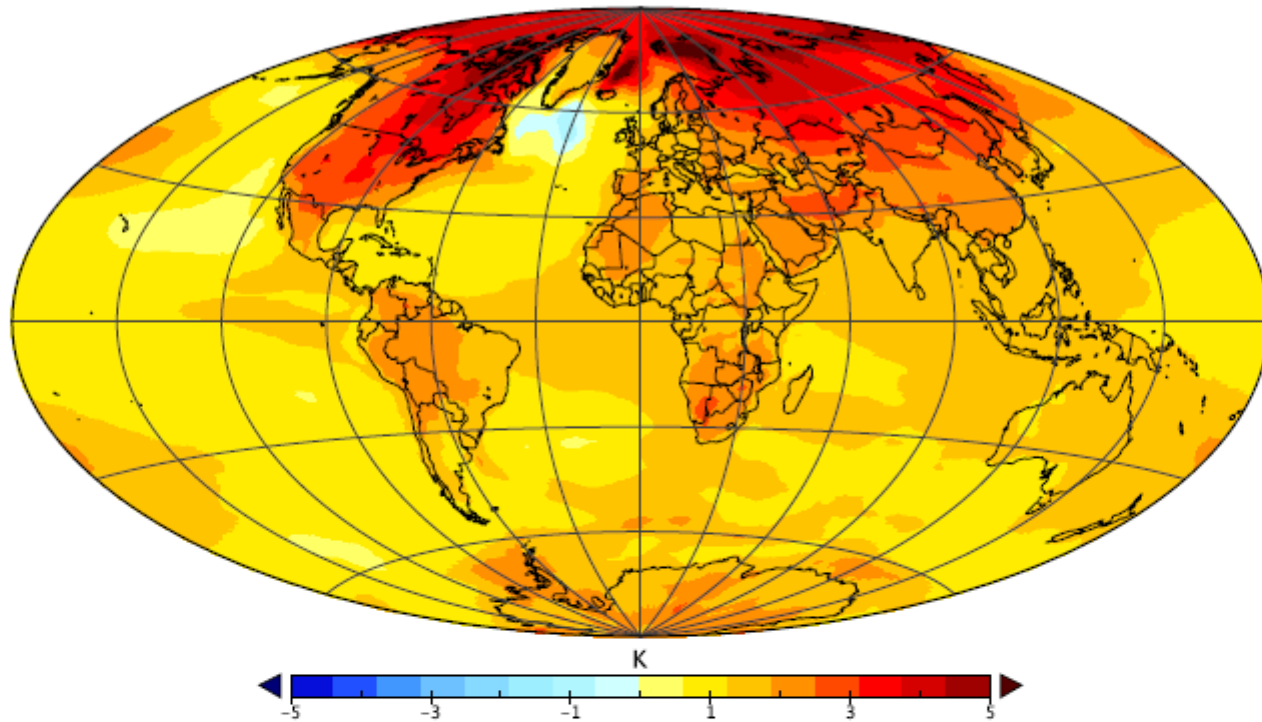




2xCO₂

Temperature:
FEEDBACK – NO FEEDBACK

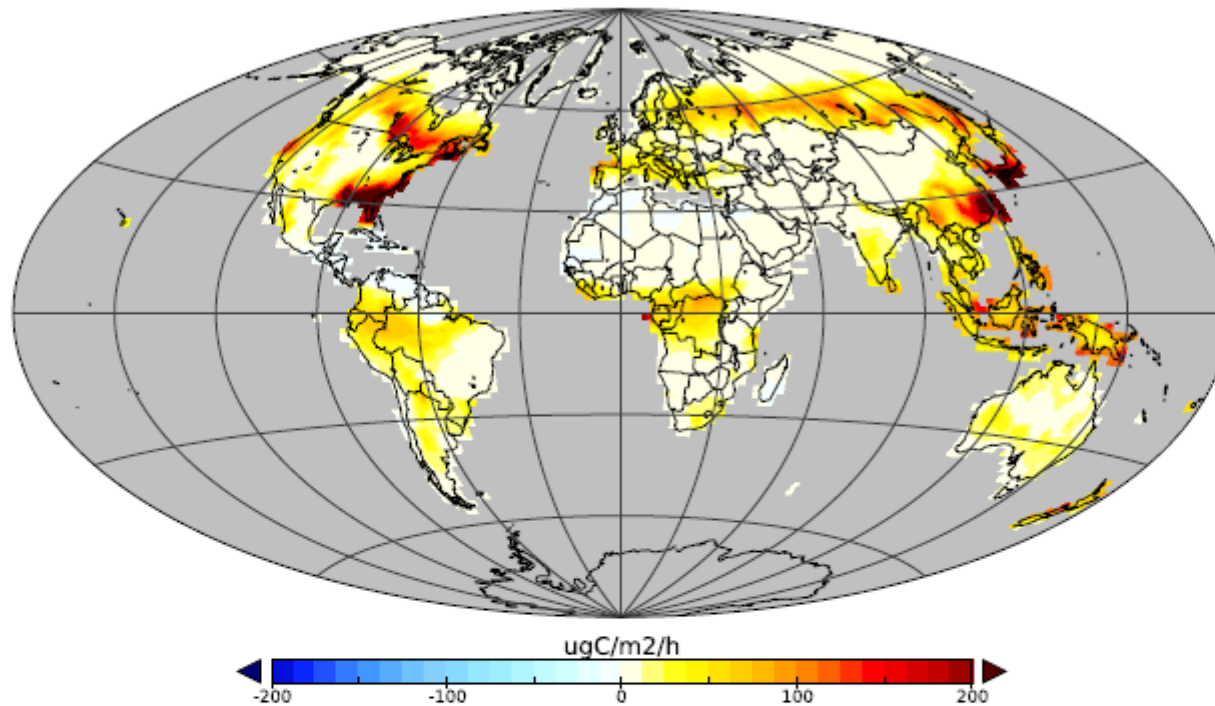




Climate change from doubled CO₂

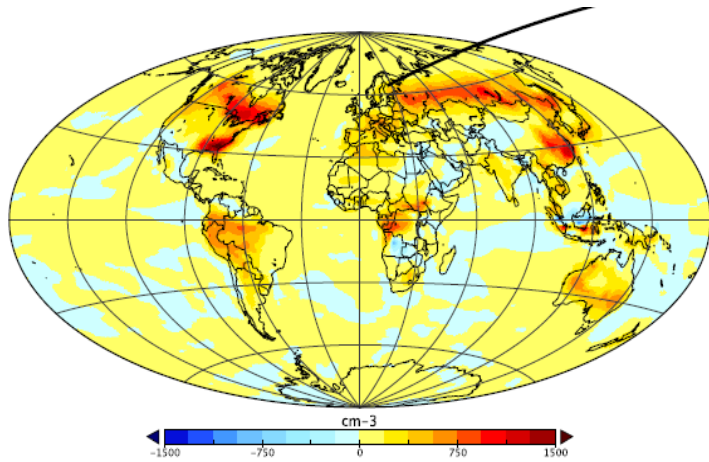
(NoFB_2xCO₂ - NoFB_1xCO₂)

The above figure show the response of the model to doubled CO₂ concentration, with BVOC feedback switched off. Global average atmospheric temperatures increase by ~3.5 K, and the change is most pronounced in the Northern Hemisphere and the Arctic.



Increase of monoterpene emission in warmer climate (FB_2xCO₂ - CTRL)

The increase in temperature and CO₂ lead to an increase in biogenic VOC emissions. Monoterpene emissions are increased globally by ~20%.



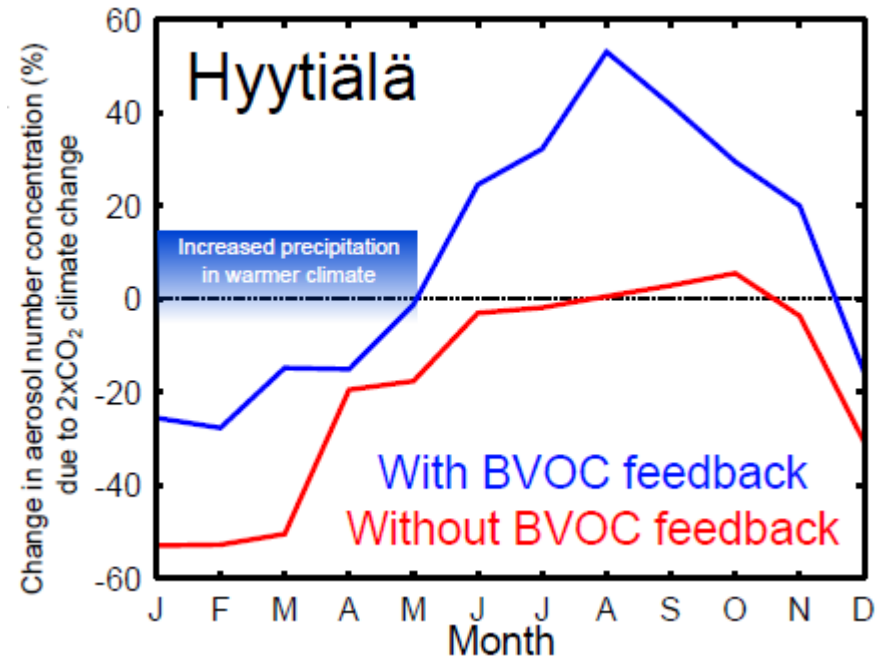
Increase in aerosol number concentration
(FB_2xCO₂ - CTRL)

The increase in VOC emission results in increased SOA formation, which both increases aerosol mass and growth of nucleated particles.

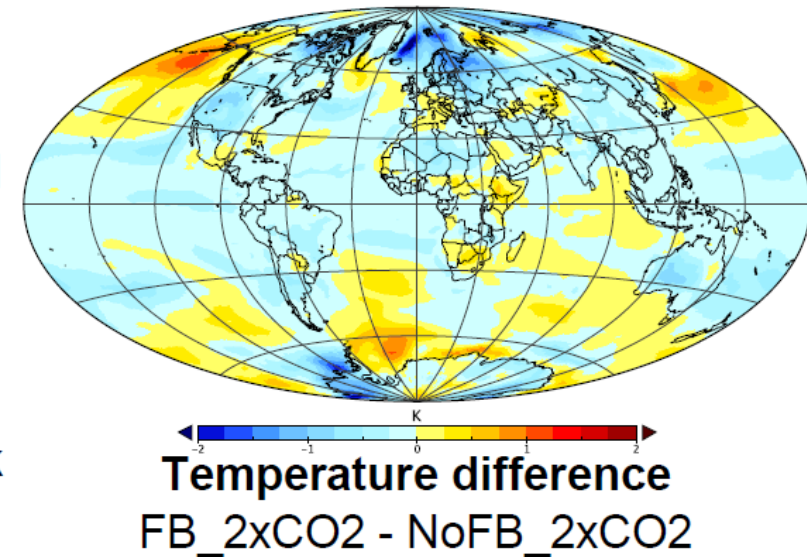
Cooling effect of increased BVOC

Increase in aerosol formation, aerosol optical depth and cloud droplet number concentrations in the 2xCO₂ climate lead to increased aerosol forcing. The cooling effect of this additional forcing can be seen by comparing the equilibrium temperatures of the experiments (figure on the right).

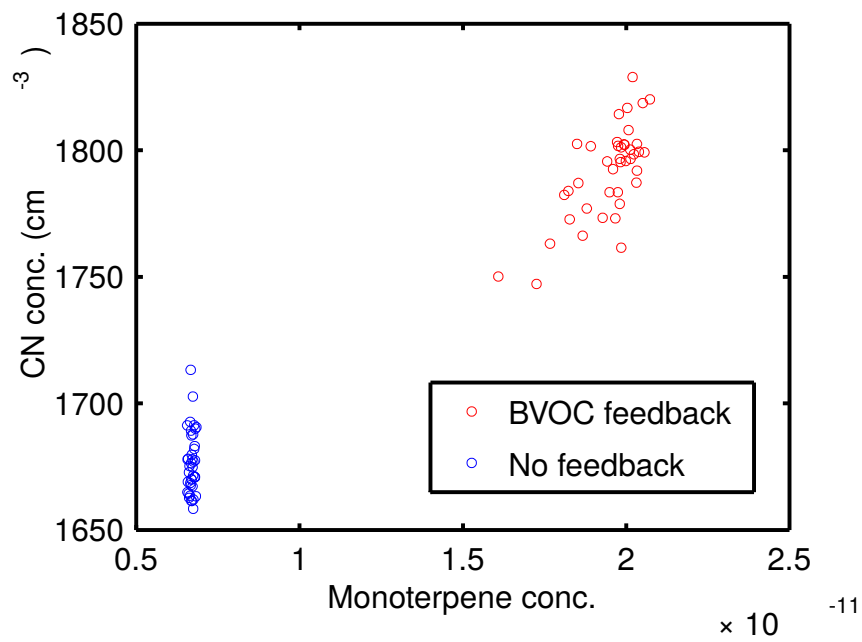
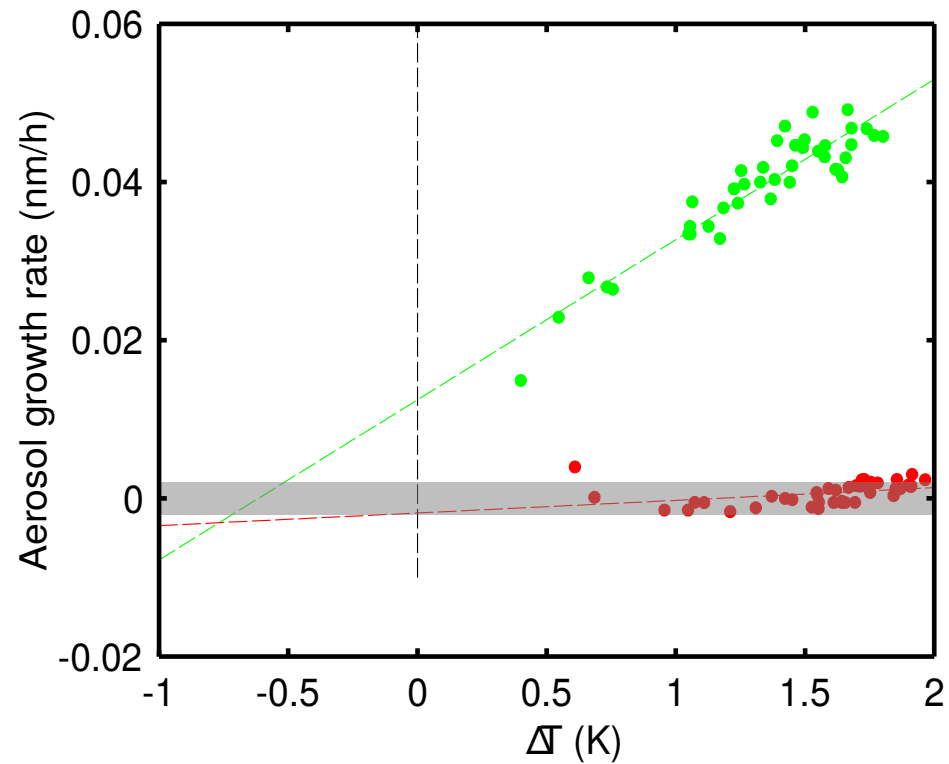
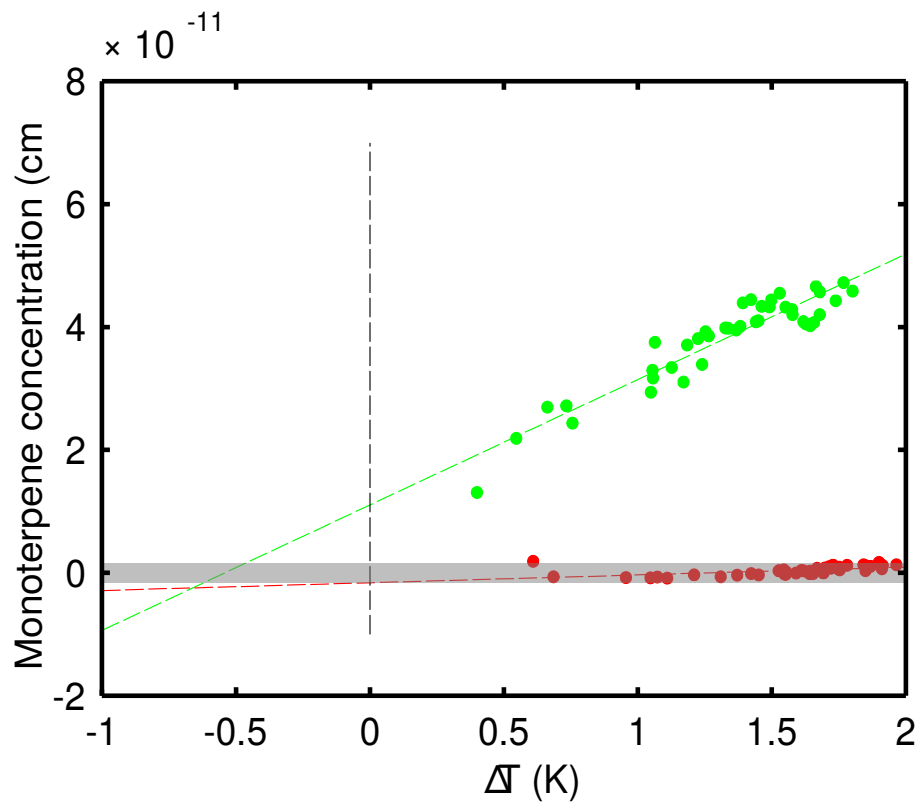
According to the simulations, BVOC-aerosol-climate feedback can act to reduce the climate warming.

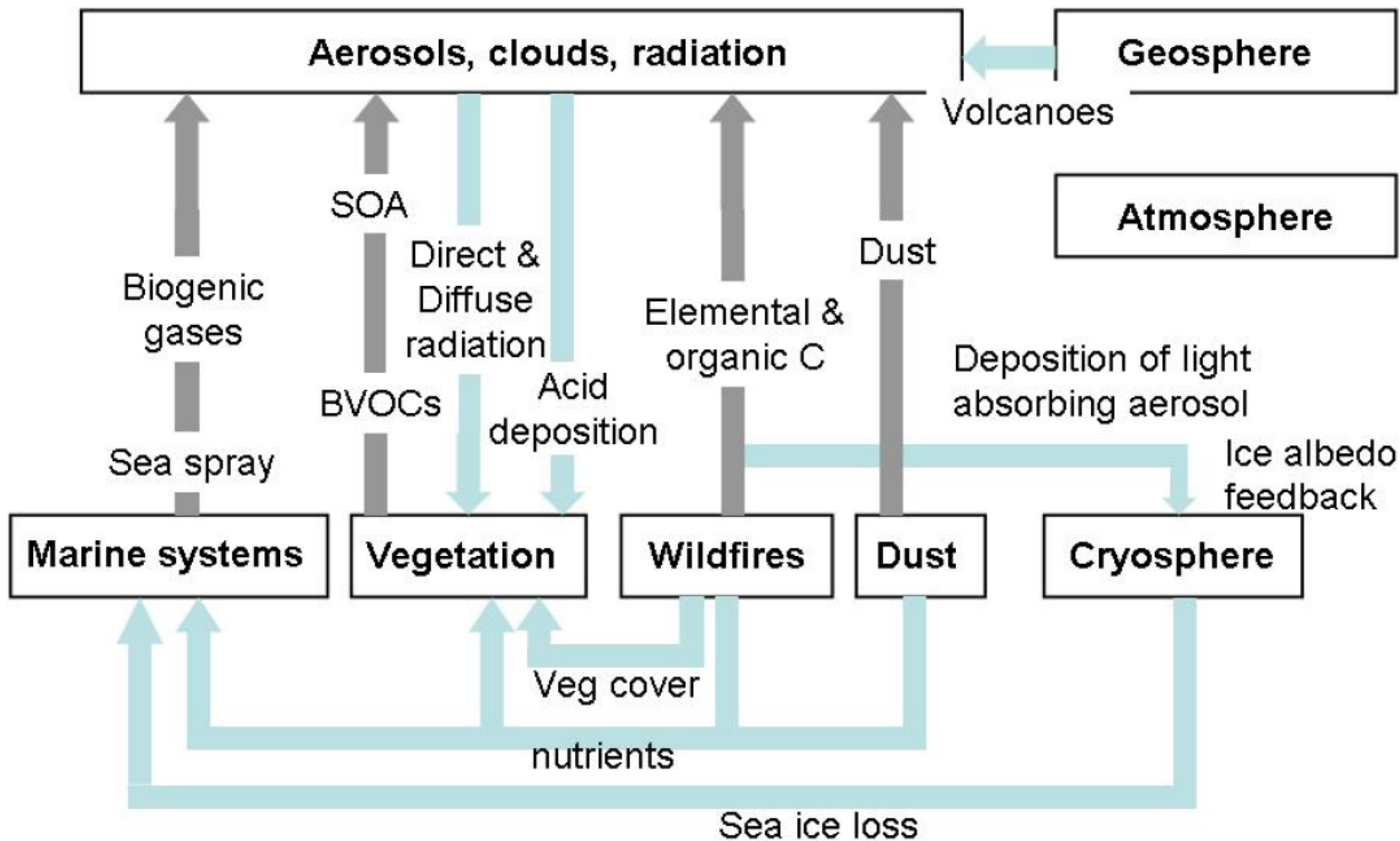


The signal from increased BVOC emission is modulated by changes in simulated climate. In Hyytiälä, for example, particle number is increasing in summer due to increased BVOCs, but decreasing in winter due to increased precipitation.

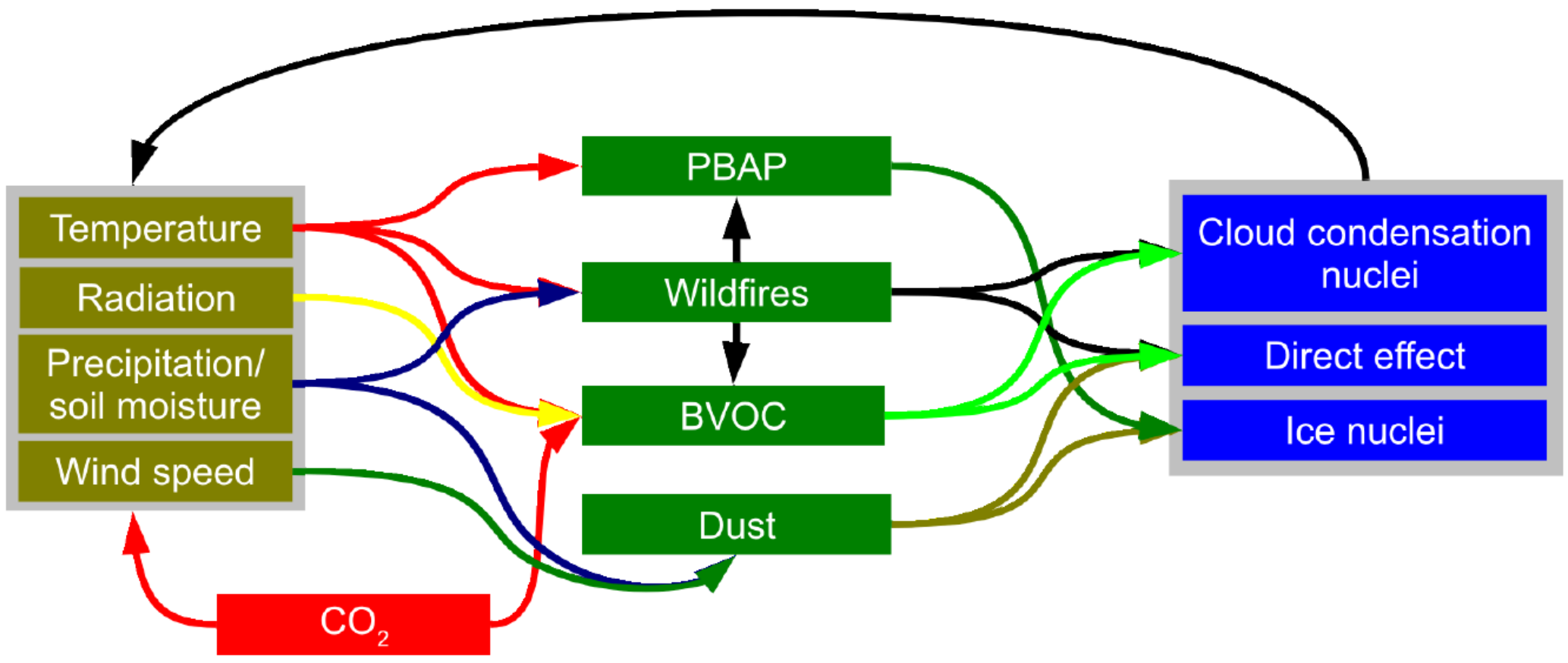


Temperature difference
FB_2xCO₂ - NoFB_2xCO₂

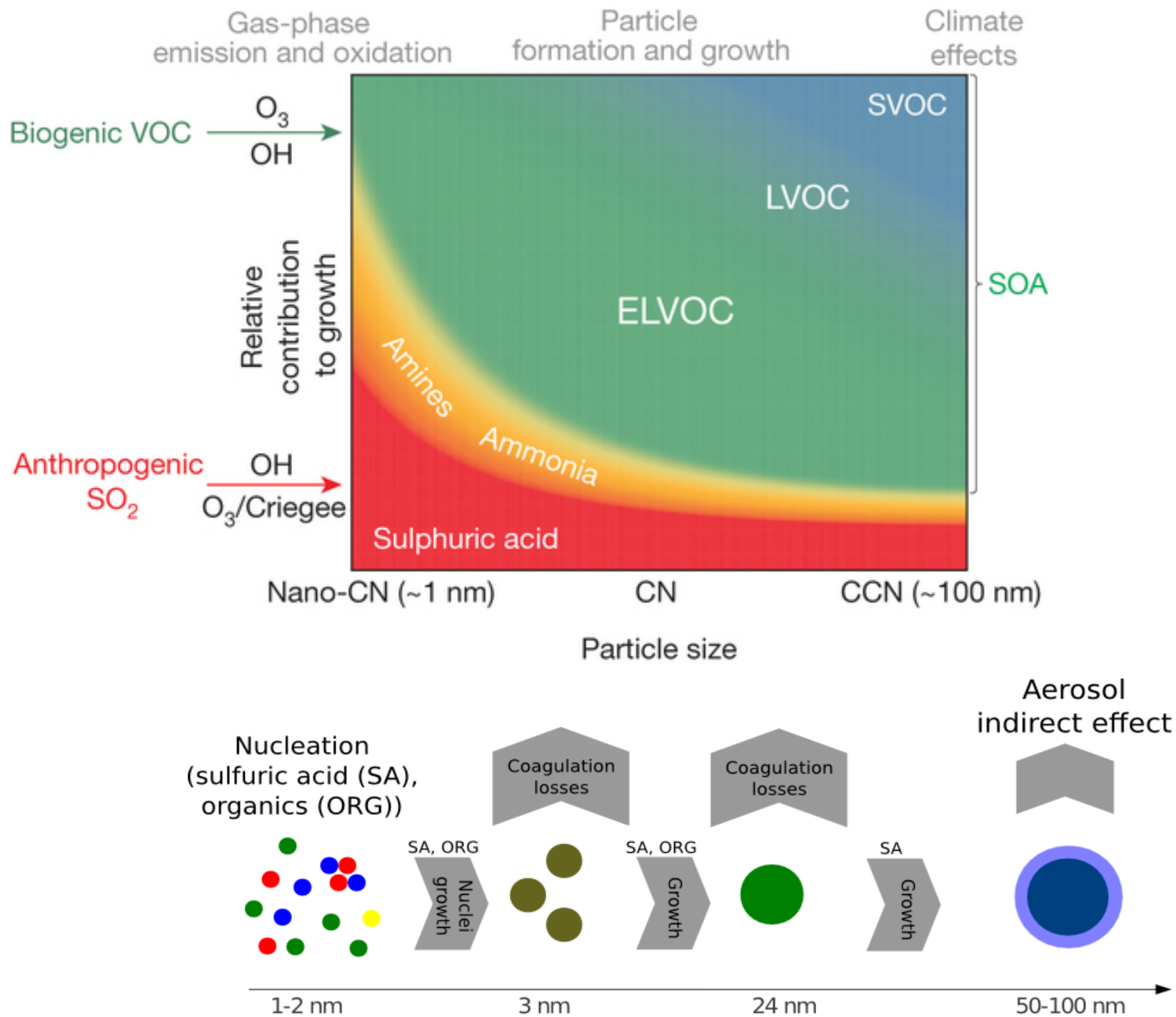




Direct aerosol effects
Indirect aerosol effects

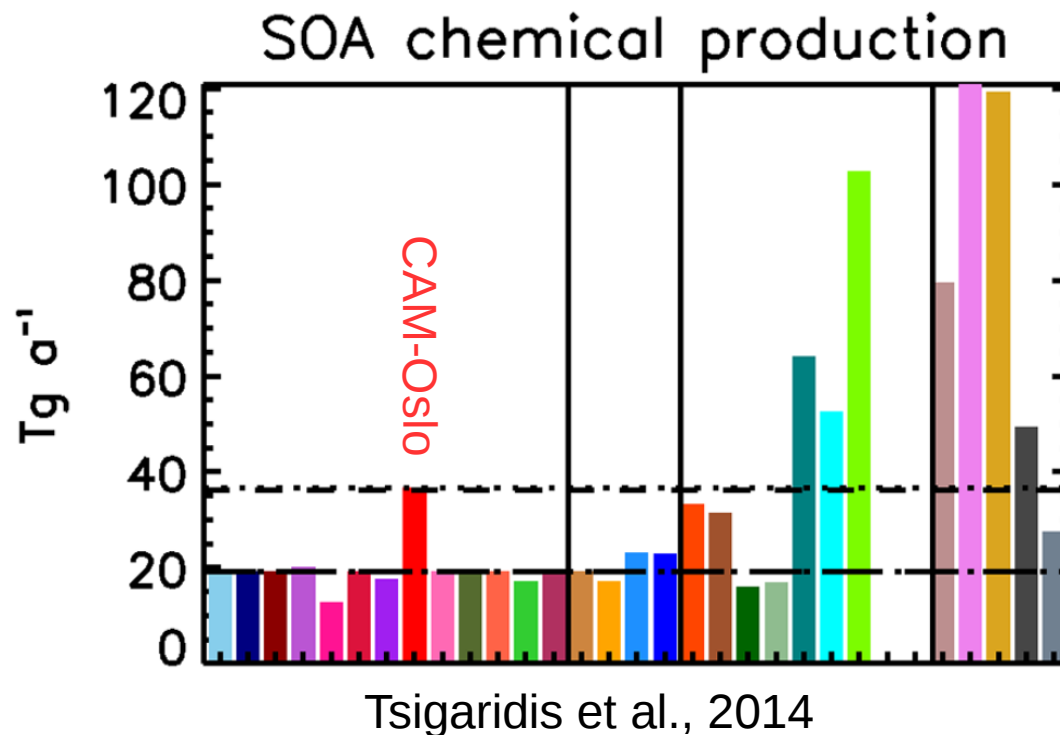


Extremely Low-volatility organic compounds (ELVOCs)



Near-future work

- (Aerosol-)climate feedback simulations for CRAICC
- Evaluation of Arctic aerosols in CRAICC/PEEX/eSTICC
- Implement new nucleation mechanisms (e.g. ion-induced)
- Implementing/improving ELVOC-chemistry in NorESM-NPF, working towards “missing SOA”



Spracklen et al., 2011:
50-380 Tg/a (140 Tg/a
best estimate)

