

Oxygenated volatile organic compounds in the remote marine troposphere: Results from the Cape Verde Atmospheric Observatory

Cape Verde Atmospheric Observatory (CVAO)
16° 52' N, 24° 52' W



THE UNIVERSITY *of York*

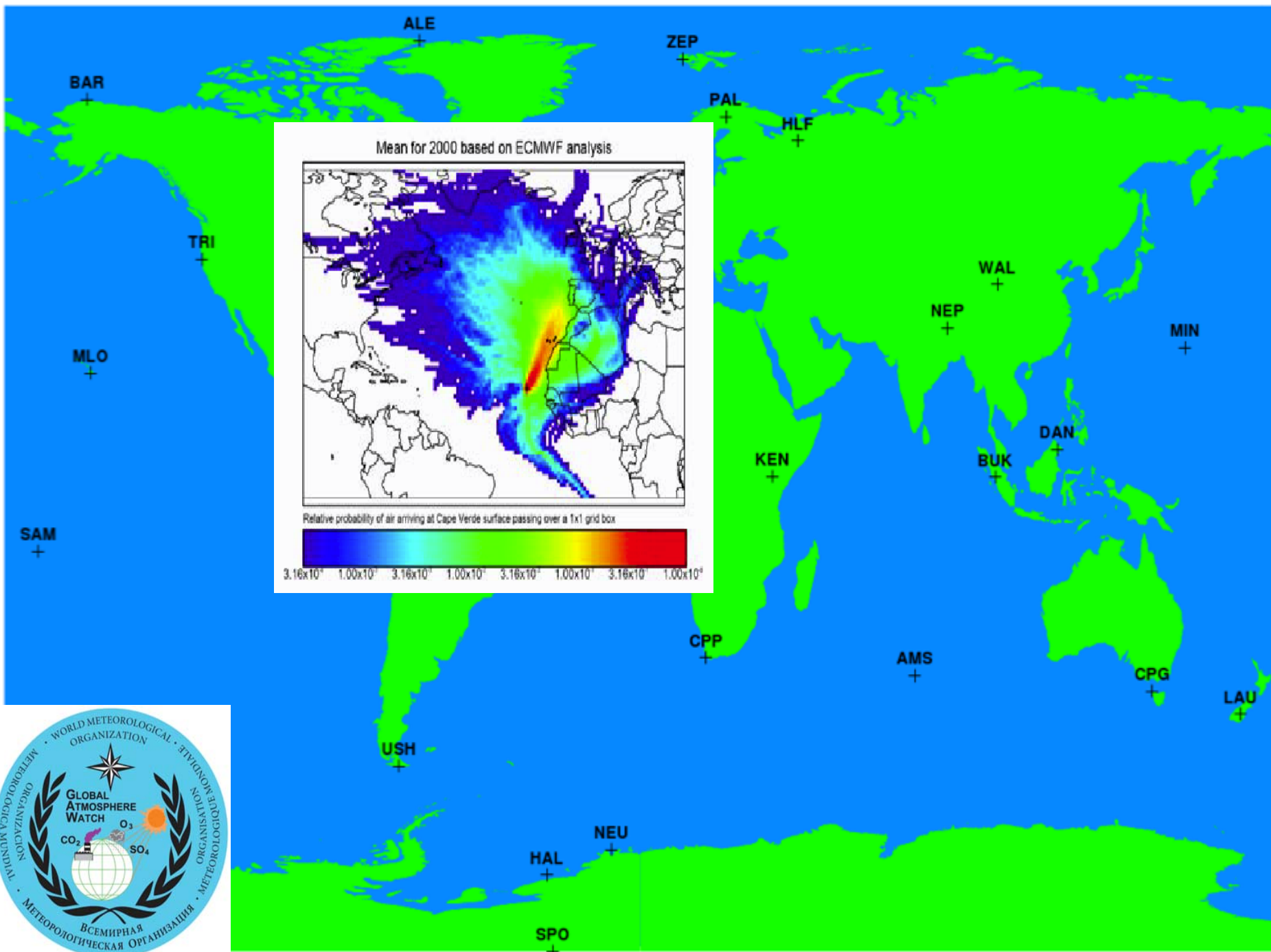
Lucy J. Carpenter, Katie Read, James Lee, Ally Lewis, James Hopkins -
NCAS, University of York

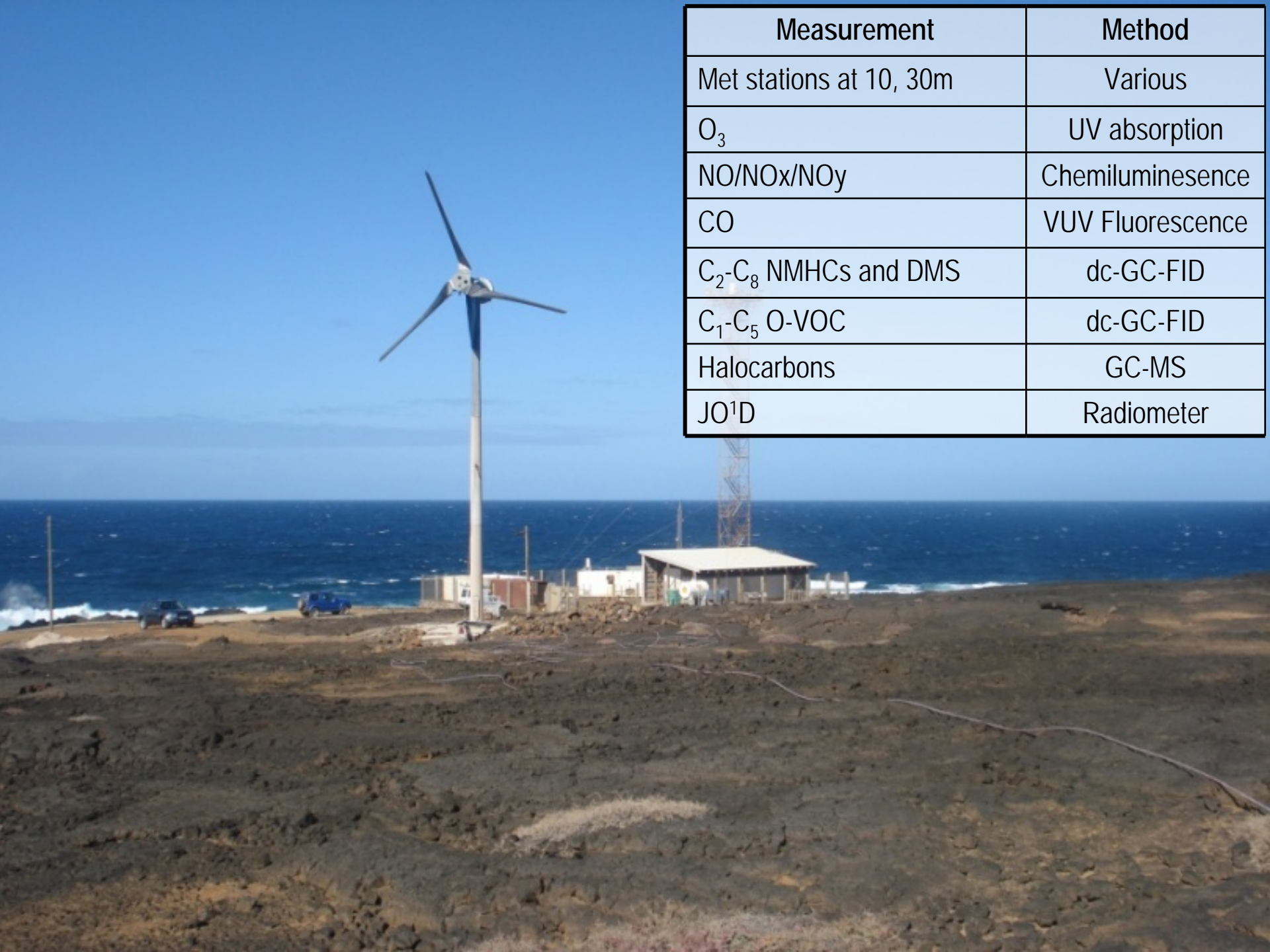
Luis Mendes, Helder Lopez - *INMG, Cape Verde*

Steve Arnold - *Earth and Environment, University of Leeds*

Rachael Beale, Phil Nightingale – *PML, UK*







Measurement	Method
Met stations at 10, 30m	Various
O ₃	UV absorption
NO/NO _x /NO _y	Chemiluminescence
CO	VUV Fluorescence
C ₂ -C ₈ NMHCs and DMS	dc-GC-FID
C ₁ -C ₅ O-VOC	dc-GC-FID
Halocarbons	GC-MS
JO ¹ D	Radiometer

Oxygenated volatile organic compounds (OVOCs)

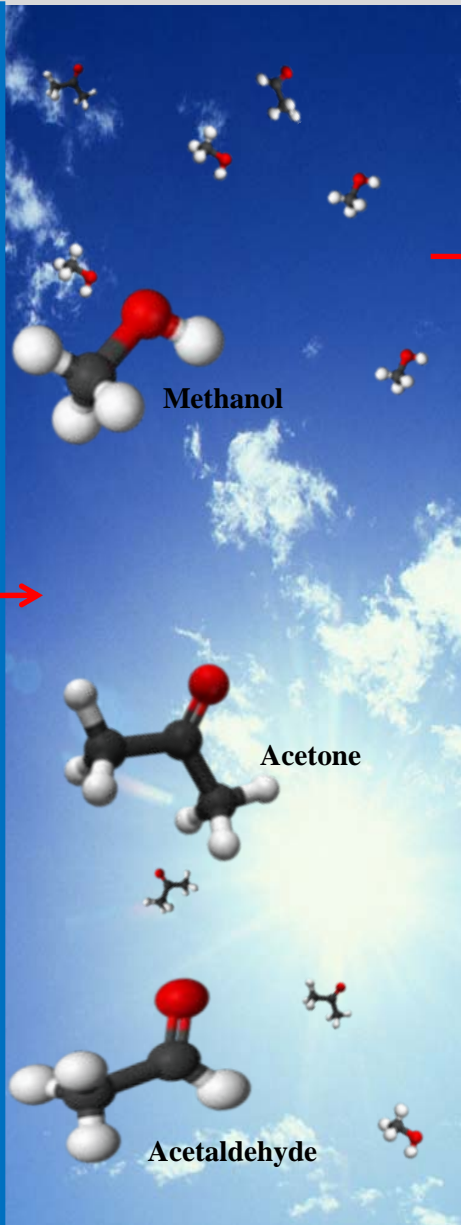
Atmospheric OVOCs

Ocean: source or sink?

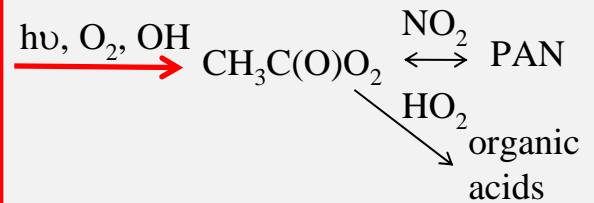
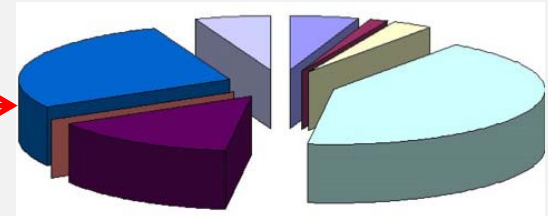
Anthropogenic and biomass burning sources

Biogenic sources

Primary and secondary sources

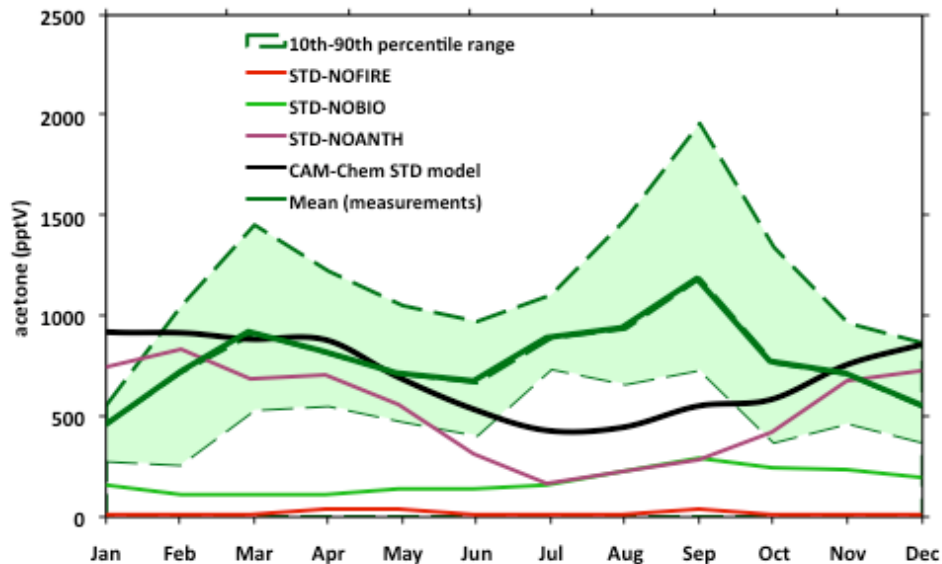
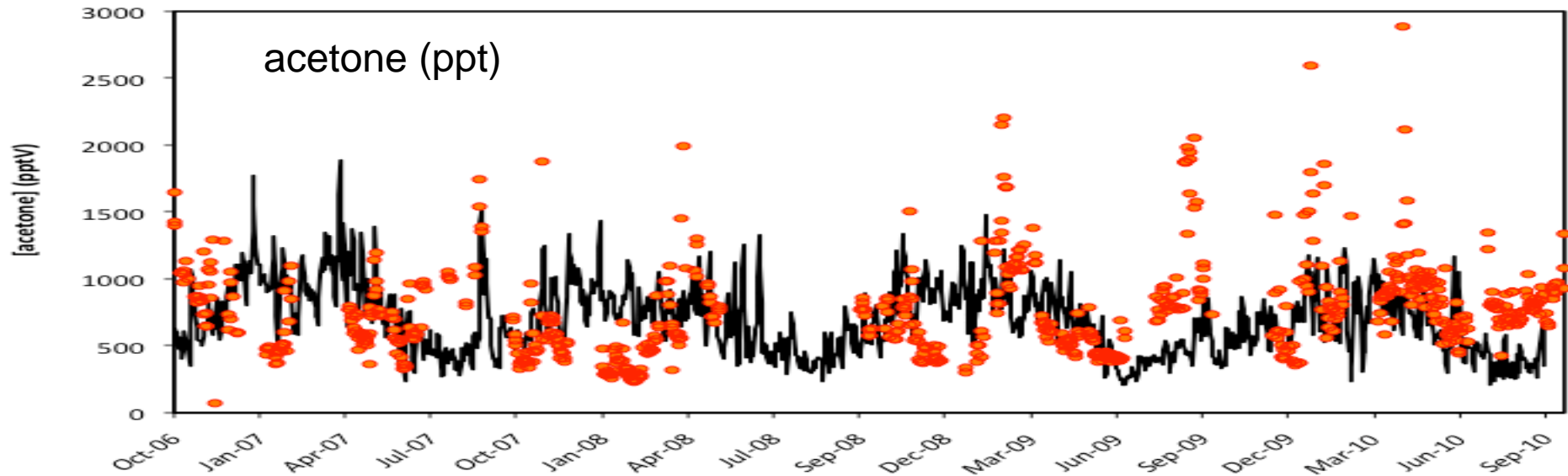


OH loss in marine boundary layer



Source of HO_x , O_3 and PAN

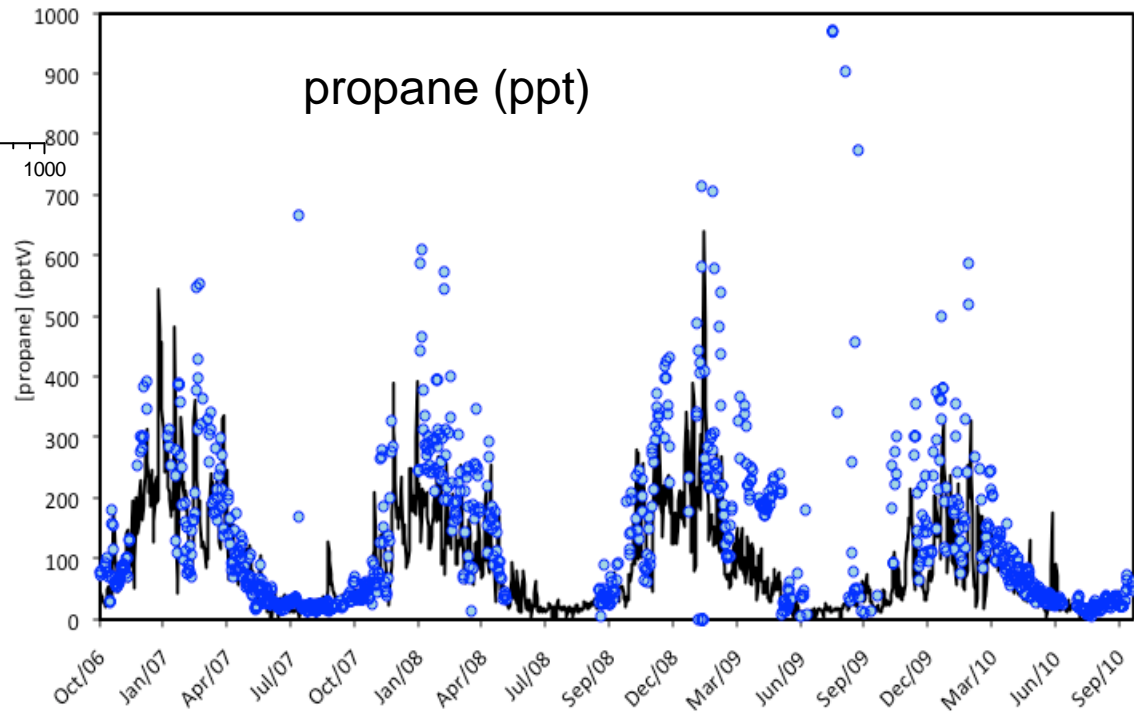
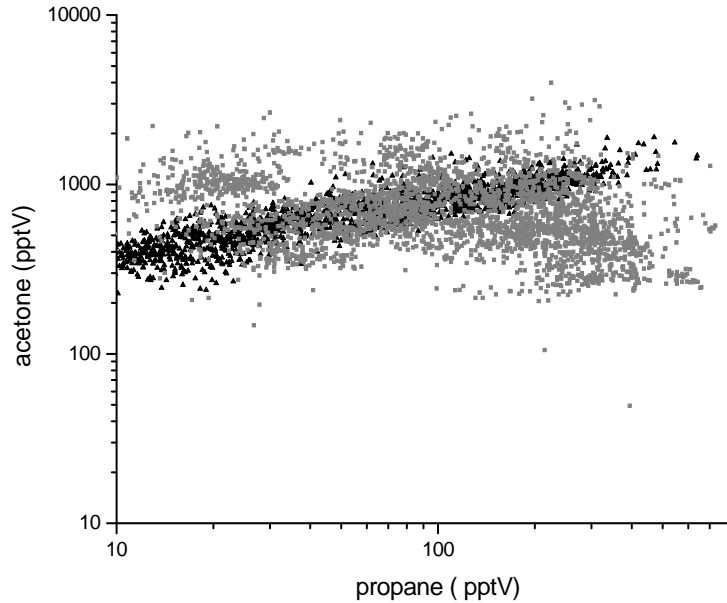
CAM-Chem vs measurements



Monthly averages

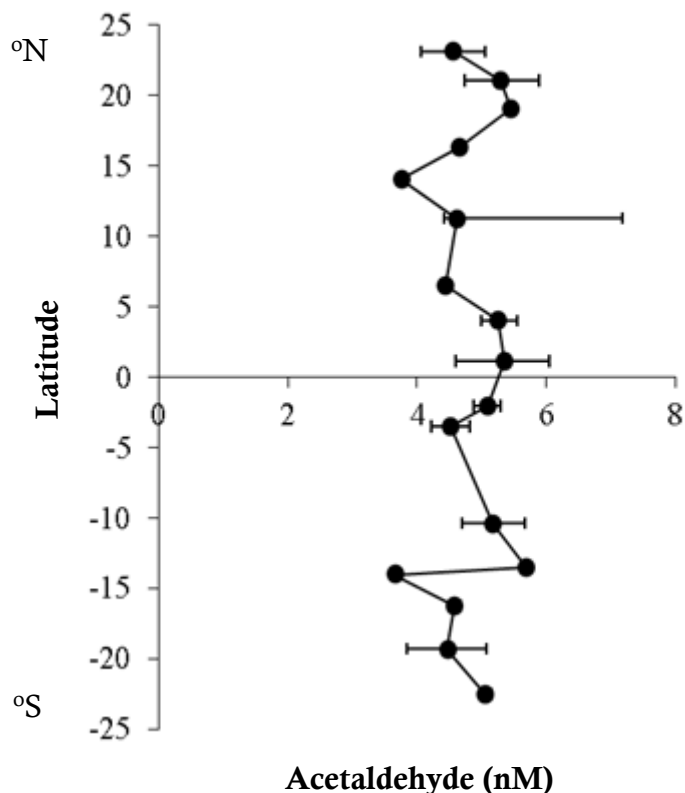
Dominated by anthropogenic emissions (39 %-91%)?

Are MBL acetone concentrations controlled by anthropogenic NMHC?

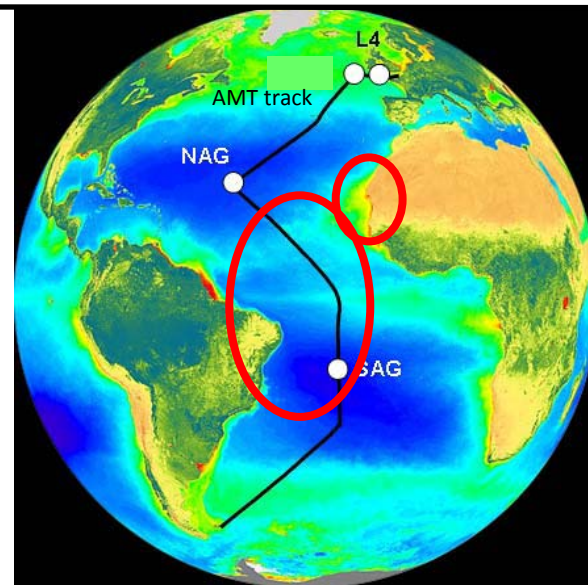


Role of the oceans?

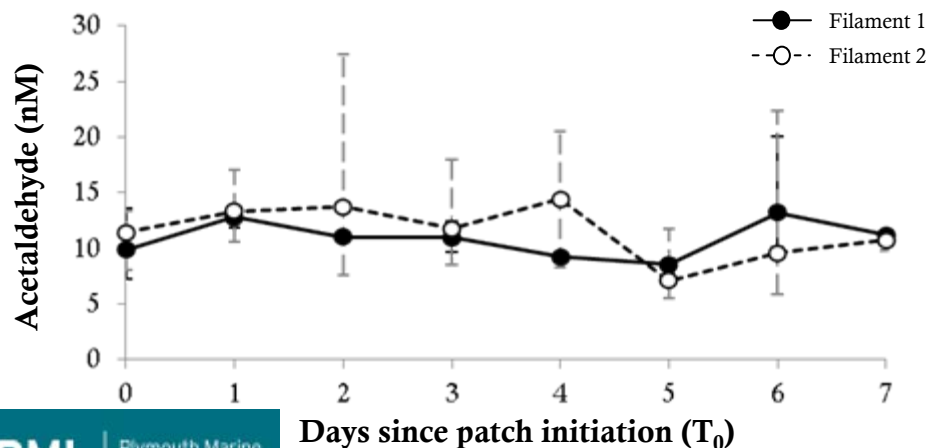
- Jacob et al. (2002)- ocean a significant source of acetone



- *Methanol, acetaldehyde and acetone quantified in seawater via MI-PTR/MS*



Acetaldehyde in Mauritanian Upwelling (ICON)

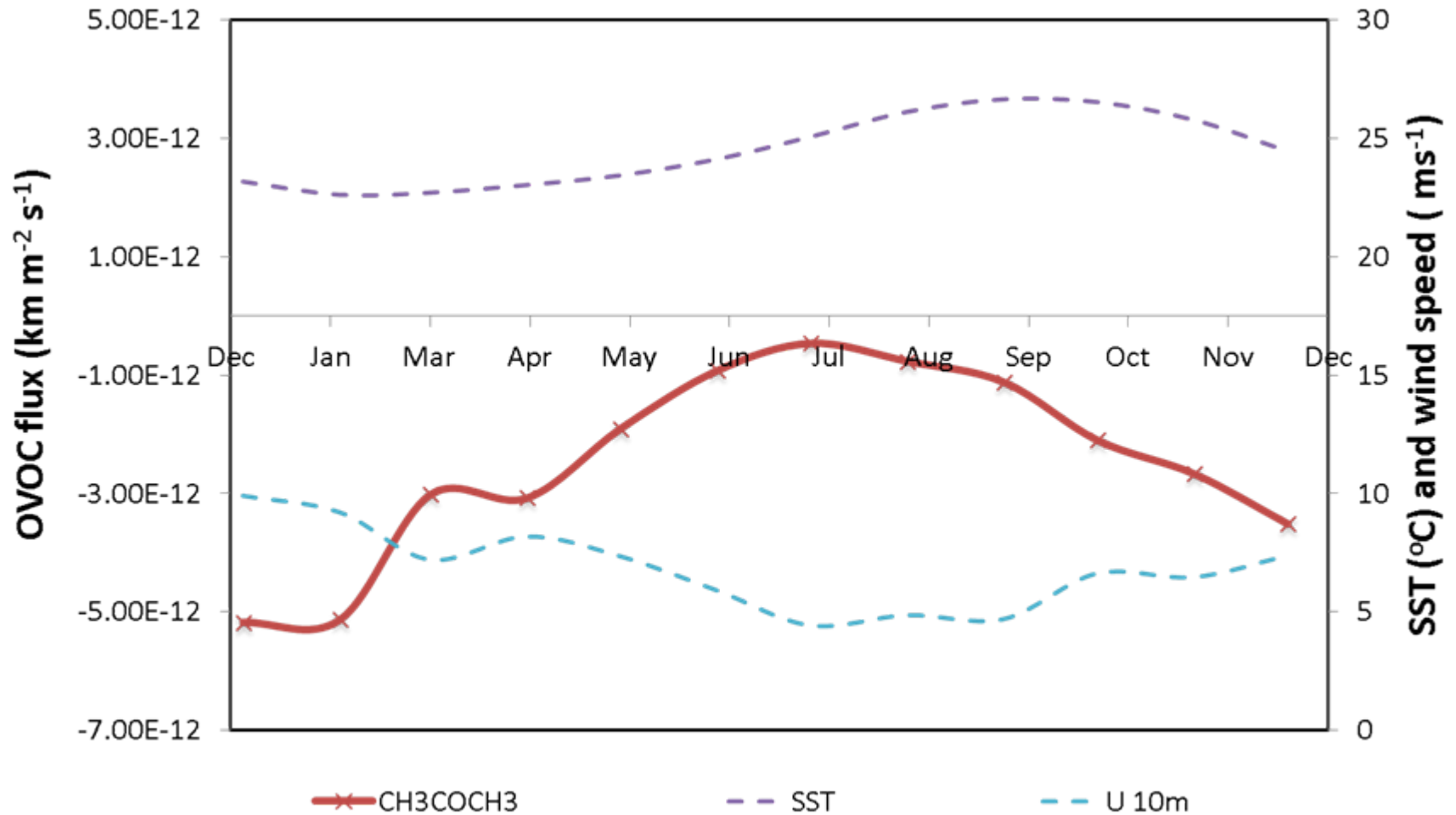


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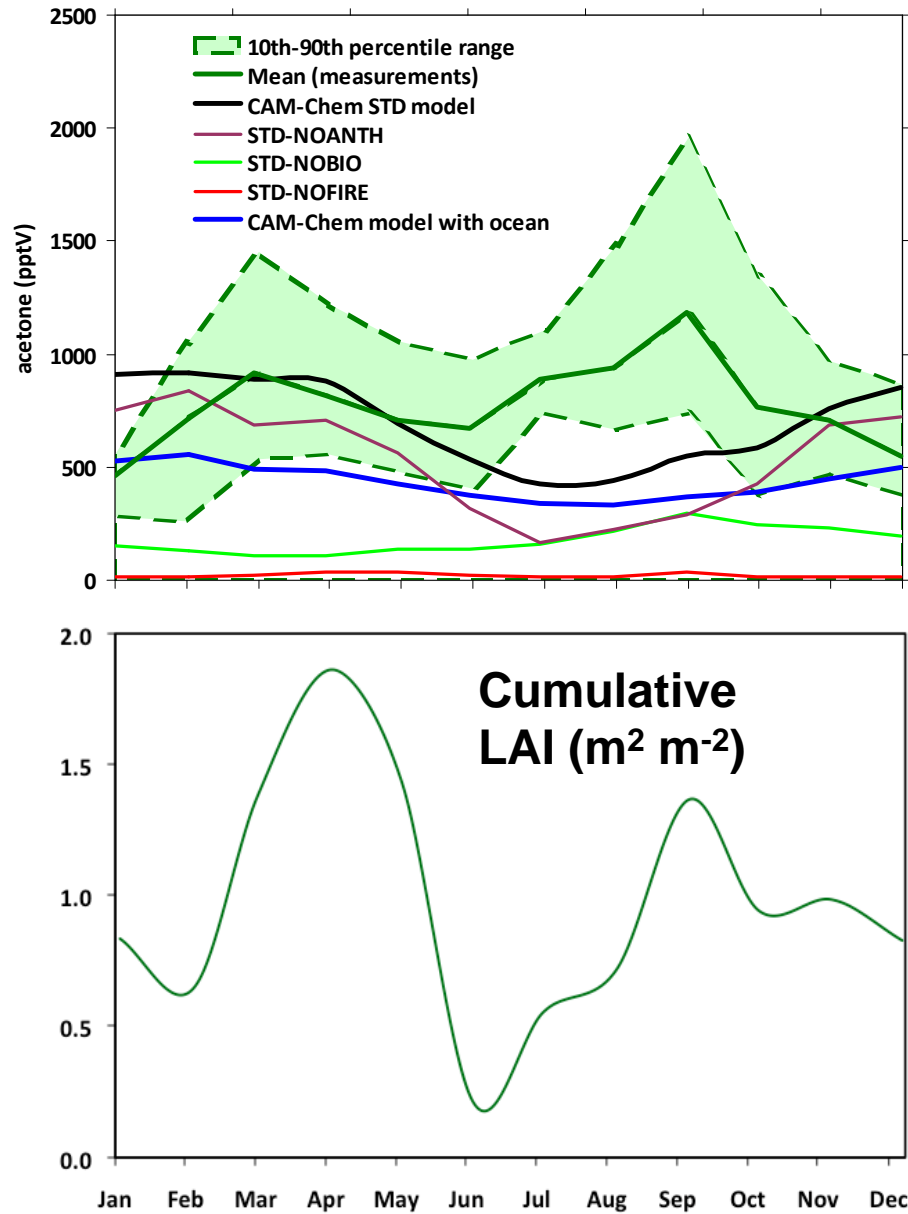
Beale, R. Quantification of oxygenated volatile organic compounds (OVOCs) in seawater, 2011, Ph.D thesis, University of East Anglia, UK. *Manuscripts in preparation.*

Modelled oceanic acetone fluxes

• Sea-air flux $F = k_t (C_w - C_a/H)$ $1/k_t = 1/k_w + 1/Hk_a$

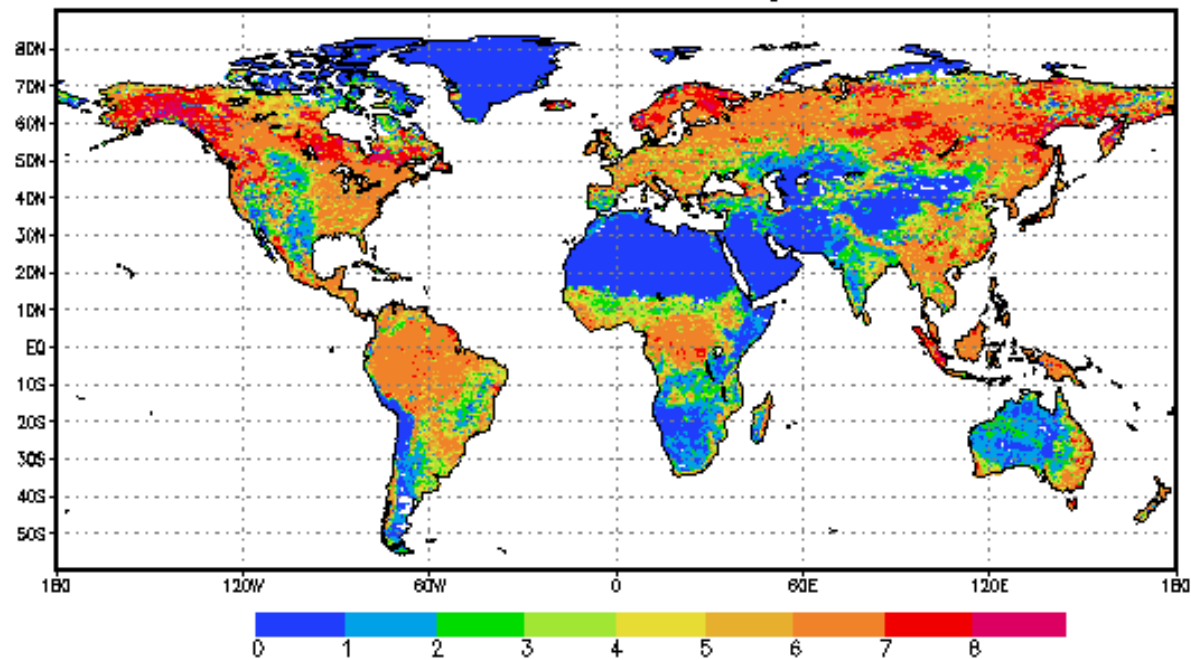


How does this change model results?

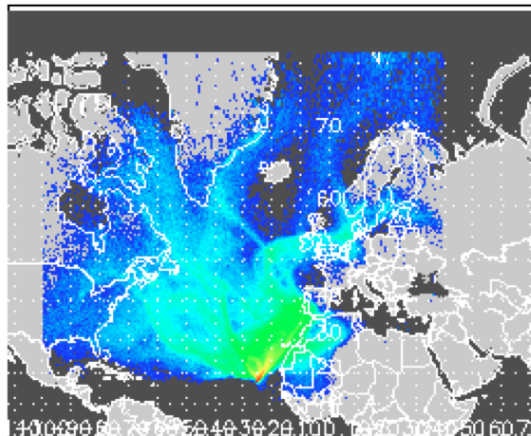


Biological (terrestrial) influences

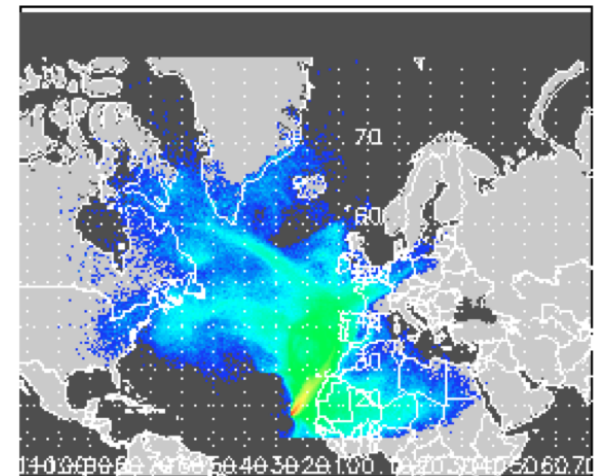
Leaf Area Index – August



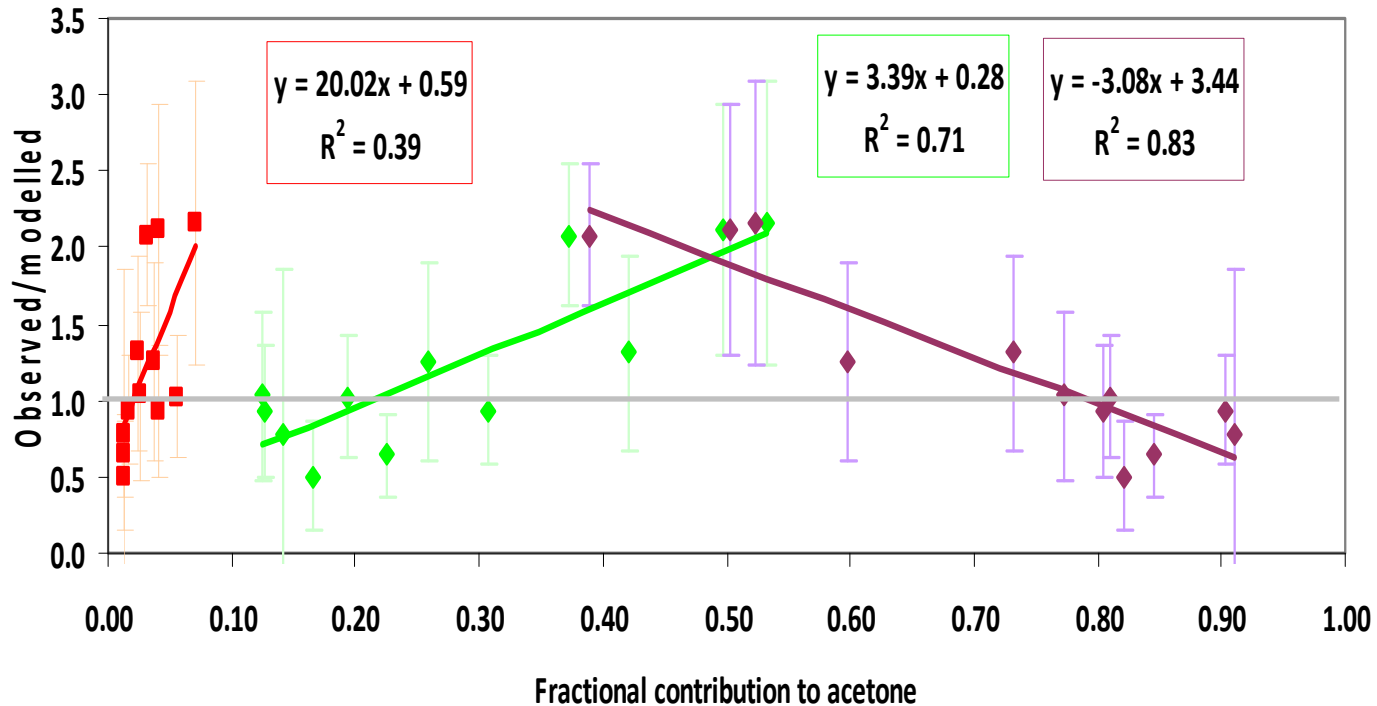
Year:2008 Month:4



Year:2008 Month:9



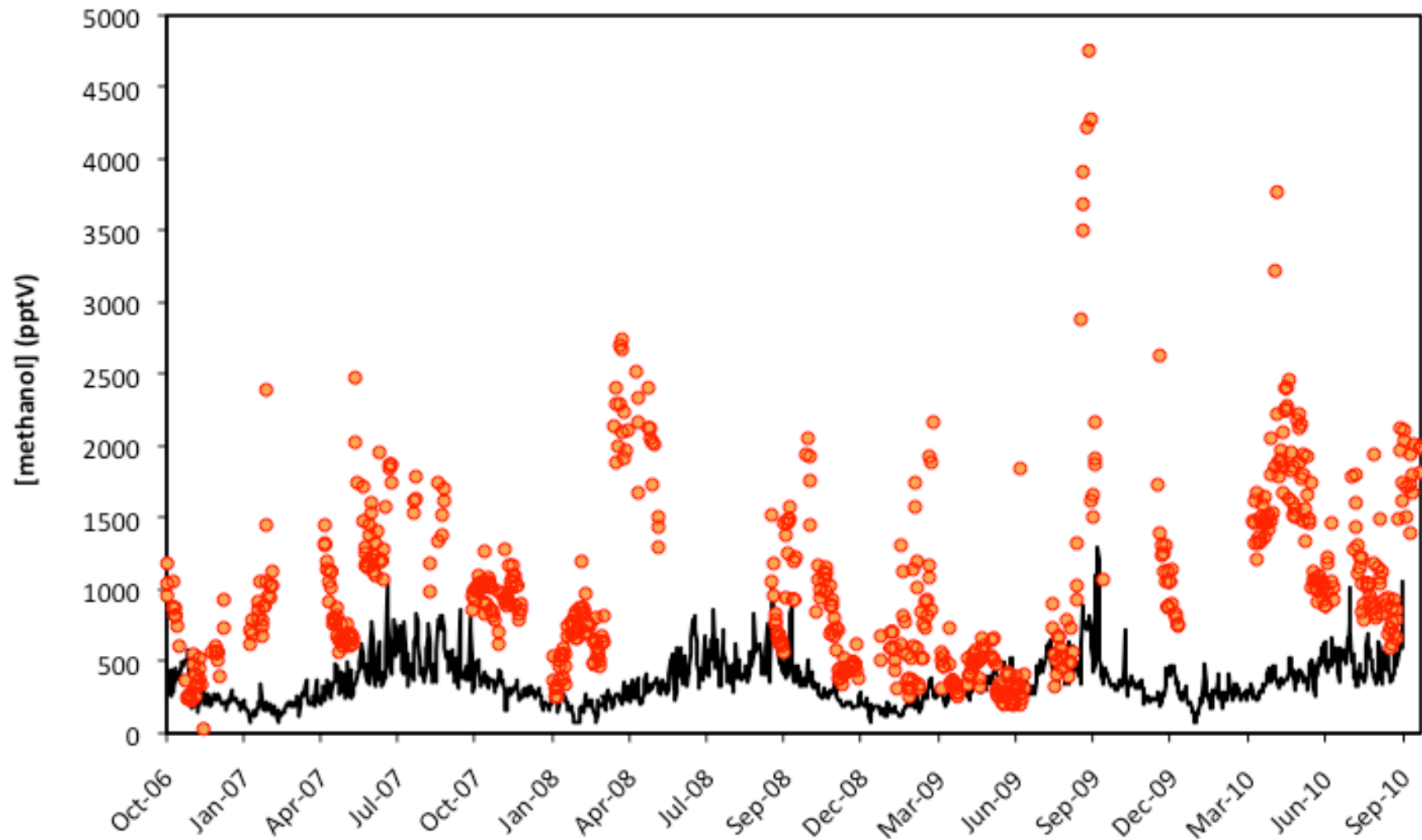
Could model bias be due to underestimated biogenic emissions?



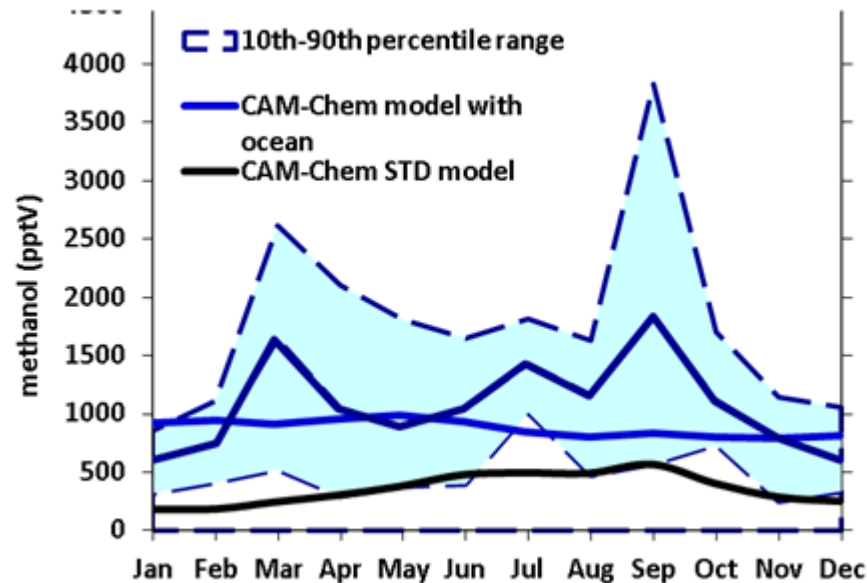
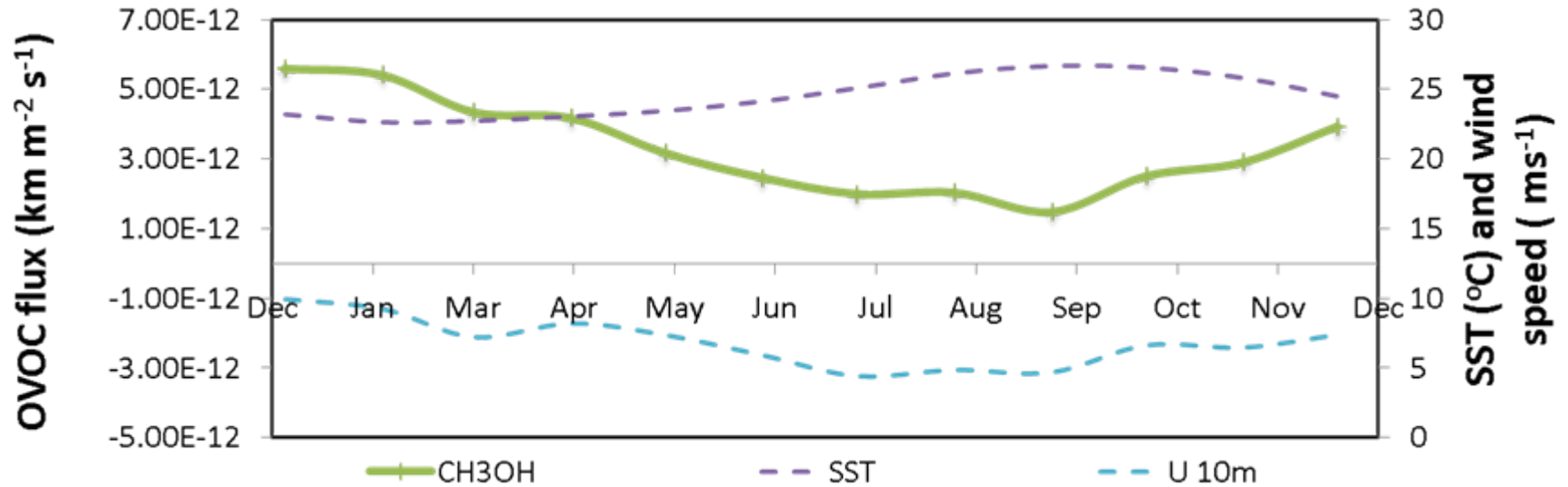
Fractional contribution from biogenic (green), anthropogenic (purple) and biomass burning (red) sources as calculated from CAM-Chem.

Grey lines indicate 1:1 observation:model agreement.

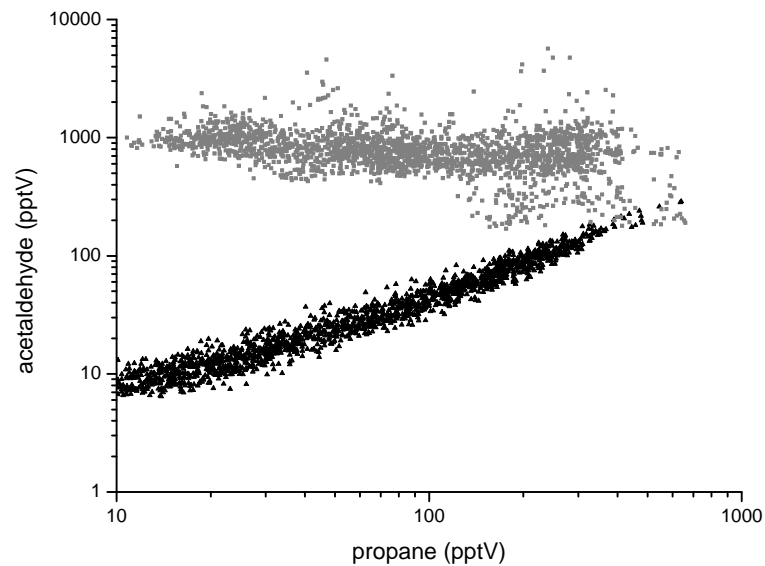
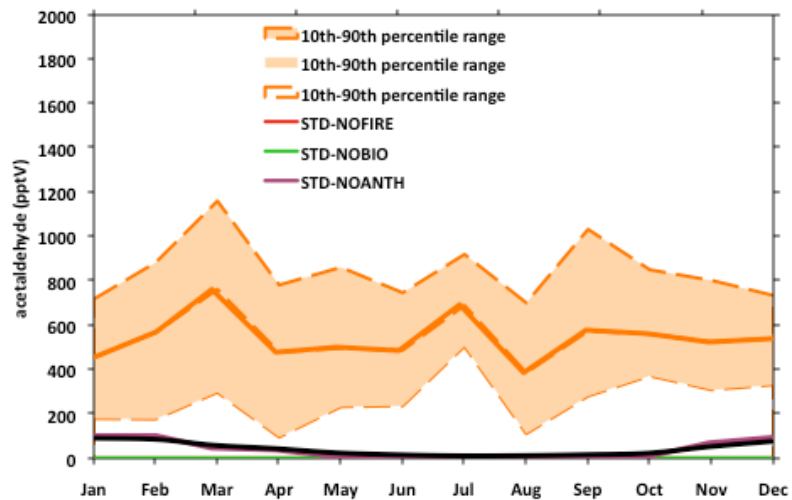
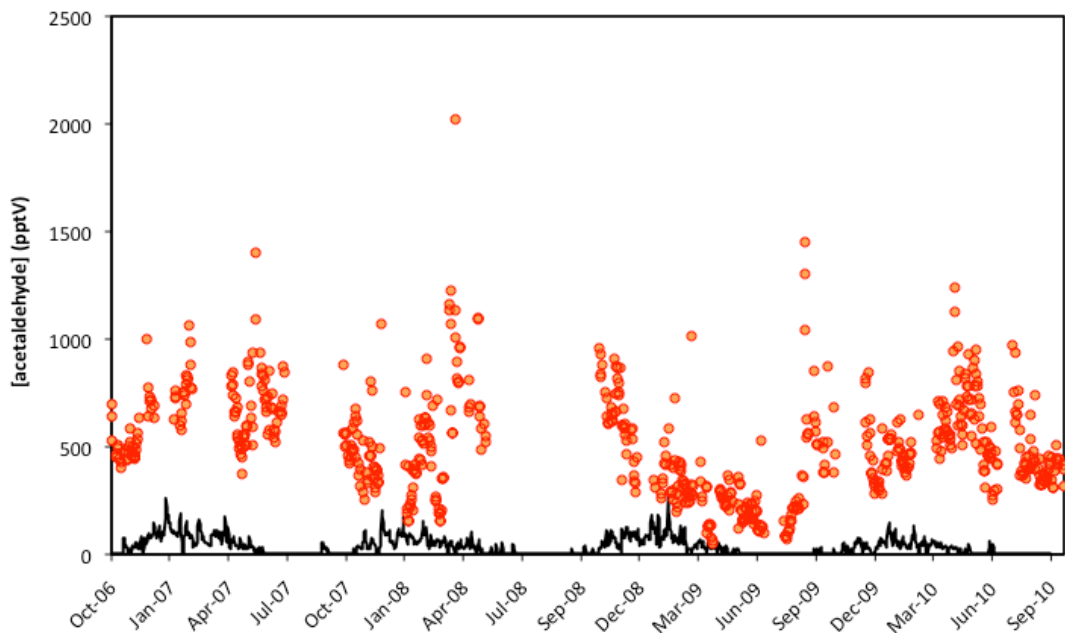
Methanol



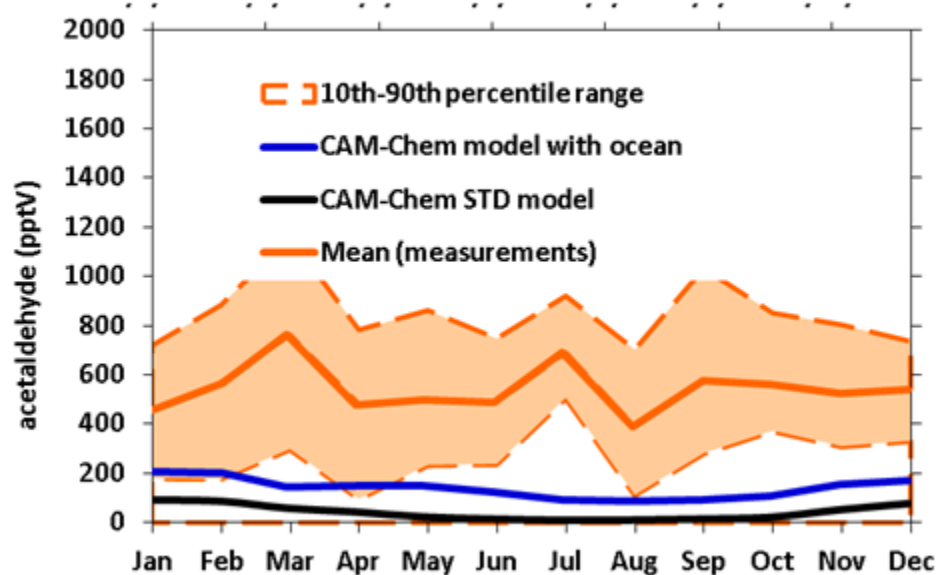
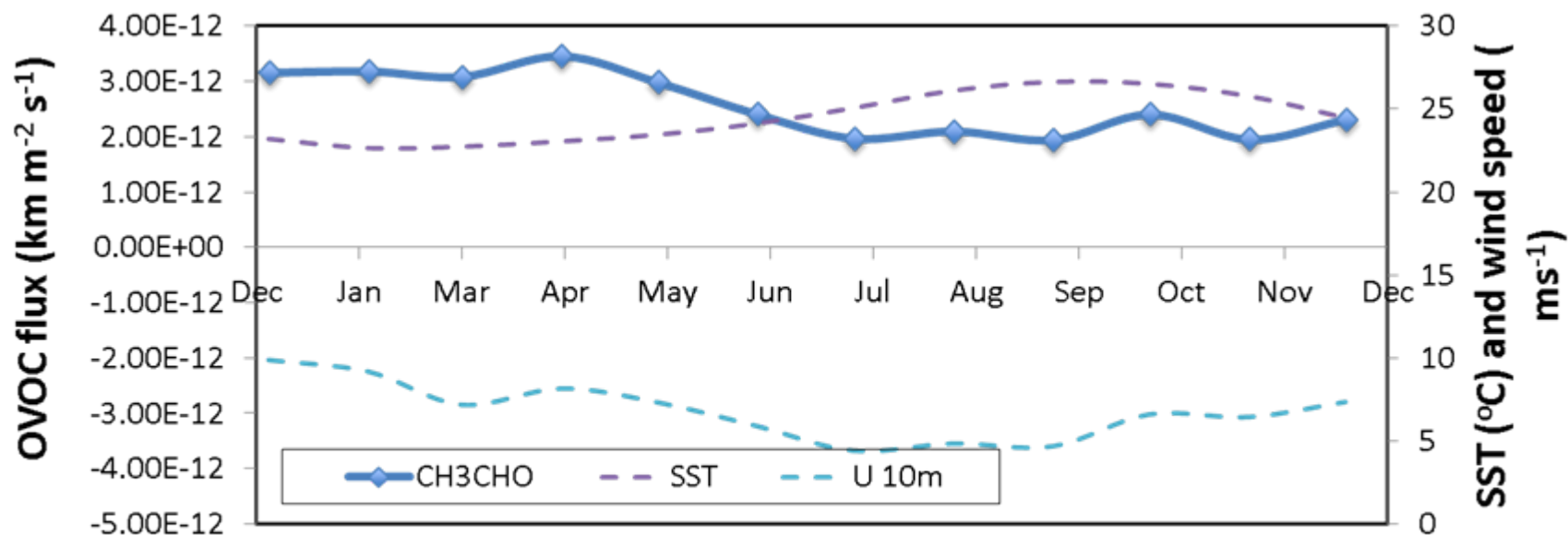
Modification of atmospheric methanol by oceans



Acetaldehyde



Acetaldehyde modification by oceans



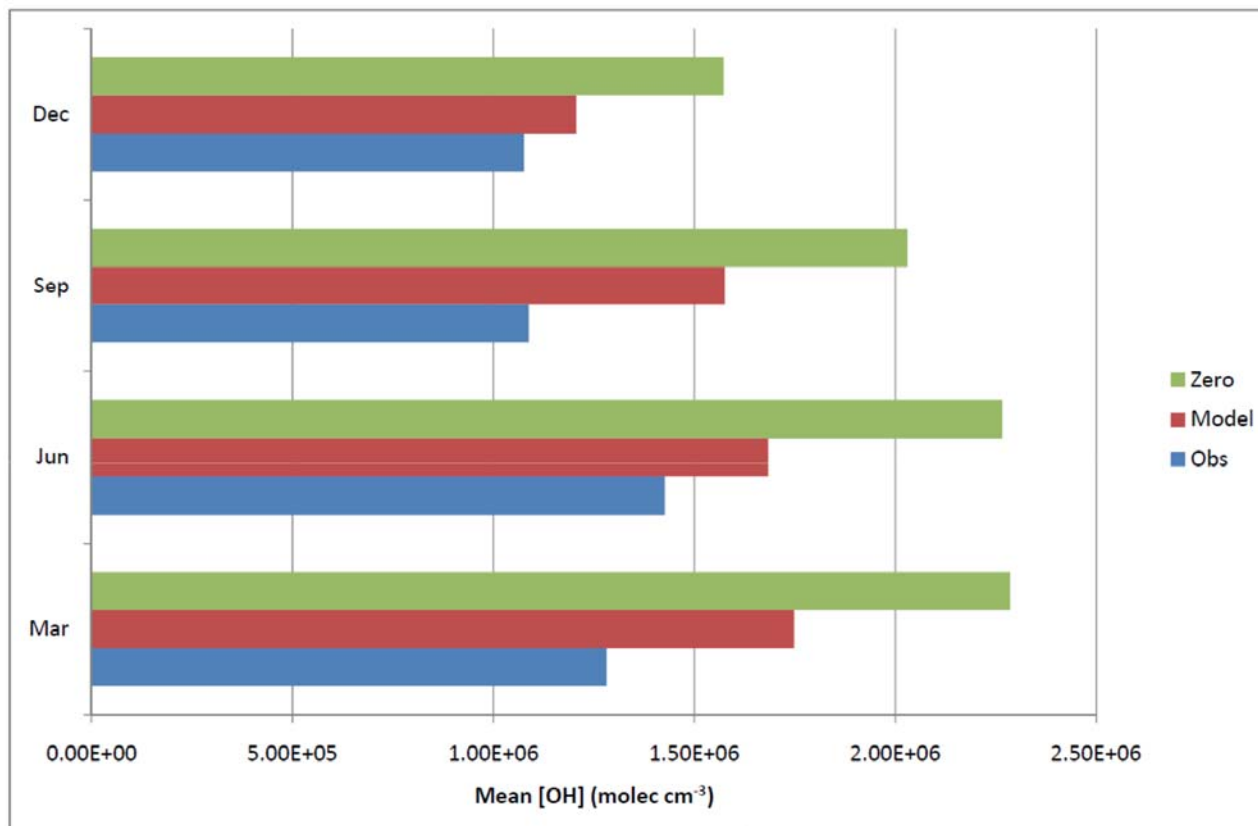
Impact of OVOCs on diurnal mean MBL [OH]

OVOC concentrations from:

(i) observations at Cape Verde

(ii) monthly-mean CAM-Chem model output including ocean fluxes

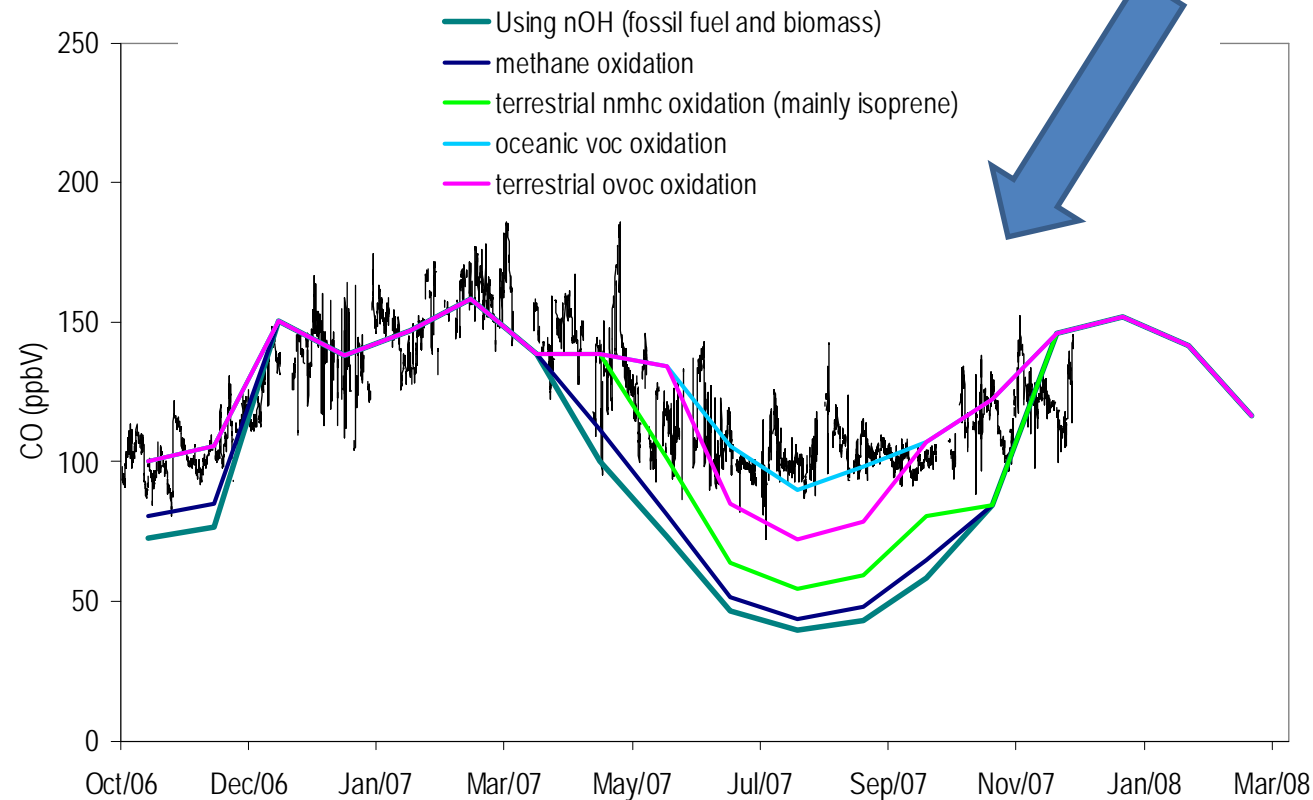
(iii) set to zero



*CittyCat box
model
simulations*

Indirect impact: OVOC as sources of CO

	Additional % contribution from summer CO emission sources not shared by ethane	Reference
Methane oxidation	5 % (May-October)	Granier et al., 2000
NMHC oxidation	12 % (May-October)	Granier et al., 2000
Terrestrial oVOC oxidation	20 % (June-November)	Miller et al., 2008
Oceanic NMVOC oxidation	20 % (July-September)	Guenther et al., 1995



Conclusions

- Oxygenated VOCs are a significant direct sink of OH in the MBL
- Their abundance in the remote marine environment is underestimated (particularly CH_3CHO)
- Marine and biological terrestrial sources of OVOCs could explain some of this model underestimation – more work required to establish emission strength and variability
- $>\text{C}_3$ alkanes and alkenes – chemistry and emissions

Acknowledgements

Funding:



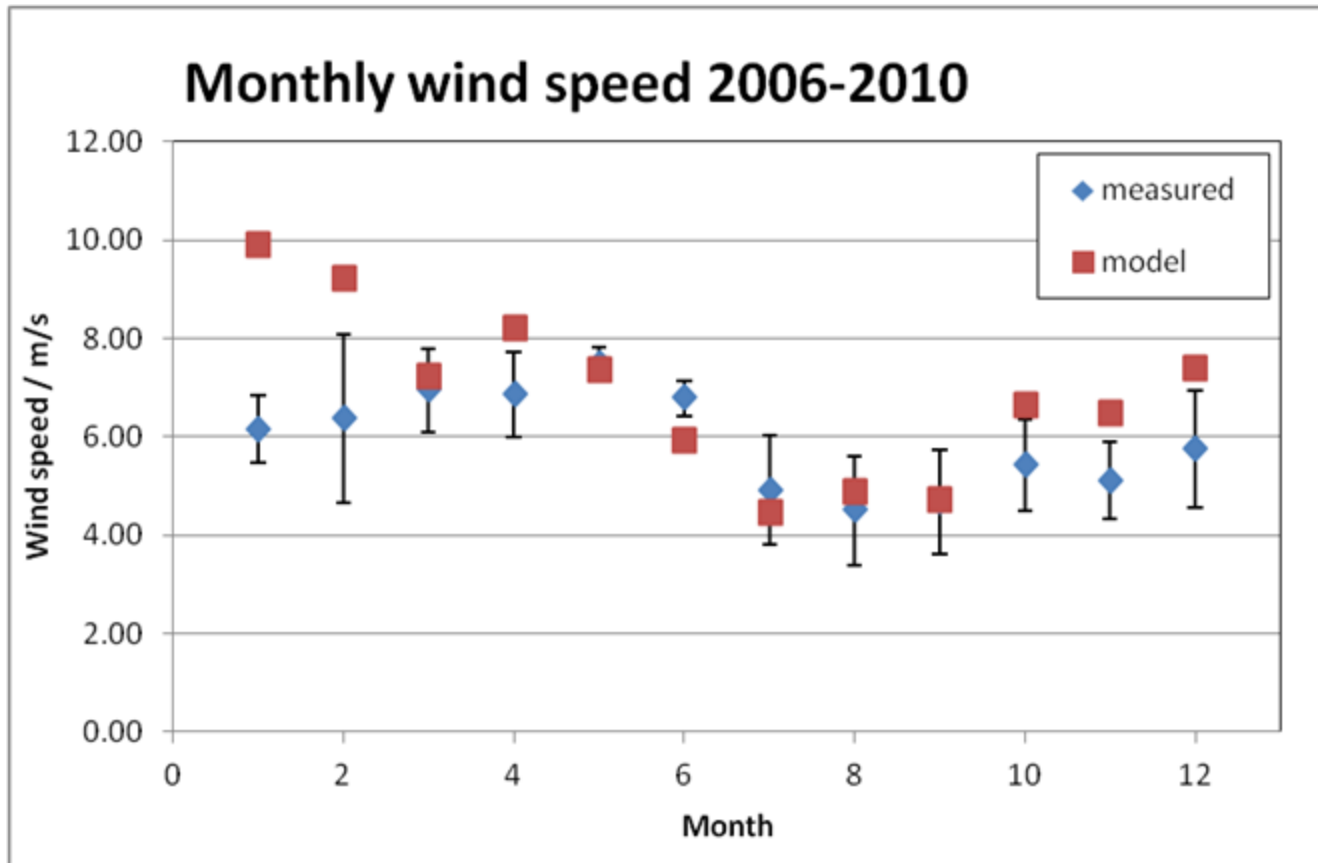
Technical support at CVAO:

Luis Mendes Neves

Helder Lopez



Modelled (GEOS-5) and measured wind speed



- With a squared wind dependence for sea-air fluxes, the difference between 10 m s^{-1} and 6 m s^{-1} is a factor ~ 3 .