

# Observational understanding of aerosols and climate

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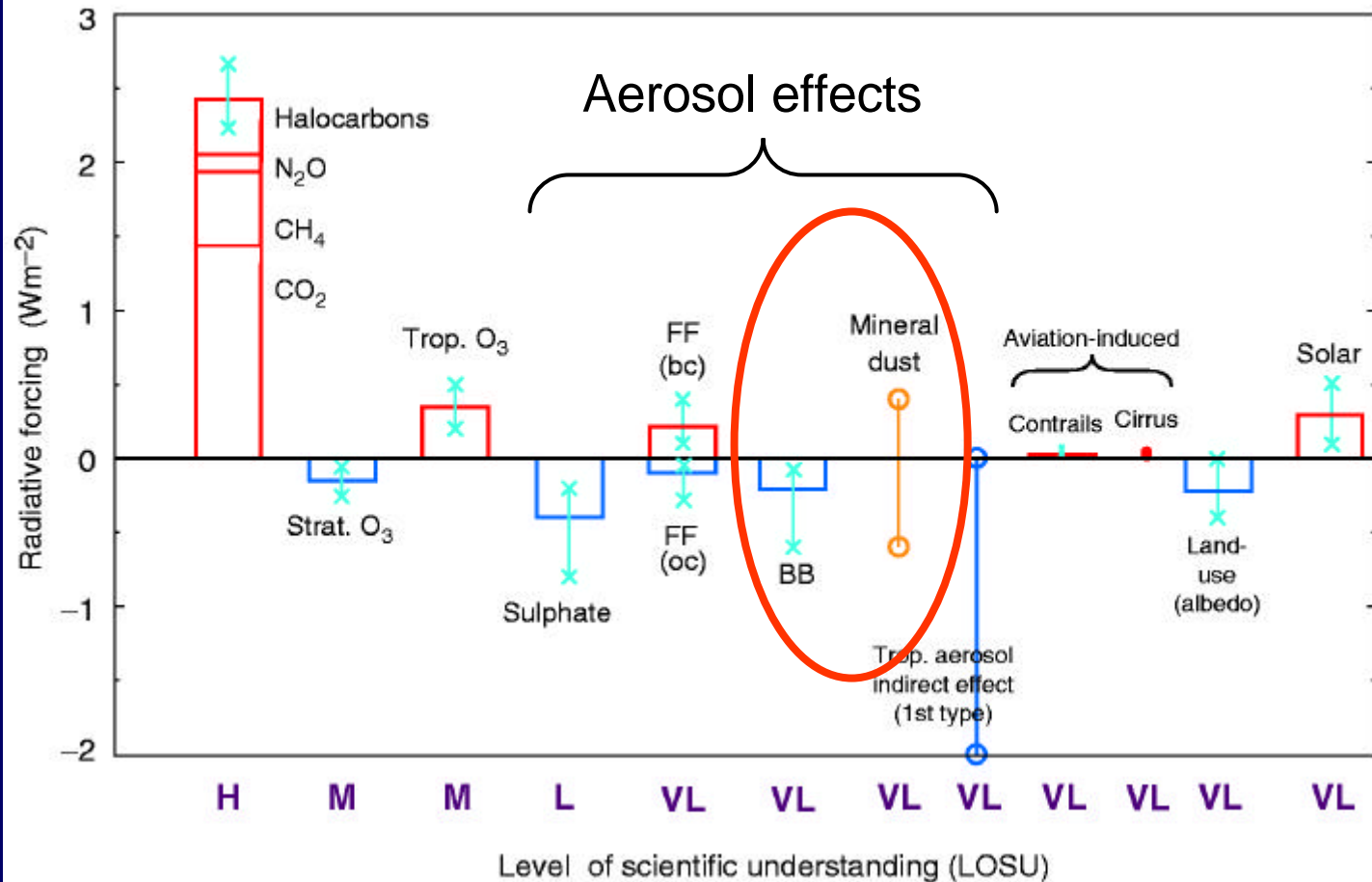
Simon Osborne, Pete Francis, Andreas Keil, Steve Abel, Ian Culverwell, Sean Milton, Andy Jones, Nicolas Bellouin, Olivier Boucher (Met Office/Hadley Centre)  
Eleanor Highwood, Rich Allan, Tony Slingo (Reading University)  
Norm Loeb, Oleg Dubovik, Brent Holben (NASA).

***ECMWF Global Earth-System Monitoring, 5<sup>th</sup> -9<sup>th</sup>***  
***September 2005***

- **Introduction: why are aerosols important in radiation budget & climate?**
- **Biomass burning aerosols**
  - **Why is the vertical profile of aerosol important?**
    - The effect of the vertical profile on the aerosol radiative forcing.
    - The effect of the vertical profile on the derivation of cloud properties.
  - **How do the biomass burning aerosol optical properties change as particles age?**
  - **Can we believe the size distributions etc from sun-photometer surface based retrievals?**
- **Saharan dust aerosols**
  - The direct solar radiative effect over ocean
  - The direct terrestrial radiative effect over ocean
  - Implications for SST retrievals
  - The direct net radiative effect over land
- **Direct forcing due to all aerosol types from observations**

Radiative forcing, DF, relates to the global mean temperature change,  $\Delta T$  via the climate sensitivity parameter  $\lambda$  :-  $\Delta T = \lambda \text{ DF}$

Global and annual mean radiative forcing (1750 to present)



# Direct effect of tropospheric aerosols (clear skies for simplicity)

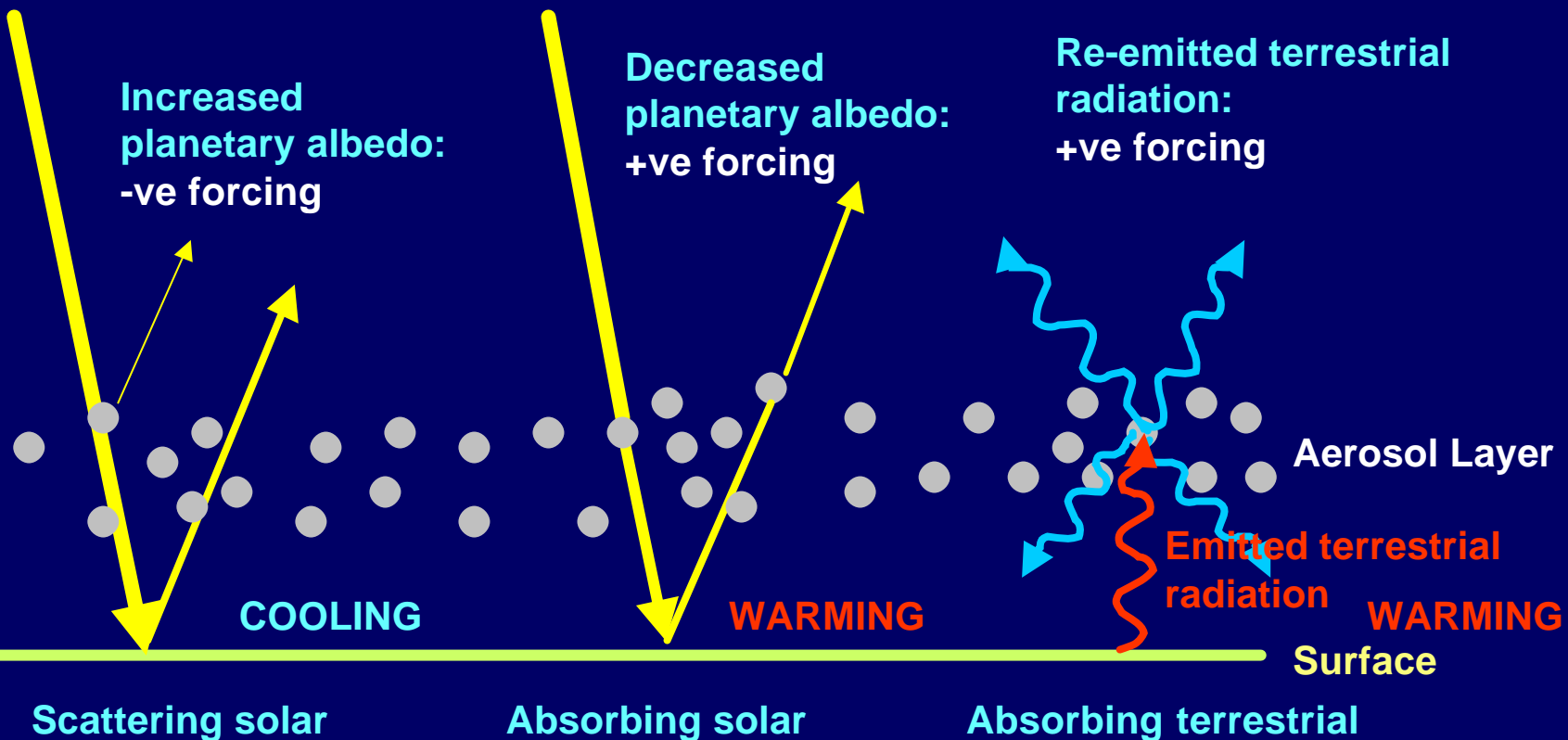
Incident solar radiation

Incident solar radiation

Increased planetary albedo:  
-ve forcing

Decreased planetary albedo:  
+ve forcing

Re-emitted terrestrial radiation:  
+ve forcing



Simple expression for the direct solar radiative effect of a scattering/absorbing tropospheric aerosol in clear skies where  $R_s$  is the surface reflectance and  $A_c$  is the cloud amount (Haywood and Shine, 1995) :-

$$\Delta F = -\frac{1}{2} S_o T_{at}^2 (1 - A_c) \left[ w_o b (1 - R_s)^2 - 2(1 - w_o) R_s \right] \tau$$

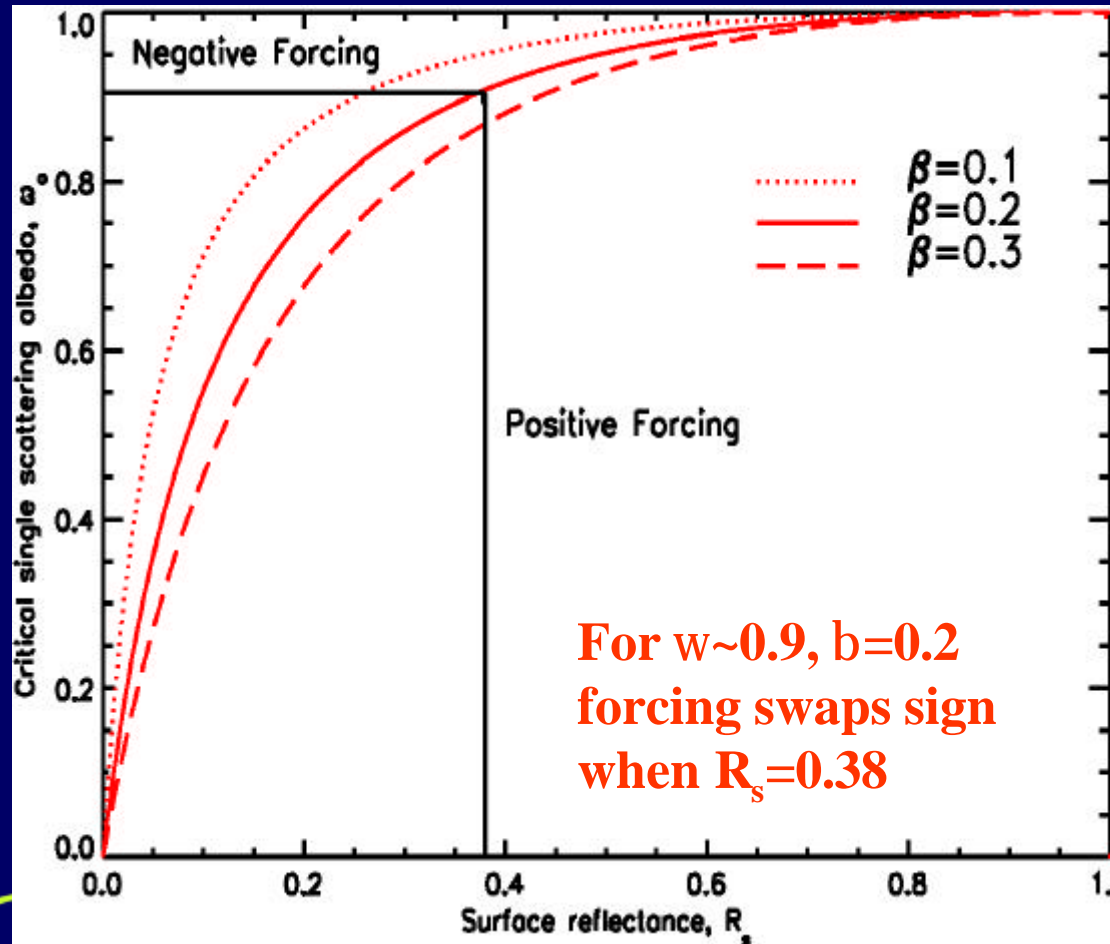
$\omega_o$ : single scattering albedo =  $Q_{sca} / (Q_{sca} + Q_{abs})$

$\beta$ : fraction backscattered to space

$\tau$  is the optical depth

$$w_o > \frac{2R_s}{b(1 - R_s)^2 + 2R_s}$$

for negative forcing



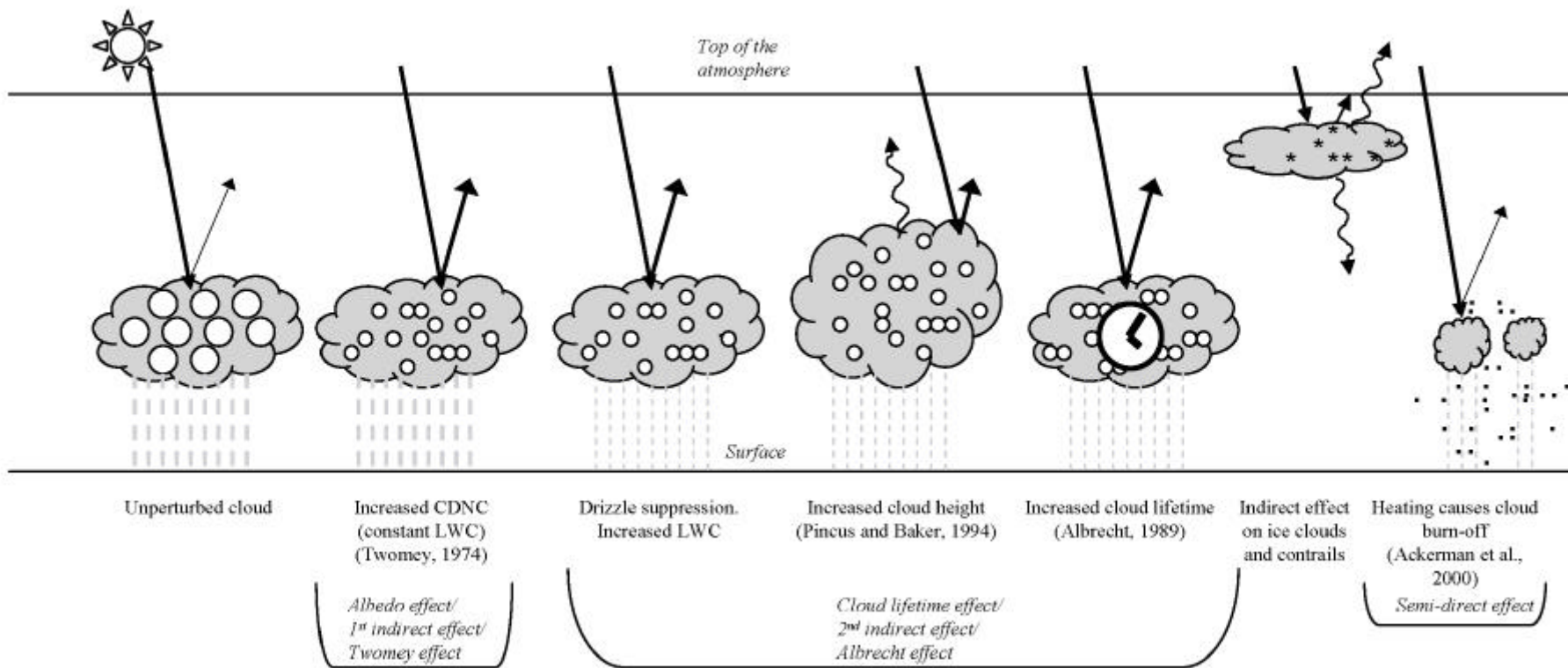
The expression is really just a complicated expression to show this:-

Dark ocean surface

Bright snow/cloud surface

<p>Dark surface appears brighter -&gt; increased planetary albedo</p>	<p>Bright surface appears darker -&gt; decreased planetary albedo</p>

# Schematic of the indirect effects - (not dealt with in detail in this talk)



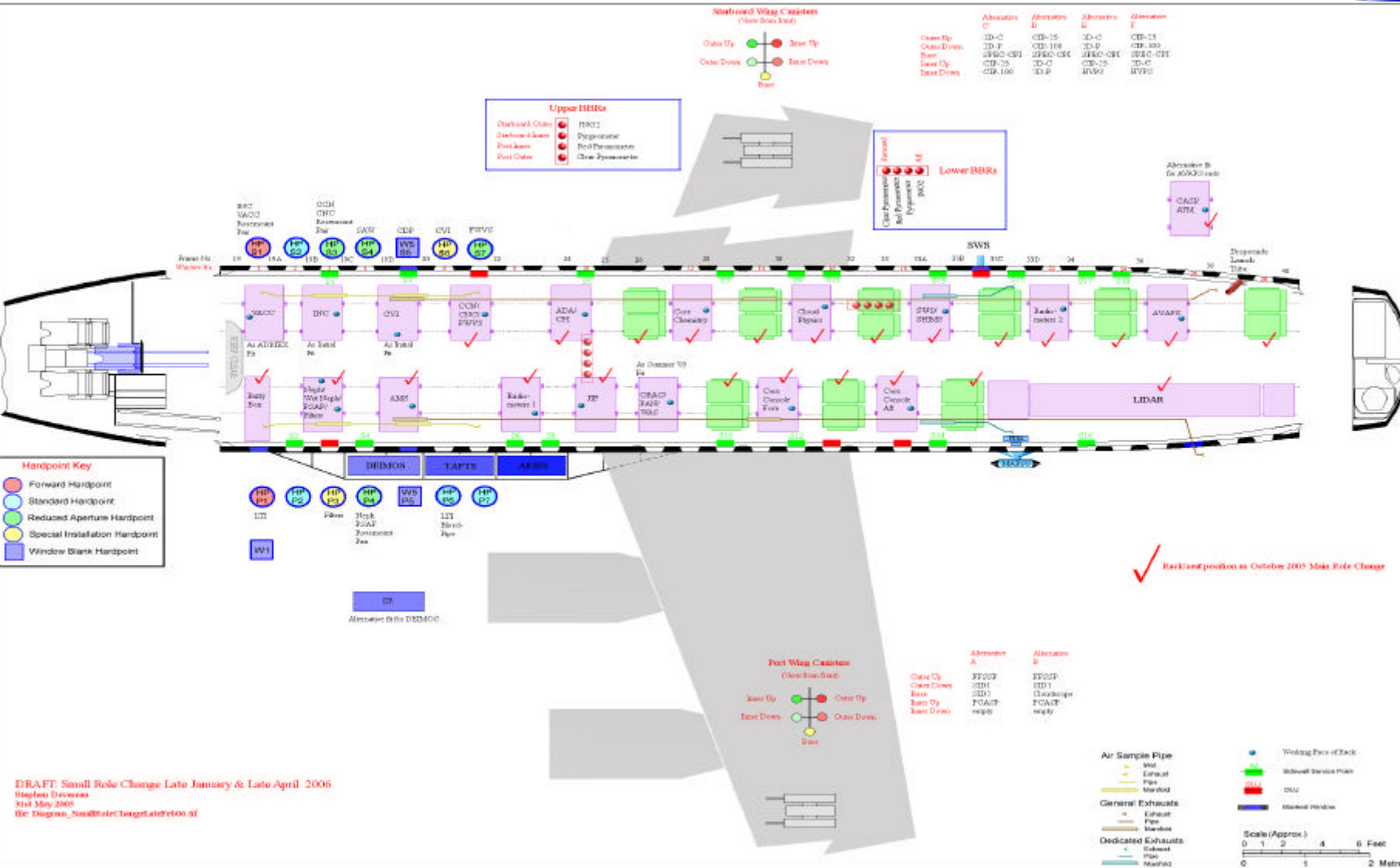
Adapted from Haywood and Boucher, 2000.

# The Met Office/NERC/FAAM aircraft



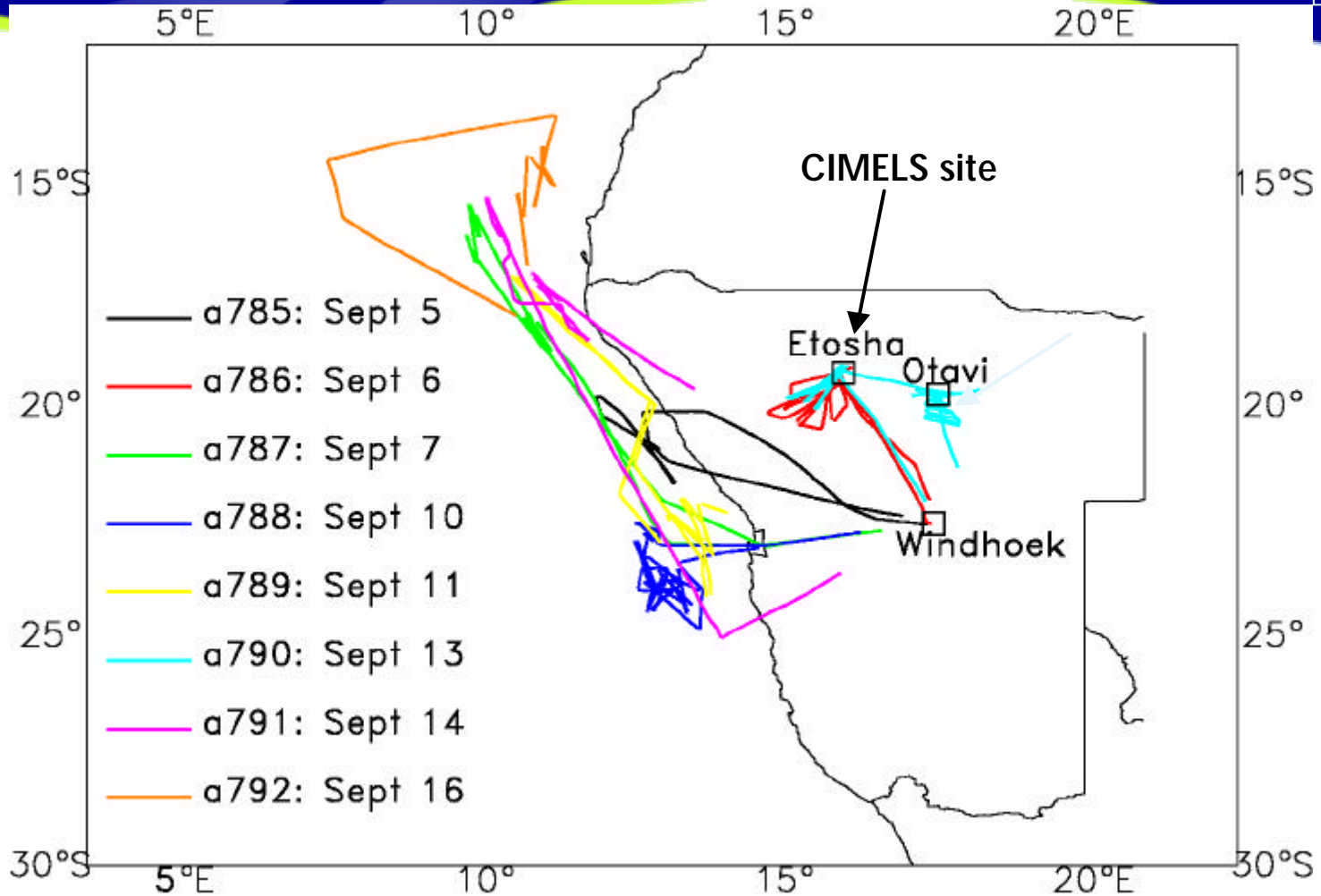


# The layout of the BAe146 (interchangeable)



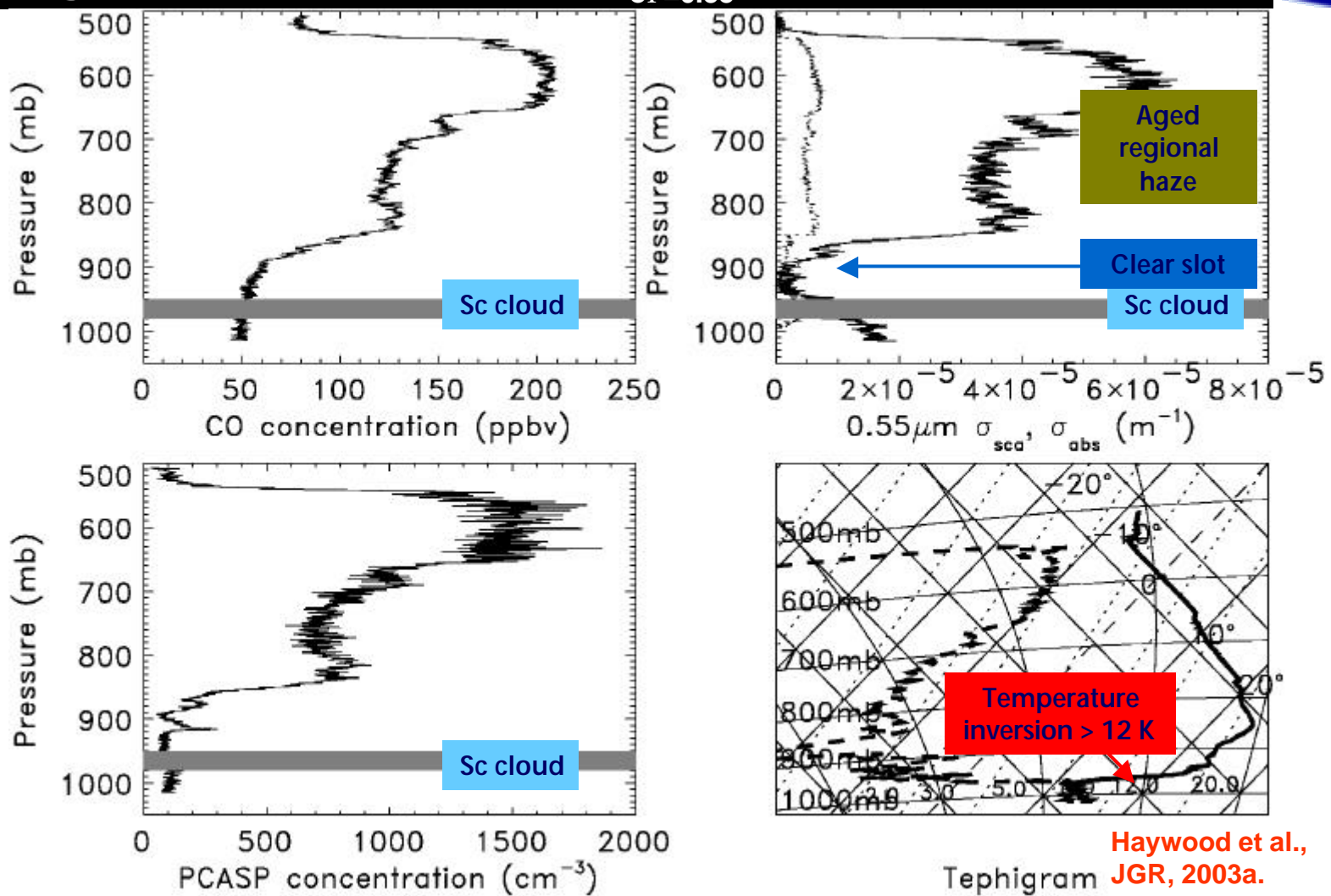
DRAFT: Small Role Change Late January & Late April 2006  
 Stephen Drevon  
 31st May 2005  
 (cc: Diagram\_SmallRoleChange@metoffice.gov.uk)

## 2. Biomass Burning Aerosols



**Figure 1: Map showing the geographical location of the flights performed by the C-130 during SAFARI 2000. The approximate positions of Windhoek, Etosha, and Otavi are marked. The geographical outline of Namibia is also shown.**

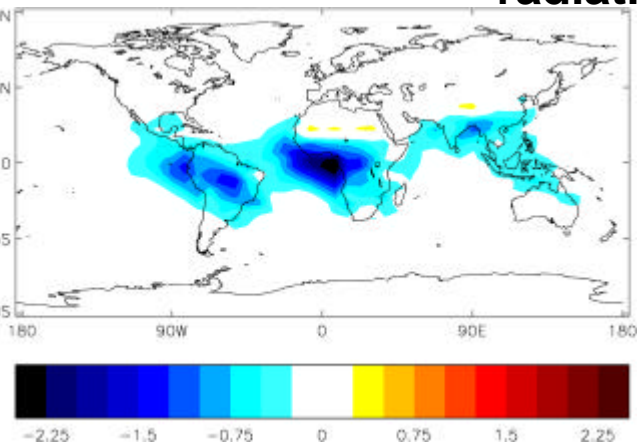
Over ocean, aerosol separated from Sc by 'clear' slot and strong temperature inversion.  $w_{ol=0.55} = 0.91$  for the aerosol.



Haywood et al., JGR, 2003a.

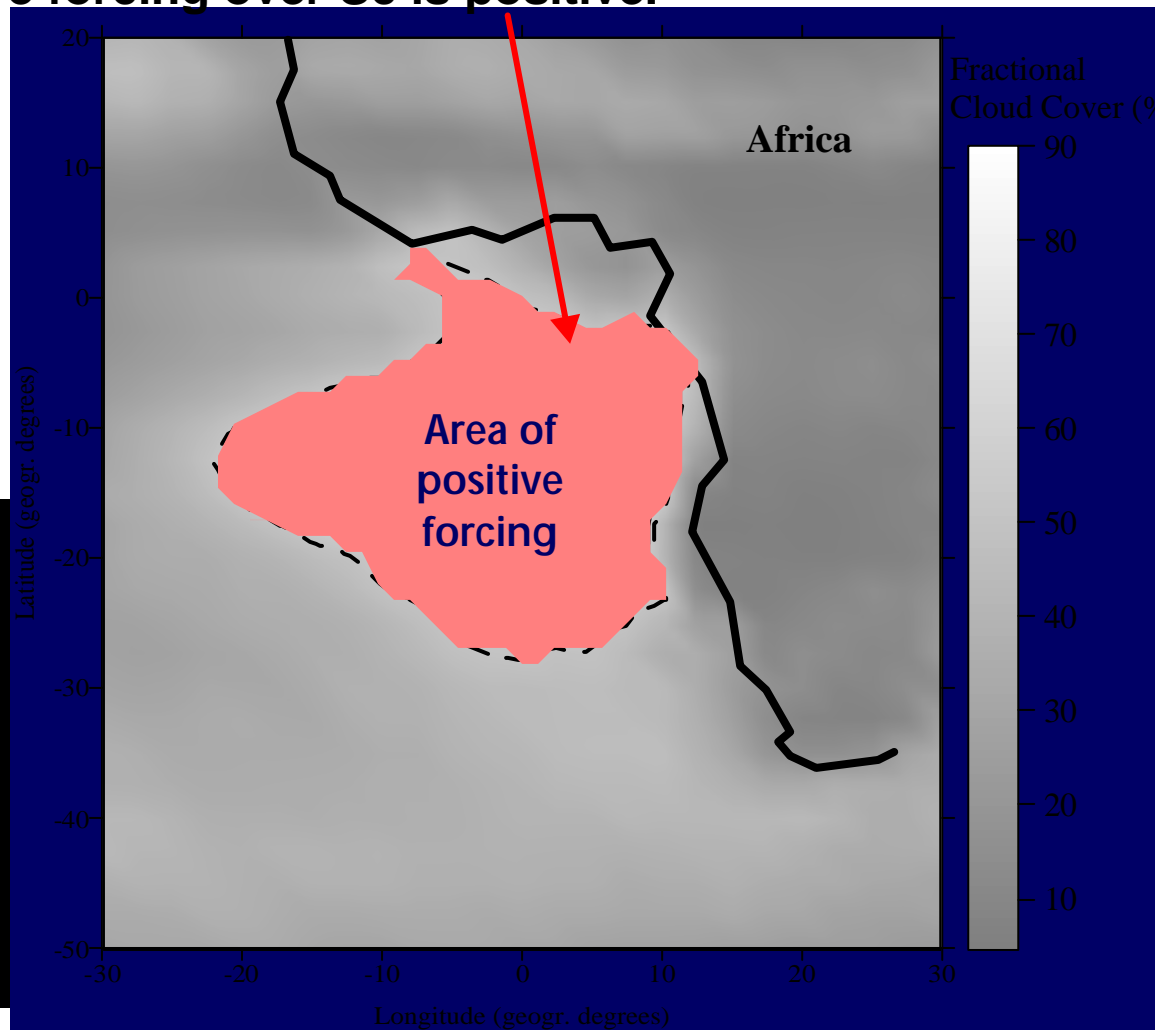
**Note the correlation between CO,  $S_{sca}$ ,  $S_{abs}$ , and PCASP conc**

Realistic vertical profiles of aged regional haze, and realistic vertical profiles of Sc (both based on aircraft measurements). Using ISCCP cloud fractions, the radiative forcing over Sc is positive.



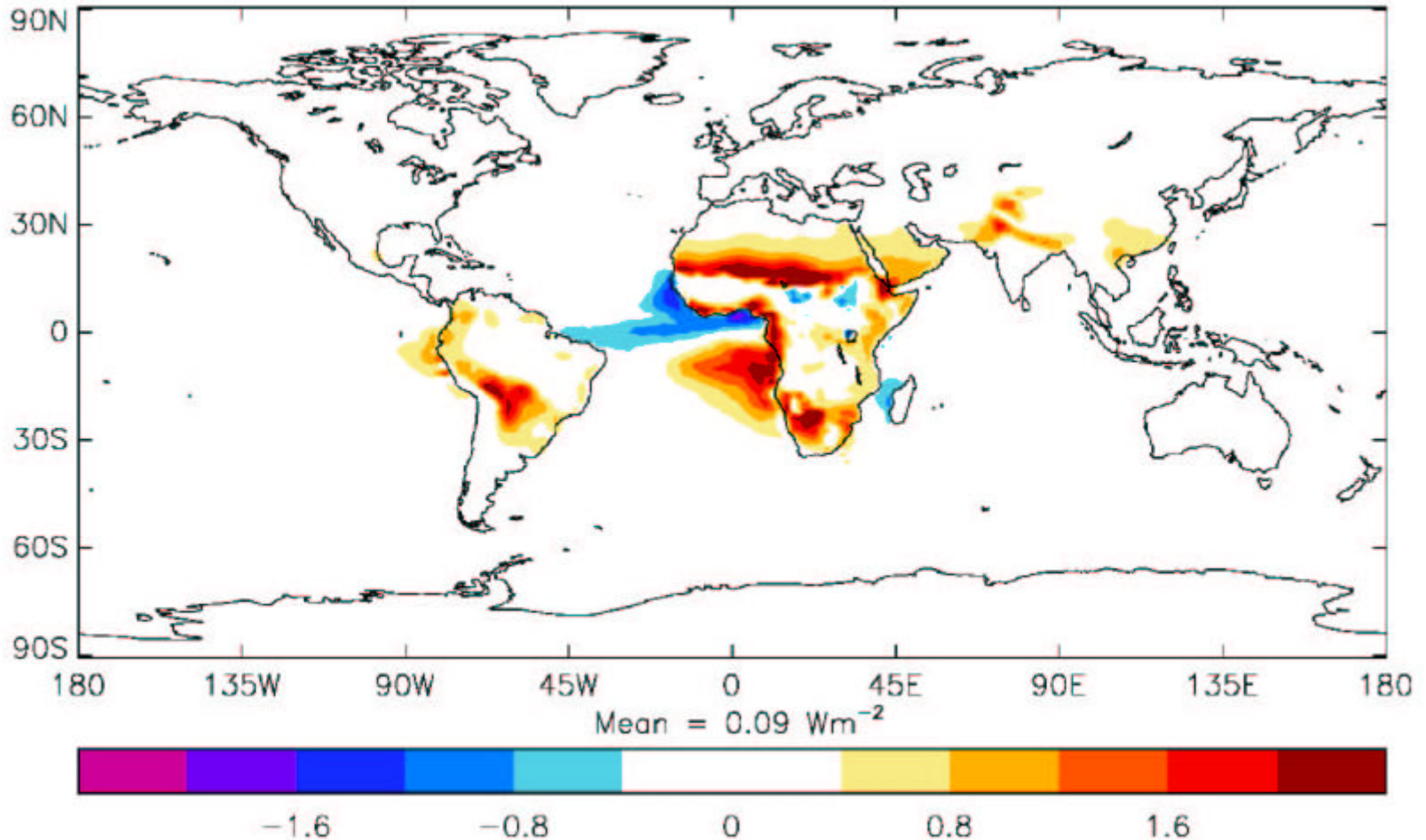
Implications - models have to get the vertical profile and cloud properties right for partially absorbing aerosol. This wasn't a concern for sulphate aerosol.

Is the sign of the global BB radiative forcing positive?!!!



# Results from the UM (Andy Jones).

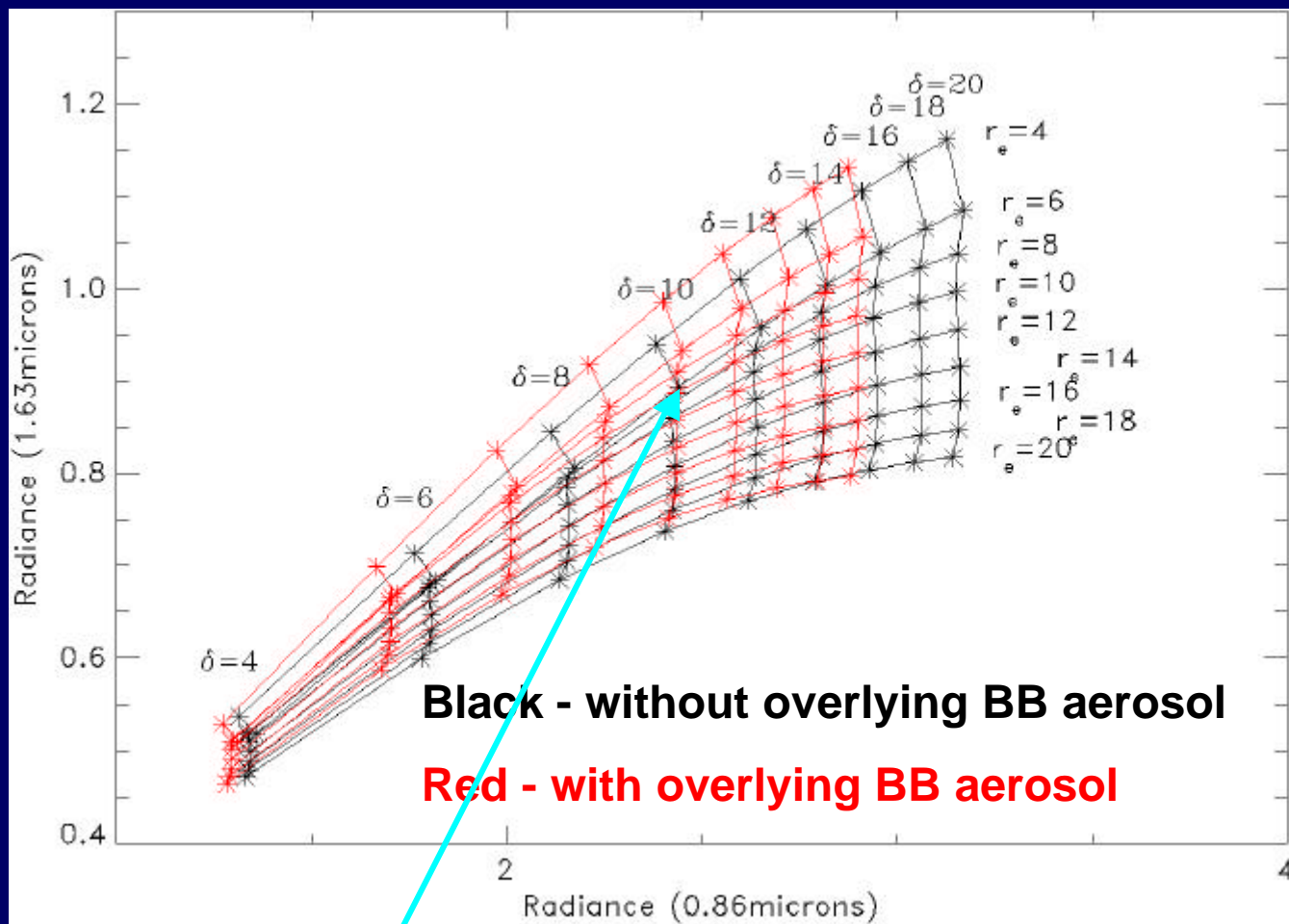
.....suggest that the TOA radiative forcing from BB aerosol may indeed be positive (but very sensitive to aerosol absorption properties). Similar results come from the AEROCOM initiative .....



# Does overlying aerosol significantly effect cloud retrievals?

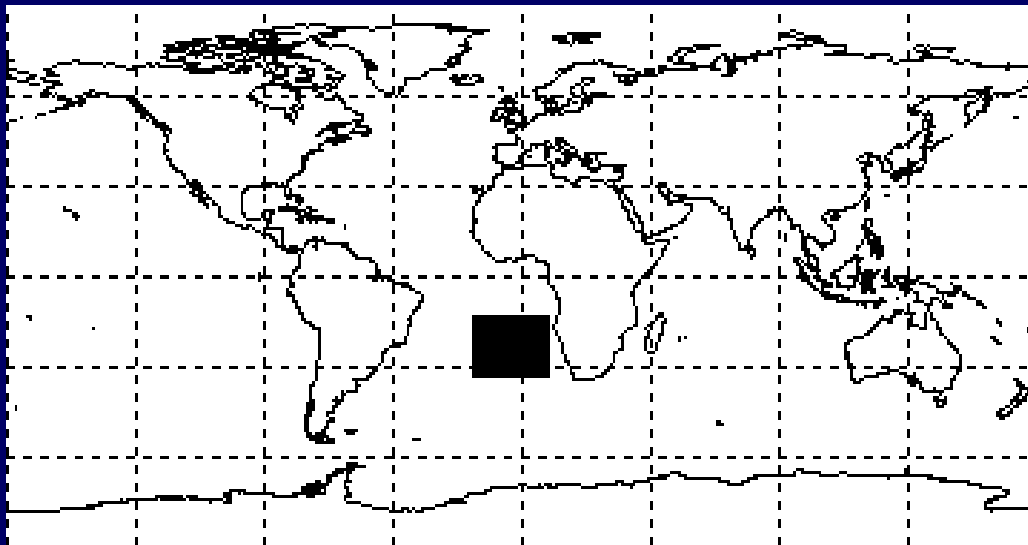
**MODIS 1.63mm:**  $d_{\text{cloud}}$  is underestimated –  $d_{\text{cloud}}^{\text{apparent}}=10$   $d_{\text{cloud}}^{\text{real}}=12$ .

$r_{\text{eff}}$  is underestimated:  $r_{\text{eff}}^{\text{apparent}}=6\text{mm}$ :  $r_{\text{eff}}^{\text{real}}=10\text{mm}$

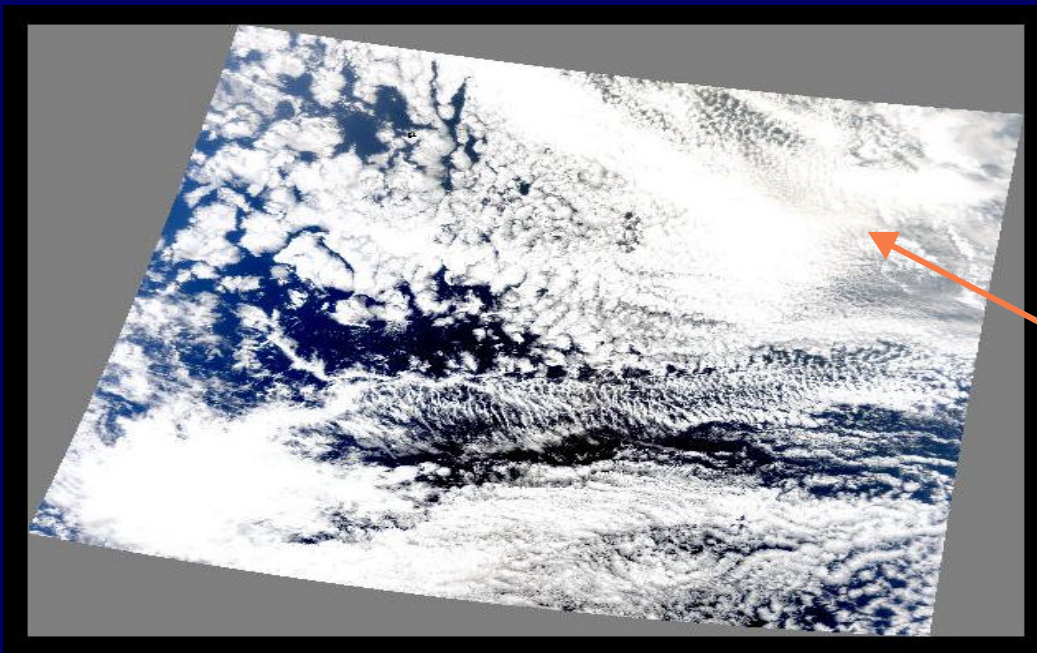


‘Apparent indirect effect’

Haywood et al., 2004, QJRMS

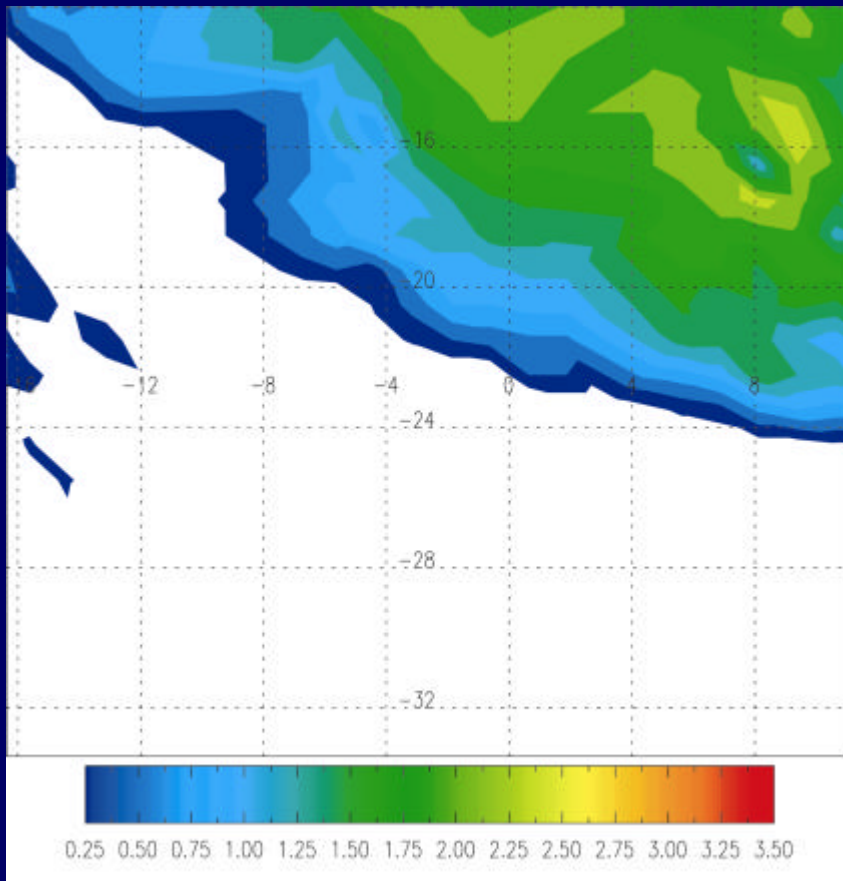


MODIS quick-look image off the coast of Namibia/Angola for 7<sup>th</sup> September 2000

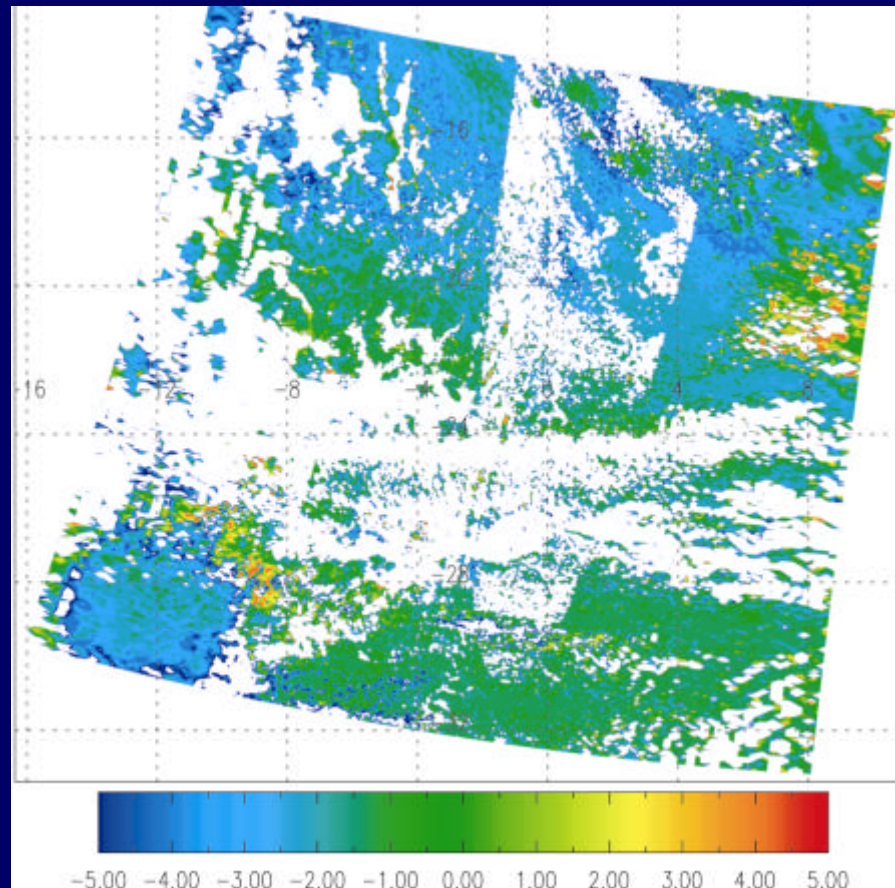


Smoke in N.E. corner

TOMS AI shows absorbing aerosol in N.E.



$r_{e1.63}$  shows negative bias in N.E.



Green colors  $r_{e1.63} = r_{e2.13}$  Blue colors  $r_{e1.63} < r_{e2.13}$

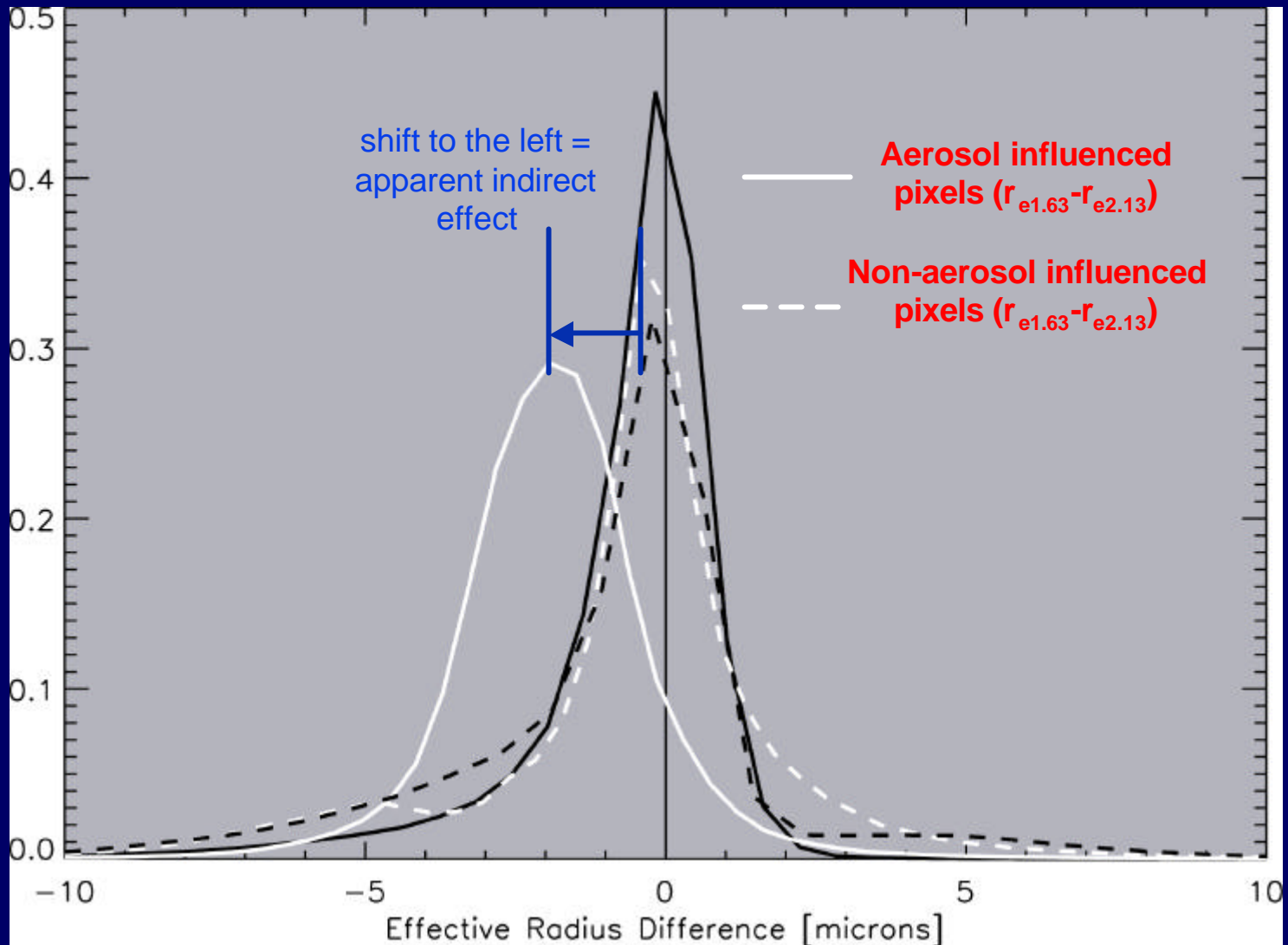
Implications –

a) ‘apparent indirect effect’

b)  $w_o$  might be derivable



# PDFs show the nature of the difference between aerosol and non-aerosol influenced pixels

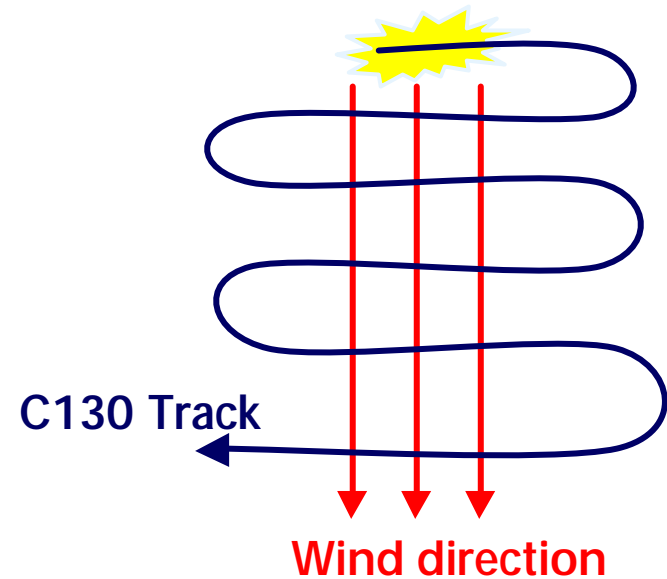
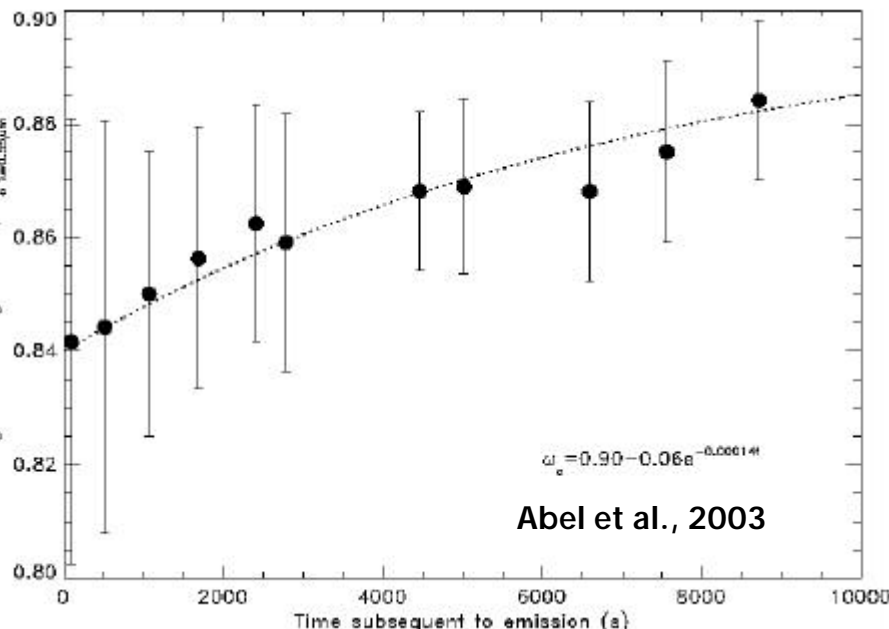


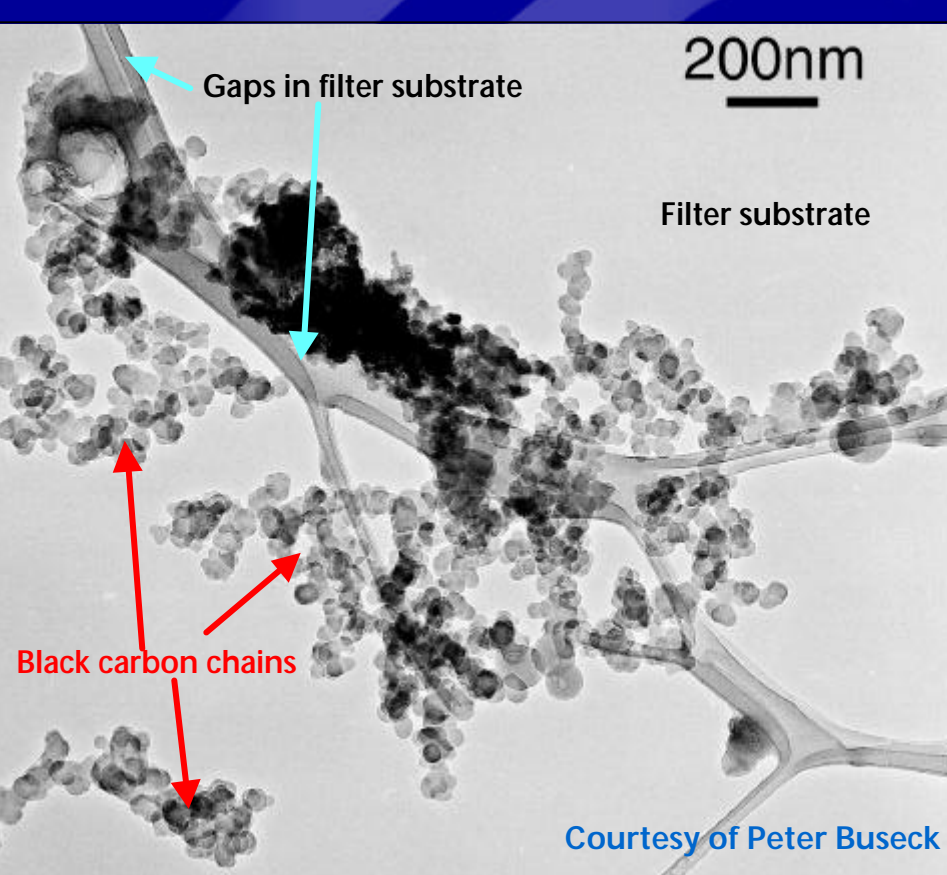


Burn scar > 5km<sup>2</sup>

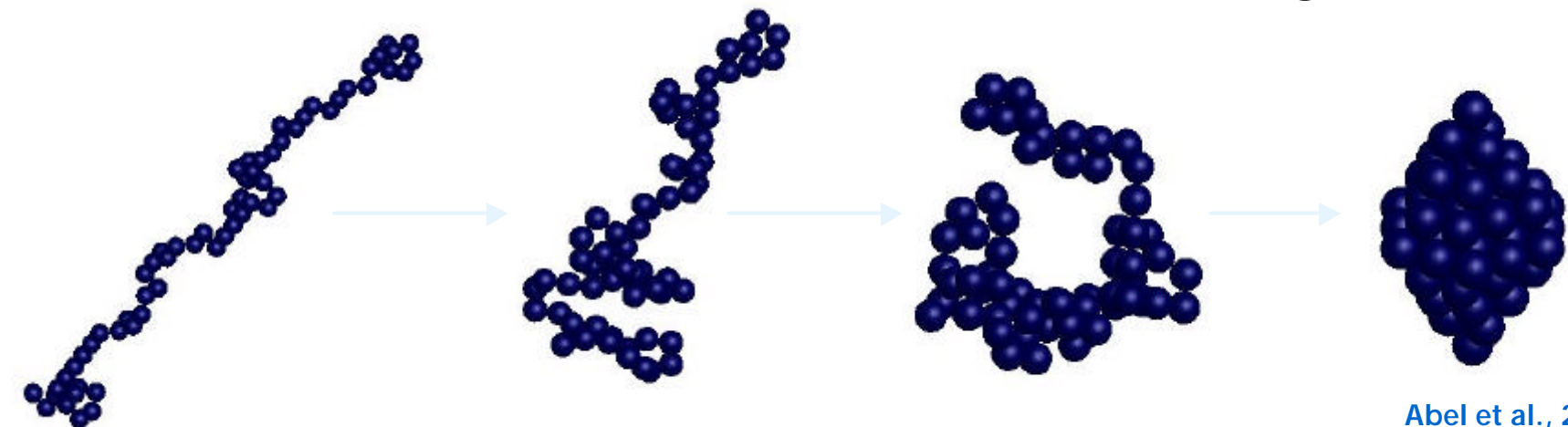
Plume easily detected 100km downwind

A raster pattern was flown downwind to determine how the single scattering albedo of the biomass burning aerosols changes:-

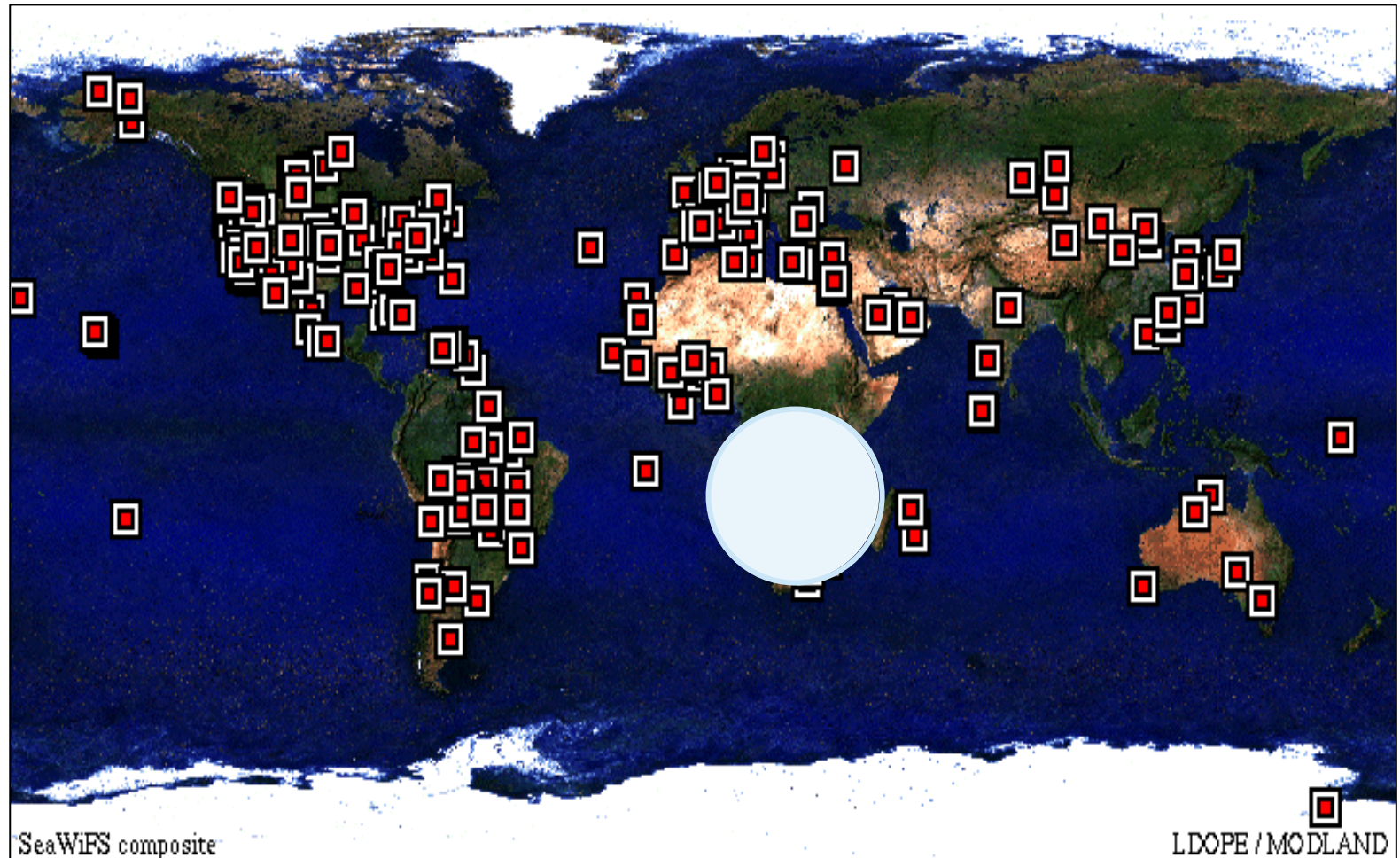




We've investigated whether the collapse of the black carbon chain structure is responsible for the change in  $w_0$ : not sufficient to explain the differences -> more likely to be the condensation of VOC gases



## Retrievals of $\tau_{\text{aer}1}$ , size distribution, absorption



**But do we believe them?**

# Flight made during SAFARI-2000 in biomass aerosol

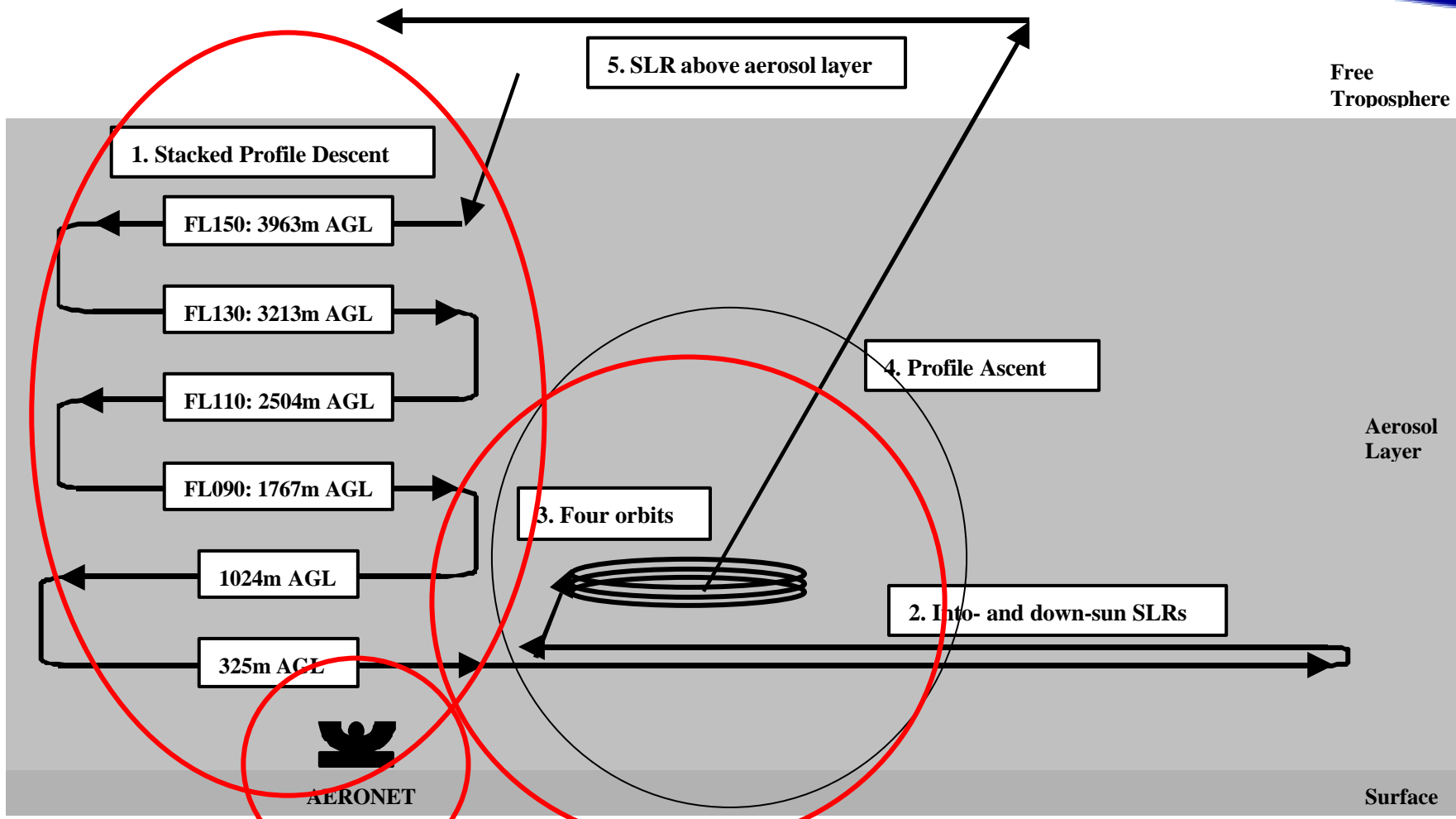
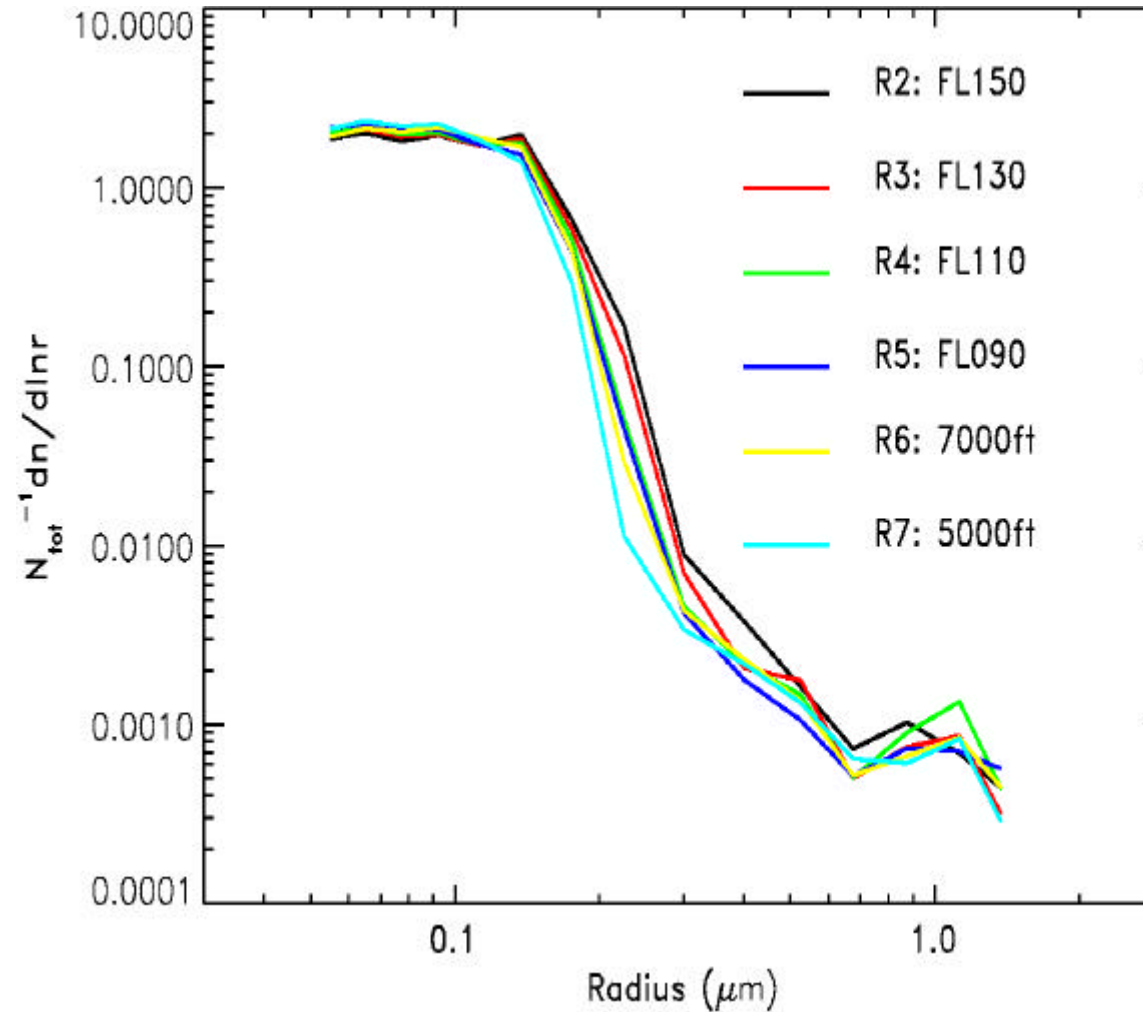
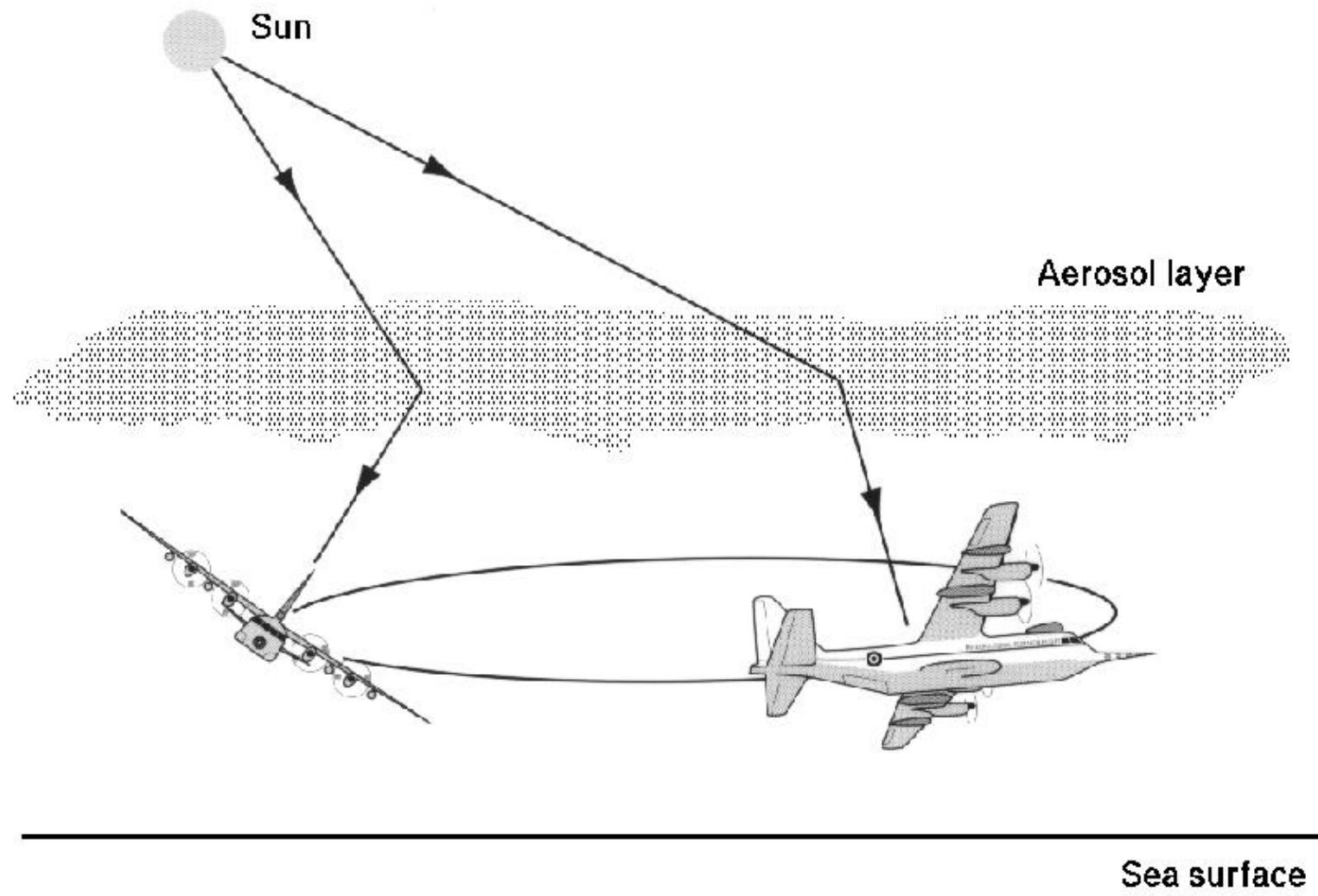


Figure 2: Schematic diagram of the flight pattern performed by the C-130 over the Etosha AERONET site on September 13, 2000. Consisting of 1) stacked profile descent, 2) into- and down- sun SLRs, 3) a series of four orbits, 4) profile ascent, 5) SLR above the aerosol layer.

**The aerosol size distribution shows little variation in the vertical due to the strong dry convective mixing.**



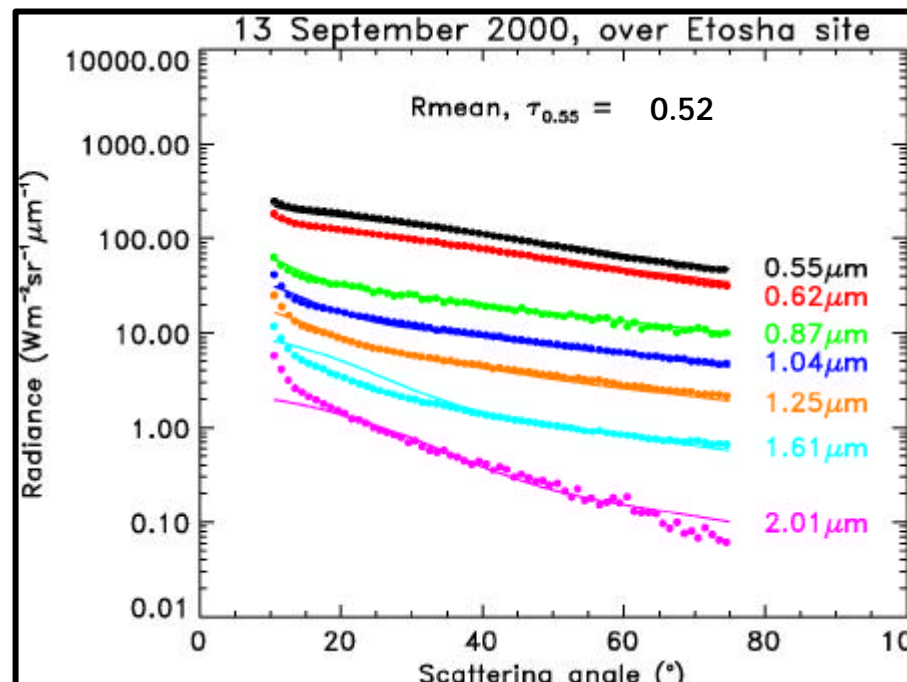
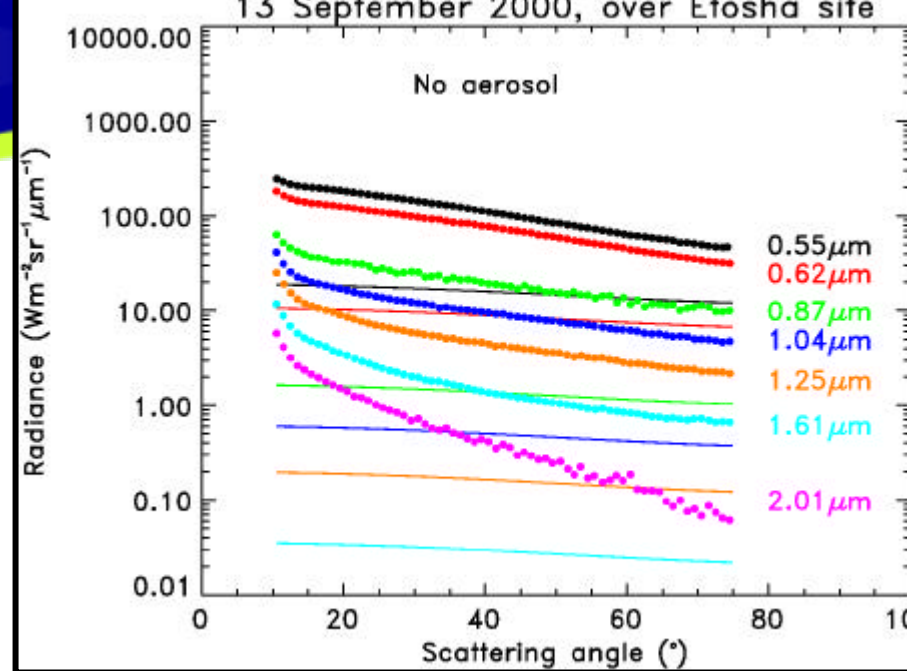
Measurements are analogous to AERONET all-sky radiance measurements by performing orbits to derive the aerosol size distribution.



Radiance as function of scattering angle.

**Circles - measurements**  
**Lines - models**

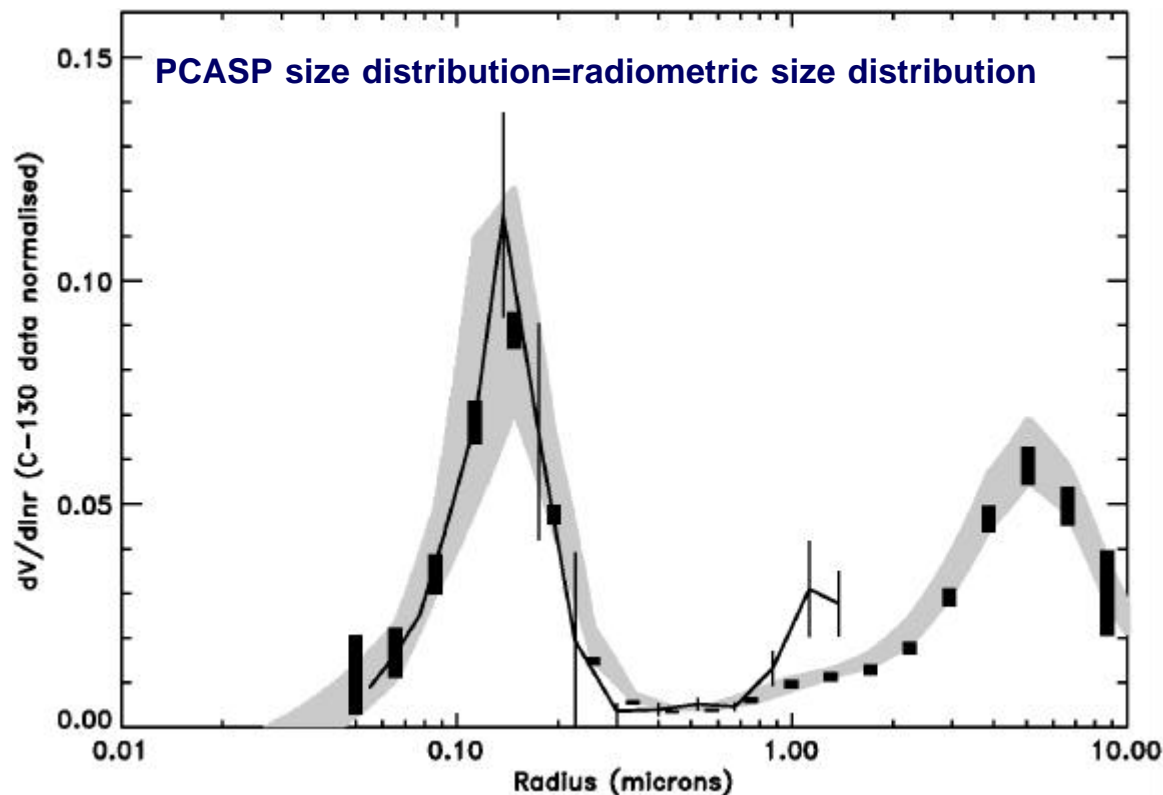
These results suggest that the sky radiances can be modelled most accurately when  $t_{\text{aer}|0.55} = 0.52$





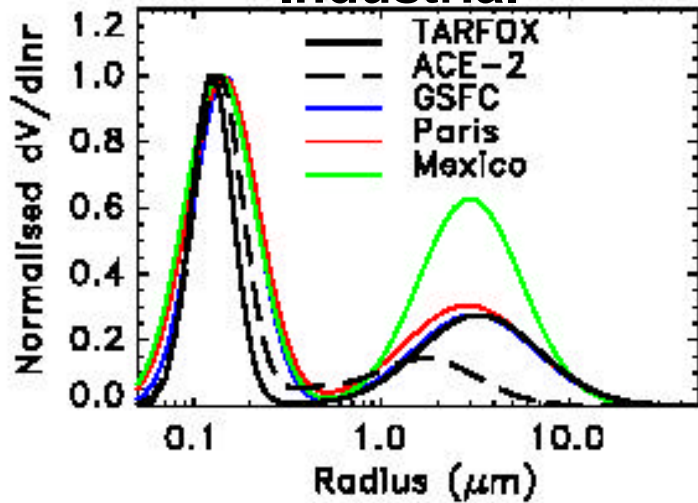
Over the radius range 0.05-1.0 $\mu$ m the size distributions are identical even though are determined completely independently.

We also found excellent agreement in  $w_0$  derived from the in-situ measurements and AERONET radiometers and hence *im*.

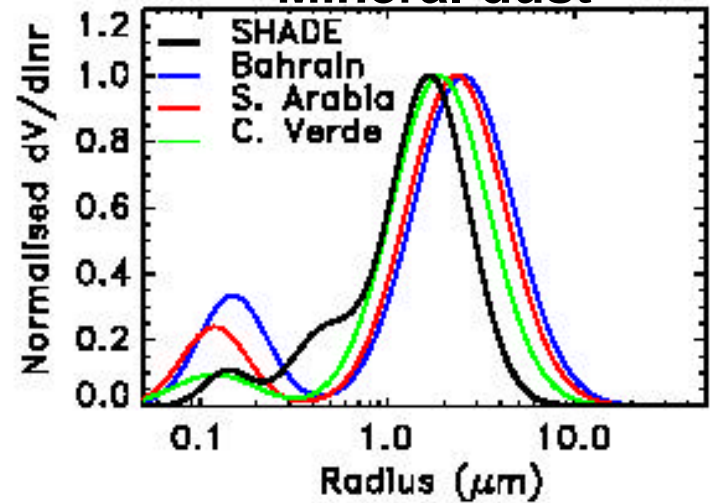


# Size distributions compare well for all aerosol types (Osborne and Haywood, 2005)

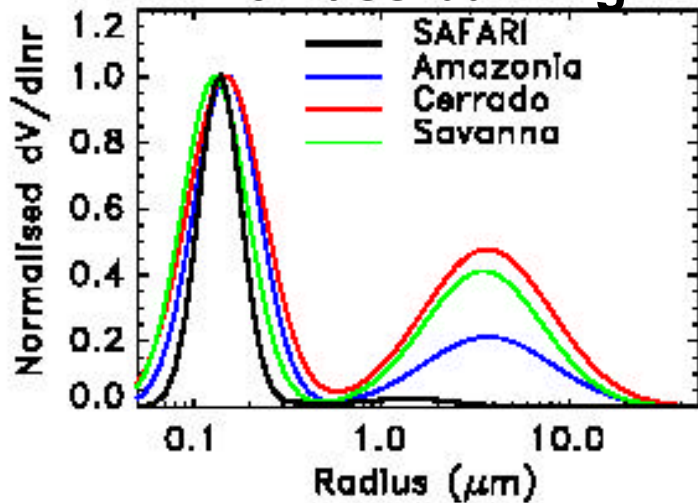
## Industrial



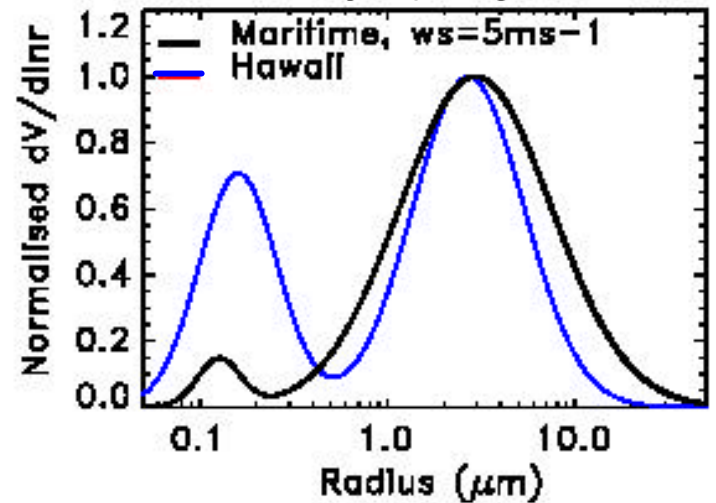
## Mineral dust



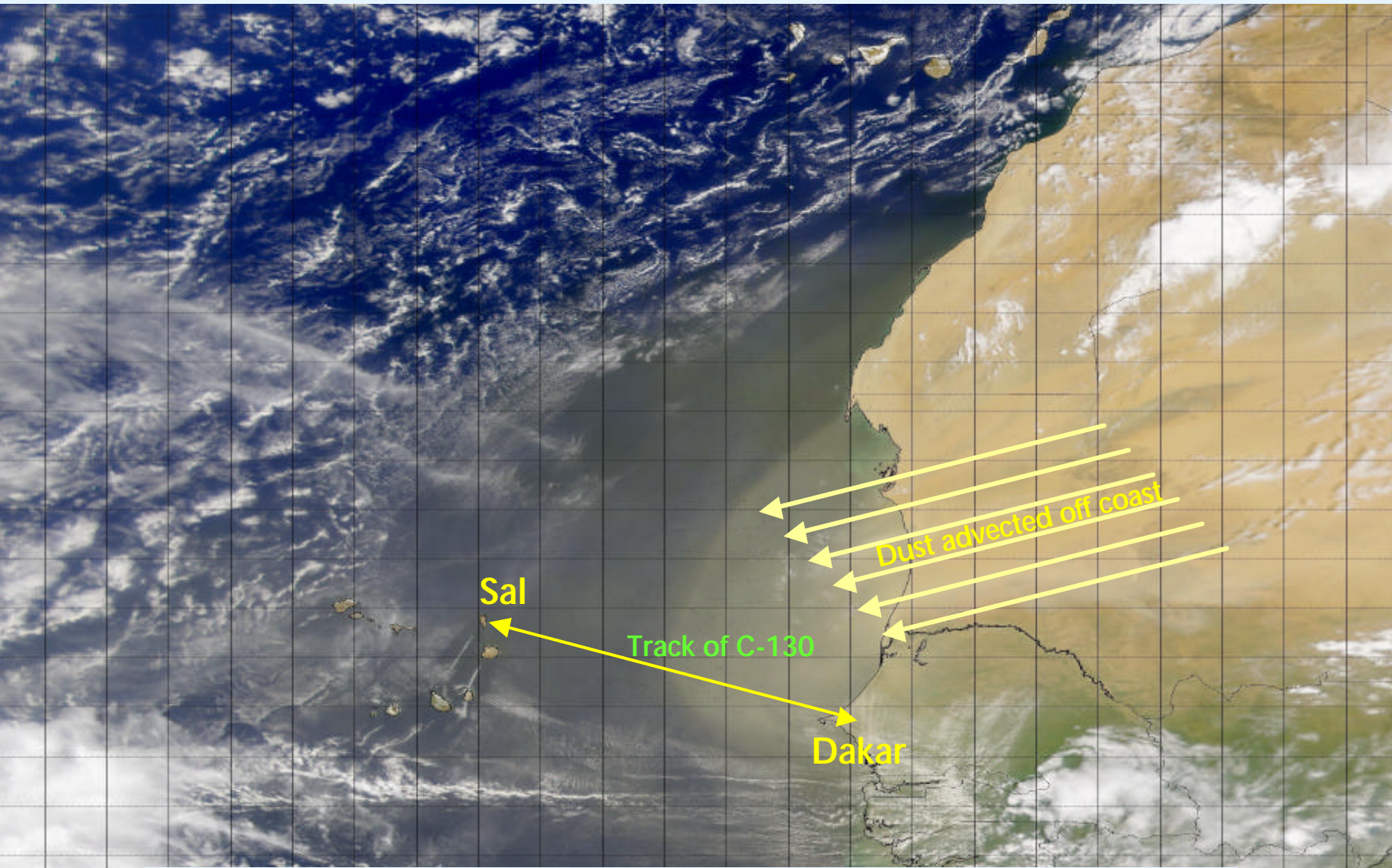
## Biomass burning



## Maritime



# C-130 measurements during the SaHAran Dust Experiment (SHADE).



SeaWiFs real-color image on 25th September 2000.

Adapted from Tanré et al., SHADE Special Issue, JGR, 2003

Colour  
2000  
and ORBIMAGE

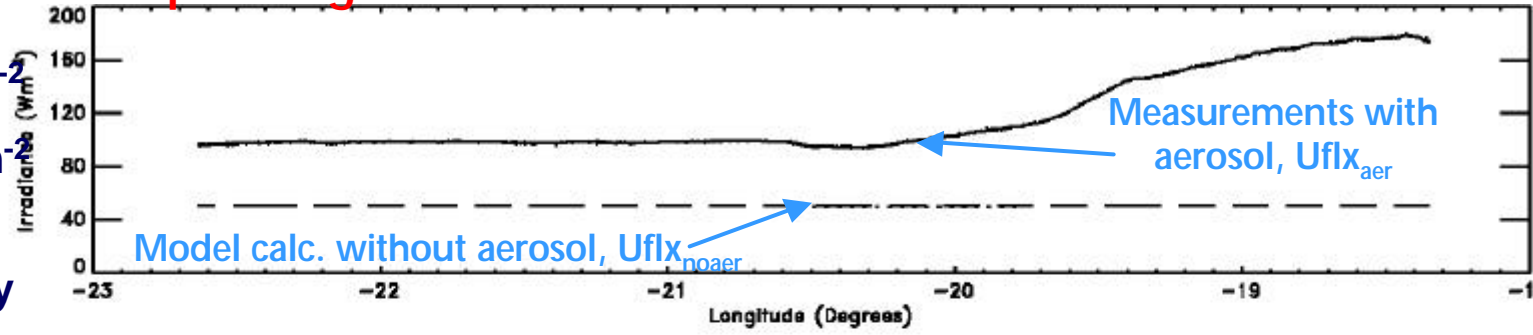
# Measurements of upwelling radiation above the aerosol



## Sal Upwelling irradiance

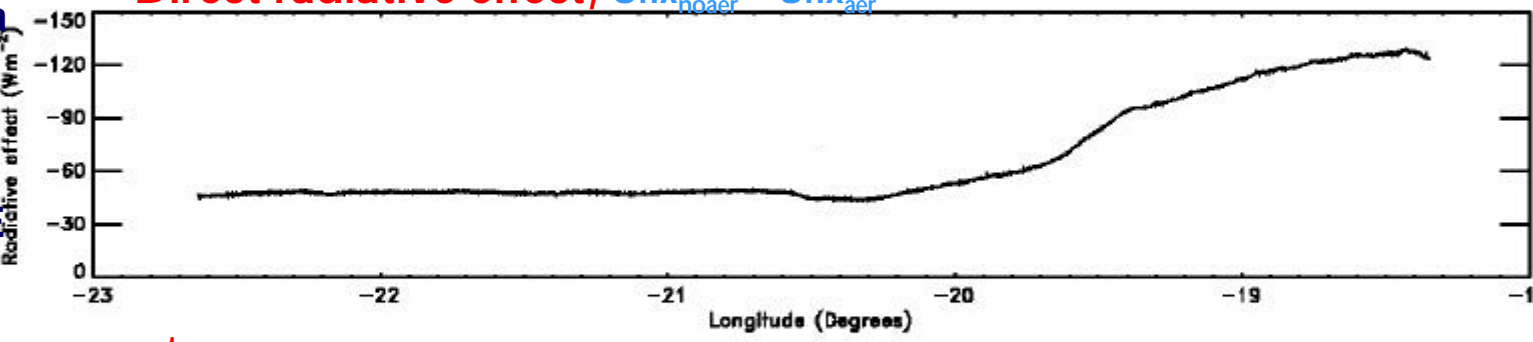
Daka

Max  $\text{Uflx}_{\text{aer}} > 179 \text{ Wm}^{-2}$   
 $\text{Uflx}_{\text{noaer}} < 50 \text{ Wm}^{-2}$   
 Local planetary albedo enhanced by a factor of ~3.  
 Max local DRE stronger than  $-120 \text{ Wm}^{-2}$



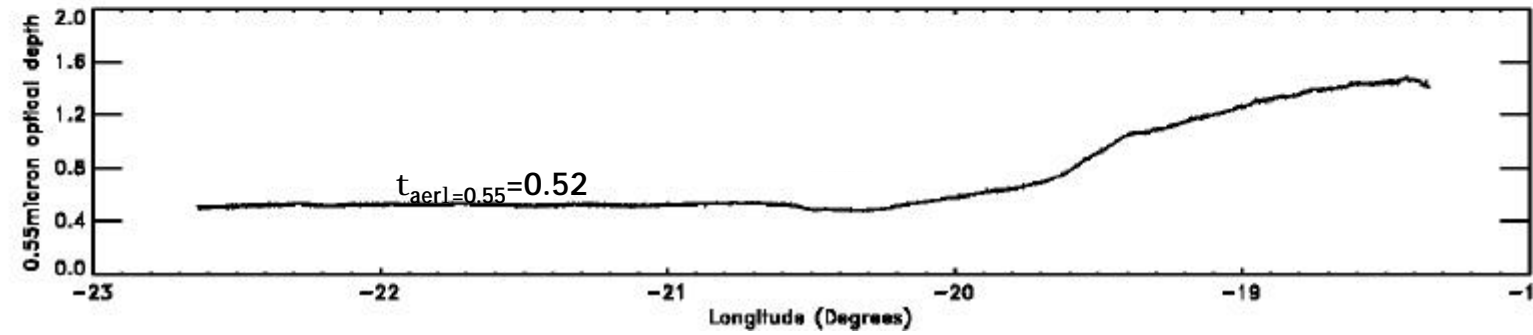
## Direct radiative effect, $\text{Uflx}_{\text{noaer}} - \text{Uflx}_{\text{aer}}$

Max local DRE stronger than  $-120 \text{ Wm}^{-2}$

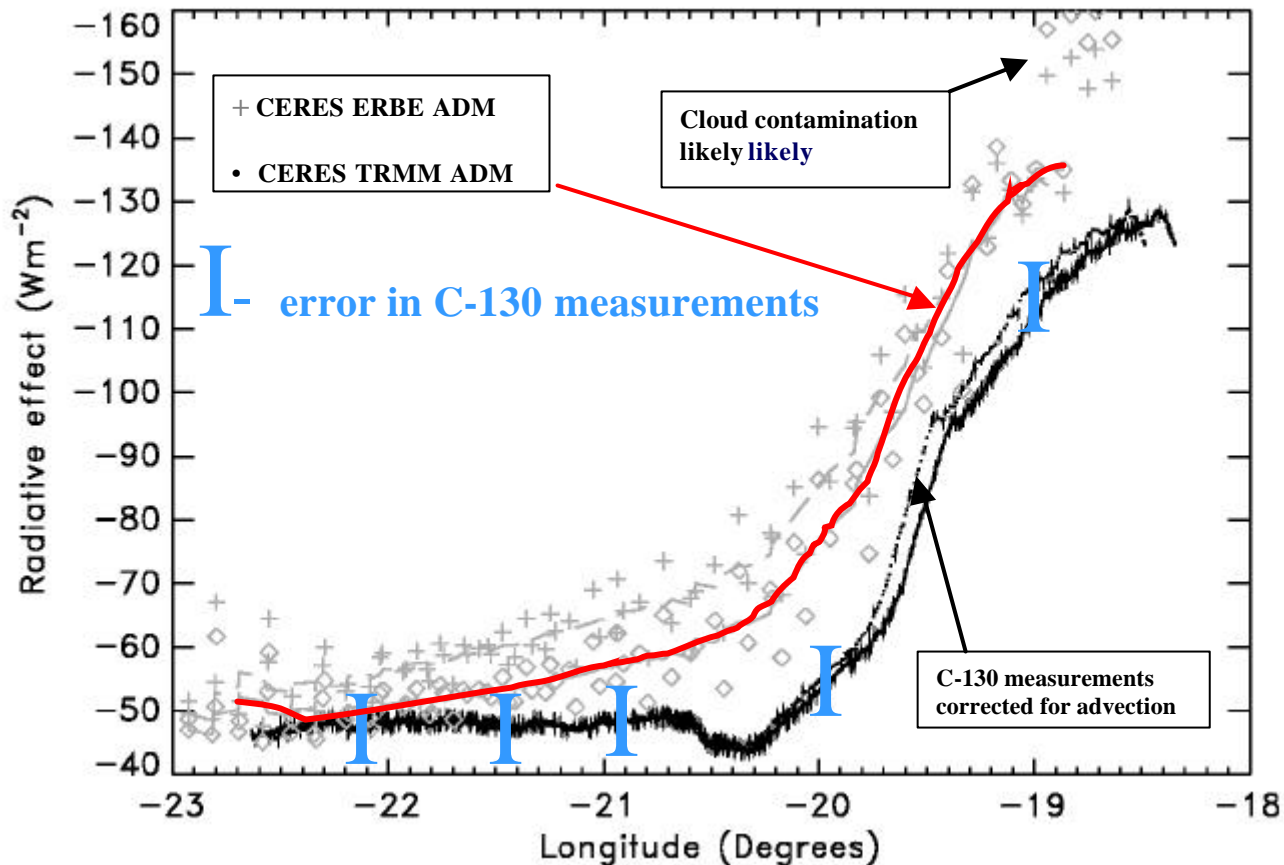


$t_{\text{aer}l} = 0.55$

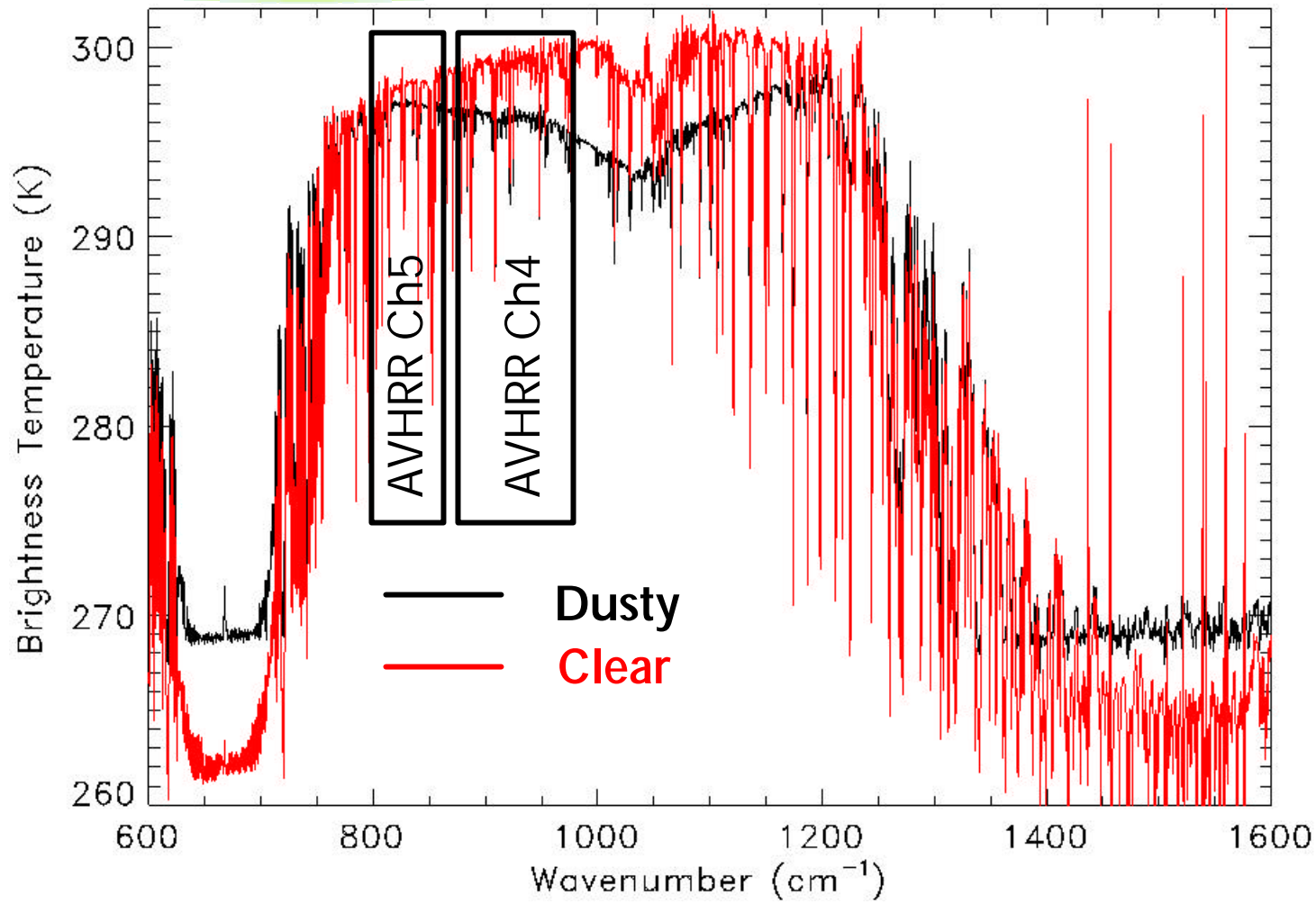
Max  $t_{\text{aer}l=0.55} > 1.5$



The C-130 and CERES direct radiative effects are in reasonable agreement, but outside the  $\pm 5 \text{ Wm}^{-2}$  error estimate for the BBRs.

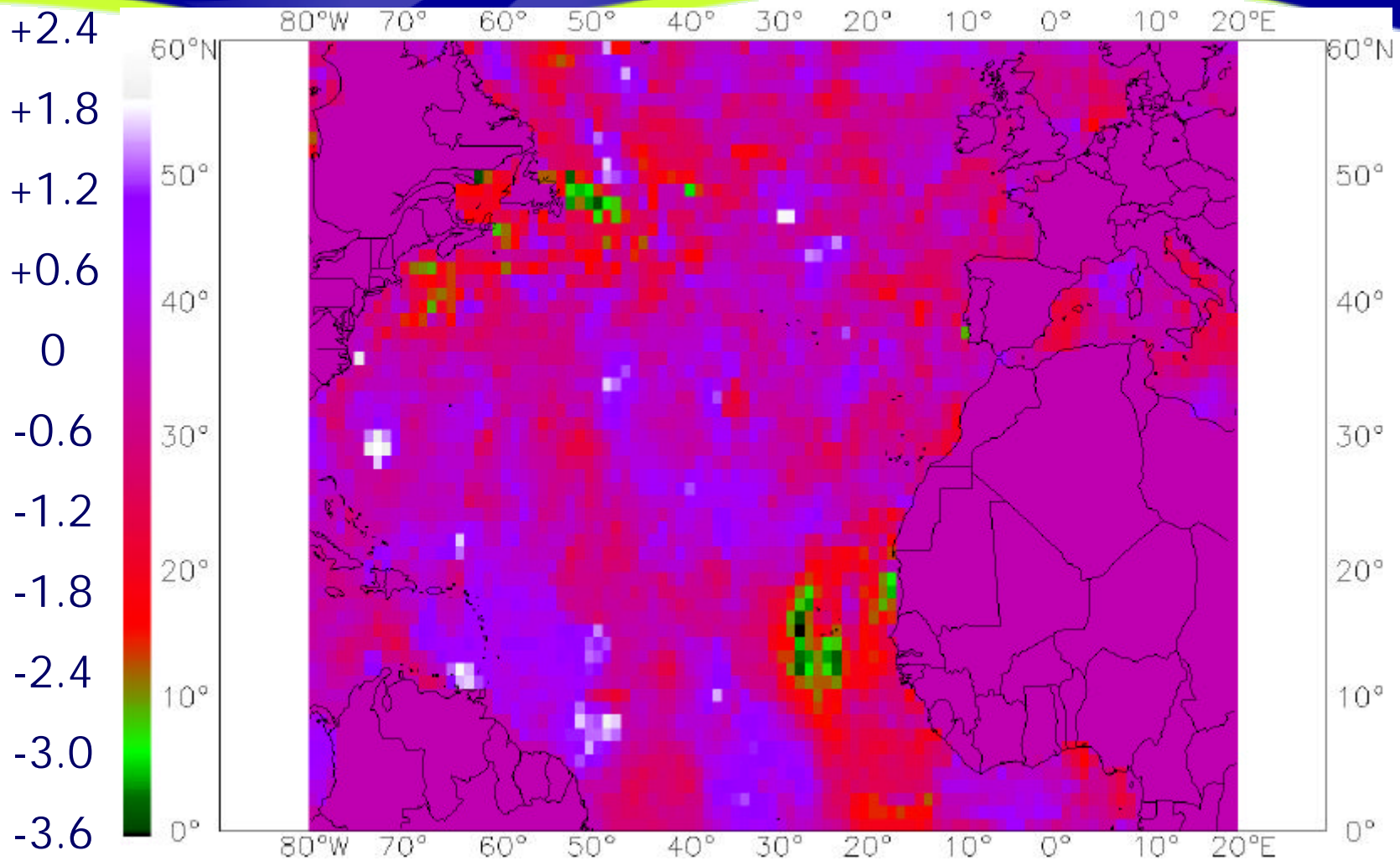


Haywood et al, 2003, JGR SHADE Special Issue



**Nadir views from 18,000ft (R6) (above aerosol).**

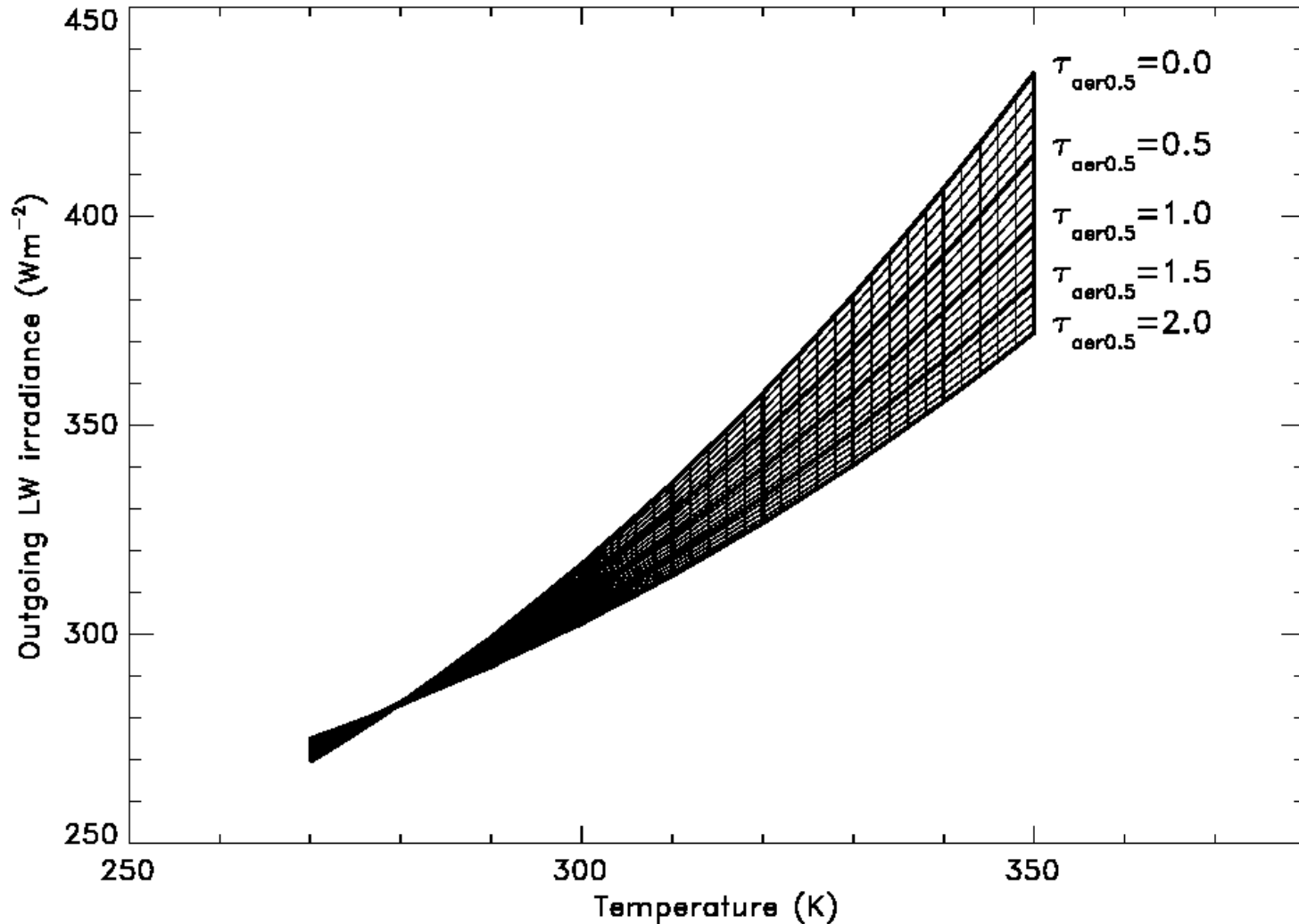
**Change in SST (K) from AVHRR data between 23rd and 27th September 2000.  
The SST anomaly over the Cape Verde Islands is evident and reaches -3.6K.**



**This is an artefact of the AVHRR retrieval algorithms which do not include mineral dust**

... the terrestrial effect is much stronger over warmer surfaces.

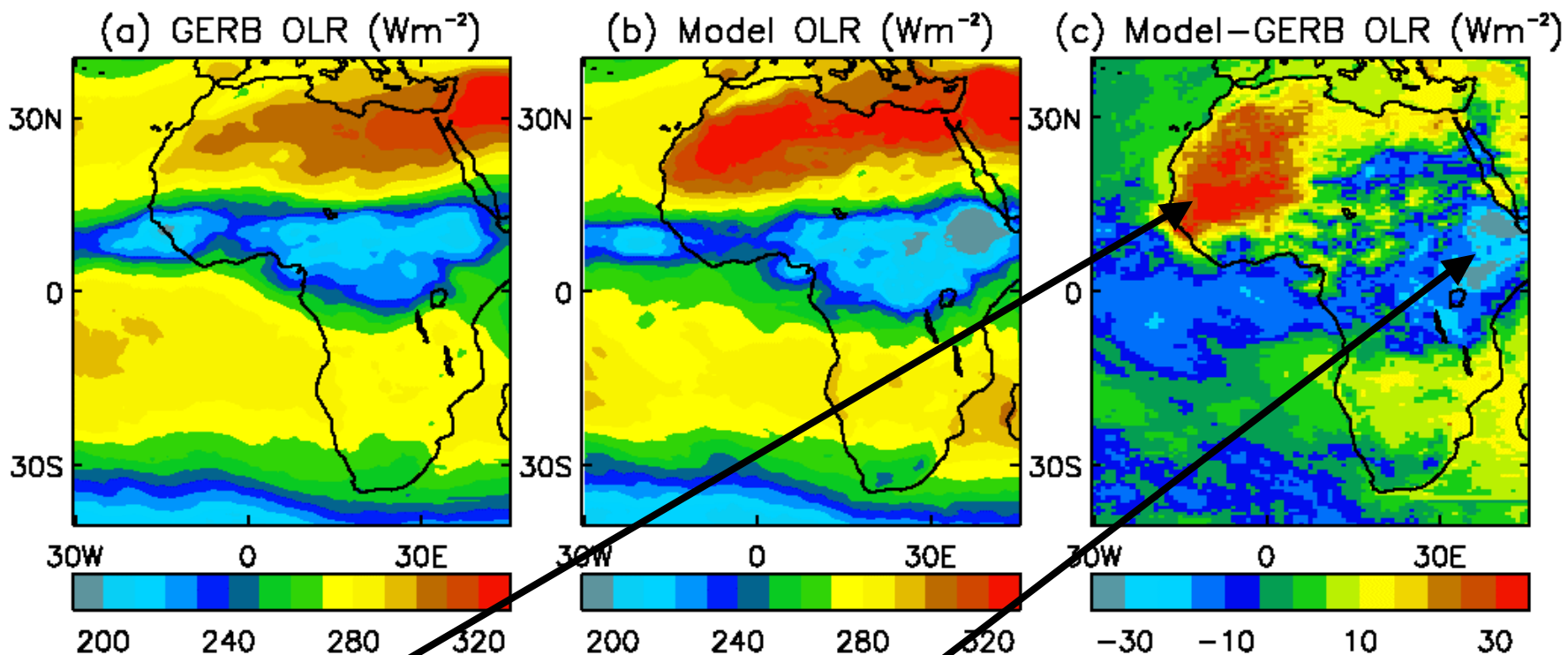
## Radiative calculations using the Edwards and Slingo radiation code.





# How does the NWP model OLR compare with new observations by the Geostationary Earth Radiation Budget instrument (GERB)?

Data from SINERGEE project using 6Z, 12Z, 18Z, 24Z, July 2003



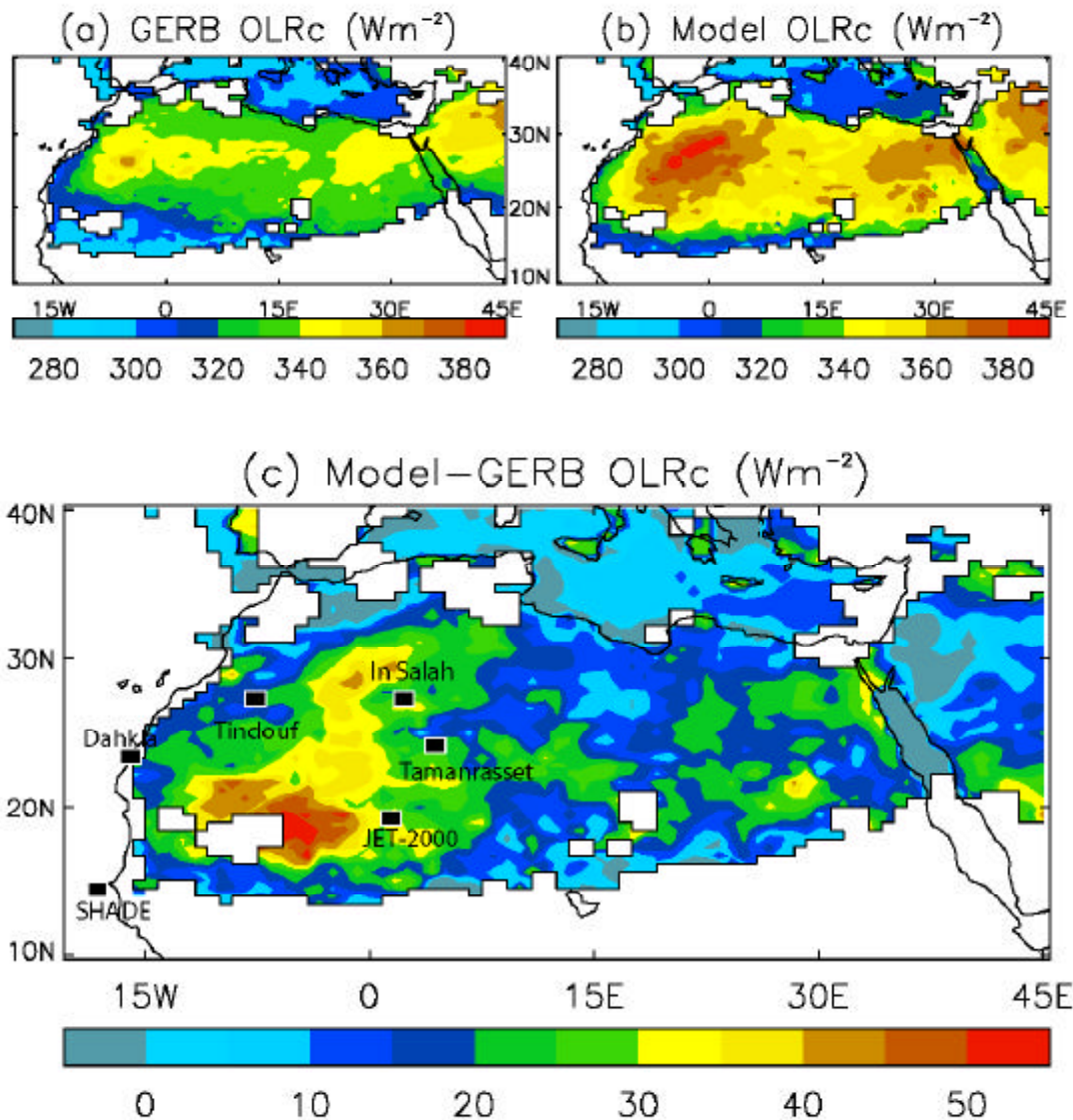
The +ve anomaly over desert is ~ -ve anomaly over ITCZ clouds

Rich Allan, Tony Slingo

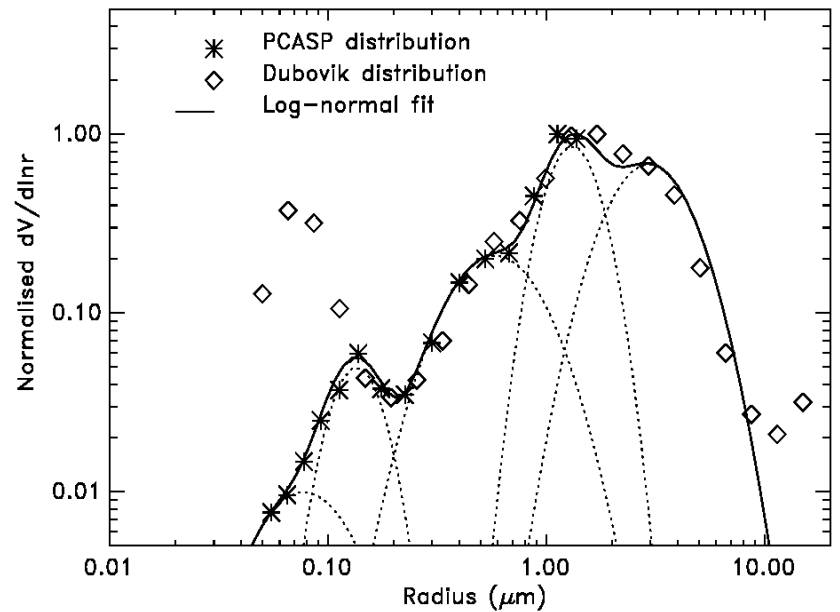
# Cloud screened data

The Geostationary Earth Radiation Budget instrument (GERB) shows significantly less OLR over regions of the desert during July 2003. What is the explanation?

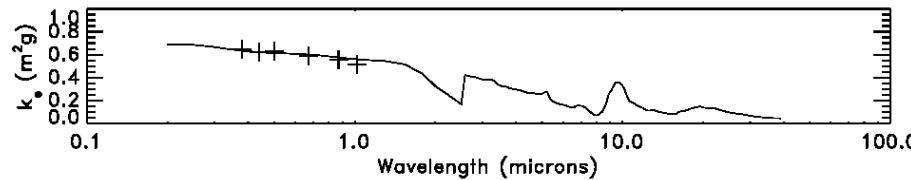
- a) Surface temperature?
- b) Emissivity?
- c) Atmospheric transmission?



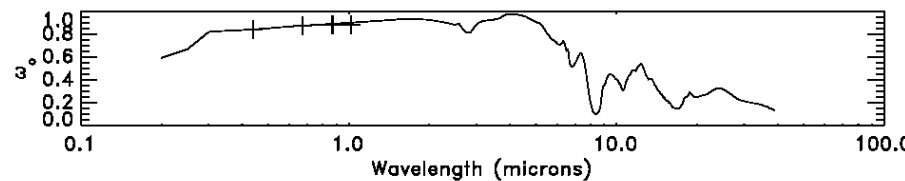
The July 2003 monthly mean aerosol size distribution from the nearby Dahkla AERONET site can be used with suitable refractive indices (Volz) to estimate the optical parameters associated with mineral dust.



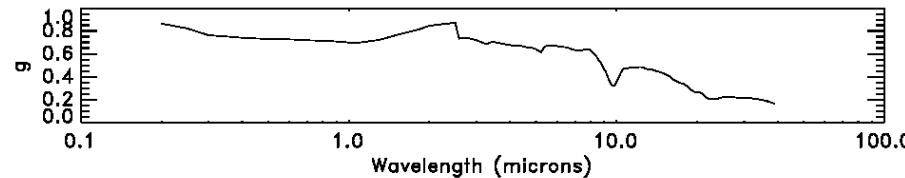
Specific extinction coefficient →



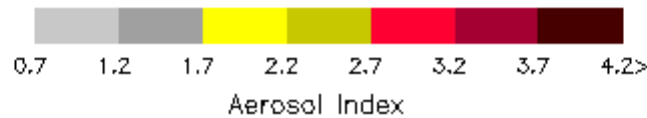
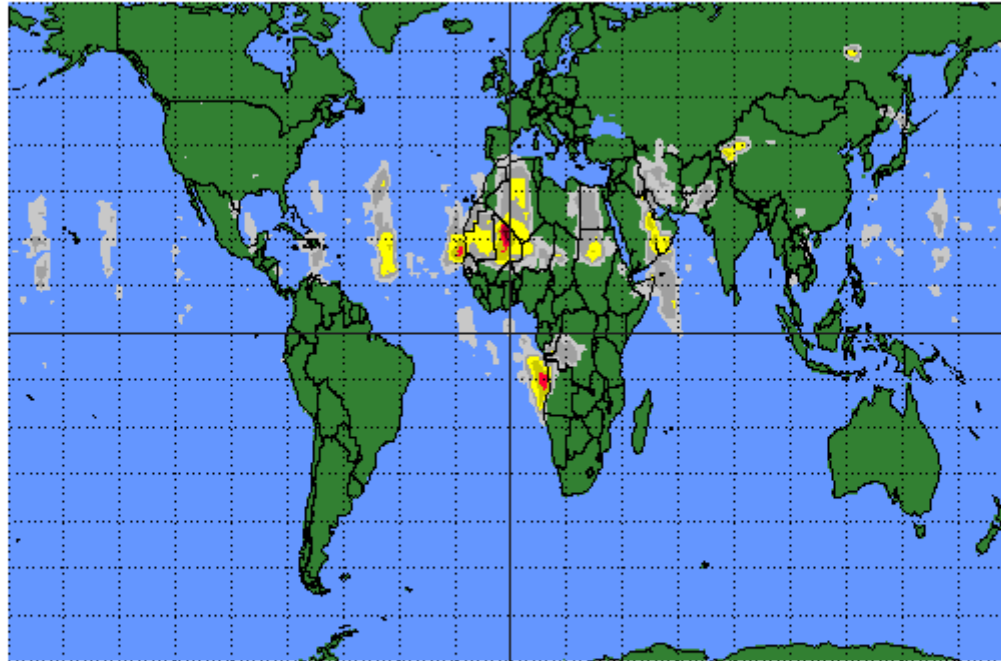
Single scattering albedo →



Asymmetry Factor →

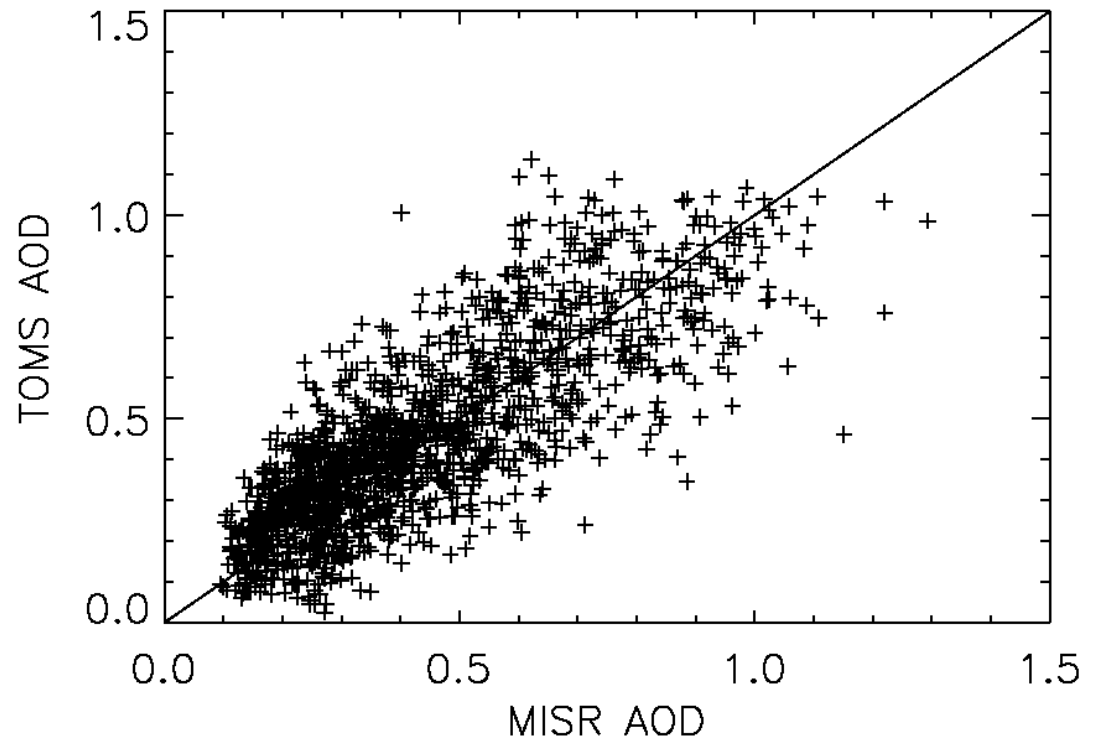
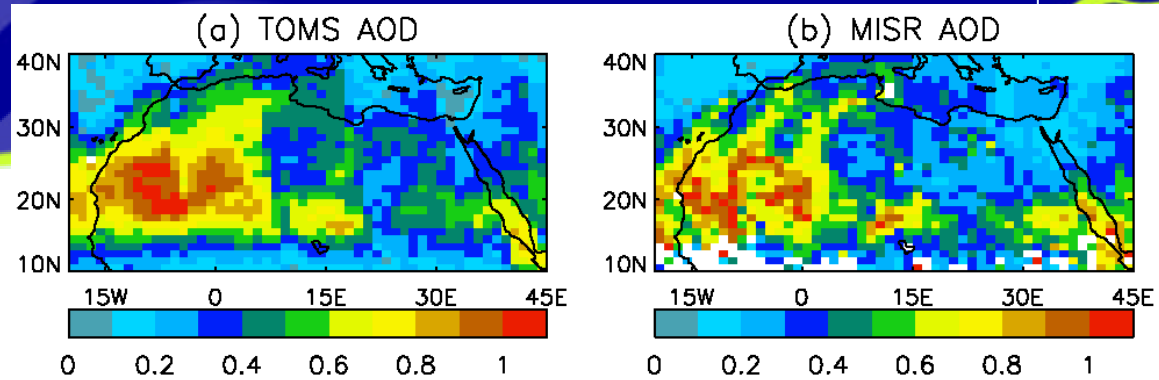


### Earth Probe TOMS Aerosol Index on July 31, 2003

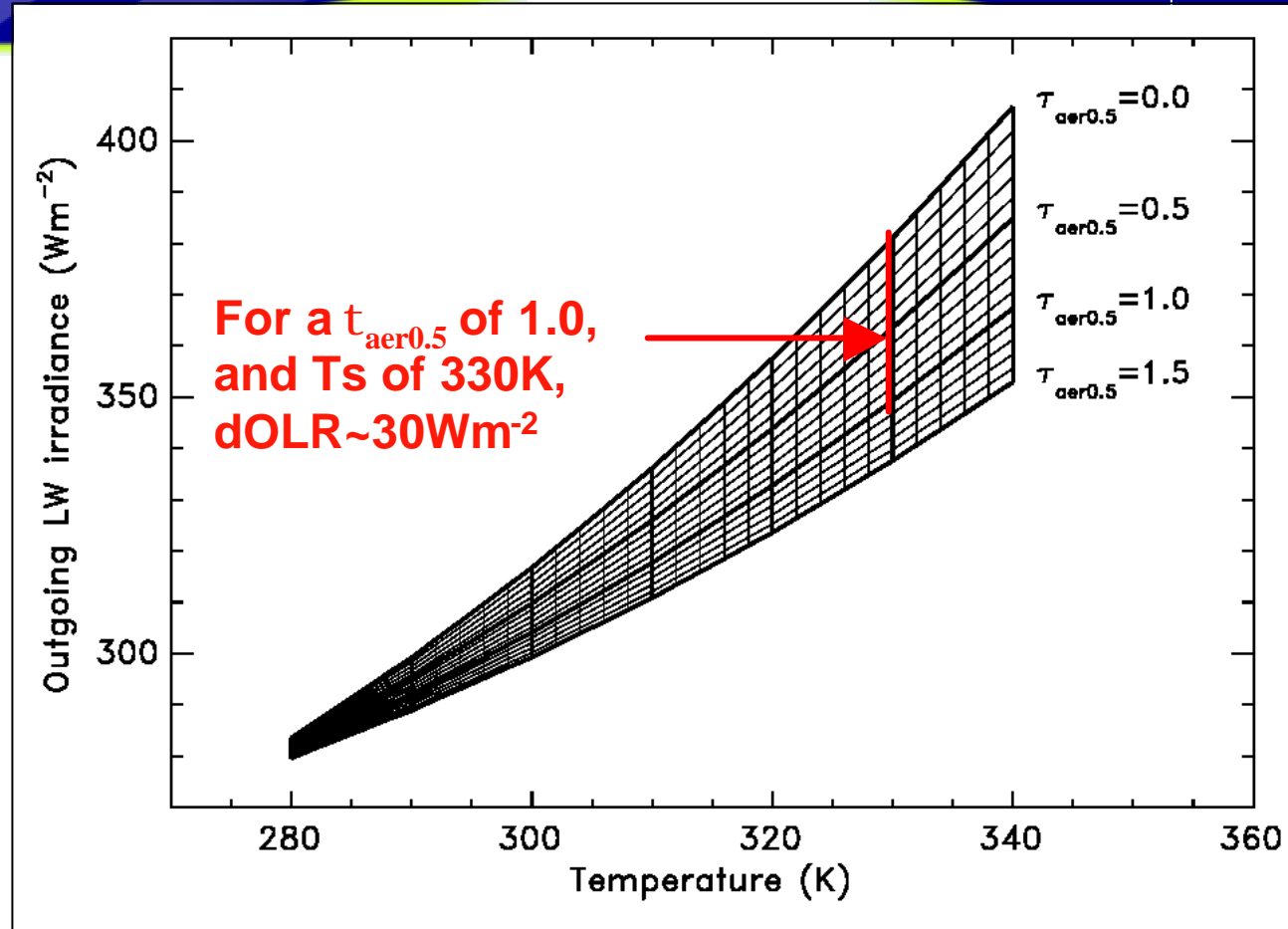


Goddard Space  
Flight Center

The monthly mean TOMS AI can be converted to a monthly mean AOD using empirical relationships based on AERONET observations. The results agree with the (v. much more) sophisticated MISR instrument.

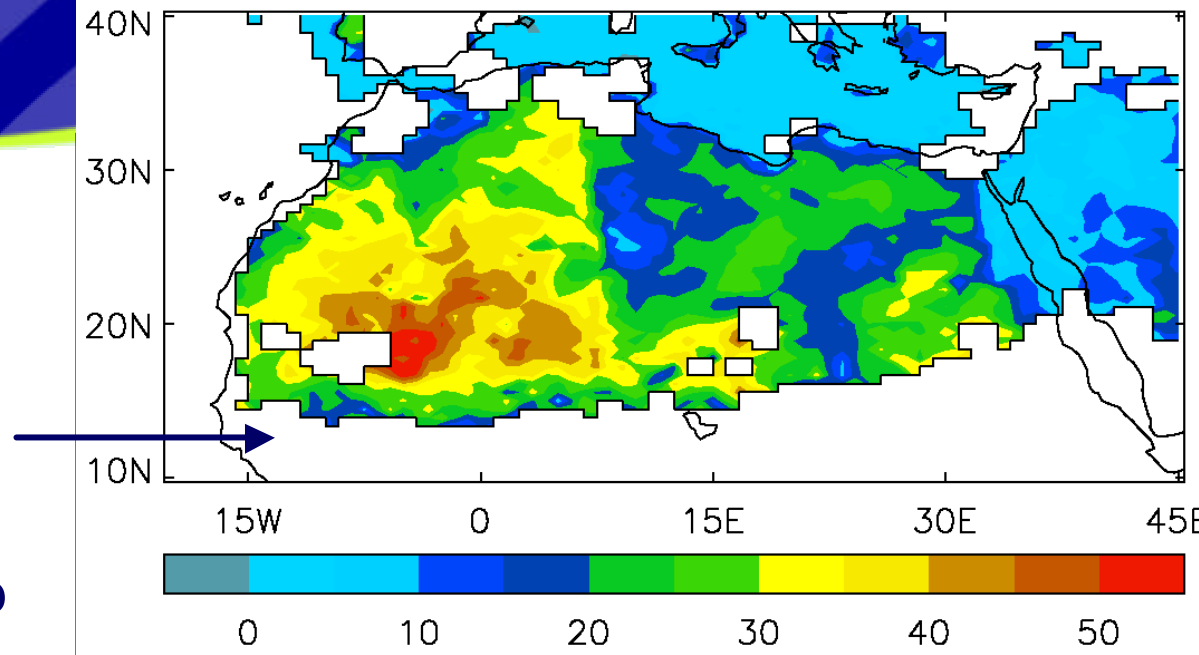


A look-up table may be produced whereby the dOLRc caused by mineral dust may be calculated as a function of aerosol optical depth and of  $T_s$ .

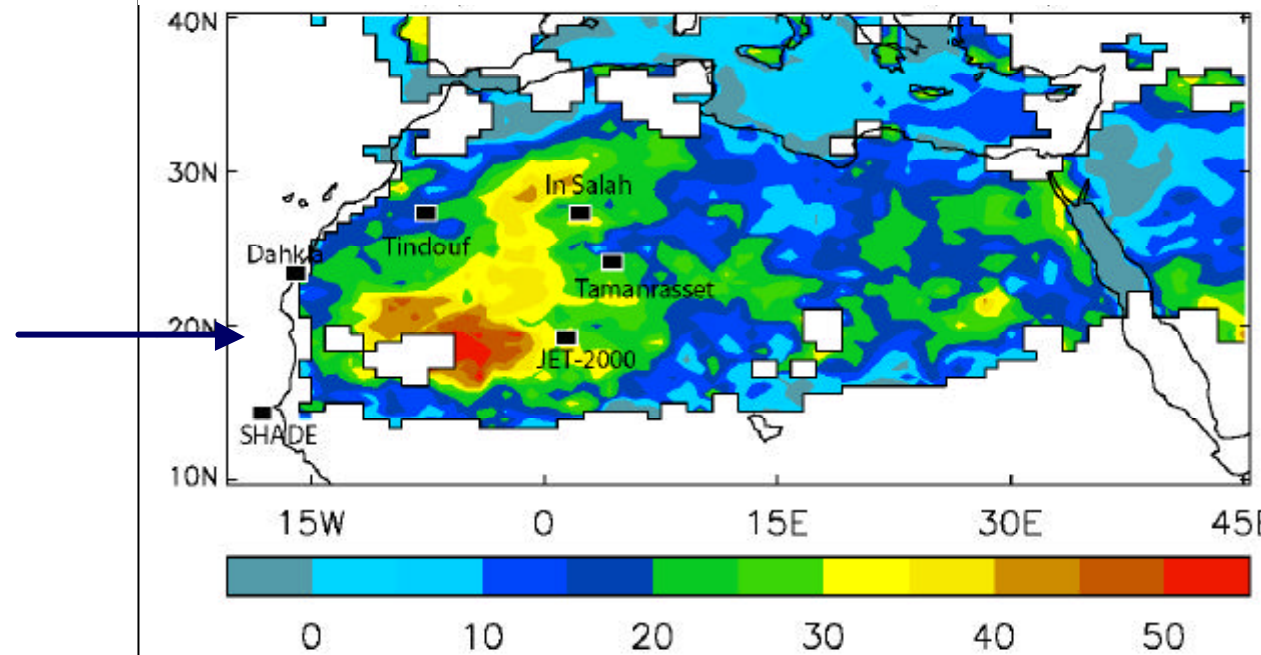


$DRE_{LW} + DRE_{LWfeedback}$

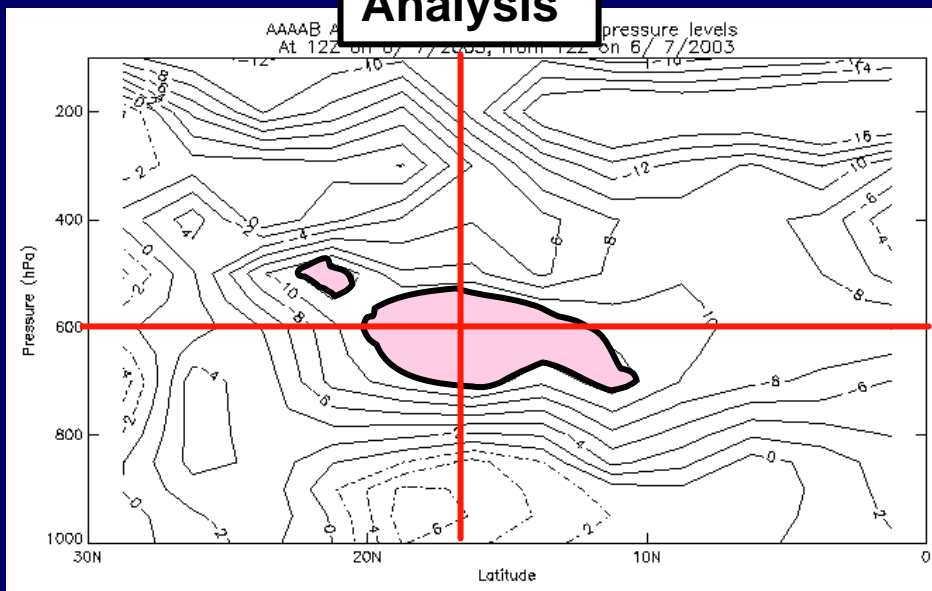
If we account for the effect of the aerosol on the SW at the surface which reduces the surface temperature and hence reduces the OLR as well, we end up with this.



Which is in good agreement with the dOLR between GERB and the UM.



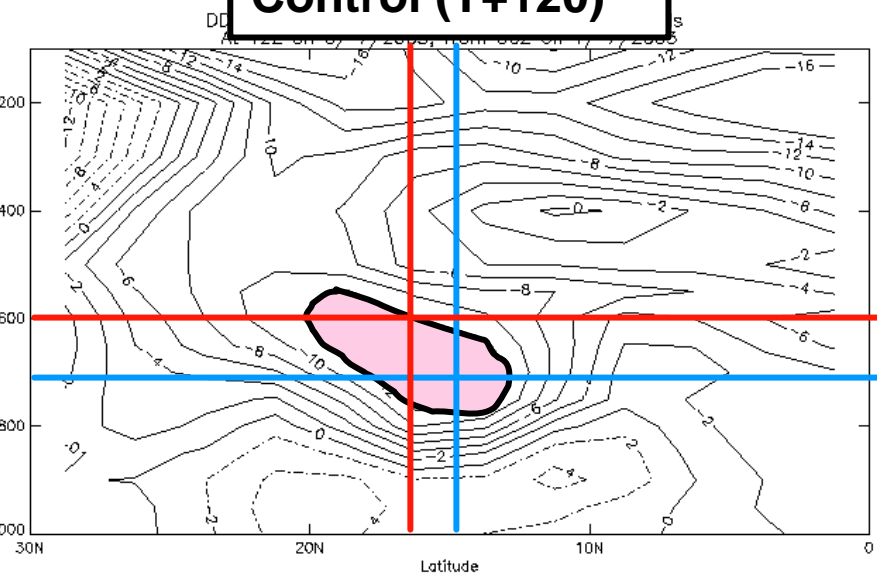
# Analysis



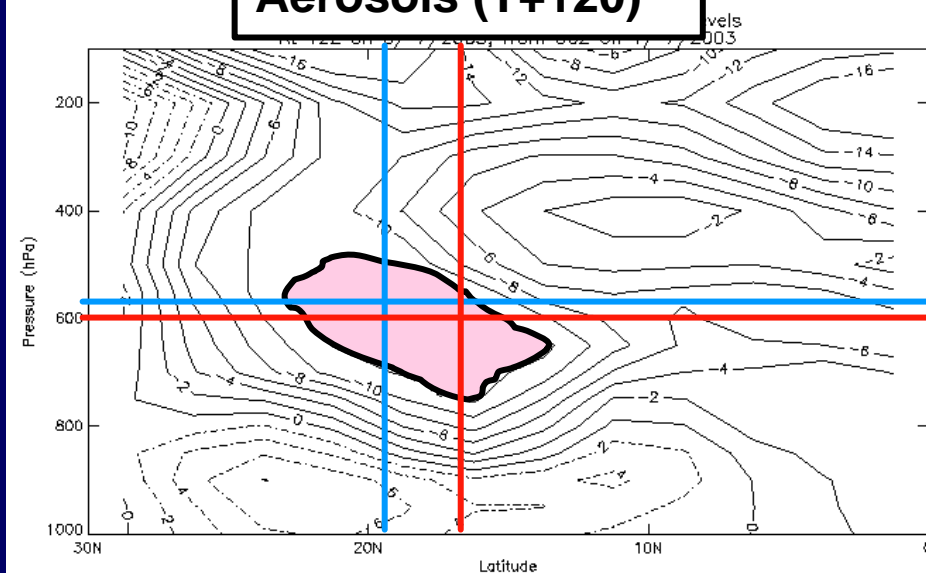
The altitude of the AEJ (and possibly the latitude) is in better agreement with the analysis when aerosols included



## Control (T+120)



## Aerosols (T+120)





Satellite retrievals are now available over land (to a lesser accuracy; problems over reflective surfaces)



**MODIS retrievals for:-**

**a) JFM**

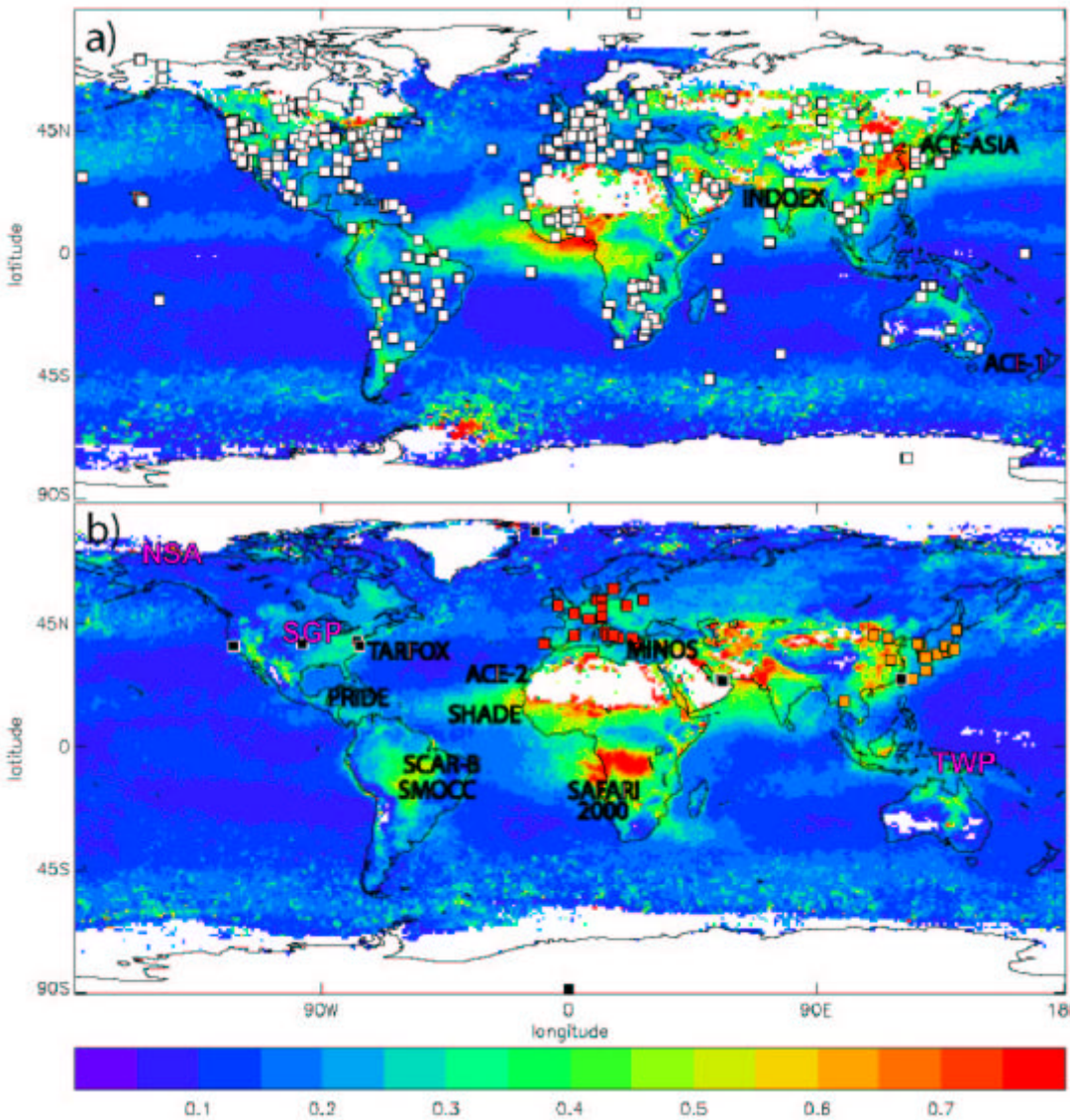
**b) ASO**

**Using a combination of MODIS Angstrom coefficient, TOMS and SSMI it is possible to break down the total aerosol optical depth into component parts:-**

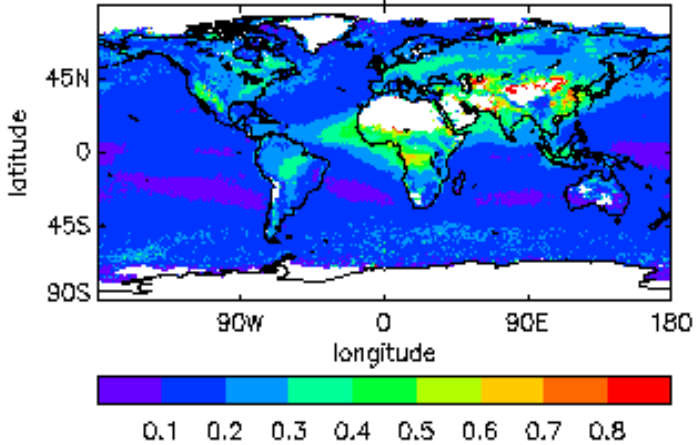
**a) Sea salt aerosol**

**b) Mineral dust**

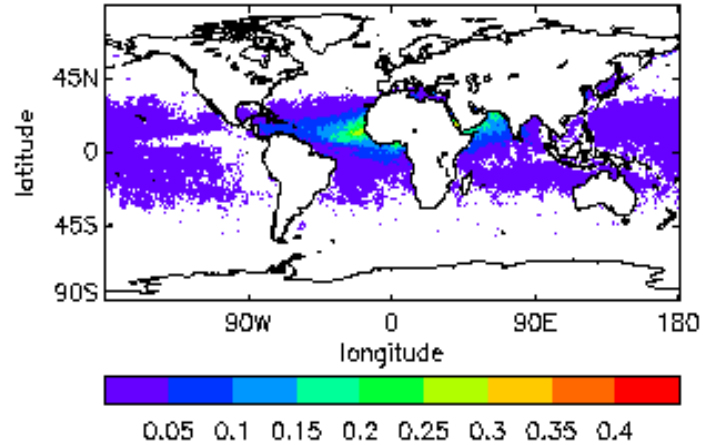
**c) Industrial aerosol/biomass burning aerosol**



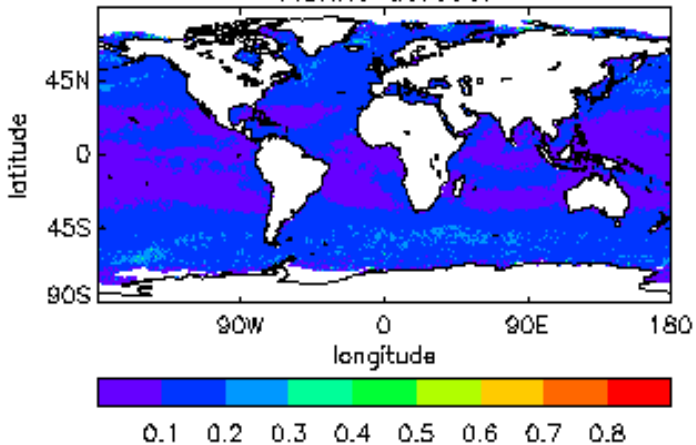
### Total aerosol optical thickness



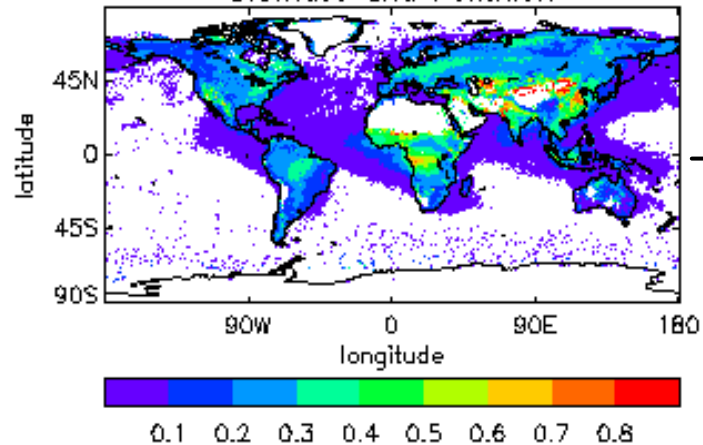
### Mineral dust



### Marine aerosol



### Biomass and Pollution



Direct radiative forcing ~  $<-1.0\text{Wm}^{-2}$

## Conclusions

1. **Observations (in-situ, surface remote sensing, satellite) are extremely useful tools in developing our understanding of the important physical processes associated with aerosols. It is important to cross calibrate these methods.**
2. **The direct radiative forcing due to aerosols derived from observational measurement methods is significant.**
3. **The radiative effects (natural component) of aerosols can be considerable particularly for thick aerosol such as mineral dust (e.g.  $-120\text{Wm}^{-2}$  in SW over ocean,  $+50\text{Wm}^{-2}$  in LW over land).**
4. **Aerosols (or the neglect of them) can cause significant problems in remote sensing methods (e.g. cloud optical depth, cloud effective radius, sea-surface temperatures, OLR etc).**