We need to reconcile the single scattering albedo inferred from satellite or AERONET measurements (0.94-0.97 @ 550nm) with what modellers have assumed (0.89 to 0.92) from refractive indices measured by Volz and Patterson

Explain discrepancies in the imaginary part of the refractive index of dust between AERONET retrievals and Volz and Patterson measurements.

We have used

1) Studies of the mineralogy since refractive indices are defined over a much broader spectral interval for minerals: Previous works include *Sokolik et al., 1999*; *Claquin et al, 1999*; *Claquin 2000 (Ph.D. thesis)* 

2) Quantified the relative volumes of the main minerals

3) Applied a dielectric model to compute the refractive indices of the minerals As internal mixtures

4) Re-compute the perturbation flux and the direct radiative effect of Mineral dust

# Retrieved Properties of Saharan DustAngstrom < 0.75</td>Dubovik et al., 2002

Average Single Scattering Albedo



### Comparison of dust refractive indices from Patterson and Volz and retrieved from AERONET





Volz (73)



Claquin et al. (1998)

#### **Three experiments**





Unique Refractive Index

Externally Mixed in Each Mineral

6 Minerals Hematite Inclusions

Pourcentage Volume OF THE 6 MINERALS From Which Refractive Index Is Derived

	LOW HEMATITE	MEDIUM HEMATITE	HIGH HEMATITE
QUARTZ	14.0	14.0	14.0
CALCITE	6.0	6.0	6.0
HEMATITE	0.9	1.5	2.7
ILLLITE	32.1	31.5	30.3
KAOLINITE	24.0	24.0	24.0
MONTMOTILLONITE	23.0	23.0	23.0



The dielectric model for refractive indices

For a two-component mixture composed of inclusions embedded In a homogeneous matrix it can be shown for spheres that the resulting Refractive index is:

$$\varepsilon_{\rm eff} = \varepsilon_{\rm m} \left\{ 1 + \left[ 3 \, v_1 \left( \varepsilon - \varepsilon_{\rm m} \right) \right/ \left( \varepsilon + 2 \, \varepsilon_{\rm m} \right) \right] / \left[ 1 - v_1 \left( \varepsilon - \varepsilon_{\rm m} \right) \right/ \left( \varepsilon + 2 \, \varepsilon_{\rm m} \right) \right] \right\}$$

Where  $\varepsilon_m$  and  $\varepsilon$  are the complex refractive indices of the matrix and of The includion

### Here the refractive index of the mineral mixture is computed With the assumption of a dielectric







"TOA RADIATIVE FORCING, ALL SKIES, 0.9% hemotite (DIELECTRIC)"



"TOA RADIATIVE FORCING, ALL SKIES, 2.7% hemotite (DIELECTRIC)"



## Retrieved Properties of Saharan DustAngstrom < 0.75</td>Dubovik et al., 2002

#### **Average Size Distributions**



### PERTURBATION TO THE RADIATIVE FLUX (TOP OF ATMOSPHERE)

]	EXTERNAL Mixture	INTERNAL VOL. WEIGHED	INTERNAL DIELECTRIC	
LOW Hen	natite			
CASE	NA	-0.68	-0.62	
MEDIUM	Hematite			
CASE	-0.54	-0.46	-0.55	
High Hem	atite			
CASE	NA	-0.23	0.41	

### DUST FORCING (TOP OF ATMOSPHERE)

EXTERI	NAL Mixture	INTERNAL VOL. WEIGHED	INTERNAL DIELECTRIC	
LOW Hematite				
CASE	NA	-0.10 to -0.34	-0,09 to -0.31	
MEDIUM Hemati	te			
CASE	-0.08 to -0.27	-0.07 to -0.23	-0.08 to -0.23	
High Hematite				
CASE	NA	-0.03 to -0.12	-0.06 to -0.21	





• We have identified why dust models assumed more absorbing aerosols than the satellite retrievals indicate

• A mineralogical approach with realistic values for the composition in iron oxides has given the basis to better constrain the direct forcing for mineral dust

• We conclude that this forcing is an overall negative forcing in the range [-0.26 to -0.05 W.m-2]. Previous uncertainty in TAR ranged from -0.7 to +0.2 W.m-2 for dust.

•We are varying the size distribution for additional sensitivity studies