

An hour for the atmosphere...

UNIVERSITY  
OF MIAMI



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7/15.

# Start with Atmospheric Optical Depth

Two basic measurement  
methods

1) Sunphotometers (handheld  
or autonomous): 2 degree field  
of view around sun (which  
subtends 0.5 degree full angle).

Most popular handheld now  
with our crowd, Microtops...you  
have seen already.



# Atmospheric Optical depth

- Another method: Shadowband radiometer

Shadowband rotates and either provides a total downwelling irradiance  $E_s$  (no shadow) or the diffuse irradiance,  $E_{ds}$ . Direct solar irradiance,  $E_d$ , is roughly the difference:

$$E_d = (E_s - E_{ds}) / \cos(\text{solar zenith})$$

J. Michalsky and J. Berndt. "Automated Multifilter Rotating Shadow-band Radiometer: an Instrument for Optical Depth and Radiation Measurements." *Applied Optics*, Vol.33, No.22. pp 5118-5125.



# Either way

$$E_d = E_o \exp(-m\tau)$$

*or*

$$\ln(E_d) = \ln(E_o) - \tau m$$

*or*

$$\tau = -\frac{1}{m} \ln\left(\frac{E_d}{E_o}\right)$$

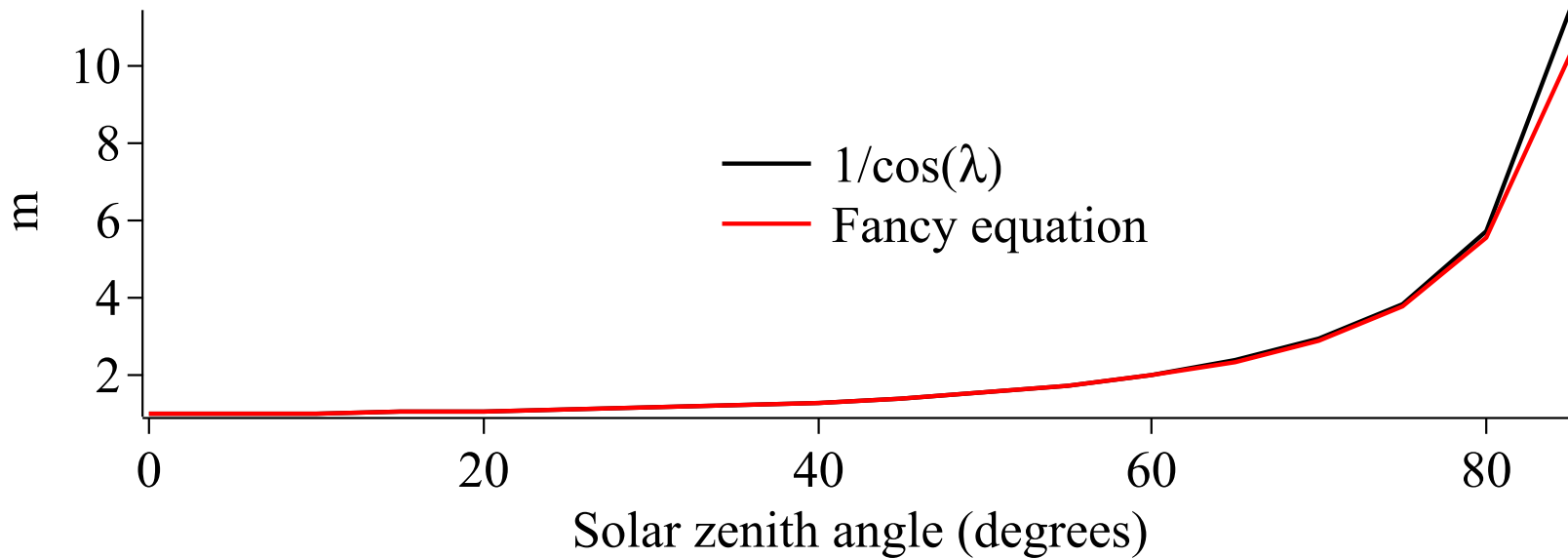
Either need  $E_o$  in terms of instrument units or have a stable day and take measurements with varying  $m$  and slope is  $-\tau$ .

$E_o$  is the extraterrestrial solar irradiance in instrument units.

# Calculate Air mass from solar ephemeris program (or tables) given date, time, location

- $m = 1.0 / [ \cos(Z) + 0.50572 * (96.07995 - Z)^{-1.6364} ]$
- Or  $m = 1/\cos(z)$  NOTE  $z$  is solar zenith angle in degrees

Only matters above 70 degrees difference is due to earth curvature



# What should you expect for Optical depth

$$\tau_{total} = \tau_R + \tau_g + \tau_a$$

$\tau_R$  is rayleigh (molecular scattering) optical depth

$\tau_g$  is absorption by molecular gases (ozone, water vapor, etc).

$\tau_a$  is aerosol optical depth (scattering and absorption....).

For aerosol information choose spectral region where most gas absorption is avoided.

water vapor, choose good water vapor absorption band...

ozone, choose good ozone absorption band....

# Rayleigh Optical depth

- Falls off as  $\lambda^{-4}$

Full blown equation, from Bodhaine et al. 1999, supposed to be accurate in the visible to within 0.02%.....

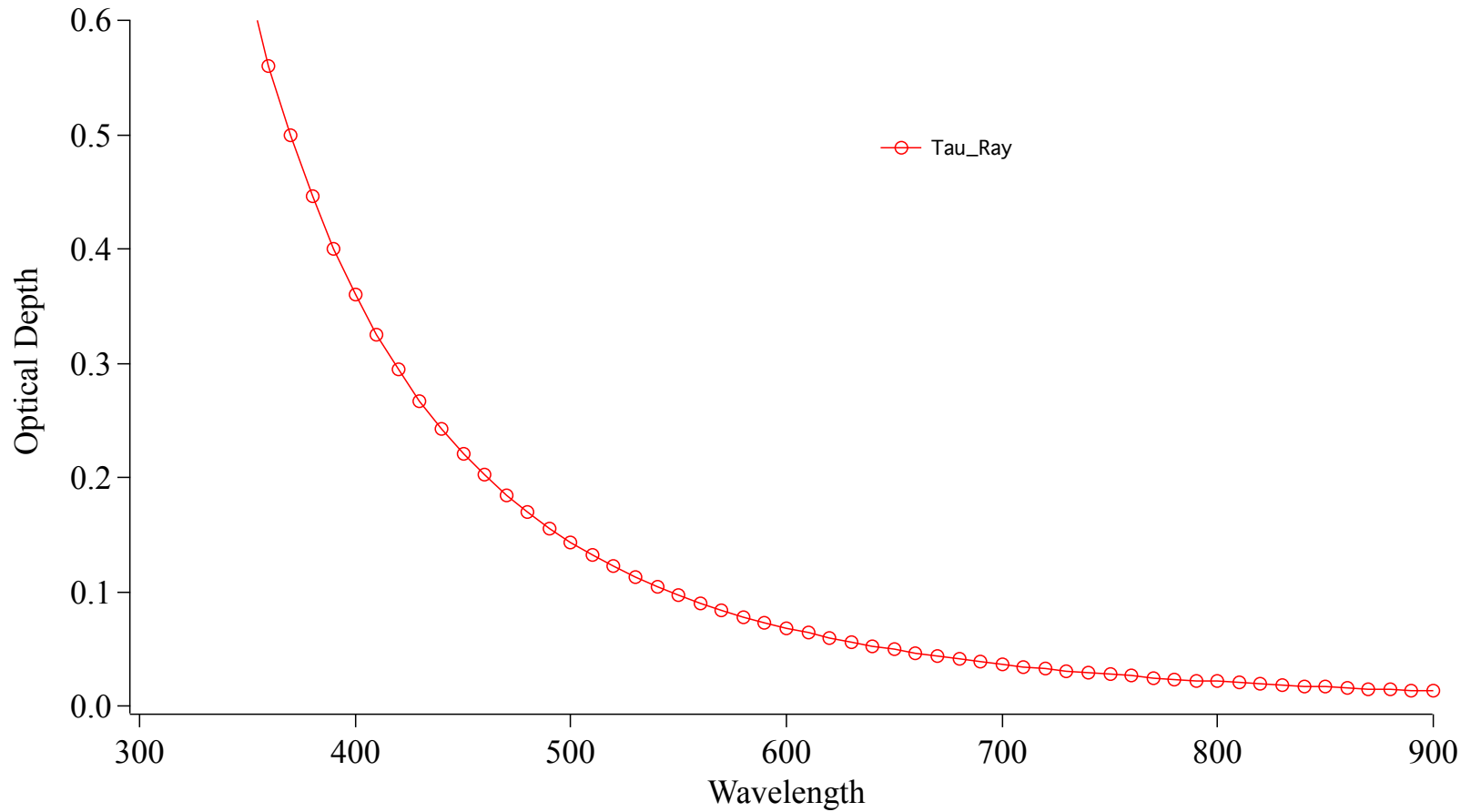
$$\tau_R = 0.0021520 \left( \frac{1.0455996 - 341.29061\lambda^{-2} - 0.90230850\lambda^2}{1 + 0.0027059889\lambda^{-2} - 85.968563\lambda^2} \right)$$

At sea level, 45 deg latitude, 1013.25 mb Scale by pressure on that day

For microtops

Wavelength	$\tau_R$
380	0.446
440	0.243
500	0.143
675	0.042
870	0.015

# Rayleigh Optical Depth





# Ozone

- In UV very strong, to measure ozone, differential UV bands are used (300, 305.5, 312.5 nm)
- In visible still evident. Ozone varies over the globe, so need to get column ozone value at measurement site from climatology or from some other source (TOMS, etc.). Typically reported in Dobson Units which are milli atm-cm. Get ozone optical depth by multiplying ozone absorption coefficient by  $DU/1000$

# Ozone climatology, Ziemke et al. 2011 (need to add tropospheric ozone, around 40DU, in paper)...MLS is Microwave Limb Sounder

**Table 3.** Global stratospheric column ozone zonal mean monthly mean climatology (in Dobson Units) derived from MLS integrated ozone profiles at 5 latitude resolution.

Latitudes	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec
85 N–90 N	311	350	352	340	324	297	269	253	243	246	267	279
75 N–80 N	319	363	363	359	330	301	272	254	248	254	275	287
65 N–70 N	330	359	359	355	332	304	280	269	262	265	280	299
55 N–60 N	332	355	354	347	329	307	291	277	270	270	282	309
50 N–55 N	329	349	348	340	325	306	290	276	267	267	279	308
45 N–50 N	321	338	338	331	318	299	280	269	262	260	273	302
40 N–45 N	307	320	324	318	304	286	266	259	254	252	265	289
35 N–40 N	284	292	300	298	286	271	256	252	248	243	252	269
30 N–35 N	257	262	272	275	269	259	251	248	244	237	237	247
25 N–30 N	234	238	248	255	256	252	248	246	242	234	229	230
20 N–25 N	221	224	234	242	246	246	245	244	240	232	224	219
15 N–20 N	215	218	227	235	240	242	244	245	241	232	223	215
10 N–15 N	214	216	224	232	237	240	244	245	242	232	223	214
5 N–10 N	215	217	223	230	234	237	240	242	240	231	224	216
0–5 N	219	221	226	231	232	233	236	238	237	229	225	219

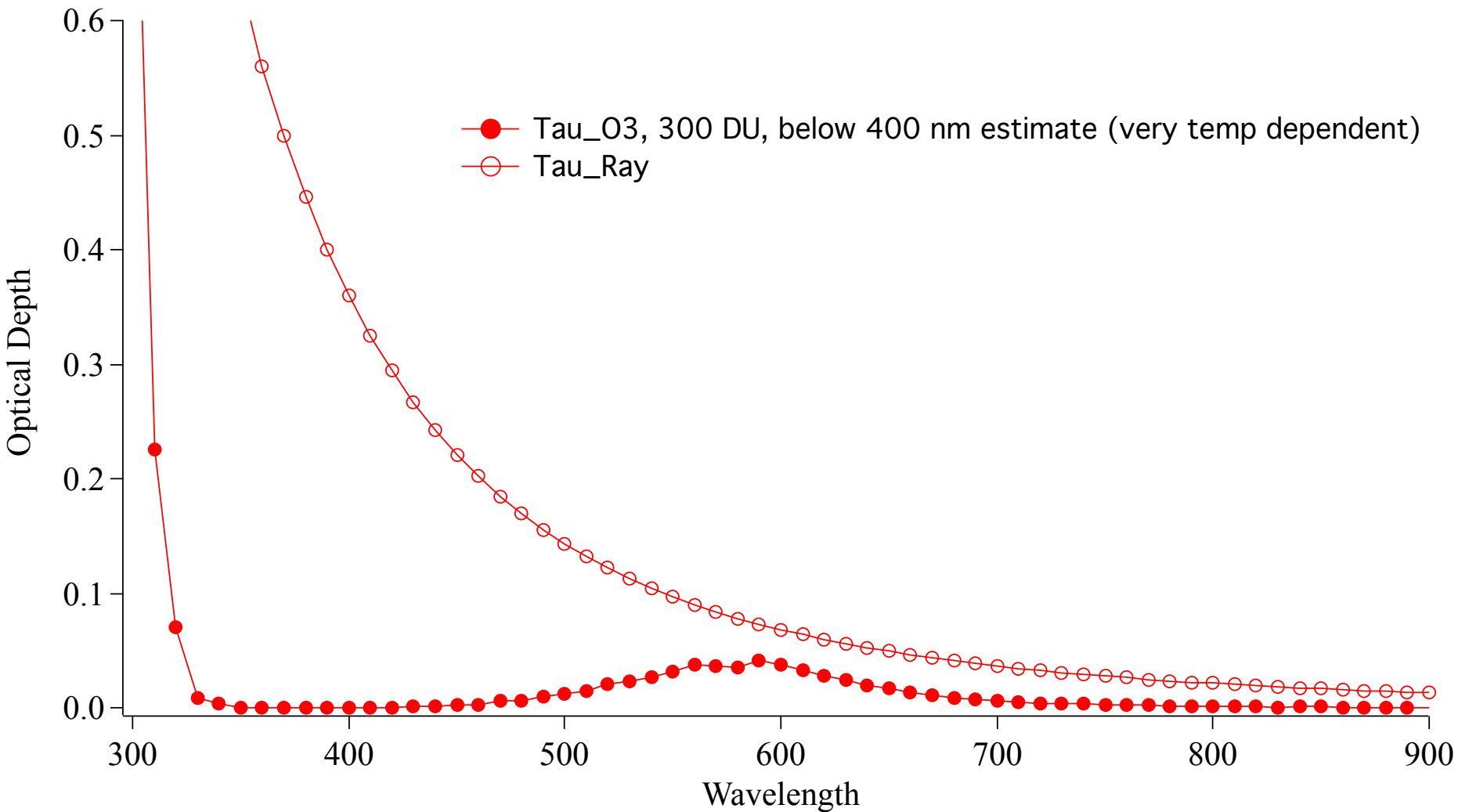
NOTE: I TRUNCATED TABLE>>>>HAS SOUTHERN HEMISPHERE ALSO

# Ozone (308 DU, estimate for our cruise)

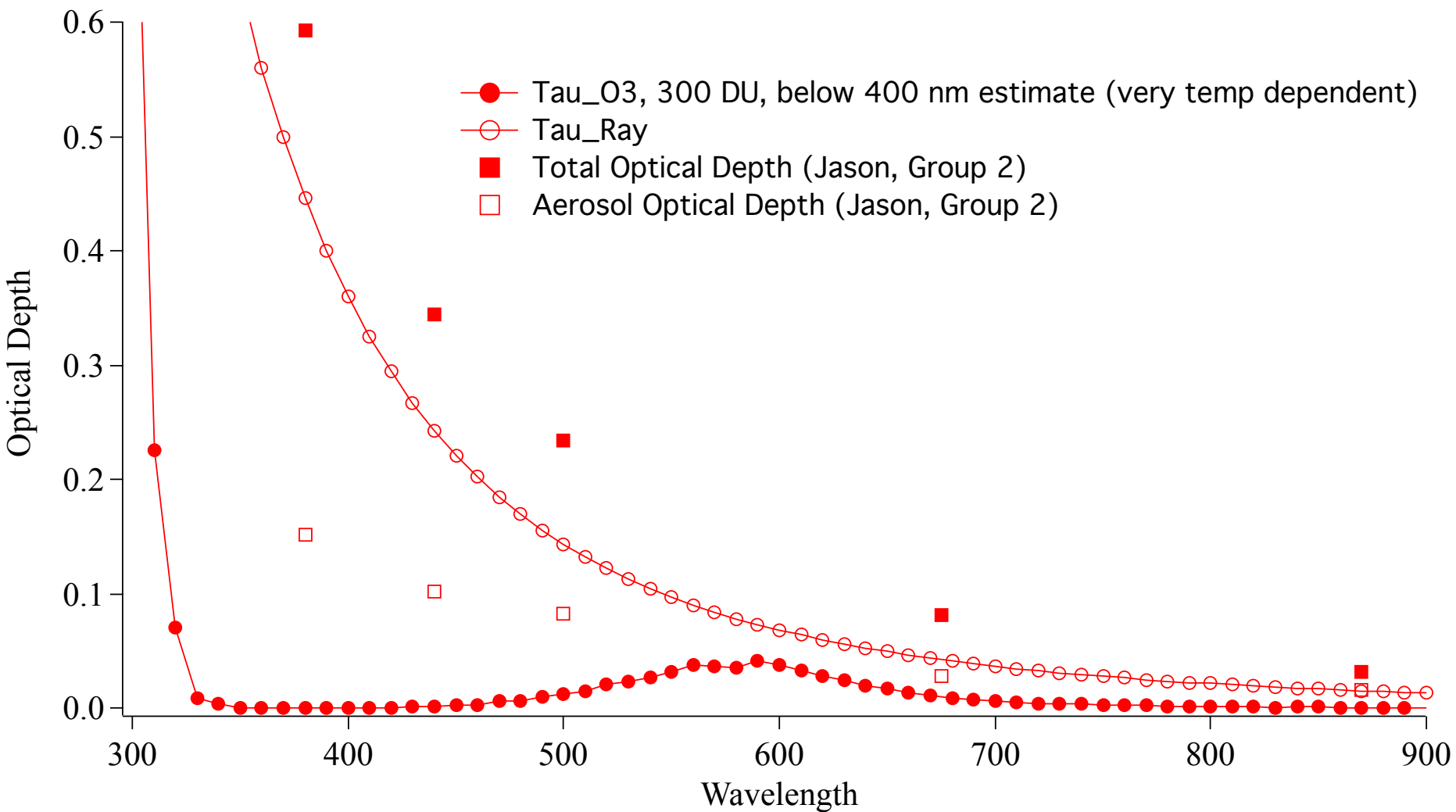
- | Wavelength | absorption coeff. | tauO3 |
|------------|-------------------|-------|
| 380        | 0                 | 0.000 |
| 440        | 0.0021            | 0.001 |
| 500        | 0.0320            | 0.010 |
| 675        | 0.0401            | 0.012 |
| 870        | 0.0013            | 0.000 |

308 DU from Ziemke et al. 2011, absorption coefficients from doe-sc-arm-tr-129.pdf and Gorshelev et al. 2014 (associated tables)

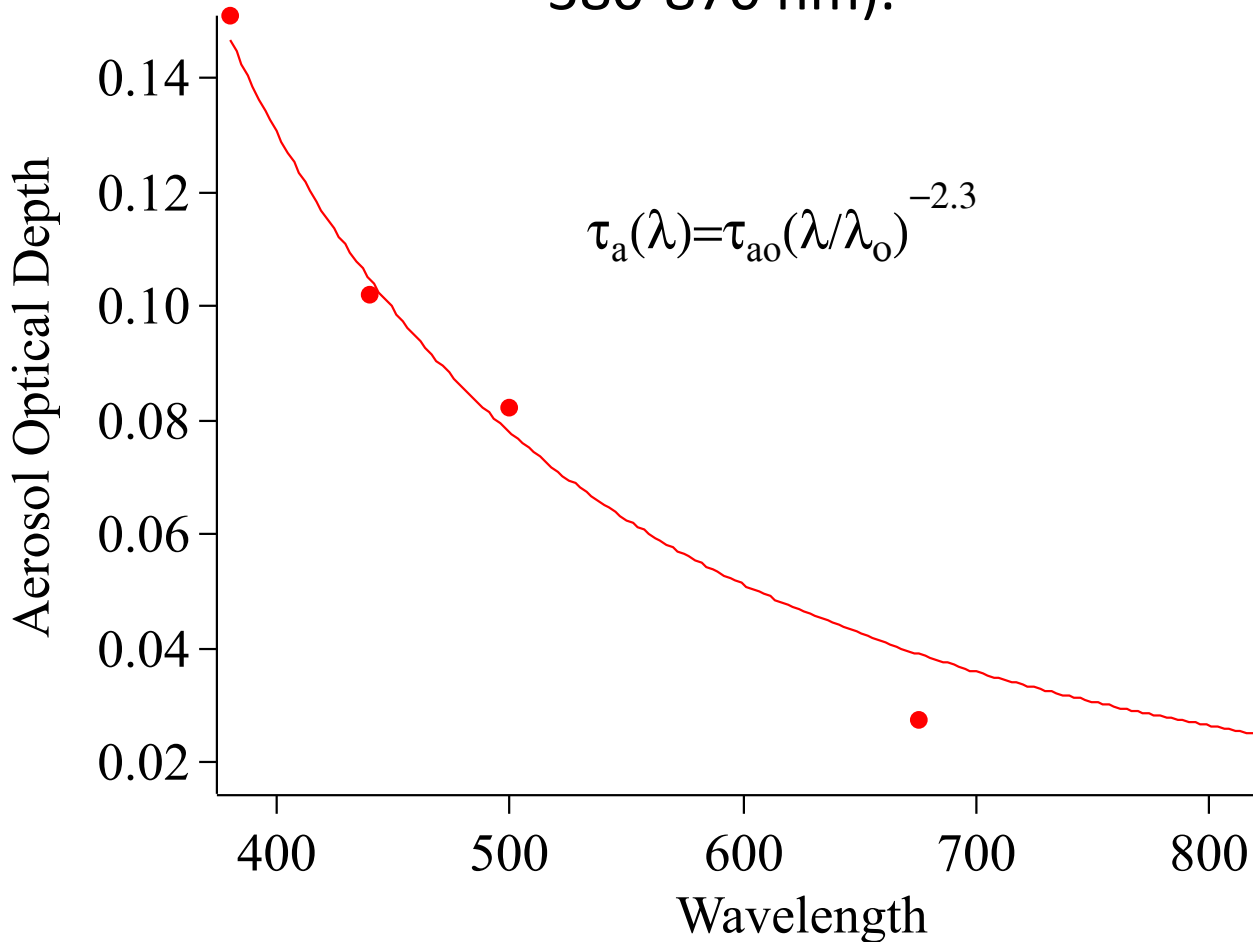
# Ozone and Rayleigh



# Your data



Expect aerosol to follow angstrom law, with angstrom coeff. around 1. (lower often for maritime atmosphere and larger in polluted atmospheres...often follows different angstrom exponent over different ranges...so need to specify (Below is 380-870 nm).



# For completeness, NO<sub>2</sub>

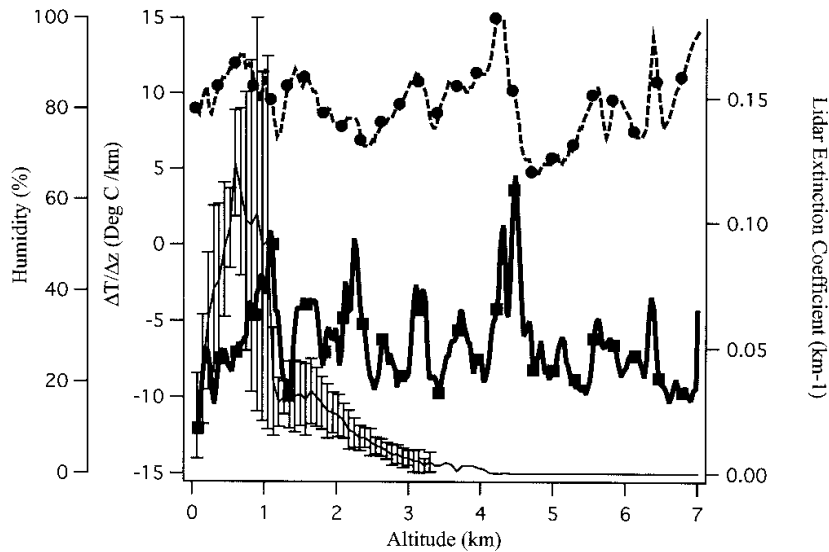
- Shaw, 1976..discusses in respect to optical depth
- Peak at 390 nm, falls off to half this value at 325 nm and 480 nm
- Shaw estimated the optical depth could vary between 0.008 and 0.087 at 390 nm for values of NO<sub>2</sub> of  $5 \times 10^{-3}$  to 0.4 atm-cm (note 0.4 atm-cm = 400 DU,  $5 \times 10^{-3}$  is 5 DU).

More recent data (derived from AERONET data) From Rublev et al 2014.

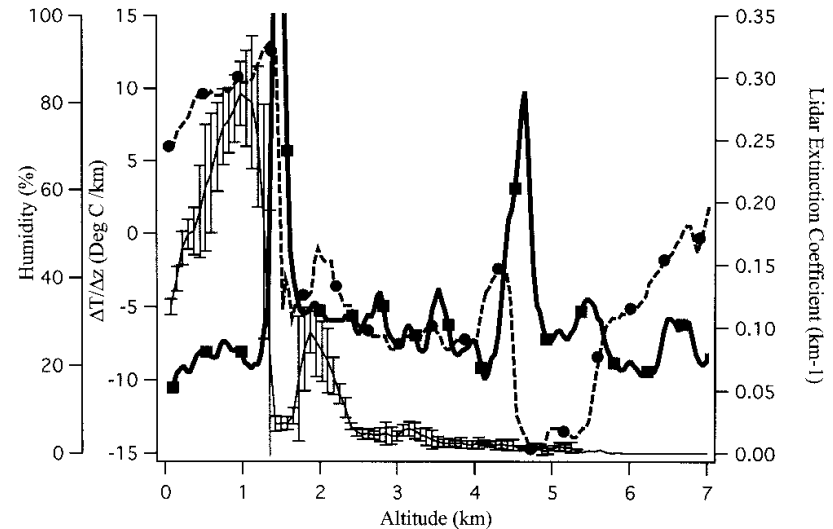
<b>Table 1. Types of Regions in Dependence on Column NO<sub>2</sub></b>						
#	Region type	Description and Examples	Q <sub>NO<sub>2</sub></sub> , DU	τ <sub>NO<sub>2</sub></sub> (380 nm)	F <sub>NO<sub>2</sub></sub> , W/m <sup>2</sup> Sum. Wint.	
1	Oceanic and Remote - continental	Without own sources, no any NO <sub>2</sub> advection due to transfer within troposphere (East Pacific; Northern Caucasia; Oklahoma, the US)	0.1 ÷ 0.4	0.002 ÷ 0.007	0.5 ÷ 1.9	0.7 ÷ 2.6
2	Continental	Without own anthropogenic sources, some NO <sub>2</sub> advection from remote industrial areas; (Northern Europe; some parts of Russia)	0.4 ÷ 0.8	0.007 ÷ 0.014	1.9 ÷ 3.8	2.6 ÷ 5.2
3	Suburban	Without own powerful anthropogenic sources of NO <sub>2</sub> , but a site is nearby the air plume of city or big industrial area (Northern part of West Siberia, nearby Norilsk steel mill)	0.8 ÷ 3.0	0.014 ÷ 0.051	3.8 ÷ 13.4	5.2 ÷ 18.5
4	Urban	There are own powerful sources of NO <sub>2</sub> (Moscow; Hamilton)	1 ÷ 15	0.017 ÷ 0.253	4.7 ÷ 58.7	6.5 ÷ 78.6



# Extinction in the atmosphere



**Figure 7.** Radiosonde and lidar-derived profiles typical for region 2, Northern Hemisphere clean (31°N–15.5°N). This specific case was for DOY 18.5, 27.7°N. LIDAR extinction is shown as the fine line with no symbols. This is an average of the nearest three profiles (each of which is a 10 min cloud-free

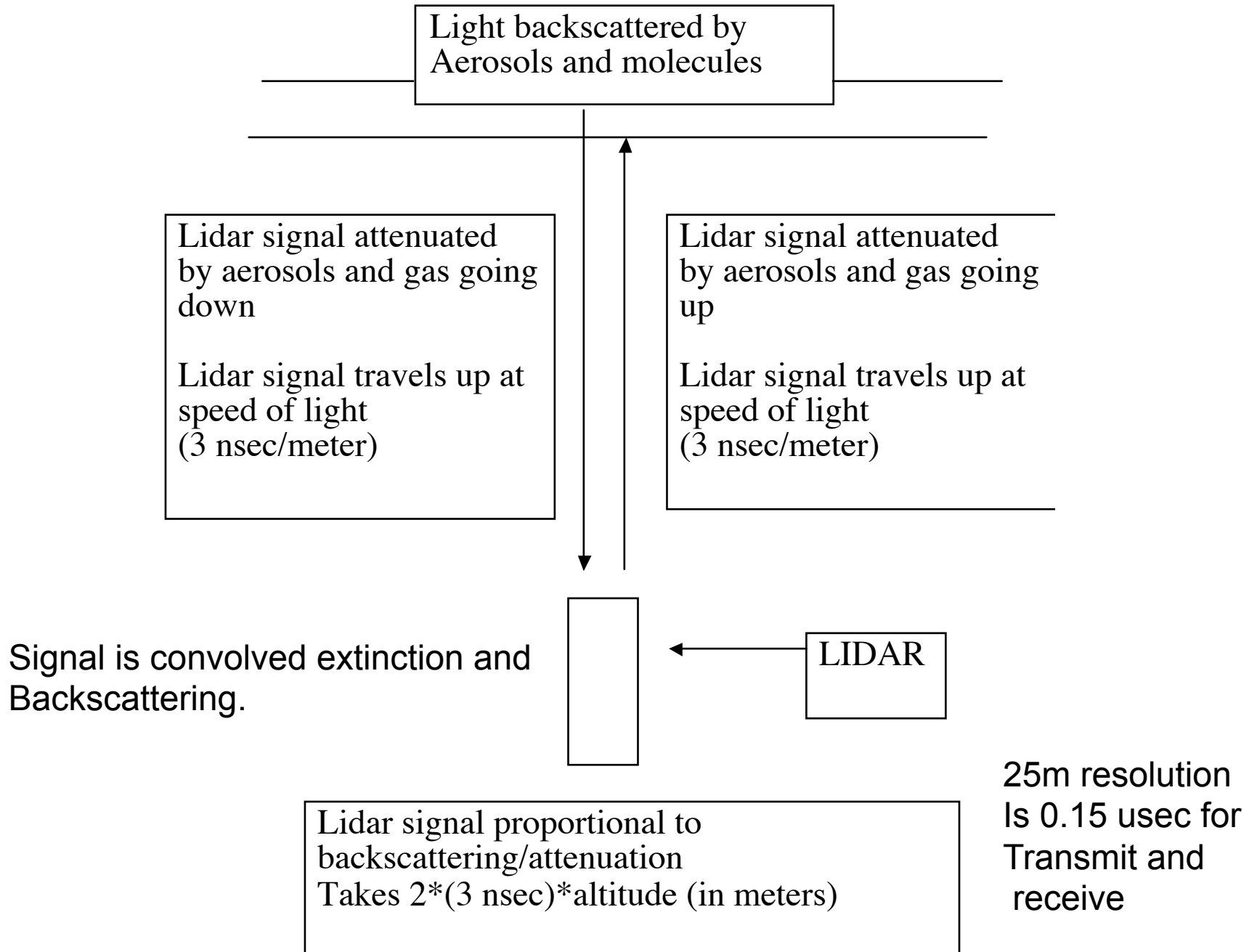


**Figure 8.** Region 3: dust (15.5°N–8°N). This specific case was for DOY 24.5, 11.2°N. Symbols and error bars are as in Figure 7. Extinction has increased from region 2, and there is another aerosol layer above the first temperature inversion (positive  $\Delta T/\Delta z$ ).

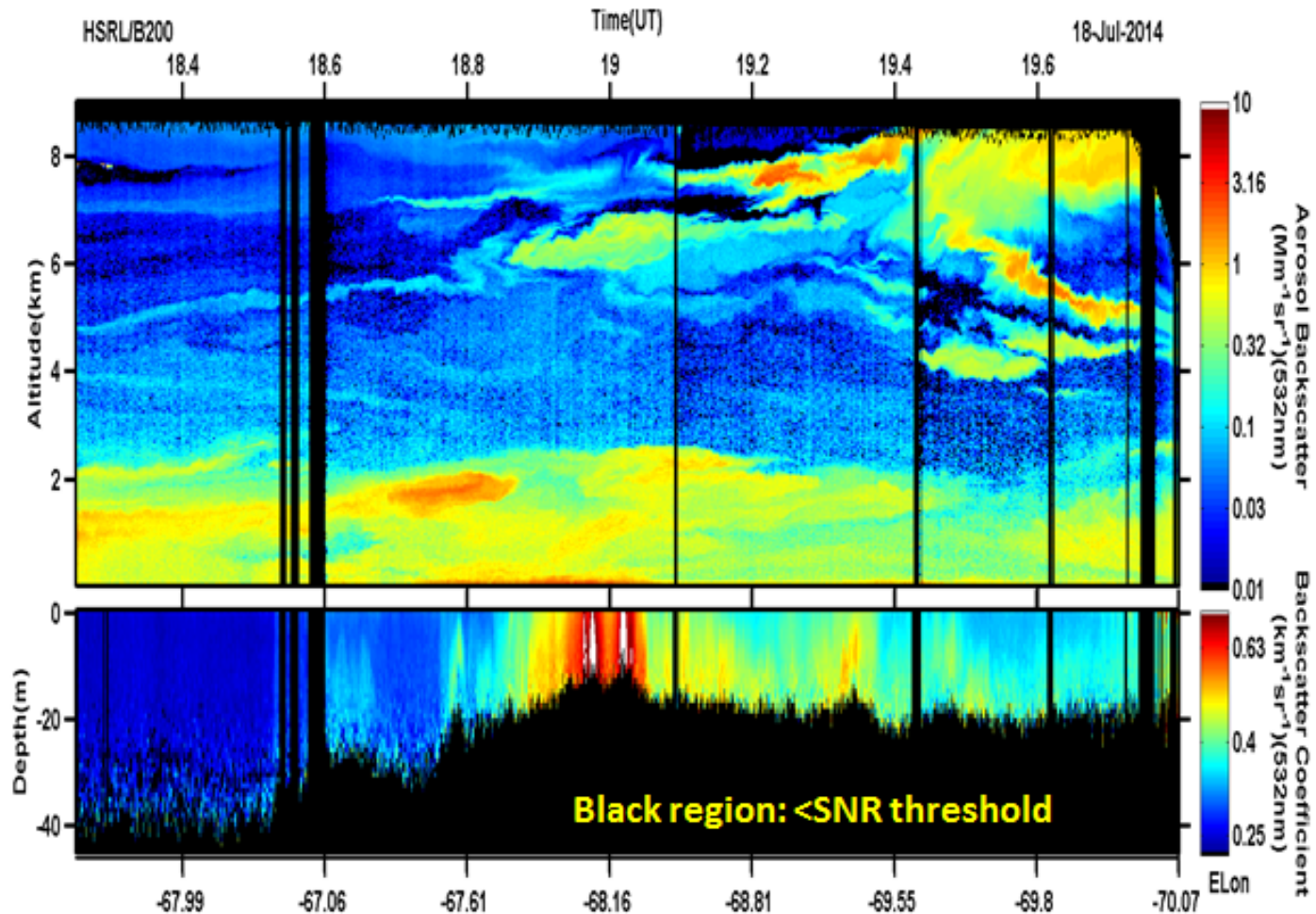
To get frame of reference, extinction is the thin line, note units...km-1! Low density allows sky radiance to contain more information (more SS, less multiple scattering).

# Methods to measure extinction

- Long pathlength (between mountains, etc) beam transmissometers.
- Lidars :Light Detection and Ranging...sort of like Radars....



# Lidars are hot topic...from SABOR, High Resolution Spectral Lidar (HRSL)

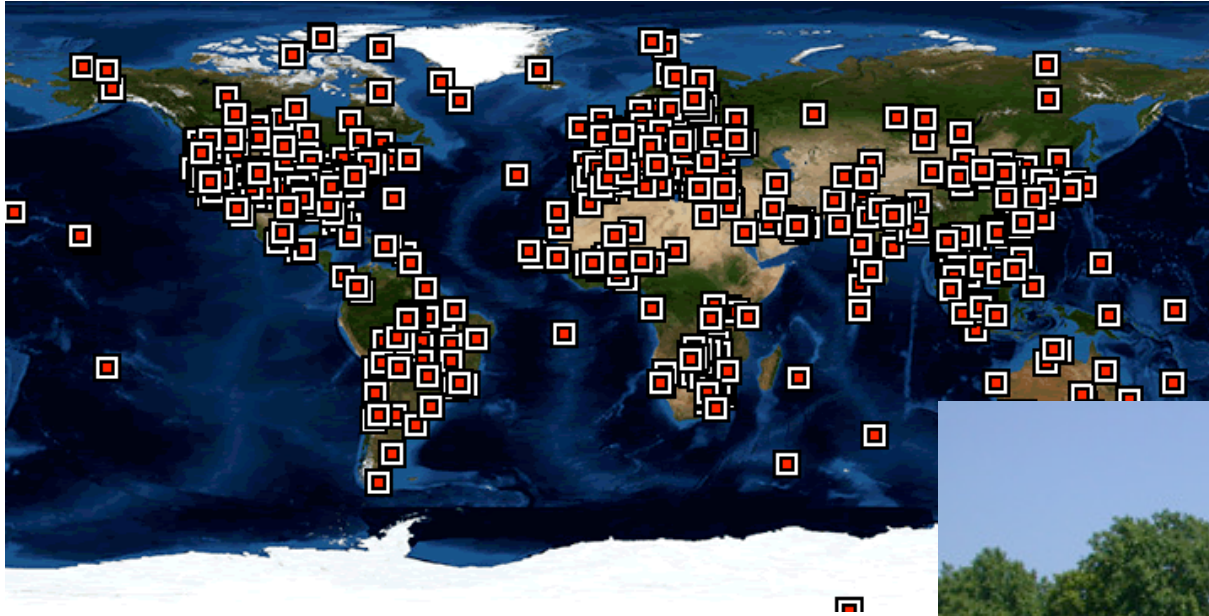


# Aerosol Absorption....really hard to measure (think about value of extinction)

- Techniques:
- Capture aerosols on filters using high volume (filter all day) and measure reflectance, or transmittance through filter....
- Measure direct/diffuse sky irradiance and calculate missing part in diffuse.....
- Ring lasers (absorbing cavity inside laser)
- Photoacoustics

Lots of variability between techniques.....really a relative mess

# AERONET (Brent Holben, GSFC/NASA)



From  
[Aeronet.gsfc.nasa.gov](http://Aeronet.gsfc.nasa.gov)

Note....not all stations are active

# Based on CIMEL sunphotometers

- Standard:  
1020-870-675-440-936-500-  
340-380 nm channels
- Measures direct solar irradiance and almucantor and principal plane sky radiance



# Measurement frequency

- Direct solar irradiance at 0.25 AM intervals at large solar zenith angles, and 15 minute intervals at small solar zenith angles (high sun) with the 8 wavelengths (to get spectral aerosol optical depth)
- Principal plane and almucantor measurements at 440, 670, 870 and 1020 nm at air mass 4,3,2,1.7 in morning and afternoon.



# Sky radiance inversions

- Does simultaneous inversion using AOD( $4 \lambda$ ) and sky radiance along almucantor at all  $4 \lambda$ .
- If solar zenith angle is 60 deg, max scattering angle is 120 deg.

Dubovik and King (2000)

Retrieves,  $m$  ( $1.33 < n < 1.6$ ,  $0.0005 < k < 0.5$ )

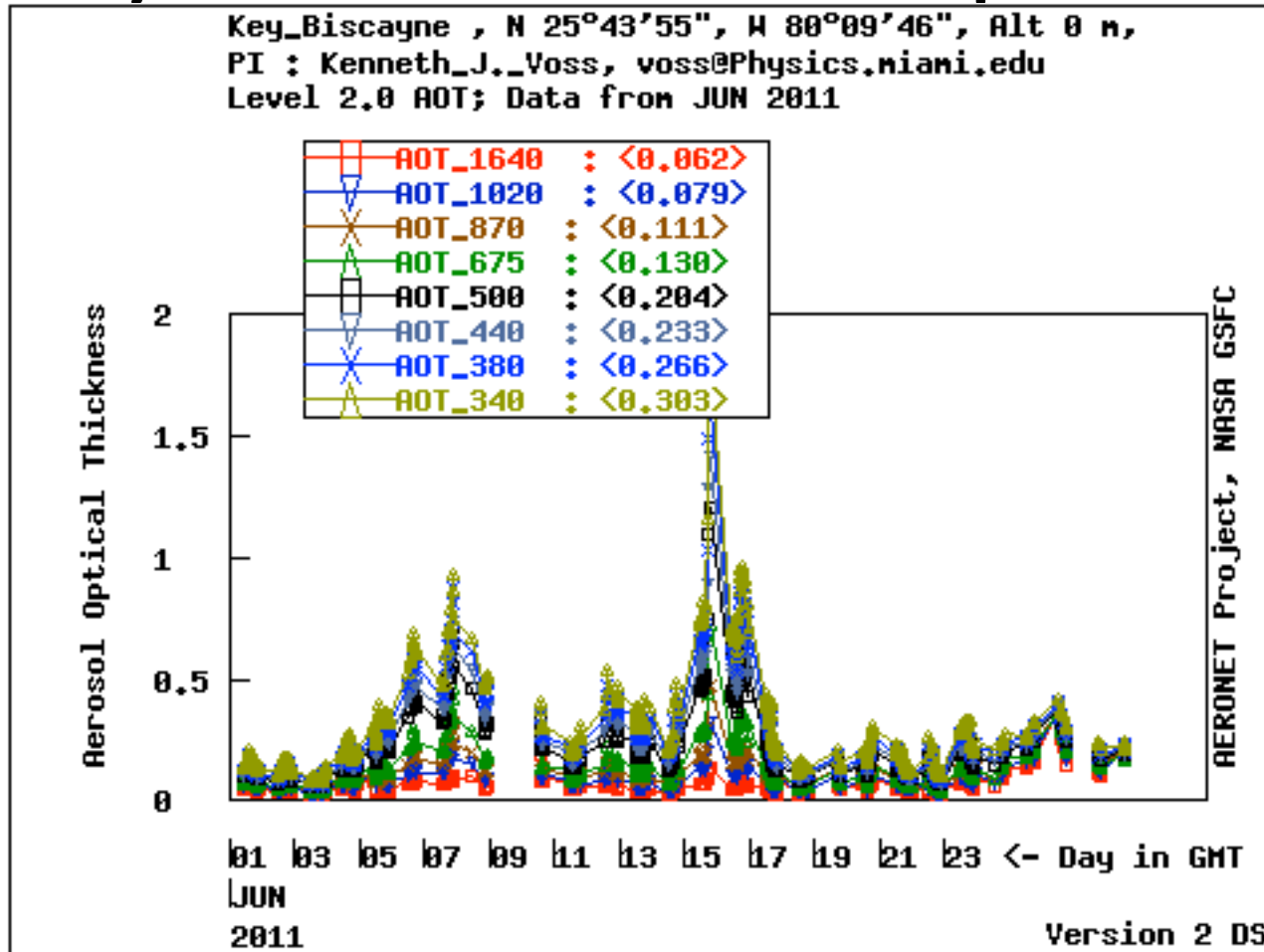
$0.05 < r < 15 \text{ um}$ , 22 sizes of  $dV/d\ln(r)$ ,  $\Delta\ln(r)$   
constant

# Estimated inversion accuracy (version 1)

**Table 4.** Errors in the Size Distribution, Complex Refractive Index, and Single-Scattering Albedo

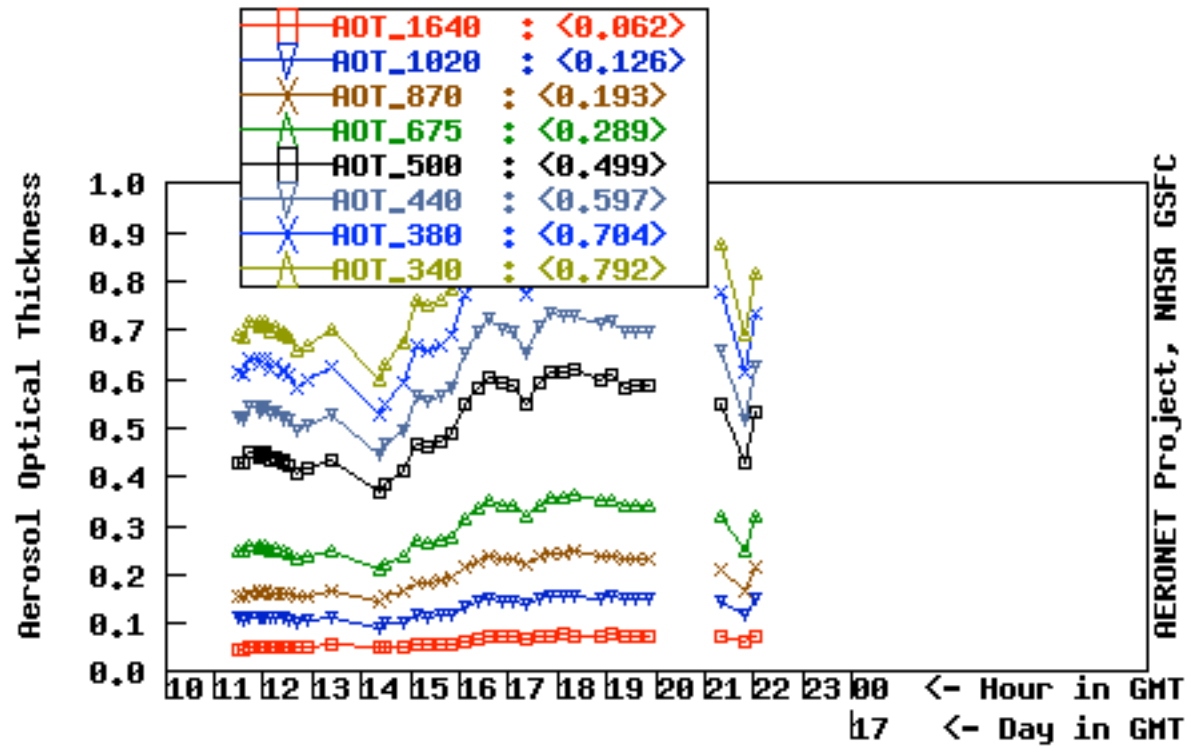
	Water-Soluble	Dust	Biomass Burning
$dV/d \ln r(r_i)$ , %			
$0.1 \mu\text{m} < r < 7 \mu\text{m}$	15	35	25
$r < 0.1 \mu\text{m}$ and $r > 7 \mu\text{m}$	15–100	35–100	25–100
$n(\lambda)$			
$\tau_a(440) \leq 0.2$	0.05		
$\tau_a(440) > 0.2$	0.025		
$\tau_a(440) \geq 0.5$		0.04	0.04
$k(\lambda)$			
$\tau_a(440) \leq 0.2$	80–100%		
$\tau_a(440) > 0.2$	50%		
$\tau_a(440) \geq 0.5$		50%	30%
$\omega_0(\lambda)$			
$\tau_a(440) \leq 0.2$	0.05–0.07		
$\tau_a(440) > 0.2$	0.03		
$\tau_a(440) \geq 0.5$		0.03	0.03

# Example inversions (from aeronet.gsfc.nasa.gov)...note always use Level 2.0 if possible!



# During Day

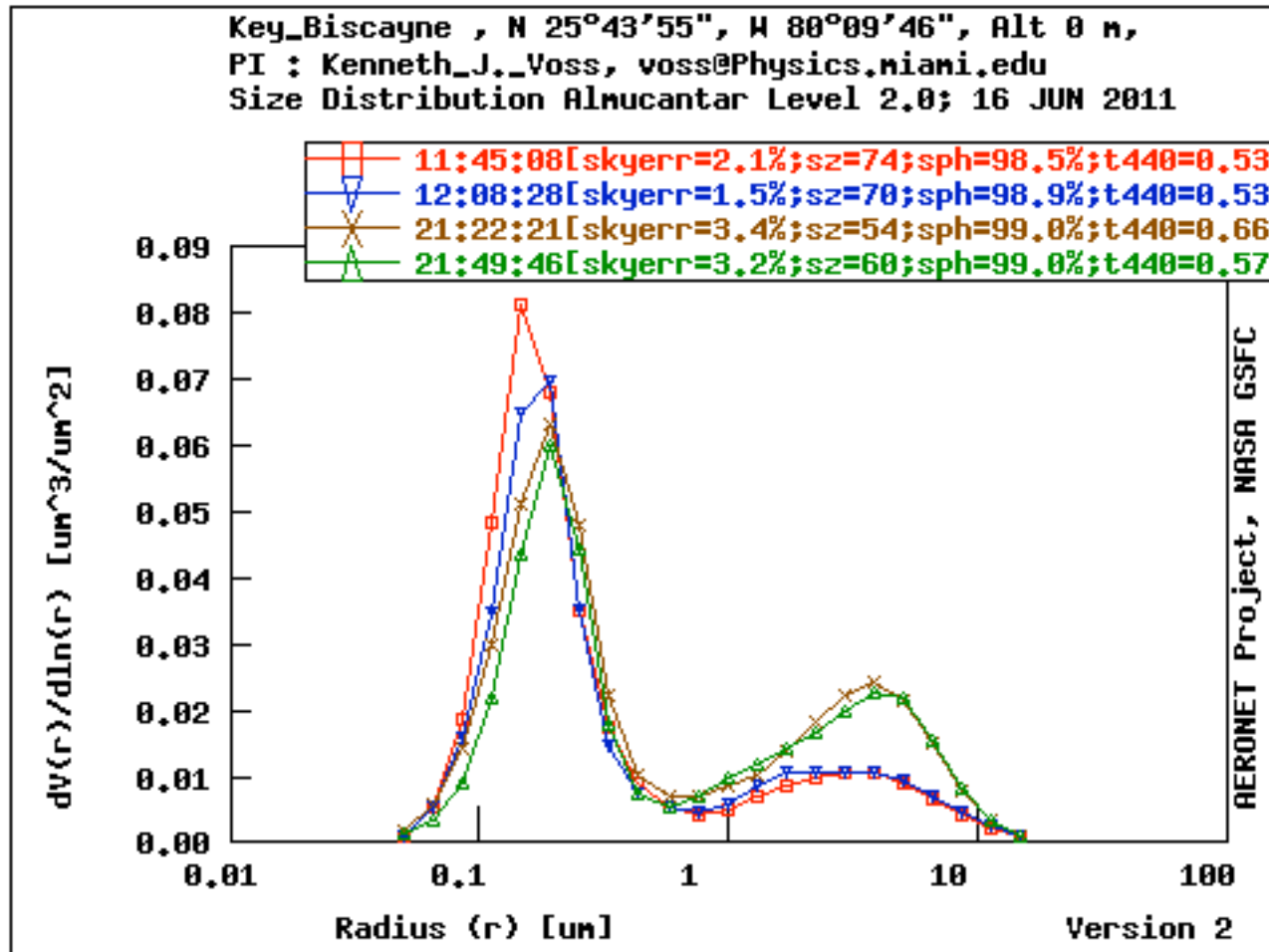
Key\_Biscayne , N 25°43'55", W 80°09'46", Alt 0 m,  
PI : Kenneth\_J.\_Voss, voss@Physics.miami.edu  
Level 2.0 AOT; Data from 16 JUN 2011



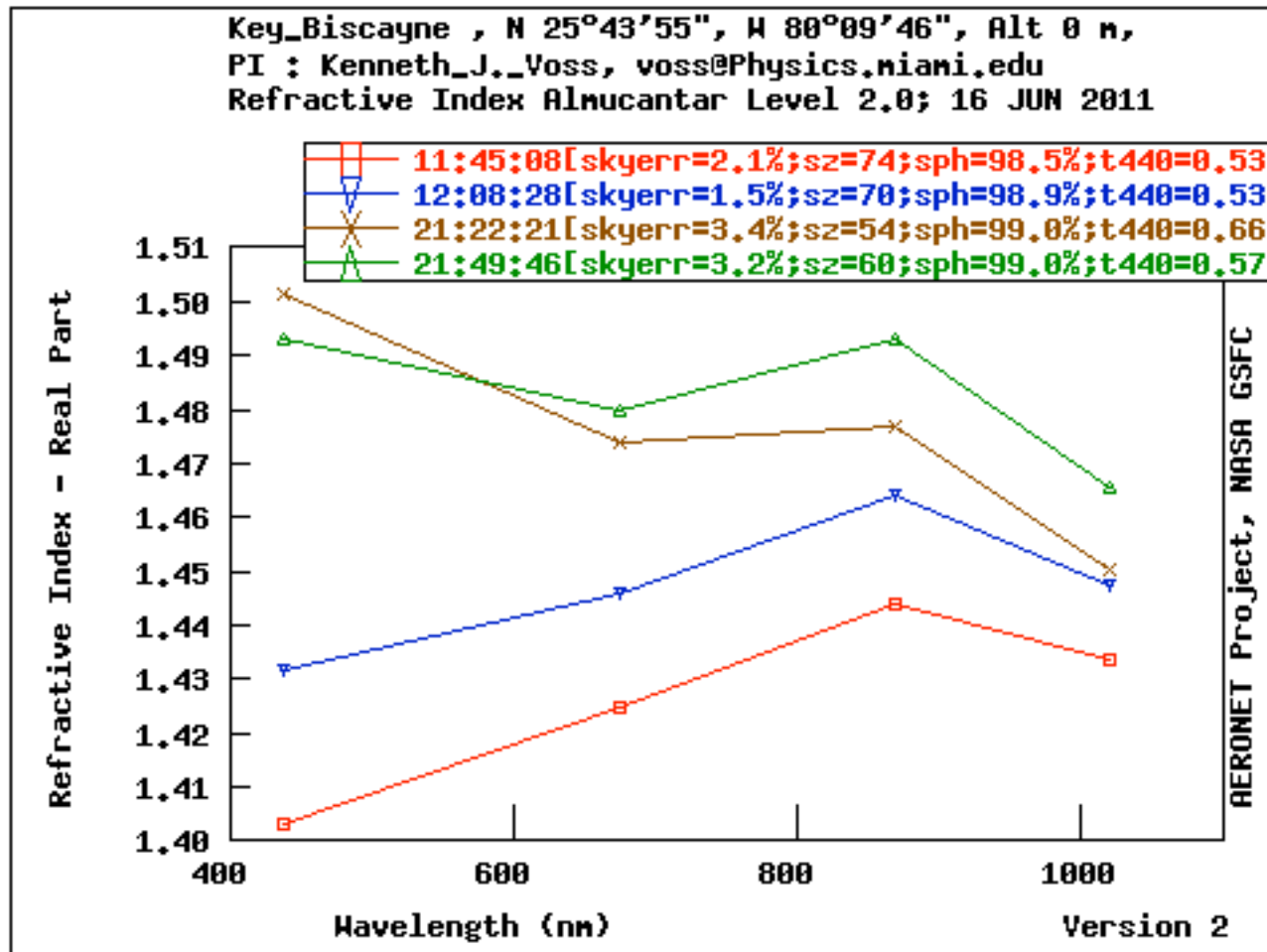
JUN  
2011

Version 2 DS

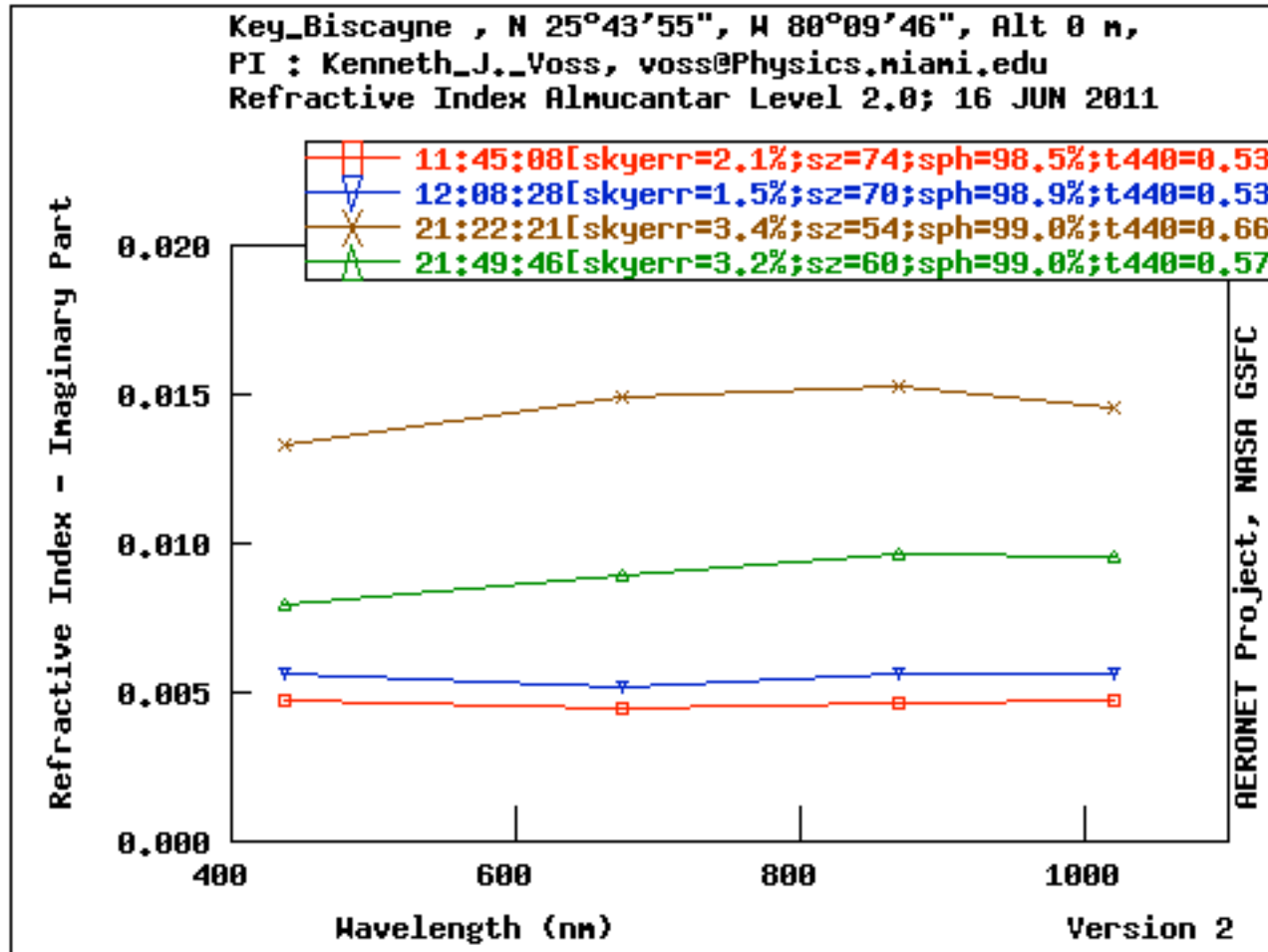
# size



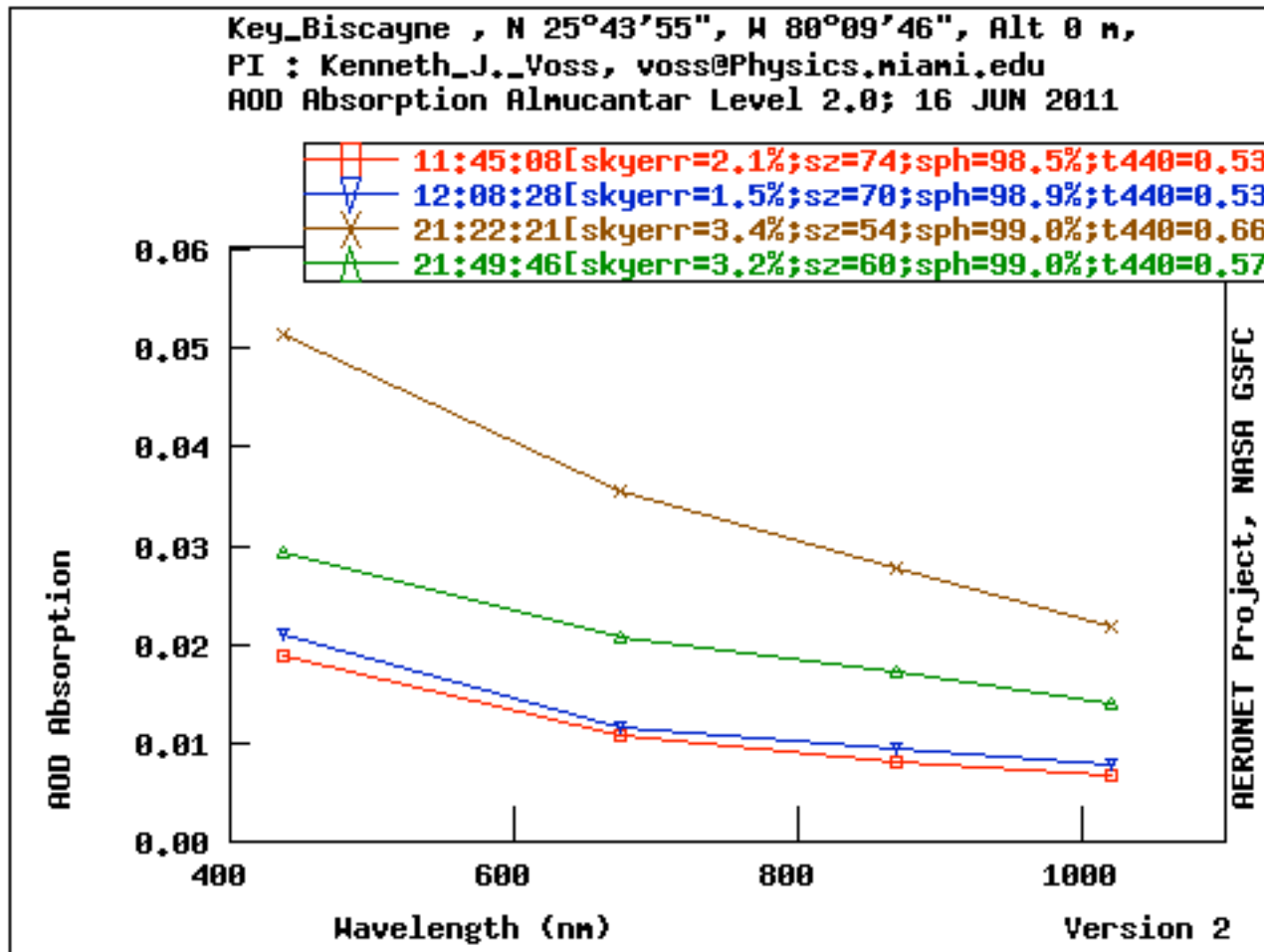
# Real index of refraction (remember accuracy 0.05)



# Imaginary index of refraction accuracy 50%

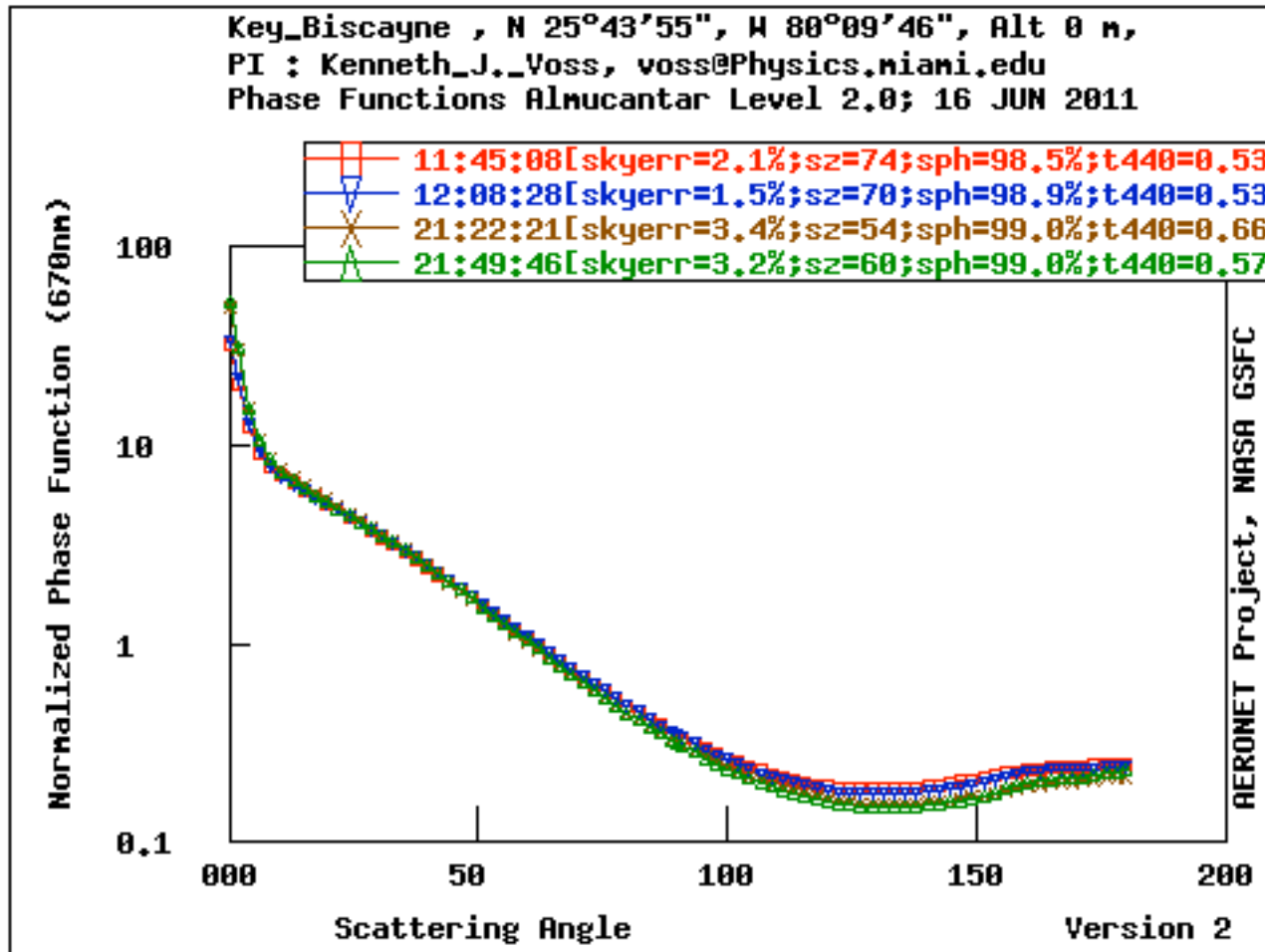


# Absorption optical depth

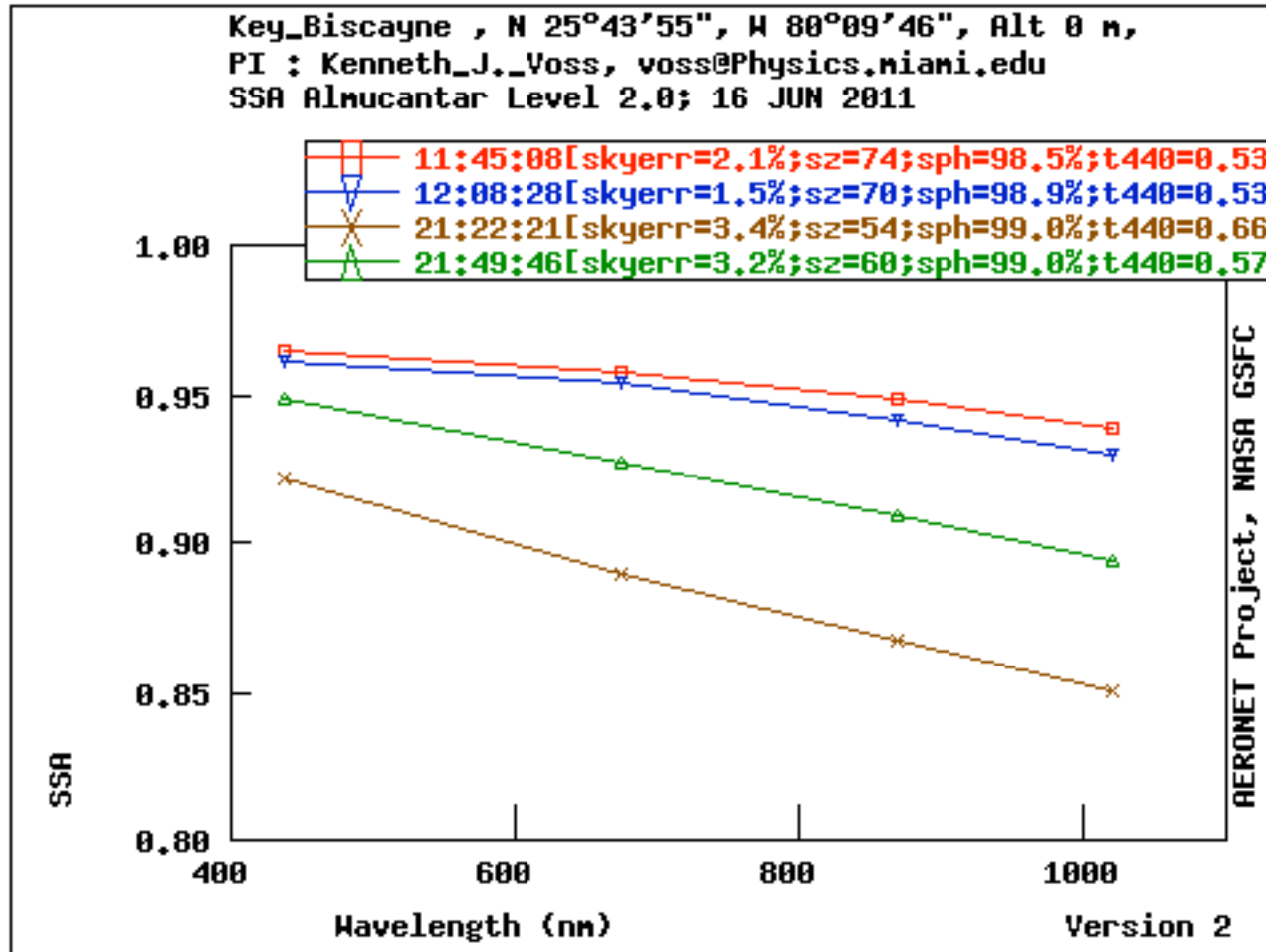




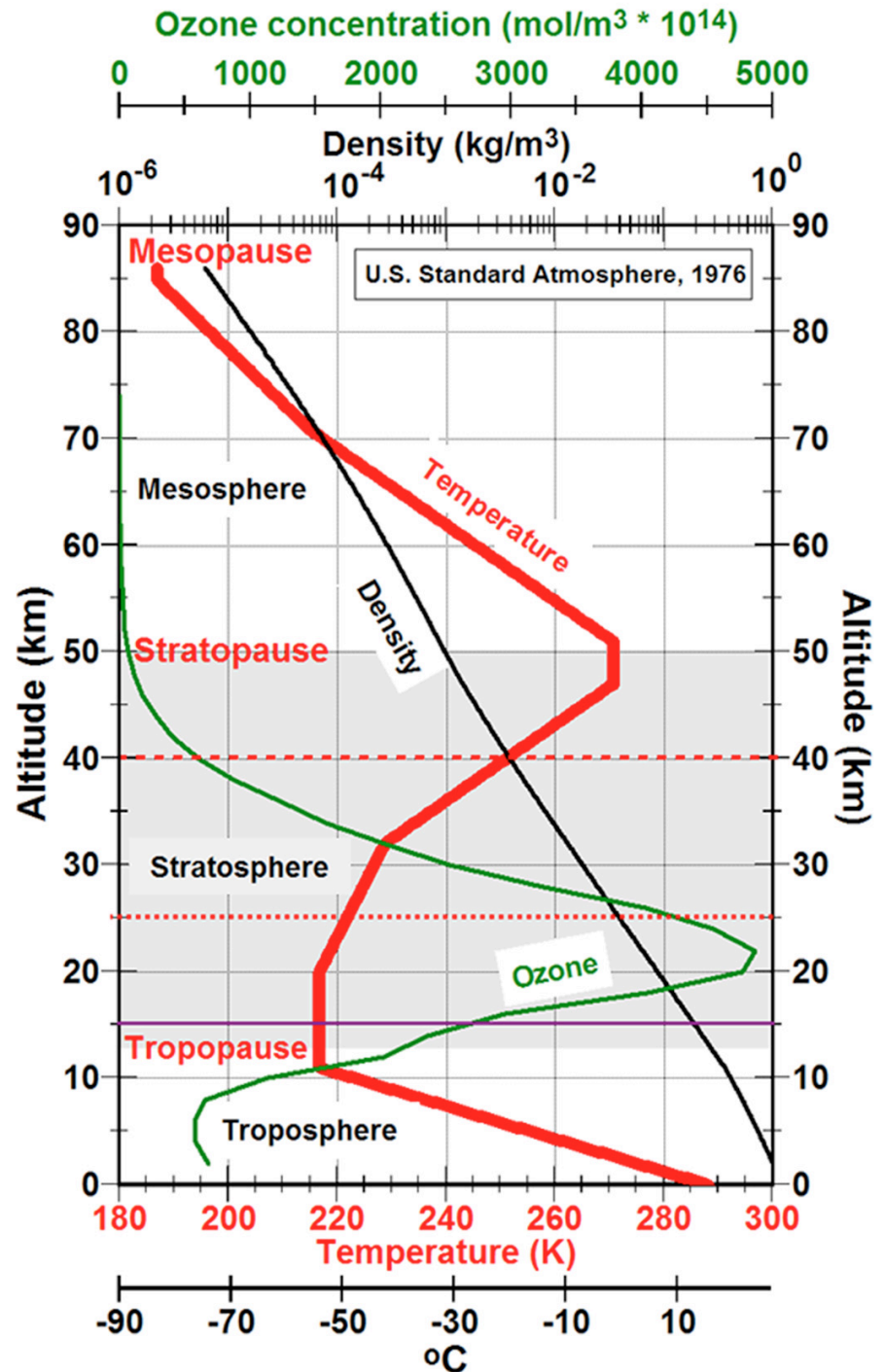
# Phase function (at 670 nm)



# Single scattering albedo



Why do we worry about absorbing aerosols for Atmospheric Correction? Rayleigh backscattering proportional to density.



# How do we currently handle vertical structure?

- Typical...2 layers...Rayleigh overlaying aerosols (aerosols packed in bottom).
- There is a rayleigh-aerosol interaction term....
- But lots of work has shown that for non absorbing aerosols vertical structure doesn't matter....
- But what about absorbing aerosols?

- For absorbing aerosols matters where they are relative to the molecular scattering optical depth.
- How about absorbing gasses?
  - Ozone above aerosols and much of molecular scattering.
  - NO<sub>2</sub>, mostly tropospheric....hence the problem....

All the techniques I know of for absorbing aerosols are spectral optimization techniques of some sort... i.e. require a model ocean and models for atmosphere. Problem in polluted coastal regions, or dusty regions.