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Comparison of in situ and columnar aerosol spectral measurements during TexAQS-GoMACCS 2006: testing parameterizations for estimating aerosol fine mode properties

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During the 2006 Texas Air Quality Study and Gulf of Mexico Atmospheric Composition and Climate Study (TexAQS-GoMACCS 2006), the optical, chemical and microphysical properties of atmospheric aerosols were measured on multiple mobile platforms and at ground based stations. In situ measurements of the aerosol light extinction coefficient (σ_{ep}) were performed by two multi-wavelength cavity ring-down (CRD) instruments, one located on board the NOAA *R/V Ronald H. Brown* (RHB) and the other located at the University of Houston, Moody Tower (UHMT). An AERONET sunphotometer was also located at the UHMT to measure the columnar aerosol optical depth (AOD). The σ_{ep} data were used to extract the extinction Ångström exponent (\mathring{a}_{ep}), a measure of the wavelength dependence of σ_{ep} . There was general agreement between the \mathring{a}_{ep} (and to a lesser degree σ_{ep}) measurements by the two spatially separated CRD instruments during multi-day periods, suggesting a regional scale consistency of the sampled aerosols. Two spectral models are applied to the σ_{ep} and AOD data to extract the fine mode fraction of extinction (η) and the fine mode effective radius ($R_{eff,f}$). These two parameters are robust measures of the fine mode contribution to total extinction and the fine mode size distribution respectively. The results of the analysis are compared to $R_{eff,f}$ values extracted using AERONET V2 retrievals and calculated from in situ particle size measurements on the RHB and at UHMT. During a time period when fine mode aerosols dominated the extinction over a large area extending from Houston/Galveston Bay and out into the Gulf of Mexico, the various methods for obtaining $R_{eff,f}$ agree qualitatively (showing the same temporal trend) and quantitatively (pooled standard deviation=28 nm).

1 Introduction

In its most current report, the Intergovernmental Panel on Climate Change continues to emphasize the importance of quantifying the connections between aerosols and cli-

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mate with better precision and accuracy at local, regional and global scales (Andreae et al., 2004; Bates et al., 2006; Lohmann and Feichter, 2005). The accurate estimate of aerosol direct effects due to particle light extinction (σ_{ep}), scattering (σ_{sp}), and absorption (σ_{ap}), where extinction is the sum of scattering and absorption, requires knowledge of the physical and radiative properties as well as of the vertical distribution of the bulk aerosol on local to global scales. Cavity ring-down (CRD) spectroscopy (Moosmuller et al., 2005; Strawa et al., 2006; Atkinson, 2003; Baynard et al., 2007) offers robust and precise measurements of the in situ σ_{ep} coefficient. The unitless aerosol optical depth (AOD), the σ_{ep} integrated along the vertical column, is a key parameter for determining the radiative effects of aerosols (Yu et al., 2006). The AOD is measured on a worldwide scale by networks of ground based spectro-radiometers and sunphotometers, such as the Aerosol Robotic Network (AERONET; (Holben et al., 1998); <http://aeronet.gsfc.nasa.gov>).

The wavelength dependence of σ_{ep} can be expressed by the Ångström exponent \mathring{a}_{ep} which can then be used to infer information about particle size: values of \mathring{a}_{ep} near to or less than zero pertain to large (coarse mode) particle sizes, while larger values are produced by fine mode particles. The \mathring{a}_{AOD} , obtained from AERONET sunphotometers, has a similar definition and properties to the in situ \mathring{a}_{ep} but is averaged along the vertical path. Recently, satellite retrieval algorithms have been developed to extract and quantify the contribution of the fine mode fraction to the column AOD (Anderson et al., 2005). Such information is critical for models to calculate the climate forcing by anthropogenic aerosols (Chin et al., 2004; Kleidman et al., 2005; Bates et al., 2006; Yu et al., 2006). However, due to the generally high uncertainties of satellite retrievals, those methods are less effective unless supported by in situ measurements.

This work examines the extent to which the spectral information from both in situ and remote instruments can be used to provide information about the fine aerosol mode in the vertical column. We process the σ_{ep} data collected by a three-wavelength cavity ring-down instrument (hereafter the NOAA CRD) onboard the *R/V Ronald H. Brown* (RHB) using two spectral analysis methods to quantify the relative contribution of the

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fine aerosol mode to the total σ_{ep} (or the AOD in the vertical column), referred to as the fine mode fraction of extinction (η). These methods also extract the effective radius $R_{\text{eff},f}$ of the fine mode aerosol, which is used to characterize the size distribution of the fine particle mode. The general definition of $R_{\text{eff},f}$ is given by Hansen and Travis (1974);

$$R_{\text{eff}} = \frac{\int_0^{\infty} r \pi r^2 \frac{dN}{d \ln r} d \ln r}{\int_0^{\infty} \pi r^2 \frac{dN}{d \ln r} d \ln r} \quad (1)$$

where r is the particle radius and $dN/d \ln r$ is the particle size distribution function. Application of Eq. (1) to a fine mode component assumes that the fine mode sub-distribution can be isolated or that one physically divides the radius regime into two segments, for example using a size selective inlet (we investigate both cases in this paper.) It has been suggested that variations in $R_{\text{eff},f}$ are indicative of oxidative aging or cloud processing of anthropogenic aerosols (usually fine mode particles) in the vertical column (Anderson et al., 2005). The $R_{\text{eff},f}$ values obtained from the spectral analyses of the CRD data are compared to those derived from sun/sky inversions using the AERONET V2 approach, and to those obtained from mobility size distribution instruments whose data were processed according to Eq. (1). Section 2 provides a detailed description of the experimental data and spectral models used in this study. In Sect. 3, comparisons of the extinction coefficient and \hat{a}_{ep} measurements from the two CRD instruments are used to investigate the spatial coherence of the aerosol air mass over the Houston/Galveston Bay area. The comparison of $R_{\text{eff},f}$ from the various methods is presented for a time period when long-range transported fine mode aerosols dominated the aerosol loading over a large area extending from Houston out in the Gulf of Mexico, and column AOD values measured on board the RHB were high (Massoli, et al., 2009). We show that the various methods for obtaining $R_{\text{eff},f}$ agree qualitatively and quantitatively, with a pooled standard deviation of 28 nm. Comparison of the \hat{a}_{ep} and η values derived from the CRD instruments is also shown for a time period when coarse aerosols originating from a long range transport of Saharan dust were present

over East Texas.

This work demonstrates that the different methods for obtaining $R_{\text{eff},f}$ show some skill in predicting aerosol physical properties. This implies that ground-based in situ optical measurements may be used in some situations to obtain information about the columnar properties of atmospheric aerosols for validating and constraining satellite retrievals when vertical distribution measurements are not available.

2 Experimental and theoretical methods

2.1 Aerosol sampling and measurement techniques

Continuous measurements of σ_{ep} were performed during TexAQS-GoMACCS 2006 using two custom CRD instruments operating on separate platforms. A first CRD from Portland State University (hereafter PSU CRD) was located at the 60 m high Moody Tower building on the University of Houston campus (UHMT) (Lefer and Rappenglück, 2008). The UHMT site housed a suite of instruments for aerosol, gas-phase and meteorological measurements as part of the TexAQS II Radical and Aerosol Monitoring Project (TRAMP). The area surrounding the site is not heavily urbanized, it is approximately one kilometer away from the nearest major traffic route to the Northeast, 20 km from the Houston ship channel (HSC) to the East/North-East, and 4.5 km from the urban core to the Northwest. Instruments were located inside temperature controlled ($24 \pm 2^\circ\text{C}$) trailers, and each instrument had its own sample inlet mounted 5 m above the roof top (~ 70 m a.s.l.) to minimize any ground or building interference. Wind speed and direction were measured by various instruments on the building and on the ground. The PSU CRD sampled aerosols through a cyclone impactor (URG Inc., 2000–30 EN), and 8 m. of copper tubing was used to transport the sample from the inlet to the CRD instrument. The cyclone has a nominal cut-point at $2.5 \mu\text{m}$ aerodynamic diameter at a flow rate of 10 l per minute (1 pm); however the aerosols were drawn into the PSU CRD at 5 lpm, giving a higher cut-point of $4 \mu\text{m}$, according to the URG product literature.

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The PSU CRD measured σ_{ep} at 532 and 1064 nm, with a maximum time resolution of 0.25 Hz, and a 95% confidence level detection limit of 4 Mm^{-1} based on previous laboratory data (Radney et al., 2008). The σ_{ep} results presented here are averaged to at least 30 min, resulting in a detection limit of less than 1 Mm^{-1} (Wright et al., 2009).

A Sequential Mobility Particle Sizer (SMPS) (GRIMM Technologies Inc.) was operated at the UHMT site throughout the project. The SMPS consists of a Differential Mobility Analyzer (Model DMA-L) and an Ultrafine Particle Counter (Model 5.403). The SMPS scanned the aerosol particle size distribution from 11.1 nm to 521 nm in 38 size bins approximately every 7 min for the majority of the study period. The size distribution data did not include corrections for variations in shape or density. In addition, a laser-based Aerosol Spectrometer (GRIMM Model 1.108 PAS) recorded a coarse mode particle distribution (15 size bins from $0.3 \mu\text{m}$ to $\sim 22 \mu\text{m}$) every 60 s. A thorough investigation of this instrument's inlet transmission as a function of particle size allowed for the correction of the raw data for particle losses, which were $\leq 10\%$ for sizes up to $4 \mu\text{m}$.

An AERONET CIMEL sun/sky radiometer was also located at the UHMT site and was continuously operated during the study period. Level 2.0 AERONET data (cloud screened and quality assured) are used in the analysis presented here. In addition to multi-wavelength AOD and pair-wise \hat{a}_{AOD} values, the AERONET V2 data products also include estimates of the particle size distribution and refractive index (Dubovik et al., 2002). This size distribution information is obtained through an inversion algorithm that operates on the full sky scan data (almucantars) combined with near-simultaneously acquired AOD data (Dubovik and King, 2000). The inversion fits the angular and spectrally resolved sky radiance and AOD data to radiative transfer computations derived for a variable (binned) size distribution and variable (pan-radius) refractive index (Dubovik and King, 2000). Non-spherical particle effects are accounted for by introducing a variable fraction of fixed-geometry spheroids (Dubovik et al., 2006; Sinyuk et al., 2007), (see http://aeronet.gsfc.nasa.gov/new_web/Documents/Inversion_products_V2.pdf). Data products of this method (hereafter referred to as “AERONET SSRI” for

“AERONET Sun-Sky Radiance Inversion”) also include the R_{eff} of the fine and coarse modes and η for the AOD.

Aerosols on the RHB were sampled through an inlet (mast) located 18 m above sea level and forward of the ship’s stack. The mast was automatically rotated into the relative wind to maintain nominally isokinetic flow and minimize the loss of supermicrometer particles. Air entered the inlet through a 5 cm diameter hole, passed through a 7° expansion cone, and then into the 20 cm inner diameter sampling mast. The flow through the mast was $1 \text{ m}^3 \text{ min}^{-1}$. The transmission efficiency of the inlet for particles with aerodynamic diameters less than $6.5 \mu\text{m}$ (the largest size tested) is greater than 95% (Bates et al., 2002). The bottom 1.5 m of the mast was heated to establish a stable reference RH for the sample air of $60 (\pm 5)\%$. On average, the aerosol was heated 2.5°C above ambient temperature. Stainless steel tubes extending into the heated portion of the mast were connected with conductive silicone tubing to downstream instrumentation for aerosol optical property measurements and particle sizing. The NOAA CRD sampled at a rate of 30 lpm off the mast and the light extinction coefficient σ_{ep} was measured at 355, 532, and 1064 nm in six independent cavities. The NOAA CRD (Massoli et al., 2009) measured the σ_{ep} coefficient for both the sub 1 and sub $10 \mu\text{m}$ sizes (aerodynamic diameter at 60% RH at mast outlet) and at different RH conditions (at 25%, 60% and 85% RH for the 532 nm, and at 25% and 85% for the 355 and 1064 nm). The relative uncertainty for σ_{ep} at any wavelength at 25% RH is approximately 1% for data averaged to 1 min, whereas the σ_{ep} coefficients measured at 85% RH are characterized by higher relative uncertainty (5%). The \dot{a}_{ep} values calculated from the 532 and 1064 nm σ_{ep} coefficients are characterized by an absolute uncertainty of 0.015 if estimated from the 25% RH σ_{ep} coefficients, and 0.07 when obtained for 85% RH conditions (Massoli et al., 2009).

On the RHB, size distributions from 20 to 200 nm and 200 to 800 nm in geometric diameter were measured with two parallel differential mobility particle sizers (DMPS, University of Vienna (Winklmayr et al., 1991) with short and medium length columns for Aitken and accumulation modes, respectively, coupled to condensation particle coun-

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ters (CPC model 3760A, TSI, Inc., St. Paul, MN.) Size distributions from 0.9 to 10 μm in aerodynamic diameter were measured with an aerodynamic particle sizer (APS model 3321, TSI). APS size distributions were converted from aerodynamic to geometric diameters using densities based on the measured chemical composition. More details regarding RHB sizing instruments and data can be found in (Bates et al., 2008; Quinn et al., 2008).

It should be pointed out that although the expressions “fine mode” and “submicron aerosols” are often treated as if they were interchangeable (likewise for “coarse mode” and “supermicron aerosols”) we are aware of the substantial differences regarding how the various measurements and methods discussed in this paper obtain the size information: the in situ instruments base such distinction on the aerodynamic particle diameter, usually determined through impaction (Bernier et al., 1979). In these cases, the fine mode fraction corresponds to a sub 1 μm fraction resulting from an 50% aerodynamic cut-off diameter within the impactor of 1 μm (Doherty et al., 2005; Yu et al., 2006) whereas all of the spectral methods base such distinction on the different optical behavior of small and large particles.

2.2 Data analysis and spectral methods

The $\hat{a}_{ep}(\lambda_1, \lambda_2)$ can be calculated using σ_{ep} values at two different wavelengths (λ_1 and λ_2) according to the logarithmic treatment of the assumed power-law expression (Ångström, 1929)

$$\hat{a}_{ep}(\lambda_1, \lambda_2) = -\ln(\sigma_{ep,\lambda_1}/\sigma_{ep,\lambda_2})/\ln(\lambda_1/\lambda_2) \quad (2)$$

The PSU CRD σ_{ep} coefficients measured at 532 and 1064 nm were used to obtain an estimate of \hat{a}_{ep} for the UHMT site (hereafter $\hat{a}_{ep,\text{UHMT}}$). Similarly, three different \hat{a}_{ep} values can be obtained pair-wise from the NOAA CRD wavelengths 355, 532 nm, 355, 1064 nm, and 532, 1064 nm (hereafter $\hat{a}_{ep,\text{RHB}}$). Direct comparison between the $\hat{a}_{ep,\text{UHMT}}$ and $\hat{a}_{ep,\text{RHB}}$ (532, 1064 nm) for specific time periods during TexAQS-GoMACCS is presented in Sect. 3.

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Next we describe the spectral methods applied to the NOAA CRD and AERONET data to extract the fine mode fraction of extinction η and the fine mode effective radius $R_{\text{eff},f}$: the Spectral Deconvolution Algorithm with Fine Mode Curvature (SDA/FMC) and the Graphical Spectral Method (GSM). The PSU CRD data were not processed in this way because application of the spectral methods requires data at 3 or more wavelengths.

The SDA/FMC approach (O'Neill et al., 2003, 2005) uses the AOD directly measured by the sunphotometer to extract η and the Van de Hulst phase shift parameter $\rho_{\text{eff},f}$ (if a particle refractive index is assumed then $\rho_{\text{eff},f}$ can be converted to an estimate of $R_{\text{eff},f}$, cf. Eq. (8) below). This spectral model takes advantage of the higher data density of the direct solar attenuation AOD measurements compared to the full sky scans (i.e. AERONET SSRI retrievals), especially with a new high frequency instrument mode which permits the acquisition of AOD data at 3 minute intervals.

The SDA/FMC procedure starts with a second order fit of $\ln(\sigma_{ep})$ vs. $\ln(\lambda)$

$$\ln(\sigma_{ep}(\lambda)) = a_0 + a_1 \ln(\lambda) + a_2 \ln(\lambda)^2. \quad (3)$$

The three fitting coefficients a_0 , a_1 and a_2 can be used to calculate the $\lambda=1 \mu\text{m}$ extinction coefficient ($\sigma_{ep}=a_0$) as well as the extinction Ångström exponent α_{ep} at any selected wavelength, and its wavelength dependence, the “spectral curvature” α'_{ep}

$$\alpha_{ep}(\lambda) = -d \ln(\sigma_{ep}) / d \ln(\lambda) = -a_1 - 2a_2 \ln(\lambda) \quad (4)$$

$$\alpha'_{ep} = d \alpha_{ep}(\lambda) / d \ln(\lambda) = -d^2 \ln(\sigma_{ep}) / d \ln(\lambda)^2 = -2a_2 \quad (5)$$

Note that although we will refer to both \hat{a} and α as Ångström exponents, they are subtly different in definition and usage: \hat{a} is an average value for a wavelength range with a proscribed lack of spectral curvature, whereas α is specific to one wavelength (in the sense of a pure derivative); its formulation accounts for higher order spectral curvature, and exploits it to extract further information about the fine and coarse modes. The spectral curvature α'_{ep} value of a fine-mode dominated distribution increases as

$R_{\text{eff},f}$ increases from 10 nm to 500 nm, and then decreases for fine-mode particles with radius >500 nm (O'Neill et al., 2005; Schuster et al., 2006). For coarse-mode dominated aerosols α'_{ep} is ≤ 0 . The α_{ep} and α'_{ep} calculated from both the NOAA CRD σ_{ep} and the UHMT AERONET sunphotometer AOD data are used to extract η and the fine mode Ångström parameters at a reference wavelength of $\lambda=500$ nm. This information is then used in the FMC procedure to extract the fine mode $\rho_{\text{eff},f}$. The FMC inversion is based on the observation that the value of $\rho_{\text{eff},f}$ is a function of the fine mode Ångström exponent $\alpha'_{ep,f}$ and its spectral derivative $\alpha'_{ep,f}$

$$\rho_{\text{eff},f} = 0.31593 + 1.25050\psi + 0.52859\psi^2 \quad (6)$$

where ψ is in radians (Eq. (5) in O'Neill et al., 2005 was incorrectly written and has since been slightly modified to correct for higher-order-polynomial artifacts at the extremes of the ψ range – O'Neill et al., 2008). From O'Neill et al. (2005) we have;

$$\tan \psi = \frac{\alpha'_f - 0.2}{\alpha_f - 0.5}. \quad (7)$$

The parameters α_f and α'_f (outputs of the coarse/fine mode separation in the SDA procedure) are typically characterized by relatively large errors which propagate into the calculation of $\rho_{\text{eff},f}$. We have found that these errors are difficult to quantify empirically. They are a significant fraction of the amplitude of $\rho_{\text{eff},f}$ but tend to decrease as $\eta \rightarrow 1$. Once the $\rho_{\text{eff},f}$ has been estimated, the $R_{\text{eff},f}$ can be calculated according to

$$\rho_{\text{eff},f} = 2(2\pi R_{\text{eff},f}/\lambda) |m - 1| \quad (8)$$

where λ and m are the reference wavelength and the complex refractive index, respectively. We obtain two $R_{\text{eff},f}$ values extracted from the SDA/FMC method, one using the AERONET AOD data (hereafter AERONET SDA/FMC) and one from the NOAA CRD data (hereafter CRD SDA/FMC). For the NOAA CRD, only the data collected at 85% RH were used to extract $R_{\text{eff},f}$ via the SDA/FMC approach to best match the

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sampling conditions at ambient RH of both PSU CRD and AERONET data. The RH measurements on the RHB showed that the ambient RH measured on RHB was between 80 and 85% during most of the study period.

The GSM approach (Gobbi et al., 2007) uses a graphical parameterization of the AERONET products to obtain $R_{\text{eff},f}$. The method relies on AOD data from three CIMEL data channels only (unlike the SDA, which uses as many wavelengths as are available) and the data is first screened to remove values below AOD=0.1. The discrete difference of specified pair-wise \hat{a}_{AOD} values computed from AERONET sunphotometer data is used to obtain $\delta \hat{a}_{\text{AOD}}$

$$\delta \hat{a}_{\text{AOD}} = \hat{a}_{\text{AOD}}(440, 675) - \hat{a}_{\text{AOD}}(675, 875) \quad (9)$$

which is also referred to as “spectral curvature”. Due to a different choice of definitions, the analogous parameter in the SDA/FMC approach (α'_{ep}) has the opposite sign, so in the GSM method more negative $\delta \hat{a}$ values will be associated with larger $R_{\text{eff},f}$ values. The derived $\delta \hat{a}_{\text{AOD}}$ data are then plotted vs. the \hat{a}_{AOD} data on a non-rectilinear grid that was constructed by modeling an extensive set of conditions (see Fig. 1 in Gobbi et al., 2007). Gobbi et al. (2007) demonstrate that the impact of using different atmospherically relevant refractive indices (i.e. different particle compositions) on the retrieval is small. O’Neill (2009) has argued that the Gobbi approach is essentially equivalent to the SDA/FMC since both rely on fundamental spectral curvature mechanisms to extract an analogous set of fine-mode parameters.

The GSM model was also applied to the NOAA CRD data by constructing the parameters $\hat{a}_{ep}(355,1064)$ and $\delta \hat{a}_{ep} = \hat{a}_{ep}(355,532) - \hat{a}_{ep}(532,1064)$ from the four channels (25% and 85% RH, sub 1 and sub 10 μm). Only the 25% RH sub 1 μm data yielded reasonable results and are presented below. As for the SDA/FMC method, we obtain two $R_{\text{eff},f}$ values from the GSM approach, one using the AERONET data (hereafter AERONET GSM) and one from the NOAA CRD data (hereafter CRD GSM).

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2.3 Meteorological conditions during the case study period

The prevailing meteorological conditions characterizing the Houston/Galveston bay area were represented by Southerly winds (ESE to WSW) until 30 August 2006, followed by Northerly flow (N to NNE) until 6 September 2006, and winds between N-NE and ESE to WSW until mid September. Detailed information about these meteorological patterns and the properties of the corresponding air masses are reported in Wright et al. (2009) for the UHMT site and in Bates et al. (2008) and Massoli et al. (2009) for the RHB. Figure 1 illustrates the overall RHB track in the area during the field study (panel 1a), with an enlarged view of the ship tracks in Houston/Galveston Bay and the Houston ship channel (panel 1b)

Under the ESE winds, long range transport of Saharan dust was an important contributor to the bulk aerosol in the Gulf of Mexico based on aerosol chemical measurements and FLEXPART back trajectory analysis (Bates et al., 2008; Stohl and Eckhardt, 2008). One of the major Saharan dust-intrusions occurred between 25 and 30 August 2006. Under prevailing Northerly flow, air masses over the Houston/Galveston Bay region and coastal Texas were a mixture of transport from the continental US and local/regional contributions. In particular, during the period 2–5 September, a long range transport event of sulfate aerosols from the Ohio River valley characterized a large region extending from Houston to coastal Texas and out in the Gulf of Mexico (Massoli et al., 2009). Examination of various datasets suggested that during both of these two particular periods, the RHB and the UHMT site were exposed to similar air masses. In particular, the Community Multiscale Air Quality (CMAQ) model, initiated with the National Center for Environmental Prediction (NCEP) weather forecast and the National Emission Inventory (NEI '99) predicts large scale vertical homogeneity in the aerosol for several extended periods in early September, including the period taken into account here (Byun and Lee, personal communication, 2009).

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3 Results and discussion

3.1 Data Overview and Investigation of Spatial Homogeneity

Figures 2 and 3 give an overview of the data that were used in the analyses conducted here. Figure 2 refers to the period 28 August (0000)–30 August (0000) 2006, when a Saharan dust intrusion was observed over the Houston / Gulf of Mexico region, as shown by trajectory analysis and particle composition information collected from impactor/filter samplers. (All times referred to in the text and figures of this paper are UTC.) We show the 30 min average time series of 532 and 1064 nm NOAA CRD σ_{ep} coefficients for the sub 1 and sub 10 μm aerosols at 85% RH (panel a); 532 and 1064 nm PSU CRD σ_{ep} data for sub 4 μm aerosols at the nominal 50% RH (panel b); the coarse ($>1 \mu\text{m}$) aerosol size distributions as measured by the GRIMM PAS at the UHMT site (panel c); the full volume size distribution from the DMPS/APS on the RHB (panel d); and the FLEXPART footprint emission sensitivity (panel e). After 27 August (18:00 UTC) a significant relative enhancement in the 1064 nm sub-10 μm σ_{ep} coefficient was observed by the NOAA CRD, suggesting a substantial increase in the coarse aerosol load. During the same time, the PSU CRD recorded higher σ_{ep} levels at both 532 and 1064 nm, although in this case the $\sigma_{ep,1064} < \sigma_{ep,532}$, perhaps due to the different size cuts of the CRD instruments (10 μm for the NOAA CRD and 4 μm for the PSU CRD). The mobility size distribution data from both the UHMT and RHB showed that the $>1 \mu\text{m}$ volume concentration peaked on 28 August with a maximum at around 3 μm , consistent with the presence of coarse dust particles in the area.

Figure 3 shows data for the period 2–9 September 2006 when the Houston/Galveston bay region was characterized by continental outflow under N-NE winds. The bottom panel shows the 532 and 1064 nm σ_{ep} time series data for the NOAA CRD sub 1 μm 85% RH channel and the PSU CRD. Overall high σ_{ep} levels were observed, with the 532 nm $\sigma_{ep} > 100 \text{ Mm}^{-1}$ and the 1064 nm σ_{ep} coefficients around 50 Mm^{-1} . The volume size distributions collected on board the RHB (top panel) show that the

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particles were mostly in the accumulation mode, peaking between 300 and 400 nm particle mobility diameter. Organic matter and sulfate were present in equal amounts for the latter half of the period, while during 2–5 September sulfate aerosols were the dominant type and a regional haze event occurred in the Gulf region (Massoli et al., 2009). The quantitative agreement between the σ_{ep} coefficients measured by the NOAA and PSU CRD instruments, with both large and small scale structures captured by the two CRDs, is noteworthy given that the physical distance between the two platforms reached 50 miles at times during this period. This confirms that the air masses probed at the two platforms (RHB and UHMT) were similar on a regional scale, as predicted by the NCEP/CMAQ models (Byun and Lee, personal communication, 2009).

Figure 4 shows the $\hat{a}_{ep,UHMT}$ and $\hat{a}_{ep,RHB}$ calculated from the PSU and the 85% RH sub 1 and sub 10 μm NOAA CRD σ_{ep} data for August 28–30 (period 1, coarse mode dominated) and 2–9 September (period 2, fine mode dominated). The indicated areas during period 2 represent times when the data are known to be affected by local pollution plume sources sampled by RHB in the Galveston area. The $\hat{a}_{ep,RHB}$ and $\hat{a}_{ep,UHMT}$ values show overall good agreement, both qualitatively and quantitatively. A scatter plot of the PSU CRD vs. the NOAA CRD data (sub 1 μm in period 1 and sub 10 μm in period 2) is shown in the lower panel of Fig. 4. The linear regression of the $\hat{a}_{ep,UHMT}$ vs. the $\hat{a}_{ep,RHB}$ data leads to a overall slope of 0.96, with an R^2 value of 0.75. Again, this implies that there was large scale aerosol air mass consistency during this part of the study.

3.2 Ångström exponent and spectral curvature analysis of the NOAA CRD data

Figure 5 shows the α_{ep} (top panel) at $\lambda=500$ nm and α'_{ep} (bottom panel) extracted from the NOAA CRD data with the SDA/FMC procedure as described in Sect. 2.2. The α_{ep} and α'_{ep} are presented for all the NOAA CRD data, i.e. sub 1 and sub 10 μm , and 25% and 85% RH, for both periods 1 and 2.

Some interesting features emerge from the analysis. For period 1 (coarse mode dominated), the average value of α_{ep} for the sub 1 μm data is 1.5, whereas the α_{ep}

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value for sub 10 μm is around 0.8. No significant difference is observed between α_{ep} extracted at different RH; the α_{ep} reported for the $<1 \mu\text{m}$ sizes at 25% RH is approximately the same as that for the sub 1 μm data at 85% RH. The minimal difference between α_{ep} values obtained at 25% and 85% RH during the dust event indicates minimal water uptake even by the sub 1 μm aerosols, as the aerosol sample was dominated by mineral dust in both size ranges. The analysis of impactor data collected on the RHB between 28 and 30 August shows that sub 1 μm dust contributed up to 50% of the $<1 \mu\text{m}$ aerosol mass (the rest being non sea-salt sulfate), whereas it was more than 80% of the total sub 10 μm aerosol mass.

During period 2, the trend in the extracted α_{ep} values is different. In this case we observe similar α_{ep} values extracted for the same RH with no distinction between inlet size cut to a first approximation: $\alpha_{ep} \sim 1.2$ for 85% RH data (both sub 1 and sub 10 μm) and $\alpha_{ep} \sim 2.0$ for 25% RH data (again both sub 1 and sub 10 μm). These results indicate the dominance of sub 1 μm aerosols during period 2, with larger sizes (corresponding to the lower α_{ep} values at 85% RH) mainly due to particle hygroscopic growth (Massoli et al., 2009).

The curvature data α'_{ep} (bottom panel of Fig. 5) reflect the trends discussed above for α_{ep} . During the coarse-mode dominated period, near-zero α'_{ep} is observed for the sub 1 μm data (25% and 85% RH), and negative values are obtained for sub 10 μm data (25% and 85% RH). For the fine-mode dominated Period 2, positive α'_{ep} values of ~ 1.8 and 2.5 (average values) are obtained for the 85% RH data (sub 1 and sub 10 μm) and for the 25% RH data (sub 1 and sub 10 μm) respectively.

3.3 Fine mode fraction (η) and $R_{\text{eff},f}$ retrievals and comparisons

Here we describe the results of the spectral analysis conducted on the NOAA CRD data to obtain the fine mode η and $R_{\text{eff},f}$ parameters. As stated earlier, for these analyses we use the sub 10 μm 85% RH CRD data because it is closest to the ambient RH values. To further facilitate comparison between the AERONET and CRD retrievals, all of the parameters are reported at 500 nm. Figure 6 shows the fine mode fraction

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η extracted from the NOAA CRD data using the SDA/FMC procedure (red trace) for the two time periods shown earlier. In addition the ratio of the NOAA CRD 85% RH sub $1\ \mu\text{m}$ to the sub $10\ \mu\text{m}$ σ_{ep} coefficients (hereafter “SMF”, or “submicron fraction”) is shown in Fig. 6 (black trace). The SDA retrieval of the fine mode contribution to the total extinction gives satisfactory agreement with the direct sub $1/\text{sub } 10\ \mu\text{m}$ σ_{ep} ratio ($\eta \sim 0.95$ and SMF ~ 0.9) during period 2 and similar agreement ($\eta \sim 0.3$ and SMF ~ 0.4) during the coarse mode event. In general one would expect η to be slightly less than SMF due to the manner in which these parameters are calculated (c.f. Kleidman et al. (2005) for example). It was noted above that the $R_{\text{eff},f}$ ($\rho_{\text{eff},f}$) calculation becomes less accurate as η decreases. Because we want to minimize the influence of retrieval results other than those related to the variation of $R_{\text{eff},f}$, we will only use the fine mode dominated period to extract the $R_{\text{eff},f}$ parameter from the CRD data.

Figure 7 depicts values of the fine mode effective mean radius $R_{\text{eff},f}$ calculated from 24 h averages of the data obtained with the various instruments, via the different methods discussed above, for the period 1–9 September 2006 slightly beyond the long range transport event (2–5 September) discussed in Massoli, (2009). We show $R_{\text{eff},f}$ extracted from: the in situ NOAA CRD data via the SDA/FMC and GSM approaches (CRD SDA/FMC and CRD GSM, respectively) and from the AOD AERONET sun-photometer data via the SDA/FMC and GSM approaches (AERONET SDA/FMC and AERONET GSM, respectively). For the SDA/FMC treatment, the $R_{\text{eff},f}$ was calculated from Eq. 8 using sub $1\ \mu\text{m}$ 85% RH $\rho_{\text{eff},f}$ and a value for the refractive index broadly applicable to ambient aerosols, $m = 1.4 - 0.001i$ (Gobbi et al., 2007). The pair-wise \hat{a} and $\delta\hat{a}$ values from the sub $1\ \mu\text{m}$ 25% RH NOAA CRD data were constructed and placed on the grid shown in Fig. 1 of Gobbi, et al., 2007 to extract the geometric fine mode radius, which was then converted to the effective radius for Fig. 7.

The last two reported $R_{\text{eff},f}$ datasets are calculated using a numerical evaluation of Eq. 1 from the in situ size distribution measurements at the UHMT site (hereafter Grimm SMPS) and on the RHB (hereafter PMEL DMPS). The particle size distribution data were averaged to 24 h periods and converted bin-wise to effective volume con-

centration ($dN \times 4/3\pi r_{\text{bin}}^3$) which were then summed and divided by the sum for effective area concentration ($dN \times \pi r_{\text{bin}}^2$) to produce the Grimm SMPS and PMEL DMPS $R_{\text{eff},f}$ estimates.

Figure 7 shows the satisfactory level of agreement between the $R_{\text{eff},f}$ values derived from different experimental data and through different methods. A pooled standard deviation of 27.9 nm was calculated, considering each day as a set. The agreement between the in situ size distribution derived $R_{\text{eff},f}$ values from the GRIMM SMPS and PMEL DMPS is better early in this period (2–5 September) when the size distributions peaked at a small radius. This provides additional evidence that a synoptic scale air-mass was being observed at both platforms. The $R_{\text{eff},f}$ values from the sunphotometer based size distribution (AERONET SSRI) seem to also agree rather well with the particle mobility size-based GRIMM SMPS and PMEL DMPS values suggesting a certain level of homogeneity in the surface size distribution throughout the column. Unfortunately the AERONET full sky scan data was missing after 3 September because of cloud contamination, hence limiting the comparisons with SSRI retrievals. The agreement between the $R_{\text{eff},f}$ values obtained from AERONET AOD and NOAA CRD σ_{ep} data via the spectral methods is also compelling. All of the in situ methods seem to have captured the rapid increase in $R_{\text{eff},f}$ during the first days of September 2006 and then the stable higher radius during the middle of the period. The low bias of the Grimm SMPS measurements may be from an intentional choice to limit the size range to a particle diameter of 521 nm because of concern that the few larger size bins may have been contaminated by small particle count data. As the particle mean radius may have approached 250 nm, this choice almost certainly excluded some particle sizes that were relevant to the evaluation of $R_{\text{eff},f}$.

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The purpose of this study was to assess the possibility of using multi-wavelength in situ aerosol σ_{ep} measurements to extract information pertaining to the fine mode fraction of extinction η and the fine mode effective radius $R_{\text{eff},f}$ in a fashion similar to that used with columnar AOD measurements (such as those from the AERONET sunphotometers or satellite borne sensors). This information is potentially useful for partitioning the contributions of anthropogenic (mostly represented by the fine modes) and natural (largely coarse sizes) aerosols to the measured σ_{ep} coefficients.

There was a reasonable agreement between the fine mode fraction and $R_{\text{eff},f}$ results obtained from the different methods. The agreement of the SDA-derived η with the SMF (Fig. 6) suggests that the method is robust, especially since it was run for a sub optimal number of wavelengths (three is the minimum usable). This is useful for helping to validate the SDA AERONET product and also suggests that the wavelength dependence of extinction from CRD can provide a reasonable estimate for the partitioning between the fine and coarse modes.

The $R_{\text{eff},f}$ retrieval from the wavelength dependent optical measurements also appears to be promising. While the GSM method is simple and provides qualitative information, it is difficult to apply in a quantitative way mainly because graphically interpreting a large amount of time-resolved data to obtain values for η and $R_{\text{eff},f}$ is impractical. The production of η and $\rho_{\text{eff},f}$ by the SDA/FMC algorithm is efficient and is probably a more reasonable approach since it properly leaves the $R_{\text{eff},f}$ and refractive index information intertwined. Although there currently are no universally accepted measurements of particle refractive index applied to ambient aerosols in field studies, there are a number of potentially useful surrogates, including using literature recommended values as used here, or the refractive index product from the AERONET SSRI (a temporally sparse output for which there actually exists little validation in the literature). The ability to extract $R_{\text{eff},f}$ would significantly expand the utility of the in situ CRD instruments, which are currently being used on land, ship, and aircraft to obtain

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3-dimensional profiles of the aerosol optical effects for use in radiative forcing calculations and indirect-effect estimates that are currently needed for global climate change models.

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References

Anderson, T. L., Charlson, R. J., Bellouin, N., Boucher, O., Chin, M., Christopher, S. A., Haywood, J., Kaufman, Y. J., Kinne, S., Ogren, J. A., Remer, L. A., Takemura, T., Tanre, D., Torres, O., Treppe, C. R., Wielicki, B. A., Winker, D. M., and Yu, H. B.: An "A-Train" strategy for quantifying direct climate forcing by anthropogenic aerosols, *B. Am. Meteorol. Soc.*, 86, 1795–1809, 2005.

Andreae, M. O., Rosenfeld, D., Artaxo, P., Costa, A. A., Frank, G. P., Longo, K. M., and Silva-Dias, M. A. F.: Smoking rain clouds over the Amazon, *Science*, 303, 1337–1342, 2004.

Ångström, A.: On the atmospheric transmission of sun radiation and on dust in the air, *Geografika Ann.*, 11, 156–166, 1929.

Atkinson, D. B.: Solving chemical problems of environmental importance using cavity ring-down spectroscopy, *The Analyst*, 128, 117–125, 2003.

Bates, T. S., Coffman, D. J., Covert, D. S., and Quinn, P. K.: Regional marine boundary layer aerosol size distributions in the Indian, Atlantic, and Pacific Oceans: A comparison of INDOEX measurements with ACE-1, ACE-2, and Aerosols99, *J. Geophys. Res.-Atmos.*, 107, 148–227, doi: 10.1029/2001J001174, 2002.

Bates, T. S., Anderson, T. L., Baynard, T., Bond, T., Boucher, O., Carmichael, G., Clarke, A., Erlick, C., Guo, H., Horowitz, L., Howell, S., Kulkarni, S., Maring, H., McComiskey, A., Middlebrook, A., Noone, K., O'Dowd, C. D., Ogren, J., Penner, J., Quinn, P. K., Ravishankara,

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A. R., Savoie, D. L., Schwartz, S. E., Shinozuka, Y., Tang, Y., Weber, R. J., and Wu, Y.: Aerosol direct radiative effects over the northwest Atlantic, northwest Pacific, and North Indian Oceans: estimates based on in-situ chemical and optical measurements and chemical transport modeling, *Atmos. Chem. Phys.*, 6, 1657–1732, 2006,

<http://www.atmos-chem-phys.net/6/1657/2006/>.

Baynard, T., Lovejoy, E. R., Pettersson, A., Brown, S. S., Lack, D., Osthoff, H., Massoli, P., Ciciora, S., Dube, W. P., and Ravishankara, A. R.: Design and application of a pulsed cavity ring-down aerosol extinction spectrometer for field measurements, *Aerosol Sci. Technol.*, 41, 447–462, 2007.

Berner, A., Lurzer, C., Pohl, F., Preining, O., and Wagner, P.: Size Distribution of the Urban Aerosol in Vienna, *Sci. Total Environ.*, 13, 245–261, 1979.

Chin, M., Chu, A., Levy, R., Remer, L., Kaufman, Y., Holben, B., Eck, T., Ginoux, P., and Gao, Q. X.: Aerosol distribution in the Northern Hemisphere during ACE-Asia: Results from global model, satellite observations, and Sun photometer measurements, *J. Geophys. Res.-Atmos.*, 109, 15, D23s90, doi:10.1029/2004jd004829, 2004.

Doherty, S. J., Quinn, P. K., Jefferson, A., Carrico, C. M., Anderson, T. L., and Hegg, D.: A comparison and summary of aerosol optical properties as observed in situ from aircraft, ship, and land during ACE-Asia, *J. Geophys. Res.-Atmos.*, 110, D04201, doi:10.1029/2004jd004964, 2005.

Dubovik, O. and King, M. D.: A flexible inversion algorithm for retrieval of aerosol optical properties from Sun and sky radiance measurements, *J. Geophys. Res.*, 105, 20673–20696, 2000.

Dubovik, O., Holben, B. N., Lapyonok, T., Sinyuk, A., Mishchenko, M. I., Yang, P., and Slutsker, I.: Non-spherical aerosol retrieval method employing light scattering by spheroids, *Geophys. Res. Lett.*, 29(4), 1415, doi: 10.1029/2001gl014506, 2002.

Dubovik, O., Sinyuk, A., Lapyonok, T., Holben, B. N., Mishchenko, M., Yang, P., Eck, T. F., Volten, H., Munoz, O., Veihelmann, B., van der Zande, W. J., Leon, J. F., Sorokin, M., and Slutsker, I.: Application of spheroid models to account for aerosol particle non-sphericity in remote sensing of desert dust, *J. Geophys. Res.-Atmos.*, 111, 34, D11208, doi: 10.1029/2005jd006619, 2006.

Gobbi, G. P., Kaufman, Y. J., Koren, I., and Eck, T. F.: Classification of aerosol properties derived from AERONET direct sun data, *Atmos. Chem. Phys.*, 7, 453–458, 2007, <http://www.atmos-chem-phys.net/7/453/2007/>.

- Hansen, J. E. and Travis, L. D.: Light Scattering in Planetary Atmospheres, *Space Science Reviews*, 16, 527–610, 1974.
- Holben, B. N., Eck, T. F., Slutsker, I., Tanré, D., Buis, J. P., Setzer, A., Vermote, E., Reagan, J. A., Kaufman, Y. J., Nakajima, T., Lavenu, F., Jankowiak, I., and Smirnov, A.: AERONET—
5 A federated instrument network and data archive for aerosol characterization, *Rem. Sens. Env.*, 66, 1–16, 1998.
- IPCC, 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change Cambridge, United Kingdom and New York, NY, USA, Climate Change 2007.
- 10 Kleidman, R. G., O'Neill, N. T., Remer, L. A., Kaufman, Y. J., Eck, T. F., Tanre, D., Dubovik, O., and Holben, B. N.: Comparison of moderate resolution Imaging spectroradiometer (MODIS) and aerosol robotic network (AERONET) remote-sensing retrievals of aerosol fine mode fraction over ocean, *J. Geophys. Res.-Atmos*, 110, 6, D22205, doi: 10.1029/2005jd005760, 2005.
- 15 Lefer, B. and Rappenglück, B.: Overview of TRAMP Texas II Study, *Atmos. Environ.*, in press, 2008.
- Lohmann, U. and Feichter, J.: Global indirect aerosol effects: a review, *Atmos. Chem. Phys.*, 5, 715–737, 2005,
<http://www.atmos-chem-phys.net/5/715/2005/>.
- 20 Massoli, P., Bates, T. S., Quinn, P. K., Lack, D. A., Baynard, T., Lerner, B. M., Tucker, S. C., Brioude, J., Stohl, A., and Williams, E. J.: Aerosol optical and hygroscopic properties during TexAQS-GoMACCS 2006 and their impact on aerosol direct radiative forcing, *J. Geophys. Res.-Atmos*, 114, 17, D00f07, doi: 10.1029/2008jd011604, 2009.
- Moosmuller, H., Varma, R., and Arnott, W. P.: Cavity ring-down and cavity-enhanced detection techniques for the measurement of aerosol extinction, *Aerosol Sci. Technol.*, 39, 30–39, doi: 10.1080/027868290903880, 2005.
- 25 O'Neill, N. T., Eck, T. F., Smirnov, A., Holben, B. N., and Thulasiraman, S.: Spectral discrimination of coarse and fine mode optical depth, *J. Geophys. Res.-Atmos.*, 108(15), 4559, doi: 10.1029/2002jd002975, 2003.
- 30 O'Neill, N. T., Thulasiraman, S., Eck, T. F., and Reid, J. S.: Robust optical features of fine mode size distributions: Application to the Quebec smoke event of 2002, *J. Geophys. Res. -Atmos.*, 110(21), D11207, doi:10.1029/2004jd005157, 2005.
- O'Neill, N. T., Thulasiraman, S., Eck, T. F., and Reid, J. S.: Correction to the effective radius

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expression in O'Neill et al. (2005), accepted for publication in J. Geophys. Res., 2008.

O'Neill, N. T.: Comment on the paper "Classification of aerosol properties derived from AERONET direct sun data" by G. P. Gobbi et al. (2007), Atmos. Chem. Phys. Discuss., 9, 175–182, 2009,

<http://www.atmos-chem-phys-discuss.net/9/175/2009/>.

Radney, J. G., Bazargan, M. H., Wright, M. E., and Atkinson, D. B.: Laboratory Validation of Aerosol Extinction Coefficient Measurements by a Field-Deployable Pulsed Cavity Ring-down Transmissometer, Aerosol Sci. Technol., accepted, 2008.

Schuster, G. L., Dubovik, O., and Holben, B. N.: Ångström exponent and bimodal aerosol size distributions, J. Geophys. Res.-Atmos., 111(14), D07207, doi: 10.1029/2005jd006328, 2006.

Sinyuk, A., Dubovik, O., Holben, B., Eck, T. F., Breon, F. M., Martonchik, J., Kahn, R., Diner, D. J., Vermote, E. F., Roger, J. C., Lapyonok, T., and Slutsker, I.: Simultaneous retrieval of aerosol and surface properties from a combination of AERONET and satellite data, Remote Sensing of Environment, 107, 90–108, doi: 10.1016/j.rse.2006.07.022, 2007.

Strawa, A. W., Elleman, R., Hallar, A. G., Covert, D., Ricci, K., Provencal, R., Owano, T. W., Jonsson, H. H., Schmid, B., Luu, A. P., Bokarius, K., and Andrews, E.: Comparison of in situ aerosol extinction and scattering coefficient measurements made during the Aerosol Intensive Operating Period, J. Geophys. Res.-Atmos., 111(17), D05s03, doi: 10.1029/2005jd006056, 2006.

Winklmayr, W., Reischl, G. P., Lindner, A. O., and Berner, A.: A New Electromobility Spectrometer for the Measurement of Aerosol Size Distributions in the Size Range from 1 to 1000 Nm, J. Aerosol Sci., 22, 289–296, 1991.

Wright, M. E., Atkinson, D.B., Ziemba, L. D., Griffin, R., Hiranuma, N., Brooks, S., Lefer, B., Flynn, J., Perna, R., Rappengluck, B., Luke, W., and Kelley, P.: Extensive Aerosol Optical Properties and Aerosol Mass Related Measurements during TRAMP/TexAQS 2006 – Implications for PM Compliance and Planning Atmospheric Environment, doi:10.1016/j.atmosenv.2008.12.055, in press, 2009.

Yu, H., Kaufman, Y. J., Chin, M., Feingold, G., Remer, L. A., Anderson, T. L., Balkanski, Y., Belouin, N., Boucher, O., Christopher, S., DeCola, P., Kahn, R., Koch, D., Loeb, N., Reddy, M. S., Schulz, M., Takemura, T., and Zhou, M.: A review of measurement-based assessments of the aerosol direct radiative effect and forcing, Atmos. Chem. Phys., 6, 613–666, 2006, <http://www.atmos-chem-phys.net/6/613/2006/>.

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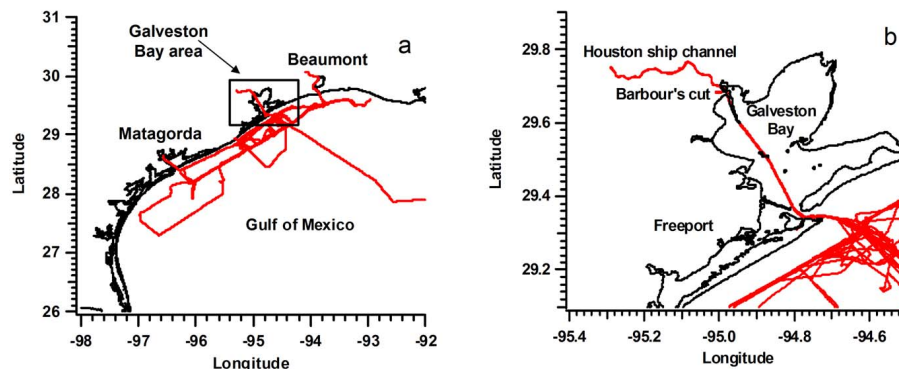


Fig. 1. RHB ship track in **(a)** the Gulf of Mexico and **(b)** Galveston bay, during the period 2 August–11 September, 2006. The UHMT site is in downtown Houston, about 5 miles east of the Houston Ship Channel (HSC).

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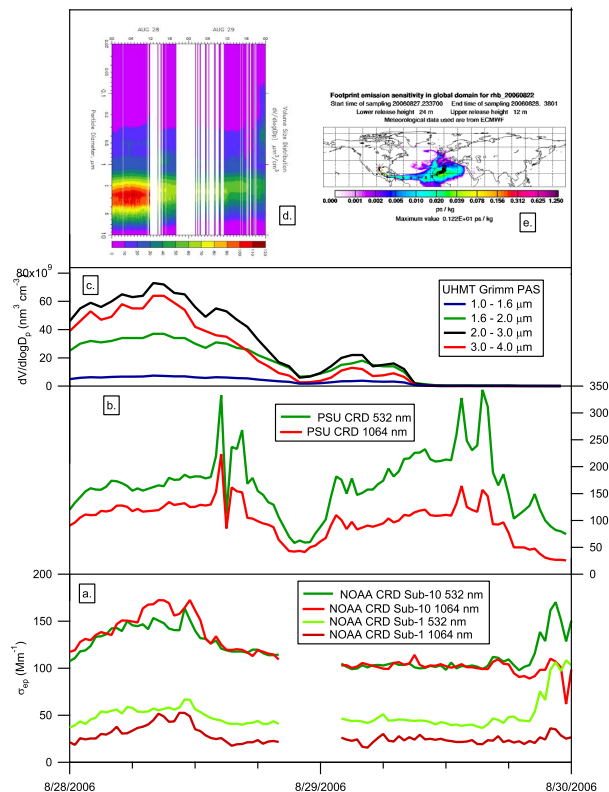


Fig. 2. Time series for the period 28 August to 30 August 2006 of: panel (a) NOAA CRD sub 1 and sub 10 μm, 85% RH, 532 and 1064 nm σ_{ep} ; panel (b) PSU CRD 532 and 1064 nm σ_{ep} ; panel (c) 1 to 4 μm size bins of the PAS at the UHMT site; panel (d) volume concentration size distribution from the DMPS/APS on the RHB; and (e) FLEXPART trajectory analysis showing transport patterns from the Saharan region to the Gulf of Mexico on 27 August.

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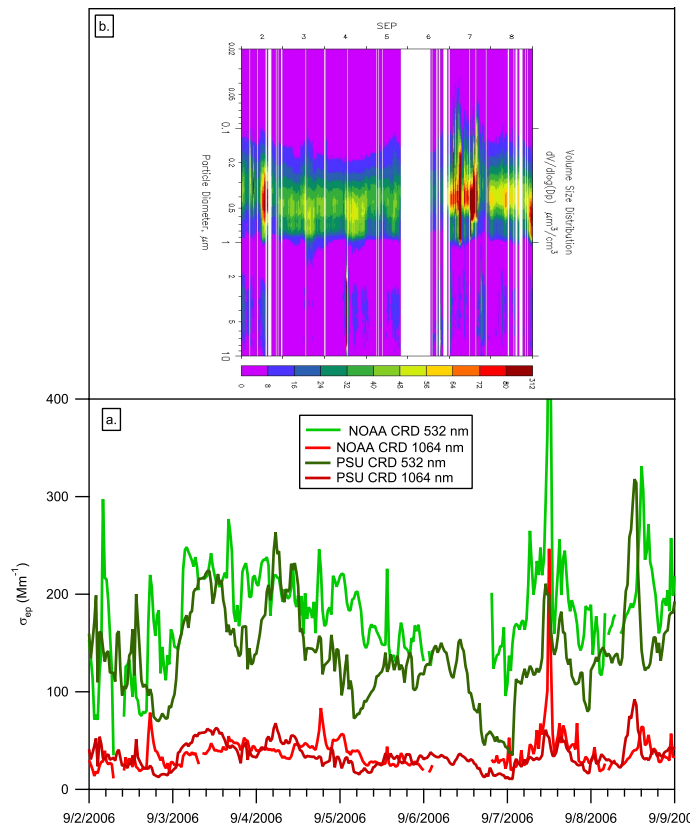


Fig. 3. Time series for the period 2 September to 9 September 2006 of: panel (a) NOAA CRD 532 and 1064 nm sub 1 micron 85% RH σ_{ep} and PSU CRD 532 and 1064 nm σ_{ep} ; and (b) volume concentration size distribution from the DMPS/APS on the RHB.

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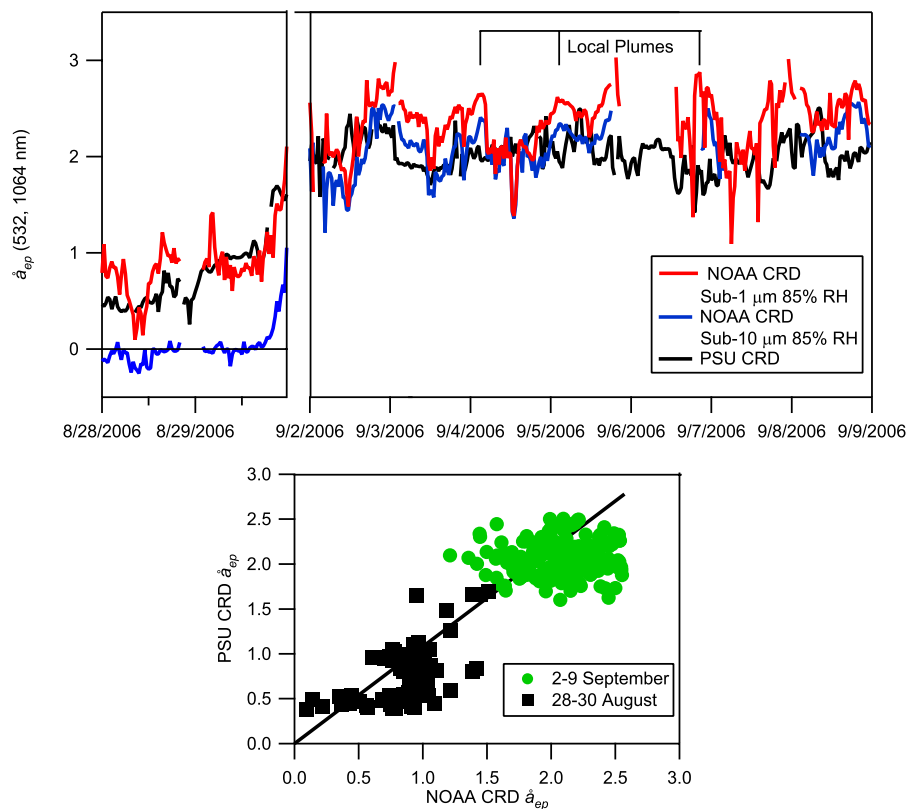


Fig. 4. Top) extinction Ångström exponents \hat{a}_{ep} obtained according to Eq. 1 using the 532 and 1064 nm σ_{ep} coefficients from the PSU CRD (black trace) and from the NOAA CRD at 85% RH for both sub $1\ \mu\text{m}$ (red trace) and sub $10\ \mu\text{m}$ (blue trace), for the two periods discussed in the text. Bottom) scatter plot of the PSU CRD $\hat{a}_{ep,UHMT}$ vs. the NOAA CRD $\hat{a}_{ep,RHB}$ for period 1 and 2.

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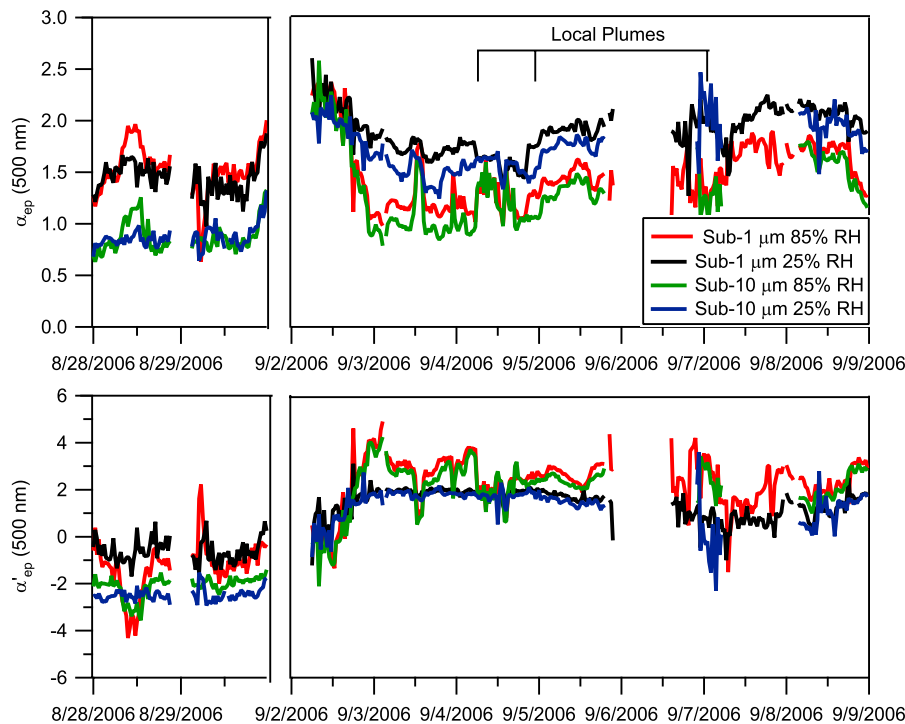


Fig. 5. The extinction Ångström exponent α_{ep} (top panel), and spectral curvature α'_{ep} (bottom panel) for the sub 1 and sub 10 μm NOAA CRD σ_{ep} coefficients at 25% and 85% RH, extracted at 500 nm via the AERONET SDA/FMC routine.

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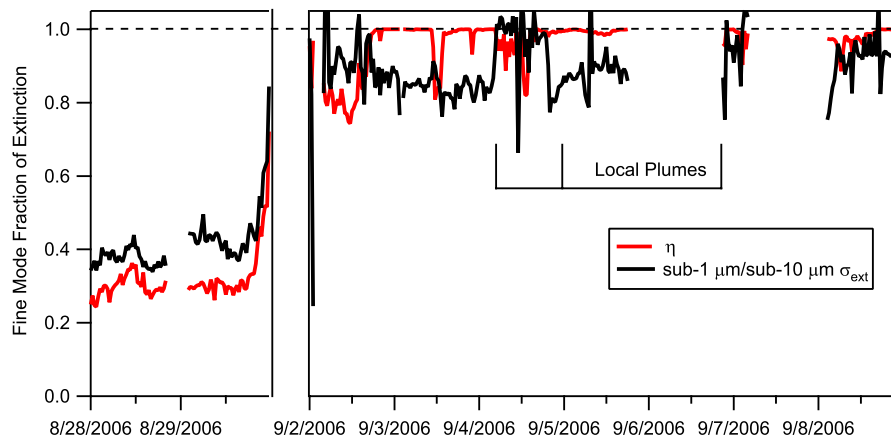


Fig. 6. Fine mode fraction of extinction η at 500 nm calculated from the NOAA CRD 85% RH σ_{ep} coefficients using the SDA/FMC method (red trace) and the ratio of the sub 1 μm to the sub 10 μm 85% RH 532 nm σ_{ep} data (black trace).

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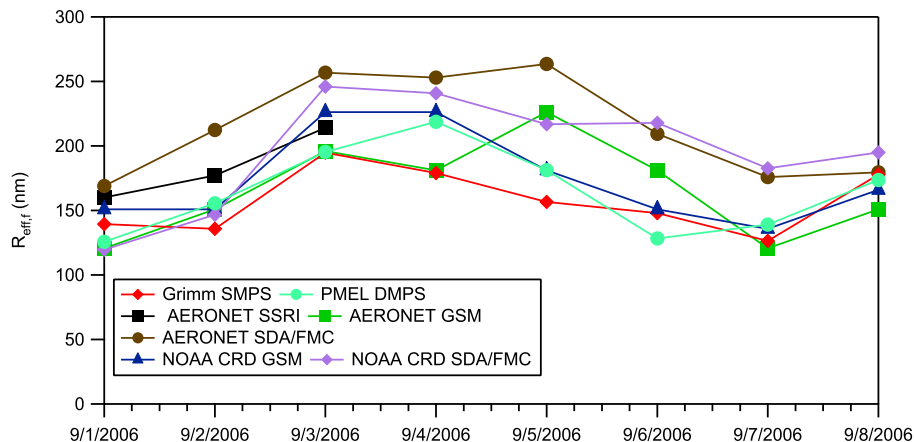


Fig. 7. Comparison of 24 h average $R_{eff,f}$ values obtained from the NOAA CRD σ_{ep} data and from the AERONET sunphotometer using the spectral methods discussed in the text, and the $R_{eff,f}$ values estimated from the size distribution instruments.

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