

**Aerosol extinction in
the UV spectral
region**

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Broadband measurements of aerosol extinction in the ultraviolet spectral region

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Abstract

Aerosols influence the Earth's radiative budget by scattering and absorbing incoming solar radiation. The optical properties of aerosols vary as a function of wavelength, but few measurements have reported the wavelength dependence of aerosol extinction cross-sections and complex refractive indices. We describe a new laboratory instrument to measure aerosol optical extinction as a function of wavelength, using cavity enhanced spectroscopy with a broadband light source. The instrument consists of two broadband channels which span the 360–390 and 385–420 nm spectral regions using two light emitting diodes (LED) and a grating spectrometer with charge-coupled device (CCD) detector. We determined aerosol extinction cross-sections and directly observed Mie scattering resonances for aerosols that are purely scattering (polystyrene latex spheres and ammonium sulfate), slightly absorbing (Suwannee River fulvic acid), and strongly absorbing (nigrosin dye). We describe an approach for retrieving refractive indices as a function of wavelength from the measured extinction cross-sections over the 360–420 nm wavelength region. The retrieved refractive indices for PSL and ammonium sulfate agree within uncertainty with literature values for this spectral region. The refractive index determined for nigrosin is $1.78(\pm 0.03) + 0.19(\pm 0.08)i$ at 360 nm and $1.53(\pm 0.03) + 0.21(\pm 0.05)i$ at 420 nm. The refractive index determined for Suwannee River fulvic acid is $1.71(\pm 0.02) + 0.07(\pm 0.06)i$ at 360 nm and $1.66(\pm 0.02) + 0.06(\pm 0.04)i$ at 420 nm. These laboratory results support the potential for a field instrument capable of determining ambient aerosol optical extinction, average aerosol extinction cross-section, and complex refractive index as a function of wavelength.

1 Introduction

Atmospheric aerosols modify the earth's energy budget directly by absorbing and scattering solar radiation and indirectly by modifying cloud properties (Solomon et al., 2007). Aerosol direct forcing remains one of the largest uncertainties in quantifying

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the role that aerosol play in the Earth's radiative budget (McComiskey et al., 2008). The interaction of light with a particle is determined by the wavelength of the incident radiation, as well as the size, shape, and complex refractive index (m) of the particle. The complex refractive index is the only intrinsic optical property of a particle, while size, shape, and composition are intrinsic physical properties. Knowing the value of m is critical in correctly modeling the optical properties of aerosols. The complex refractive index of different aerosol types is directly used in regional and global aerosol models to predict aerosol radiative forcing.

Depending on their refractive index, different aerosol types have different influences on the Earth's radiative budget. Most common inorganic aerosols, including sulfate and sea salt, are purely scattering at visible and ultraviolet wavelengths, and thus exert a net cooling effect (Toon et al., 1976; Nilsson, 1979). Alternatively, black carbon aerosols derived from combustion sources strongly absorb light at all visible and ultraviolet wavelengths (Bond and Bergstrom, 2006). Between these two extremes lie brown carbon aerosols, which comprise a potentially large and variable group of organic compounds, including secondary organic and biomass burning aerosols, and which are characterized by a wavelength-dependent absorption that increases toward the ultraviolet spectral region (Andreae and Gelencser, 2006; Sun et al., 2007; Dinar et al., 2008; Hecobian et al., 2010; Adler et al., 2010; Moosmuller et al., 2011; Zhang et al., 2011). Brown carbon may dominate the aerosol composition in some geographical regions, yet its short-wavelength optical properties are poorly characterized (Chung et al., 2012).

Conventional techniques for the measurement of aerosol scattering and absorption coefficients include integrating nephelometry (Heintzenberg and Charlson, 1996) and filter-based photometry (Bond et al., 1999), respectively. Wavelength-dependent measurements of aerosol optical properties have typically used filter-based collection or extraction methods, rather than direct, in situ methods. For example, commercial aethalometer instruments have provided filter-based measurements over a wide spectral range of discrete wavelengths (e.g., Rizzo et al., 2011), but may be subject to

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artifacts, for example due to particles on the filter (Weingartner et al., 2003). Recent advances in photo-acoustic spectroscopy (PAS) (Arnott et al., 1999; Lack et al., 2006; Lack et al., 2012) and cavity ring-down spectroscopy (CRDS) (Strawa et al., 2003; Baynard et al., 2007; Abo Riziq et al., 2008; Langridge et al., 2011) allow rapid, accurate, in situ measurements of optical absorption and extinction, but have so far been limited to single, discrete wavelengths. Similarly, cavity attenuated phase shift spectroscopy (CAPS) instruments may employ broadband LED light sources, but do not resolve the extinction as a function of wavelength (Kebabian et al., 2007; Massoli et al., 2010). Due in part to these limitations, there have been few investigations of ambient aerosol optical properties at the blue end of the visible spectrum and into the ultraviolet, where some aerosol types both scatter and absorb light (Bond, 2001; Sun et al., 2007). One existing instrument combines a White-type multipass gas cell with a grating and a linear diode array detector to measure aerosol extinction as a function of wavelength from the mid-infrared to near-ultraviolet spectral region (Chartier and Greenslade, 2012). Further development of sensitive and accurate instruments able to measure extinction continuously across a broad wavelength region, particularly in the near ultraviolet and blue end of the visible spectrum, provides a useful complement to existing PAS, CRDS, and CAPS instruments.

Recent advances in cavity enhanced spectroscopy have combined high-intensity, broadband light sources with optical cavities. The optical output from the cavity is resolved with a spectrometer and recorded using a multichannel detector. This approach allows sensitive measurements of optical extinction as a function of wavelength (Fiedler et al., 2003). For gas-phase absorption measurements, the technique is referred to as incoherent broadband cavity enhanced absorption spectroscopy (IB-BCEAS or BBCEAS), or alternatively as cavity enhanced differential optical absorption spectroscopy (CE-DOAS) (Platt et al., 2009). IBCEAS is a sensitive and selective detection method for atmospheric trace gases with broad, structured absorptions in the visible and ultraviolet spectral region. When the optical cross-section of the analyte species is known, the gas-phase concentration of the species may be calculated

directly from the optical extinction. IBBCEAS has been used successfully to measure gas phase concentrations of NO_2 , NO_3 , CHOCHO , CH_3COCHO , HONO , O_3 , I_2 , IO , and OIO (Ball et al., 2004; Venables et al., 2006; Washenfelder et al., 2008; Gherman et al., 2008; Vaughan et al., 2008; Thalman and Volkamer, 2010). Initial cavity measurements have reported aerosol extinction for ambient aerosol (Washenfelder et al., 2008; Varma et al., 2009; Thalman and Volkamer, 2010).

In this study, we further extend the use of broadband cavity enhanced spectroscopy to measure aerosol extinction as a function of wavelength in a high-finesse cavity. By size-selecting different aerosol types, it is possible to measure their extinction cross-section as a function of wavelength, and directly observe the Mie scattering resonances. For aerosols whose extinction cross-section has minimal spectral dependence, the selection and measurement of a single particle diameter allows the retrieval of the complex refractive index (RI). Alternatively, for aerosols whose extinction cross-section varies with wavelength, the measurement of several particle sizes allows the retrieval of the complex RI as a function of wavelength. This paper demonstrates this technique for aerosols that are purely scattering (polystyrene latex spheres and ammonium sulfate), slightly absorbing (Suwannee River fulvic acid), and highly absorbing (nigrosin dye). We refer to our new method and instrument as the Broadband Aerosol Extinction Spectrometer (BAES).

2 Experimental

The laboratory instrument consists of two BAES channels and one CRDS channel, which measure aerosol extinction at 360–390, 385–425, and 407 nm respectively. The optics, aerosol generation, and flow system are described in detail below and presented schematically in Fig. 1. All of the instrumental components are mounted on a standard laboratory optical bench.

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2.1 Description of the broadband aerosol extinction spectrometer (BAES) channels

Each of the two BAES channels consist of an LED, collimating lens, optical cavity, bandpass filter, and shared grating spectrometer with CCD array detector as shown in Fig. 1a. The laboratory instrument is similar to an instrument for gas phase measurements of glyoxal and nitrogen dioxide described previously in Washenfelder et al. (2008). Notable changes include the addition of a second channel, replacement of the Xenon arc lamp with LEDs, and changes to the collimating and filtering optics.

The first BAES channel covers the spectral region from 360–390 nm, using an LED with manufacturer-specified center wavelength between 360–370 nm (NCSU033A, Nichia Corp., Tokyo, Japan). The experimentally measured center wavelength and optical power output at 0.5 A and 20.0 °C are 370.2 nm and 0.210 W, respectively. The second BAES channel covers the spectral region from 385–425 nm, using an LED with manufacturer-specified center wavelength between 390–410 nm (LZ1-00UA05, LEDEngin Inc., San Jose, CA, USA). The experimentally measured center wavelength and optical power output at 0.7 A and 20.0 °C are 407.1 nm and 0.450 W, respectively. To maintain a constant optical power output, the temperature of each LED is controlled at 20.0(±0.3) °C and powered by a constant-current power supply.

The output from each LED is collimated using a single F/1.2 fused silica lens and passively coupled into a cavity consisting of two 2.5 cm, 1 m radius of curvature mirrors (Advanced Thin Films, Boulder, CO, USA). The manufacturer's reported transmission for the two sets of mirrors is 82 ppm at the nominal center wavelength of 362 nm and 35 ppm at the nominal center wavelength of 405 nm. After exiting the cavity, the light is optically filtered (XB05, Omega Optical Inc, Brattleboro, VT, USA; FB400-40, Thorlabs, Newton, NJ, USA) to block stray light and out-of-band light from the LEDs. The filtered light is imaged by a 2.5 cm F/3.1 lens onto a 0.5 cm F/2 lens that couples the light into a fiber bundle (BFB-455-7, Princeton Instruments, Trenton, NJ, USA) containing seven

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200 μm diameter UV-VIS fibers which are then oriented linearly along the 100 μm input slit for the grating spectrometer.

Spectra are acquired using a 150 mm focal length Czerny-Turner spectrometer (InSpectrum 150, Princeton Instruments) with a 16-bit back-illuminated 250×1024 pixel charge coupled device (CCD) array detector cooled to -20°C (Langford et al., 2007). The spectrometer contains a $1200 \text{ groove mm}^{-1}$ (500 nm blaze) grating rotated to allow spectral measurements between 333–451 nm. The grating spectrometer is temperature controlled at $33.0(\pm 0.2)^\circ\text{C}$. Dark spectra are acquired with the InSpectrum input shutter closed. The wavelength calibration was determined using narrow Hg emission lines from a Hg/Ar lamp, and was consistent within 0.03 nm over the course of the experiments.

The spectra from the InSpectrum 150 are measured as two vertical regions on the CCD, and digitized to produce two spectra. Custom data acquisition software (Labview, National Instruments Corp., Austin, TX, USA) is used to control the spectrometer and acquire data.

2.2 Description of the cavity ring-down spectroscopy (CRDS) channel

The CRDS channel consists of a diode laser, optical cavity, bandpass filter, and photomultiplier tube, following the design in Fuchs et al. (2009). Light from a continuous wave diode laser with temperature-controlled diode (Fabry-Perot diode model IQ μ series, optical power output 80 mW, Power Technology Inc., Little Rock, AR, USA) is amplitude modulated with a square-wave signal at a repetition of 2 kHz and a duty cycle of 50%. The laser is optically isolated and aligned on-axis to the cavity with no active mode matching. The cavity is formed by a pair of 2.54 cm diameter, 1 m radius of curvature mirrors (Advanced Thin Films, Boulder, CO, USA). The center wavelength and full-width at half-maximum of the diode laser were measured with a grating spectrometer to be $407.0(\pm 0.1) \text{ nm}$ and $0.6(\pm 0.15) \text{ nm}$, respectively. The observed ring-down time constant at 407 nm was approximately 33 μs at ambient pressure (840 hPa in Boulder, CO; 1600 m above mean sea level). The light transmitted through the end mirror

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of the cavity is collected with a 200 μm diameter fiber optic, optically filtered with a bandpass filter (FB405-10, Thorlabs), and then measured by a photomultiplier tube module (HC120, Hamamatsu Photonics, Hamamatsu, Japan). Ring-down transients are recorded at a repetition rate of 1 kHz, coadded, and fitted with a single exponential following the method in Fuchs et al. (2009).

2.3 Aerosol generation and measurement

Four different aerosol types were generated: polystyrene latex spheres (PSL; Nanosphere size standards, Thermo Fisher Scientific Inc., Waltham, MA, USA), ammonium sulfate (AS; a component of anthropogenic aerosol; Sigma Aldrich, St. Louis, MO, USA), Suwannee River fulvic acid (SRFA; a model compound for humic-like substances; 1S101F, International Humic Substances Society, Saint Paul, MN, USA), and water-soluble nigrosin (an organic black dye used as a model for highly absorbing substances; CAS 8005-03-6, part number 198285, Sigma Aldrich). As shown in Fig. 1b, an aerosol was atomized from the aqueous solutions of pure compounds in distilled water (2–6 g L^{-1}) using a custom-built Collison-type atomizer (Liu and Lee, 1975) with dry nitrogen flow. The atomizer output was diluted with dry, particle-free nitrogen, and dried with a silica gel diffusion dryer, which reduced the relative humidity to less than 10%. Subsequently, particles were size-selected using a custom-built differential mobility analyzer (DMA) (Knutson and Whitby, 1975). Prior to entering the DMA column, a steady-state charge distribution was achieved using a 20 mCi Po 210 source. The DMA was operated with a fixed applied voltage, a sample flow of 0.3–0.5 volumetric liters per minute (vLpm), and a ratio of 10 : 1 between the sheath flow and sample flow. The particle concentration after the three optical channels was measured continuously using a condensation particle counter (CPC; 3022A, TSI Inc., Shoreview, MN, USA).

Measurements were performed for particle diameters from 200 to 800 nm in 100 nm steps (excluding 700 nm) for PSL and in 50 nm steps for AS, SRFA, and nigrosin. For each measured PSL diameter, the particle loss through the system was both measured and modeled (von der Weiden et al., 2009), with agreement of 1% between the two

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methods. The particle loss was included in calculations of the extinction cross-section for each cavity, with the assumption that particle loss was evenly divided among the three cavities, which were connected in series.

The uncertainty in the diameter of spherical particles selected by the DMA is $\leq 1\%$, as determined by measuring the DMA transfer function for five sizes of NIST-traceable PSL standards. Treatment of multiply-charged particles in the size distribution produced by the DMA selection of aerosols generated from solution is described in Sect. 3.3.2.

2.4 Flow system

The size-selected aerosol from the DMA was sampled by the three cavities in series, as shown in Fig. 1b. The flow system consisted entirely of aluminum, stainless steel, and conductive silicone tubing to minimize electrostatic particle losses. A constant flow of $1.35(\pm 0.01)$ vLpm through the three cavities was provided by the condensation particle counter. The measured pressure drop through the three cells was less than 0.5 hPa. Small N_2 purge flows of $35\text{--}50$ volumetric $\text{cm}^3 \text{min}^{-1}$ (vccm) were added at the mirrors to maintain their cleanliness. These flows were produced using six critical orifices supplied by a regulated 70 kPa of N_2 . The dilution of the ambient sample by the mirror purge flows was accounted for in the data analysis. Temperature (K-type thermocouple) and pressure (PPT, Honeywell, International Inc., Plymouth, MN, USA) were measured at the CRDS cell.

Calibration gases were added by pushing through the system. This method was used to fill the cells with He and N_2 , which were used to calculate the mirror reflectivity.

2.5 Operation

During normal laboratory operation, the BAES, CRDS, and aerosol generation and counting instruments were turned on at least 1 h prior to beginning measurements to allow the temperatures and flows to stabilize. The first step in the measurement

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sequence was to determine the dark noise of the CCD detector by acquiring spectra under dark conditions, which were then scaled to the integration time and subtracted from subsequent spectra. Next, He and N₂ spectra were separately measured for one minute each to calculate the mirror reflectivity of the BAES channels, as described below. Next, aerosol samples were generated, and pulled through the flow system using the CPC. N₂ spectra were recorded regularly, usually after three or four different aerosol sizes were measured. Each aerosol and N₂ measurement reported is an average of 150 spectra integrated for 0.4 s each, for a total integration time of 1 min. The reflectivity measurement and calculation was typically repeated hourly.

3 Data analysis

3.1 Determination of aerosol optical extinction from BAES

The expression for the optical extinction in the BAES cavities may be derived using either an infinite sum for the light transmission through the cavity (Fiedler et al., 2003) or a differential equation for light input and output (Washenfelder et al., 2008). These two approaches give an equivalent, general expression that relates the optical extinction, $\alpha(\lambda)$, to the observed change in transmitted light intensity through the cavity as

$$\alpha(\lambda) = \left(\frac{1 - R(\lambda)}{d} \right) \left(\frac{I_0(\lambda) - I(\lambda)}{I(\lambda)} \right) \quad (1)$$

where λ is the wavelength of light, d is the cavity length, $R(\lambda)$ is the mirror reflectivity, $I_0(\lambda)$ is the reference spectrum (an empty cavity for the expression above), and $I(\lambda)$ is the measured spectrum at each wavelength. For our experimental conditions, we more conveniently define $I_0(\lambda)$ as a spectrum of N₂ and explicitly account for its Rayleigh scattering. Additionally, for aerosol measurements, we also account for the portion of the cell volume that is occupied by mirror purge flow. We define the ratio of the total cell length to the sample cell length as R_L . Incorporating these changes, we derive the

expression:

$$\alpha(\lambda) = R_L \left(\frac{(1 - R(\lambda))}{d} + \alpha_{\text{Rayleigh } N_2}(\lambda) \right) \left(\frac{I_{N_2}(\lambda) - I(\lambda)}{I(\lambda)} \right) \quad (2)$$

where $I_{N_2}(\lambda)$ is the spectrum of N_2 and $\alpha_{\text{Rayleigh } N_2}(\lambda)$ is the extinction due to Rayleigh scattering by N_2 .

For the aerosol measurements, R_L may either be measured using a known extinction, such as a dilute concentration of NO_2 , or defined geometrically from the assumption that aerosols follow the gas flow path and are not present in the purge volumes. In this analysis, we used R_L values measured from the optical extinction of ~ 70 ppbv of NO_2 with and without the mirror purges (i.e., with 70 ppbv of NO_2 filling only the central portion of the cell, or its entire length).

In order to use Eq. (2), it is necessary to first determine $R(\lambda)$. Mirror reflectivity (or effective cavity extinction, defined as $(1 - R(\lambda))/d$) can be determined by introducing a known extinction into the cavity. Rayleigh extinction is useful for this calculation because it varies slowly with wavelength in a well-defined manner across the entire region of interest, and requires only measurements of temperature and pressure. In this work, we calculated $R(\lambda)$ using Rayleigh scattering cross-sections for N_2 fitted to Bodhaine et al. (1999) data ($\sigma_{\text{Rayleigh } N_2} = 1.2577 \times 10^{-15} \times \lambda^{-4.1814}$) and He fitted to Shardanand and Rao (1977) data ($\sigma_{\text{Rayleigh He}} = 1.336 \times 10^{-17} \times \lambda^{-4.1287}$).

3.2 Determination of aerosol optical cross-section from BAES

For the cavity containing only particles in N_2 gas, the relationship between aerosol optical cross-section and aerosol optical extinction is equal to:

$$\sigma(\lambda, D_p, m) = \frac{\alpha(\lambda, D_p, m)}{N(D_p)} \quad (3)$$

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where $\alpha(\lambda, D_p, m)$ is the aerosol extinction cross-section and $N(D_p)$ is the aerosol number density.

The aerosol cross-section in Eq. (3) is a function of the wavelength of incident radiation (λ), the diameter of the particle (D_p), and the complex refractive index (m). The complex refractive index, m , for a particle relative to the surrounding medium is further described as:

$$m = n + ik \quad (4)$$

where the real part, n , represents the scattering component and the imaginary part, k , represents the absorbing component. Particles with $k = 0$ are purely scattering, while particles with $k > \sim 0.25$ are typically described as strongly absorbing. Slightly absorbing particles fall between these two extremes. The generalized expression for σ is:

$$\sigma = f(\lambda, D_p, n, k) \quad (5)$$

3.3 Determination of the complex refractive index from BAES

The simplified expression for σ in Eq. (5) conceals a complex dependence on the variables λ , D_p , n , and k . However, for given values of λ , D_p , n , and k and particles that are assumed to be spherical, Eq. (5) may be readily solved using numerical codes to determine the optical cross-section (Bohren and Huffman, 1983) as demonstrated previously (Pettersson et al., 2004; Lack et al., 2006; Abo Riziq et al., 2007; Flores et al., 2009; Lang-Yona et al., 2009; Nakayama et al., 2010; Miles et al., 2010; Flores et al., 2012; Chartier and Greenslade, 2012; Bluvshstein et al., 2012).

The goal of this work is to determine n and k for different aerosol types by measuring the wavelength dependence of σ for particles of known N and D_p . As Eq. (5) indicates, n and k can be determined using two approaches: (1) for particles with a slight spectral dependence of the refractive index (e.g. PSL, ammonium sulfate), measurement of a single diameter or size distribution at multiple wavelengths can be used to determine

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n and k ; (2) for aerosols with substantial spectral dependence of the refractive index (e.g. Suwannee River fulvic acid), measurement of multiple diameters can be used to determine n and k for each wavelength. This is similar to the approach used in conventional CRDS measurements (Pettersson et al., 2004; Abo Riziq et al., 2007; Flores et al., 2012), although used for multiple wavelengths in this case. The implementation of these approaches is described in further detail below.

3.3.1 Determination of the complex refractive index for monodisperse aerosols neglecting spectral dependence

For particles whose refractive index does not vary strongly with wavelength over the selected wavelength region, the extinction cross-section given in Eq. (5) is a function of only D_p , n , and k . The refractive index may therefore be retrieved from the extinction cross-section measured at multiple wavelengths for a single particle diameter. We apply this retrieval to monodisperse aerosol (PSL) and aerosol whose refractive index is sufficiently well-known to correct for multiply-charged particles (e.g. ammonium sulfate).

In this approach, the measured extinction cross-section for each diameter is compared to values calculated from Mie theory. The real and imaginary components of the complex refractive index for the calculated cross section are varied to minimize the expression

$$\chi^2 = \sum_{i=1}^{N_\lambda} \frac{(\sigma_{\text{measured}} - \sigma_{\text{calculated}})_i^2}{\varepsilon_i^2} \quad (6)$$

where ε is the estimated error in the individual measurement and N_λ is the number of wavelengths. The uncertainties are calculated from the deviation of χ^2 in the vicinity of the minimum. Assuming the errors of the measurement to be normally distributed, the values of χ^2 for different measurements follow a χ^2 distribution for the two degrees of freedom n and k . The quantile for the 1σ (68.3 %) confidence level for this parameter

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set is 2.298. Any measurement that falls within the 1σ error bound of the best measurement is considered acceptable if $\chi^2 < \chi_{\min}^2 + 2.298$ (Spindler et al., 2007; Dinar et al., 2008).

3.3.2 Determination of complex refractive index as a function of wavelength for polydisperse aerosol

A more general approach to retrieve the complex refractive index uses multiple aerosol diameters and does not assume that n and k are invariant with wavelength. In this case, $\sigma(\lambda, D_p, n, k)$ is determined from $\alpha(\lambda, D_p, n, k) / N(D_p)$. PSL have narrow, monodisperse, size distributions, and $N(D_p)$ may be accurately represented as a single value, N . In contrast, an aerosol produced from liquid solution (or sampled from the ambient atmosphere) and classified with a DMA has a broader and multi-peaked population that consists of singly-, doubly-, and triply-charged particles, each having a finite dispersion. The larger multiply-charged particles will generally have a greater extinction cross-section than the population of singly-charged particles, leading to an overestimation of the extinction cross-section for the selected, primary diameter.

A common method used in CRDS retrievals of complex refractive index is to calculate and subtract the extinction due to doubly- and triply-charged particles from the total measured extinction (Fierz-Schmidhauser et al., 2010; Cappa et al., 2011; Flores et al., 2012). The disadvantage of this method is that it requires an assumed refractive index for the multiply charged particles. To avoid this, we used the full aerosol size distribution to determine the best fit n and k values. We measured the aerosol size distribution atomized for each solute concentration using scanning electrical mobility spectroscopy (Wang and Flagan, 1990, using Markowski's, 1987, version of the Twomey, 1975, algorithm). We then used DMA transfer theory (Knutson and Whitby, 1975) with Weidensohler's steady-state charge distribution approximation (1988, with subsequent erratum) to quantify the full size distribution of singly-, doubly-, and triply-charged particles exiting the DMA for each primary diameter setting.

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For each cross-section measurement, we calculated “effective” Mie cross-sections as a function of n and k corresponding to the measured size distribution. The best fit values of n and k are determined by minimizing the difference between σ_{measured} and $\sigma_{\text{calculated}}$. This method requires measurements for at least two different diameter setpoints, because a single measurement of $\sigma(\lambda, D_p, n, k)$ for one wavelength and one aerosol size distribution represents one equation and two unknowns (n and k). The solution is determined by measuring the cross-section for multiple diameter setpoints and minimizing the expression:

$$\chi^2 = \sum_{i=1}^{N_{D_p}} \left(\frac{\sigma_{\text{measured}} - \sigma_{\text{calculated}}}{\sigma_{\text{measured}}} \right)_i^2 \quad (7)$$

This expression appears similar to Eq. (6), but represents a separate, independent calculation for each wavelength element. We determine the absolute uncertainty in the retrieved n and k empirically by varying σ_{measured} by the uncertainty of the measurement ($\pm 4.1\%$ as described in Sect. 3.4).

3.4 Precision and accuracy of the optical extinction cross-sections and refractive indices

The precision of the $\alpha(\lambda, D_p, m)$ measurement is 0.21 Mm^{-1} and 0.17 Mm^{-1} ($1 \text{ Mm}^{-1} = 10^{-8} \text{ cm}^{-1}$) for 1 min BAES measurements at the center wavelengths of 365 nm and 405 nm. This precision is limited by noise in the spectra, as the system is shot-noise limited at the CCD detector. The precision of the $\sigma(\lambda, D_p, m)$ value additionally includes the sampling statistics of the aerosol in the optical beam, although this makes a minor contribution. This is an important difference between the BAES and CRDS, because the broadband beam has a much greater diameter than a laser, and thus intersects a larger sample volume (see discussion of CRDS in Pettersson et al., 2004; Baynard et al., 2007).

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The absolute accuracy of the $\sigma(\lambda, D_p, m)$ cross-section values is limited by the uncertainty in the Rayleigh scattering cross-sections for N_2 and He, sample pressure and temperature, total flow, individual mirror purge flows, R_L , particle concentration, particle size distribution, and particle losses in the cavities. The uncertainty in the Rayleigh scattering cross-sections of N_2 is $\pm 2\%$. Although the uncertainty in the Rayleigh scattering cross-section for He is similar, its cross-section is small and makes a minor contribution to the total uncertainty. The uncertainty in the sample pressure is $\pm 0.3\%$ as determined by comparison to a Hg manometer and by confirming that the pressure drop for the three cells is less than 0.5 hPa. The uncertainty in the sample temperature is approximately $\pm 0.2\%$. The uncertainties in the total flow and mirror purge flows contribute to an uncertainty in the sample dilution correction of 0.3%, 0.2%, and 0.1% respectively for the BAES 405 nm, BAES 365 nm, and CRDS, with the largest error for the cell farthest from the CPC because it requires the greatest flow dilution correction. We estimate an uncertainty in R_L of 2%, because R_L for aerosol may differ from the value determined using NO_2 . Particle losses through the system were both measured and modeled, and we estimate a 2% uncertainty in the particle loss correction. Uncertainty in the particle size distribution is neglected here. The manufacturer's reported uncertainty for the TSI 3022A CPC is $\pm 10\%$ for particle concentrations less than $50\,000\text{ cm}^{-3}$. However, we estimate the actual uncertainty in particle concentration to be limited by the CPC sample flow rate and digital counting statistics. We calibrated the CPC sample flow rate with an uncertainty of 1%. The uncertainty in the digital counting is expected to follow Poisson statistics, with a 60 s sample time giving an uncertainty of 1.8% for $10\text{ particles cm}^{-3}$ and 0.3% for $300\text{ particles cm}^{-3}$ (for a $5.3\text{ cm}^3\text{ s}^{-1}$ detector flow rate). The addition of all uncertainties in quadrature, with the omission of the uncertainty in particle size distribution, gives a total uncertainty of 3.6–4.1% for the aerosol extinction cross-sections.

4 Results and discussion

4.1 Polystyrene latex spheres

We measured six different diameters of PSL (199, 299, 404, 499, 596, and 802 nm; see Table 1), with five or more concentration measurements for each diameter, to test the two refractive index retrieval methods described in Sect. 3. These measurements allow us to calculate the extinction cross-section for every wavelength in both BAES channels and for the single wavelength of the CRDS. As Eq. (3) shows, we expect a linear relationship between $\alpha(\lambda, D_p, m)$ and $N(D_p)$, with a slope equal to $\sigma(\lambda, D_p, m)$ (Pettersson et al., 2004; Abo Riziq et al., 2007). For single measurements of $\alpha(\lambda, D_p, m)$ at a given $N(D_p)$, we may determine $\sigma(\lambda, D_p, m)$ from the ratio of $\alpha(\lambda, D_p, m) / N(D_p)$.

Figure 2 shows the determination of $\sigma(\lambda, D_p, m)$ for multiple measurements of $\alpha(\lambda, D_p, m)$ at different $N(D_p)$ for a single wavelength element of the BAES (407.025 nm). The CRDS at 407.0 nm is also plotted, and the BAES and CRDS cross-sections agree within 2% for 600 nm diameter particles, and to better than 1% for smaller particles.

For each of the six PSL diameters, the value of σ was determined from the slope for each wavelength element of the BAES 360–390 and 385–420 nm channels. The resulting σ values are plotted in Fig. 3, together with the CRDS value at 407.0 nm. The agreement at 407 nm between the CRDS and BAES measurements in Fig. 3a is an important validation of the BAES measurements. Furthermore, there is agreement between the two independent BAES instruments in the region from 385–390 nm where they overlap spectrally, with a maximum difference in σ of 2.6%.

4.1.1 Determination of complex refractive index for PSL neglecting spectral dependence

Refractive indices were retrieved from extinction cross-sections for each PSL diameter using the method described in Sect. 3.3.1. This analysis was independently performed

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for PSL cross-section measurements by each of the two BAES channels, and for their combined spectral range from 360 to 420 nm (Fig. 3a). The results are summarized in Table 1, which shows consistency in the complex RI retrieved for different PSL sizes. The complex RI for the PSL describes an aerosol that is strongly scattering, with an absorbing component that is not significantly different from zero. As a result, the fits in Fig. 3a clearly show Mie resonances for the different particle sizes, and these are more visible for the expanded cross-sections shown in Fig. 3b and c for 299 and 802 nm diameter particles.

4.1.2 Determination of complex refractive index for PSL as a function of wavelength

The complex RI was also determined using the full retrieval (described in Sect. 3.3.2), which combines measurements made at several particle diameters and allows n and k to vary with wavelength. For the calculations, we assumed a Gaussian size distribution with center diameter and standard deviation given by the PSL manufacturer. DMA-classified PSL particles are nearly monodisperse, without multiply-charged particles.

The results for the retrievals of PSL refractive index as a function of wavelength are shown in Table 1 and Fig. 4. The error bars in Fig. 4 were calculated by scaling the measured extinction cross-section by an uncertainty of $\pm 4.1\%$. The real part of the refractive index (Fig. 4a) is $n = 1.657(\pm 0.002)$ (Table 1) at 360 nm, and decreases to $1.637(\pm 0.005)$ at 420 nm. The imaginary part of the refractive index (Fig. 4b) does not differ significantly from zero across the measured spectral region, with the exception of retrieved values reaching 0.01 (± 0.002) at 410 nm.

4.1.3 Comparison of PSL refractive index to published values

Polystyrene latex spheres are widely used to calibrate optical aerosol instruments. Refractive indices have been reported for bulk samples (Matheson and Saunderson, 1952; Nikolov and Ivanov, 2000; Sultanova et al., 2003), particles in suspension

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(Bateman et al., 1959; Smart and Willis, 1967; Ma et al., 2003; French et al., 2007), and aerosol particles (Pettersson et al., 2004; Abo Riziq et al., 2007; Baynard et al., 2007; Lang-Yona et al., 2009; Chartier and Greenslade, 2012). These measurements are nicely summarized by Miles et al. (2010), with the exception of the more recent Chartier and Greenslade (2012) result.

Only two prior studies have reported the complex refractive index of PSL particles at ultraviolet wavelengths. Baynard et al. (2007) measured four PSL diameters using a pulsed CRDS system at $\lambda = 355$ nm. The authors reported an agreement of 5% with the refractive index values in Nikolov and Ivanov (2000), but did not independently retrieve an RI. Chartier and Greenslade (2012) measured refractive indices at $\lambda = 248$ and 335 nm. To our knowledge, ours is the first study to measure the refractive index for PSL as a function of wavelength between $\lambda = 360$ –420 nm.

As shown in Table 1 and Fig. 4, the BAES measurements give a real part of the refractive index of $n = 1.64$ –1.66 at $\lambda = 360$ –420 nm, increasing with decreasing wavelength. The studies of bulk and suspended polystyrene summarized by Miles et al. (2010) also indicate that the real part of the refractive index increases slightly with decreasing wavelength, with a value of $n = 1.60$ –1.62 at 400 nm. These values are lower than the retrieved values reported here. However, the values reported here are more consistent with Chartier and Greenslade (2012), who recently measured a real part of the refractive index $n = 1.81(\pm 0.03)$ at $\lambda = 248$ nm and $n = 1.72(\pm 0.02)$ at $\lambda = 335$ nm, which is greater than the results reported for particles in suspension by French et al. (2007) and Ma et al. (2003).

As shown in Table 1 and Fig. 4, the BAES measurements give an imaginary part of the refractive index that is not significantly different from zero at $\lambda = 360$ –404, but shows an increase to $k = 0.01$ centered at 410 nm. Chartier and Greenslade (2012) reported values of $k = 0.05$ and 0.005 for PSL aerosol at $\lambda = 248$ nm and $\lambda = 355$ nm, respectively. Studies of PSL aerosol at $\lambda = 532$ nm have reported imaginary components of $k = 0.01$ and $k = 0.0$ (Abo Riziq et al., 2007; Lang-Yona et al., 2009). Overall, our n and k values are consistent with the prior literature values.

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4.2 Ammonium sulfate aerosol

To test the performance of the BAES with broader size distributions and an atmospherically-relevant purely scattering substance, we measured ammonium sulfate (AS) at DMA diameter setpoints from 200 to 700 nm in 50 nm steps. The extinction cross-section (corrected for multiply charged particles; see next section) for all the measured sizes for both BAES and the CRDS at 407 nm are shown in Fig. 5. The open squares and circles show the results for the BAES at 360–390 nm and 385–420 nm, respectively. The CRDS measurements are shown as open triangles. Figure 5 shows agreement in the overlap region of the two BAES channels, and for the BAES and CRDS at $\lambda = 407$ nm.

4.2.1 Determination of complex refractive index for ammonium sulfate neglecting spectral dependence

Using the wavelength-independent approach (Sect. 3.3.1), we retrieved the complex refractive index from extinction measurements shown in Fig. 5, after correcting for multiply-charged particles. The assumed refractive index used for the subtraction of extinction due to multiply-charged particles was $1.52 + 0i$ (Abo Riziq et al., 2008), with no wavelength dependence. The refractive index was independently determined for 13 aerosol diameters from 200–800 nm, with an average value of $1.51(\pm 0.01) + 0.02(\pm 0.01)i$.

4.2.2 Determination of complex refractive index for ammonium sulfate as a function of wavelength

Results from the full retrieval of complex refractive indices as a function of wavelength, derived using the full size distribution, are shown in Fig. 6 for the two BAES channels. The retrieval gives a value of $n = 1.513(\pm 0.004)$ at 360 nm, which increases to

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$n = 1.540(\pm 0.007)$ at 420 nm. The retrieved imaginary part of the refractive index is $0.000(\pm 0.001)$ for all wavelengths.

4.2.3 Comparison of ammonium sulfate refractive index to published values

Aerosols containing ammonium sulfate are some of the most abundant in the troposphere, and have a significant contribution to Earth's radiative budget (Forster et al., 2007). Hence, the optical properties of AS are important to atmospheric radiative transfer models and to calculations of climatic forcing. CRDS studies have retrieved complex RI values at $\lambda = 532$ nm with values for the real component between $n = 1.51$ – 1.53 and an imaginary component of $k = 0$ (Abo Riziq et al., 2007; Spindler et al., 2007; Lang-Yona et al., 2009; Flores et al., 2009). There are few measurements in the near UV region of the complex refractive index of AS, and the relevant values are all included in Fig. 6. At 355 nm, Flores et al. (2012) and Trainic et al. (2011) reported values of $m = 1.507(\pm 0.024) + 0.005(\pm 0.025)i$, and $m = 1.553(\pm 0.005) + 0.002(\pm 0.013)i$, respectively. At 390 nm, Dinar et al. (2008) reported a value of $m = 1.525(\pm 0.002) + 0.0001(\pm 0.0002)i$. At 405 nm, Toon et al. (1976) reported a real value of $n = 1.53$ – 1.55 and an imaginary value of $k < 1 \times 10^{-7}$ determined from the reflectivity and transmission of crystalline ammonium sulfate samples. Our results are consistent with these prior measurements, with the exception of Trainic et al. (2011), as shown in Fig. 6.

4.3 Nigrosin aerosol

Nigrosin aerosol is an organic black dye that is commonly used as a model for absorbing particles. We measured extinction by nigrosin particles at DMA diameter setpoints from 200 to 700 nm in 50 nm steps. Extinction cross-sections for measured sizes between 200 and 500 nm from both BAES and the CRDS at 407 nm are shown in Fig. 7 and used in the refractive index retrievals. At larger diameters, fewer nigrosin particles were generated and accurately quantifying the fraction of triply-charged particles with the DMA was difficult. Unlike the ammonium sulfate cross-sections shown in Fig. 5,

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the plotted nigrosin cross-sections have not been corrected for doubly- and triply-charged particles. This is because there are few reported values for the refractive index of nigrosin in the ultraviolet and near-visible spectral regions which would allow the contribution of doubly- and triply- charged particles to be calculated and subtracted (see Sect. 4.3.3 for a summary of published values). Instead, Fig. 7 shows the size-distribution-weighted cross-sections for each DMA diameter setpoint, calculated from $\alpha(\lambda, D_p, n, k)/N$.

4.3.1 Determination of complex refractive index for nigrosin as a function of wavelength

A wavelength-independent retrieval is not appropriate for nigrosin, because the spectral dependence of its refractive index for particles is poorly known and the absorbance of its bulk solution varies strongly with wavelength. Refractive index values determined using the full measured particle size distribution, are shown in Fig. 8a and b. The retrieval gives a value of $n = 1.78(\pm 0.03)$ at 360 nm, which decreases to $n = 1.63(\pm 0.03)$ at 420 nm. The imaginary part of the refractive index is $k = 0.19(\pm 0.08)$ at 360 nm, and $k = 0.21(\pm 0.05)$ at 420 nm, with no wavelength dependence detectable within the measurement precision.

4.3.2 Comparison of nigrosin refractive index to published values and UV-visible absorption spectra

Prior studies have reported the refractive index for nigrosin aerosol in the visible spectral region (Garvey and Pinnick, 1983; Lack et al., 2006; Flores et al., 2009; Lang-Yona et al., 2009; Bluvshstein et al., 2012). The reported values are generally consistent: $1.67 + 0.26i$ at 633 nm (Garvey and Pinnick, 1983); $1.70(\pm 0.04) + 0.31(\pm 0.05)i$ at 532 nm (Lack et al., 2006); $1.649(\pm 0.007) + 0.238(\pm 0.008)i$ at 532 nm (Dinar et al., 2008); $1.72(\pm 0.01) + 0.28(\pm 0.08)i$ and $1.65(\pm 0.01) + 0.24(\pm 0.01)i$ at 532 nm (Lang-Yona et al., 2009); and $1.64(\pm 0.04) + 0.20(\pm 0.03)i$ for a white light source at

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370–780 nm (Flores et al., 2009). No trend with wavelength is evident from the reported data. Although no refractive index values have been reported in the blue or ultraviolet spectral regions, the n and k values reported here for 360–420 nm are consistent within ± 0.01 of the values at 532 nm.

5 4.4 Suwannee River fulvic acid aerosol

Suwannee River fulvic acid is common model for humic-like substances in aged aerosols, and was selected as a surrogate for brown carbon aerosol because it is thought to have strongly wavelength-dependent absorption. We measured extinction by Suwannee River fulvic acid particles at DMA diameter setpoints from 200 to 700 nm in 50 nm steps. Extinction cross-sections for measured sizes between 200 and 500 nm from both BAES and the CRDS at 407 nm are shown in Fig. 9 and used in the refractive index retrievals. Similarly to the nigrosin cross-sections in Fig. 7, the plotted SRFA cross-sections have not been corrected for doubly- and triply-charged particles, as there are few reported values for the refractive index of SRFA in the ultraviolet and near-visible spectral regions which would allow the contribution of doubly- and triply-charged particles to be accurately calculated and subtracted, and also because the refractive index for SRFA varies with wavelength (see Sect. 4.4.3 for a summary of published values). Instead, Fig. 9 shows the size-distribution-weighted cross-sections for each DMA diameter setpoint, calculated from $\alpha(\lambda, D_p, n, k)/N$.

20 4.4.1 Determination of complex refractive index for Suwannee River fulvic acid as a function of wavelength

Similar to nigrosin, a wavelength-independent retrieval is not appropriate for SRFA, because its absorption varies as a function of wavelength (Dinar et al., 2008) and is not well known in the ultraviolet spectral region. Refractive index values determined using the full measured particle size distribution are shown in Fig. 10. The retrieval gives a value of $n = 1.71(\pm 0.02)$ at 360 nm, which decreases to $1.66(\pm 0.02)$ at 420 nm.

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The imaginary part of the refractive index is $k = 0.07(\pm 0.06)$ at 360 nm, decreasing slightly to a value of $0.06(\pm 0.04)$ at 420 nm.

4.4.2 Comparison of Suwannee River fulvic acid refractive index to published values

5 Prior studies have reported the refractive index for Suwannee River fulvic acid aerosol in the visible spectral region (Dinar et al., 2008; Flores et al., 2009; Lang-Yona et al., 2009). However, it is important to note that different SRFA samples have been supplied by the International Humic Substances Society, and that we have used SRFA 1S101F for this work. For sample 1R101F, (Dinar et al., 2008) reports values of $1.602(\pm 0.005) + 0.098(\pm 0.005)i$ at 390 nm and $1.634(\pm 0.004) + 0.021(\pm 0.005)i$ at 532 nm. Previous studies have reported $1.520(\pm 0.003) + 0.02(\pm 0.01)i$ and $1.65(\pm 0.01) + 0.02(\pm 0.01)i$ for cavity ringdown measurements (Lang-Yona et al., 2009) at 532 nm and $1.53(\pm 0.01) + 0.0i$ for a white light source at 370–870 nm (Flores et al., 2009). Figure 10 shows the only measurement previously reported in the near-ultraviolet spectral region, with a value of $m = 1.602(\pm 0.005) + 0.098(\pm 0.005)i$ at 390 nm for SRFA 1R101F (Dinar et al., 2008).

5 Discussion of uncertainty in retrievals of refractive index

The error bars for each of the wavelength-dependent retrievals were calculated by scaling the measured extinction cross-section by an uncertainty of $\pm 4.1\%$. However, the resulting error bars differ widely for the four compounds. The uncertainty in n is 0.001–0.002 for ammonium sulfate, while it is 0.01–0.03 for nigrosin. For k , these uncertainty values are 0.001 and 0.05–0.09, respectively. Nigrosin has much greater uncertainty in its refractive index because it is a strongly absorbing aerosol and a relatively large change in k for an absorbing aerosol causes only a small change in the aerosol's

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extinction cross-section, making it more difficult to accurately retrieve the refractive index for absorbing compounds.

The greater difficulty in retrieving refractive indices for absorbing aerosol is demonstrated in Fig. 11, which shows the theoretical refractive indices for a series of purely scattering and strongly absorbing aerosol at three discrete diameters. The three extinction cross-sections of the purely scattering aerosol ($m = 1.5 + 0i$, $1.6 + 0i$, and $1.7 + 0i$) can be easily distinguished. By contrast, the differences between the extinction cross-sections of the strongly absorbing aerosol ($m = 1.5 + 0.3i$, $1.6 + 0.3i$, and $1.7 + 0.3i$) are much smaller, particularly for 600 nm diameter particles.

6 Summary and conclusions

We have constructed a laboratory instrument that accurately measures aerosol optical extinction in the 360–420 nm spectral range. At each aerosol diameter selected by the DMA, we measure aerosol optical extinction and the concentration of particles, which can be used to calculate the aerosol extinction cross-section. By combining multiple measurements of aerosol extinction cross-section at different diameters, we are able to retrieve a refractive index as a function of wavelength.

We have applied this method to measure aerosol cross-sections for different aerosols generated in the laboratory, including PSL, ammonium sulfate, nigrosin, and SRFA aerosol. For each of these compounds, we derive refractive indices that are consistent with results in the literature. Further, these results are the first reported refractive index measurements as a function of wavelength in the ultraviolet and near-visible spectral regions for each of these species. For SRFA we observe a mild spectral dependence in the imaginary part of the refractive index, as has been predicted by bulk solution studies. The refractive index values are directly useful for global radiative models, and this method may be used to measure other unknown aerosol types.

The results presented here suggest three possibilities for a field instrument to measure aerosol optical properties. First, BAES cavities could be constructed to measure

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the optical extinction of an ambient aerosol sample as a function of wavelength. This follows the approach of existing CRDS field instruments that operate at a single wavelength and report optical extinction (Strawa et al., 2003; Moosmuller et al., 2005; Baynard et al., 2007; Langridge et al., 2011). The strong absorption by gases in the ultraviolet and visible spectral region, particularly by NO₂, requires either a correction based on independent measurements of the absorbing species (e.g. Langridge et al., 2011) or selective removal of gas-phase absorbers (e.g. using an activated charcoal filter). Second, by combining the BAES cavities with a DMA and CPC as demonstrated in this work, optical cross-sections could be determined for selected aerosol sizes as a function of wavelength. Third, by measuring multiple optical cross-sections at different aerosol sizes, the refractive index could be determined as a function of wavelength. Careful consideration of optimum D_p setpoints to determine the refractive index, as described in Bluvshstein et al. (2012), would allow an improved time response for this approach. Our current refractive index retrieval assumes that the singly-, doubly-, and triply-charged particles exiting the DMA share the same composition and optical properties, and that they are approximately spherical. In the ambient atmosphere, particles of a single diameter may have different compositions, composition may vary strongly with particle diameter, particles may not be internally-well mixed, and particles may not be spherical. These challenges could be minimized by selecting particles of a single size and density (Beranek et al., 2012). Further, bulk aerosol composition as a function of size and single-particle composition could be quantified with laser mass spectrometry instruments (Pratt and Prather, 2012) to evaluate how these affect the refractive index retrieval.

Each of the three experimental approaches described above provides wavelength-resolved measurements of aerosol optical properties, and could potentially be combined with additional measurements of aerosol composition, aging, hygroscopicity, and other properties to more fully describe aerosol properties and processes in the atmosphere.

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Table 1. Complex refractive index determined for polystyrene latex spheres using two methods.

Diameter (nm)	Retrieval with no wavelength dependence		
	360–390 nm	385–420 nm	360–420 nm (combined)
199 ± 6	1.656(±0.01) + 0.0(±0.04) <i>i</i>	1.652(±0.007) + 0.0(±0.011) <i>i</i>	1.653(±0.008) + 0.0(±0.014) <i>i</i>
299 ± 6	1.653(±0.012) + 0.0(±0.014) <i>i</i>	1.644(±0.008) + 0.0(±0.068) <i>i</i>	1.646(±0.009) + 0.0(±0.029) <i>i</i>
404 ± 4	1.651(±0.029) + 0.008(±0.009) <i>i</i>	1.641(±0.011) + 0.003(±0.009) <i>i</i>	1.647(±0.015) + 0.008(±0.009) <i>i</i>
499 ± 5	1.642(±0.006) + 0.0(±0.013) <i>i</i>	1.624(±0.029) + 0.007(±0.008) <i>i</i>	1.639(±0.009) + 0.004(±0.009) <i>i</i>
596 ± 6	1.651(±0.005) + 0.0(±0.011) <i>i</i>	1.624(±0.009) + 0.016(±0.015) <i>i</i>	1.644(±0.010) + 0.0(±0.011) <i>i</i>
802 ± 6	1.643(±0.01) + 0.007(±0.009) <i>i</i>	1.628(±0.011) + 0.004(±0.006) <i>i</i>	1.634(±0.012) + 0.006(±0.008) <i>i</i>

Wavelength (nm)	Retrieval with wavelength dependence
360	1.651(±0.002) + 0.000(±0.001) <i>i</i>
370	1.653(±0.002) + 0.003(±0.003) <i>i</i>
380	1.643(±0.002) + 0.000(±0.001) <i>i</i>
390	1.639(±0.004) + 0.006(±0.004) <i>i</i>
400	1.63(±0.01) + 0.001(±0.001) <i>i</i>
410	1.63(±0.01) + 0.007(±0.002) <i>i</i>
420	1.633(±0.005) + 0.000(±0.005) <i>i</i>

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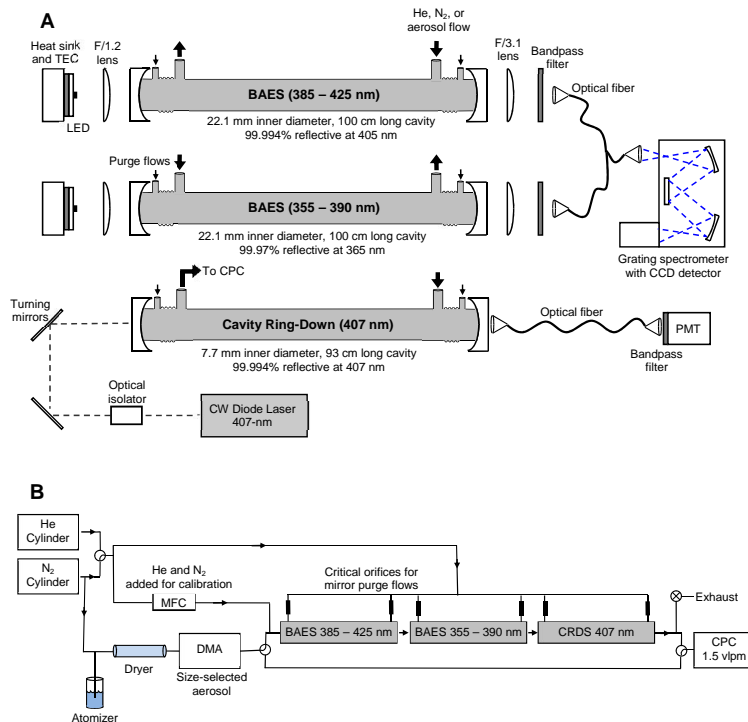


Fig. 1. (a) Schematic of the broadband aerosol extinction spectrometer, with channels for 360–390 and 385–420 nm. Each BAES channel consists of a temperature-controlled LED and optical cavity, with a shared grating spectrometer. Details of the 407 nm CRDS are given in Fuchs et al. (2009). **(b)** Block diagram showing aerosol generation and gas delivery. Aerosols are generated with an atomizer, and subsequently dried and size-selected using a differential mobility analyzer (DMA). The aerosol concentration is measured using a condensation particle counter (CPC).

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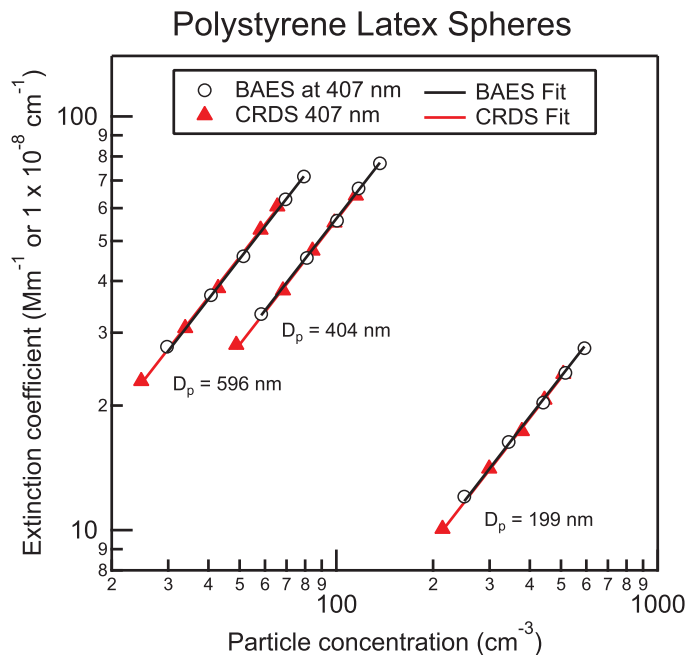


Fig. 2. Extinction coefficients as a function of particle concentration for polystyrene latex spheres (PSL) at 199, 404, and 596 nm diameters. CRDS measurements at $\lambda = 407$ nm (red triangles) are shown together with the measurements from a single wavelength element ($\lambda = 407.025$ nm) of the BAES (black circles). The extinction cross-sections calculated from linear fits to each diameter are $\sigma_{\text{CRDS}} = 4.73(\pm 0.05) \times 10^{-10} \text{ cm}^2$ and $\sigma_{\text{BAES}} = 4.70(\pm 0.02) \times 10^{-10} \text{ cm}^2$ at $D_p = 199$ nm; $\sigma_{\text{CRDS}} = 5.68(\pm 0.04) \times 10^{-9} \text{ cm}^2$ and $\sigma_{\text{BAES}} = 5.66(\pm 0.01) \times 10^{-9} \text{ cm}^2$ at $D_p = 404$ nm; and $\sigma_{\text{CRDS}} = 9.26(\pm 0.07) \times 10^{-9} \text{ cm}^2$ and $\sigma_{\text{BAES}} = 9.05(\pm 0.03) \times 10^{-9} \text{ cm}^2$ at $D_p = 596$ nm.

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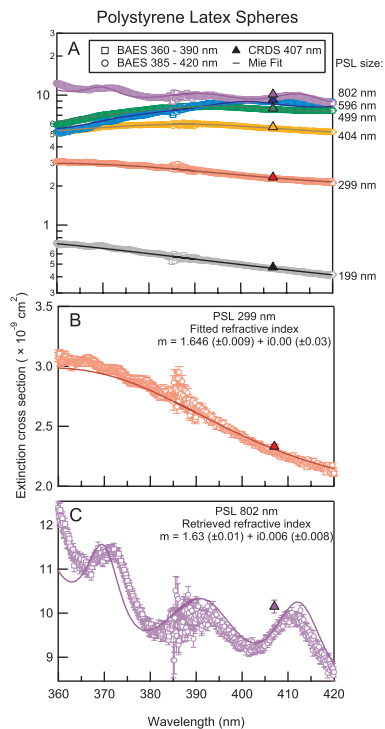


Fig. 3. (a) Extinction cross-sections as a function of wavelength for 199–802 nm diameter PSL measured with the BAES 360–390 nm (squares), BAES 385–420 nm (circles), and CRDS at $\lambda = 407$ nm (triangles). The solid lines show the best Mie fit. **(b)** Expanded view of the 299 nm PSL extinction cross-section measurements and Mie fit. The retrieved complex refractive index is given in the upper right hand corner. **(c)** Similar expanded view for 802 nm PSL.

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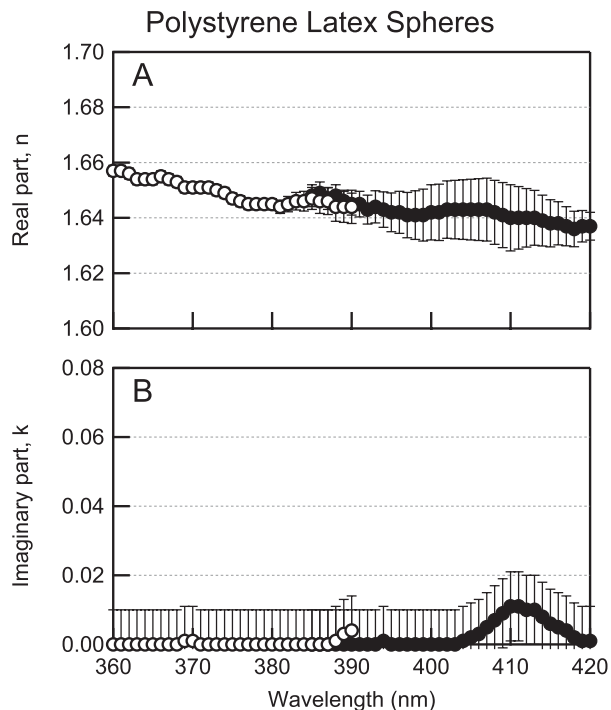


Fig. 4. (a) The real part of the complex refractive index (RI) as a function of wavelength for the BAES 360–390 nm (open circles) and the BAES 385–420 nm (filled circles). (b) The imaginary part of the RI.

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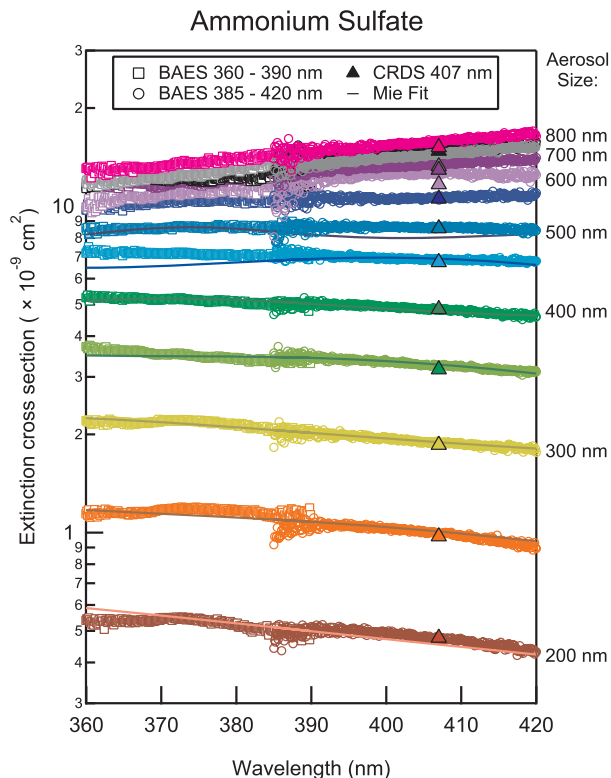


Fig. 5. Extinction cross-sections as a function of wavelength for ammonium sulfate aerosol at 200–800 nm diameter measured with the BAES 360–390 nm, BAES 385–420 nm, and CRDS at $\lambda = 407$ nm. The solid lines show the best Mie fit. The extinction cross-sections shown in this plot have been corrected for doubly- and triply-charged particles, using an assumed refractive index of $m = 1.52 + 0i$.

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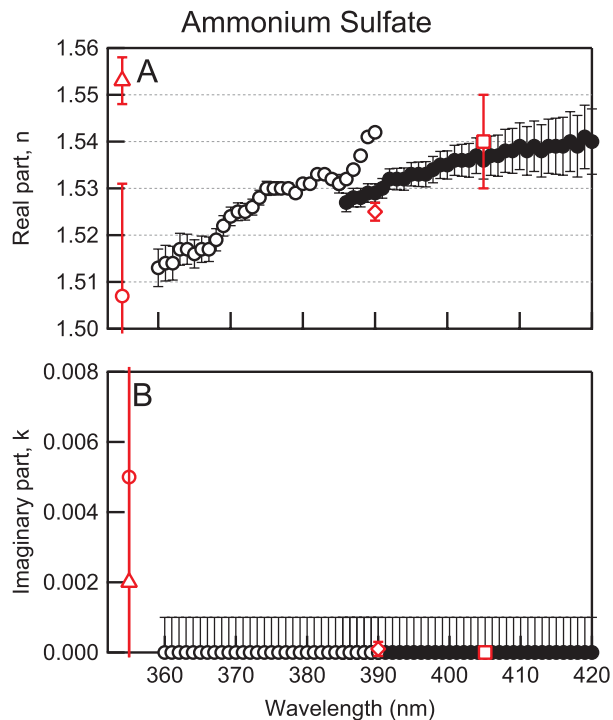


Fig. 6. (a) The real part of the complex refractive index (RI) as a function of wavelength for each wavelength from the BAES for ammonium sulfate aerosol. Red markers show values reported by Trainic et al. (2011, triangle), Flores et al. (2012, circle), Dinar et al. (2008, diamond), and Toon et al. (1976, square). **(b)** Similarly, the imaginary part of the RI.

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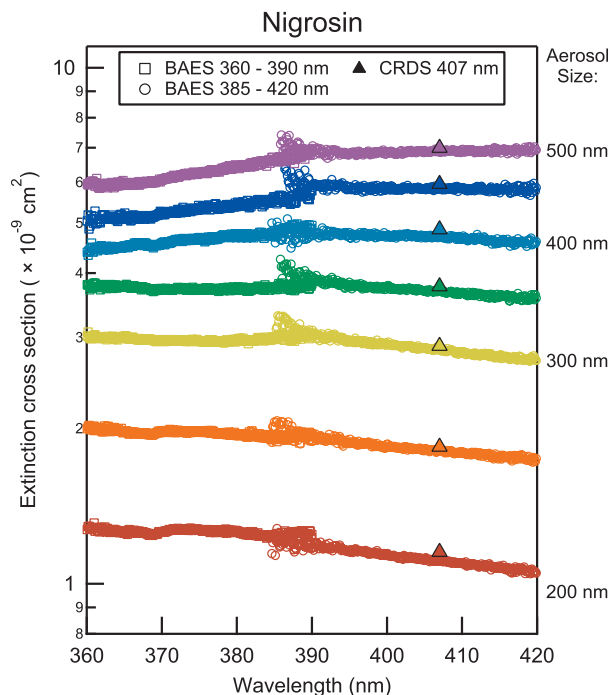


Fig. 7. Extinction cross-sections as a function of wavelength for nigrosin aerosol at 200–500 nm diameter measured with the BAES 360–390 nm, BAES 385–420 nm, and CRDS at $\lambda = 407$ nm. The extinction cross-sections shown in this plot have not been corrected for doubly- and triply-charged particles, because the refractive index of nigrosin as a function of wavelength is not well-known.

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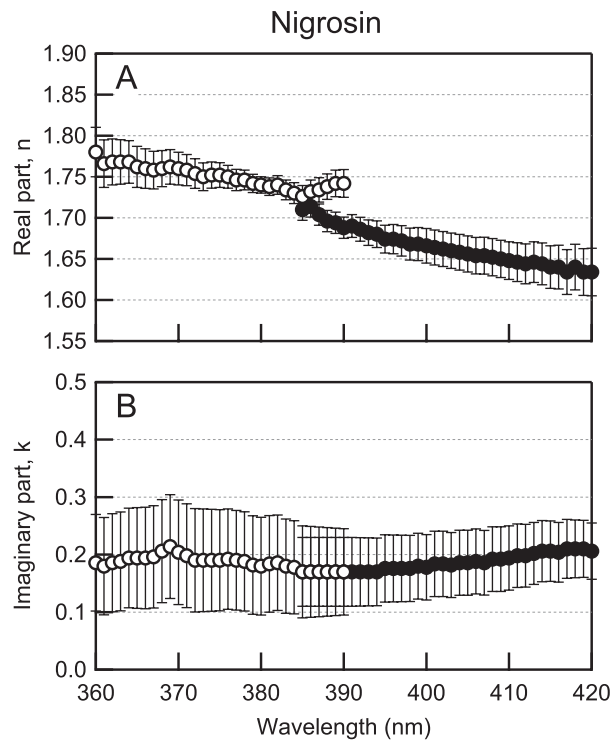


Fig. 8. (a) The real part of the complex refractive index (RI) as a function of wavelength for nigrosin aerosol. (b) The imaginary part of the RI.

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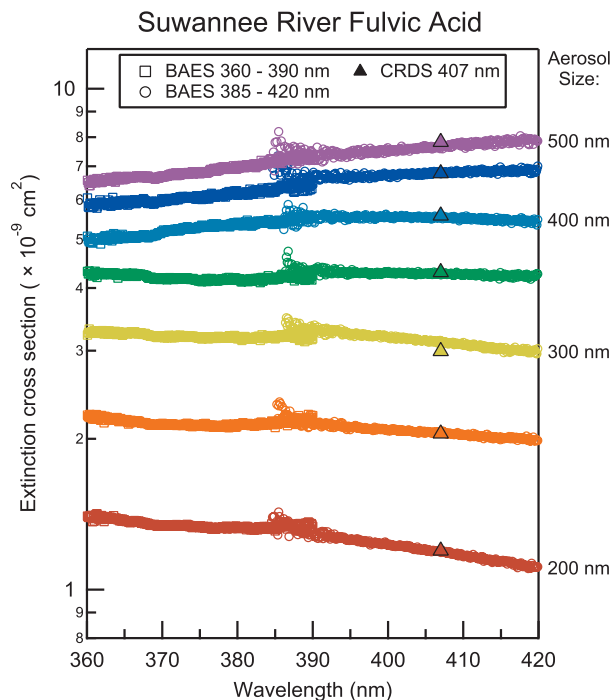


Fig. 9. Extinction cross-sections as a function of wavelength for Suwannee River fulvic acid aerosol at 200–500 nm diameter measured with the BAES 360–390 nm, BAES 385–420 nm, and CRDS at $\lambda = 407$ nm. Similarly to Fig. 7, these cross-sections include the population of singly-, doubly-, and triply-charged particles at each DMA setpoint.

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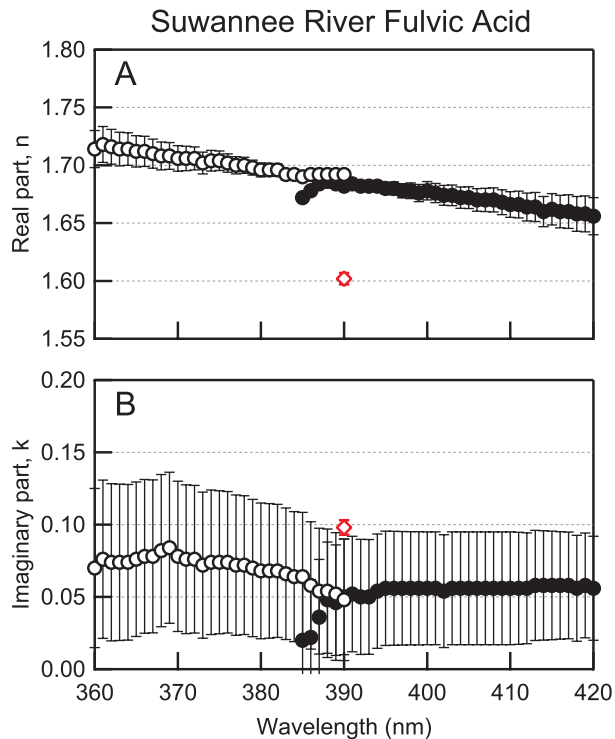


Fig. 10. (a) The real part of the complex refractive index (RI) as a function of wavelength for SRFA aerosol. Red marker shows values reported by Dinar et al. (2008, diamond). (b) Similarly, the imaginary part of the RI.

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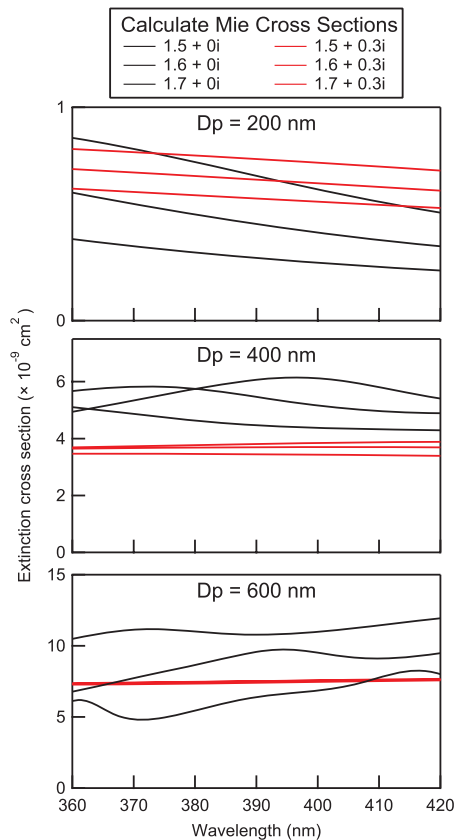


Fig. 11. Theoretical aerosol extinction cross-sections for six refractive indices at aerosol diameters of 200, 400, and 600 nm.

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