



Supplement of

The "dual-spot" Aethalometer: an improved measurement of aerosol black carbon with real-time loading compensation

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24 Filter loading effect for different measurement sites and seasons

The filter loading effect is determined by the *BC*(*ATN*) method for measurement campaigns conducted at different type of measurement site (roadside, urban background, rural background, regional background) in Klagenfurt (Austria), Anaheim (USA), Payerne (Switzerland), Sonnblick (Austria), Kathmandu (Nepal). Uncompensated *BC* data as measured by the AE33 with the detector under spot S1 was fitted using a linear function in the *ATN* range from 4.4 to 45. The filter loading effect is determined as the relative slope *RS* (Eq. 2).

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Figure S1. *BC*(*ATN*) analysis of the filter loading effect for the campaign in Klagenfurt (Austria) from 1-12 March 2012. The Klagenfurt station is located at Völkermarkter Strasse (46°37'32.1"N 14°19'05.7"E, 446 m ASL). The station is located in the middle of a crossroad and is strongly influenced by local traffic. During winter the site is strongly influenced by biomass burning from residential heating. Average BC data for each ATN bin (1 ATN unit wide) and the linear fit of the raw data are presented.





Figure S2. *BC*(*ATN*) analysis of the filter loading effect for a campaign in Anaheim (California, USA)
from 1-9 July 2012. The Anaheim station is located in an urban environment (33° 49' 50" N, 117°
56' 18"W, 39 m ASL). Average BC data for each ATN bin (1 ATN unit wide) and the linear fit of the
raw data are presented.



Figure S3. *BC(ATN)* analysis of the filter loading effect for campaign in Payerne (Switzerland) from
15 June – 5 July 2012. The Payerne aerological station (46.82 ° N, 6.95 ° E, 491 m ASL) is a rural
background air quality monitoring station located in Southwestern Switzerland, on the Swiss
plateau between the Jura and the Alps. It lies about 1 km south-east of the small rural town of
Payerne. The site is surrounded by agricultural fields (grassland and crops), forests and small
villages. Average BC data for each ATN bin (1 ATN unit wide) and the linear fit of the raw data are
presented.





Figure S4. *BC*(*ATN*) analysis of the filter loading effect for campaign in Sonnblick (Austria) from 1 July to 31 August 2013. The Sonnblick observatory is a background station located in the high alpine environment (47°03'15"N, 12°57'27"E, 3106 m ASL). Measurements at this location allow the determination of the composition of the mid-troposphere, frequently reaching into the free troposphere. Average BC data for each ATN bin (1 ATN unit wide) and the linear fit of the raw data are presented.







73 Table S1. Average filter loading effect as determined with the *BC*(ATN) analysis for different

74 measurement sites and seasons.

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Location	BC(ATN) relative slope RS	Site description	Season
Klagenfurt (Austria)	- 0.0069 +/- 0.0007	Roadside	Winter
Anaheim (USA)	- 0.0033 +/- 0.0003	Urban	Summer
	,		
Paverne (Switzerland)	- 0.0013 +/- 0.0004	Rural background	Summer
- / /	,	8	
Sonnblick (Austria)	- 0.0017 +/- 0.0002	Regional background	Summer
			••••••
Kathmandu (Nepal)	- 0.0047 +/- 0.0003	Urban	Winter
		0.001	



80 Figure S6. Time-series of the ACTRIS inter-comparison, showing AE33 and MAAP measurements of

- 81 BC.
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83 Table S2. Mass absorption cross-section values used in the AE33.

Channel	Wavelength (nm)	σ_{air} (m ² /g)
1	370	18.47
2	470	14.54
3	520	13.14
4	590	11.58
5	660	10.35
6	880	7.77
7	950	7.19

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85 The mass absorption cross-section values σ_{air} used in the Aethalometer assume an inverse dependence on the wavelength – that is, a "black" sample with an Ångström exponent equal to unity 86 87 (Moosmüller et al., 2011). The relationship between the absorption and mass was determined by optical measurements of transmission and thermal measurements of samples, where the non-88 89 refractory constituents of the carbonaceous sample were removed (Gundel et al., 1984) -90 measurements in the infra-red part of the spectrum should therefore be used to convert the optical 91 measurement into the mass concentration, as the contribution of sample components other than 92 black carbon is negligible at these wavelengths (Sandradewi et al., 2008a; Sandradewi et al., 2008b; 93 Fialho et al., 2005; Yang et al., 2009; and references therein). The relationship between the mass 94 concentration of BC and the optical absorption can be determined by comparing the filter photometer measurements with those obtained by thermal-optical analysis (Sciare, 2011). However, 95 96 as the determination of elemental carbon (EC) depends on the thermal-optical analysis method, 97 sometimes with large differences (Bae et al., 2009), the determination of the mass absorption crosssection also depends on the thermal-optical method employed. Additionally, the season and the 98 99 sample composition (Bae et al., 2009; Chiappini et al., 2014) may influence the determination of EC; and the mass absorption cross-section may depend on the aerosol mixing state and size (Bond and Bergstrom, 2006). Concurrent determination of EC in filter samples with an thermal-optical method, and the Aethalometer measurement of optical absorption allow the site-specific determination of the mass absorption cross-section, which is specific to the thermal-optical method employed – this procedure is often employed in source apportionment campaigns, where mass closure is attempted (Sciare, 2011).

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