



Supplement of

Comparison of co-located refractory black carbon (rBC) and elemental carbon (EC) mass concentration measurements during field campaigns at several European sites

Rosaria E. Pileci et al.

Correspondence to: Robin L. Modini (robin.modini@psi.ch) and Martin Gysel-Beer (martin.gysel@psi.ch)

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Supplementary information (SI)

S1. Further details concerning the optical correction in thermal optical analysis (TOA)

During TOA analysis a fraction of the OC can pyrolyze in the He step to form pyrolytic carbon (PC), which is thermally stable and only desorbs in the O₂ step, thereby causing a charring artefact in the mutual quantification of OC and EC. To correct for this latter effect a laser at 658 nm can be used to monitor the light transmission through the loaded filter before and during the analysis. PC is strongly light absorbing, thus leading to a decrease of the transmission signal when it forms upon heating in the inert atmosphere. Later, in the oxidizing atmosphere, both PC and EC are released from the filter resulting in an increase of the transmission signal. The time at which the transmission equals again the initial pre-pyrolysis value is used to separate OC and EC, depending on whether the carbon evolved before or after this “split point”, respectively. This thermal–optical transmittance (TOT) approach to correct for PC eliminates potential charring artefacts if the PC has the same mass–specific attenuation cross section as the atmospheric native EC (Yang and Yu, 2002), and if no other light–absorbing material evolves from the sample.

Instead of using light transmission, the charring correction can also be done with light reflectance (i.e., thermal-optical reflectance, TOR). EC values determined using TOT can be up to 30 – 70 % lower than those determined with TOR (Karanasiou et al., 2015), because the evaporation of non–absorbing particulate matter during heating affects the reflectance to a greater extent than the transmission signal. Furthermore, high loadings of EC result in saturation effects of both optical signals, again to a greater extent for the reflection compared to the transmission method (Chiappini et al., 2014). These two effects result in better reproducibility and accuracy of the TOT based OC/EC split compared to the TOR approach.

S2. Calculation of variability and bias

The variability (Q_{AV}) is defined as the relative standard deviation given by the 95 % confidence limit, thus:

$$Q_{AV} = \frac{n}{\sqrt{6}} \left[\max \left(\frac{RD_i}{T_i} \right) - \min \left(\frac{RD_i}{T_i} \right) \right] \quad (S1)$$

Where $RD_i = L_i - T_i$, with L_i and T_i representing the laboratory and expected concentrations, respectively.

The bias (Q_{AB}) is defined as the median of the percentage of the ratio between RD_i and T_i

$$Q_{AB} = \text{median} \left[\frac{RD_i}{T_i} \% \right]. \quad (S2)$$

S3. The Cunningham slip correction

The Cunningham slip correction factor, C_C , is used to account for non–continuum effects when calculating the drag force on small particles. C_C depends on the particle diameter, D , the mean free path of the surrounding gas, λ , and on the experimental coefficients α , β and γ (Cunningham, 1910; Seinfeld and Pandis, 2006).

$$C_C(D) = 1 + \frac{2\lambda}{D} \left[\alpha + \beta e^{-\frac{\gamma D}{\lambda}} \right] \quad (S3)$$

with $\alpha = 1.257$, $\beta = 0.4$, $\gamma = 1.1$, $\lambda = 6.5 \cdot 10^{-8} \text{ m}$

35 S4. Hygroscopic growth factor

Hygroscopic growth affects the cut-off imposed by impactors operated at ambient RH. Here we provide simplified equations to calculate the volume equivalent diameter growth factor, GF, of BC-containing particles coated with a mixture of organic and inorganic matter. The GF is calculated using κ -Köhler theory (Petters and Kreidenweis, 2007):

$$GF(RH) = \left(1 + \kappa_{\text{tot}} \frac{RH}{1-RH}\right)^{1/3} \quad (S4)$$

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where the hygroscopicity parameter of the mixed particle, κ_{tot} , is obtained with the ZSR-mixing rule written as (Petters and Kreidenweis, 2007):

$$\kappa_{\text{tot}} = \varepsilon_{\text{BC}} \kappa_{\text{BC}} + \varepsilon_{\text{org}} \kappa_{\text{org}} + \varepsilon_{\text{inorg}} \kappa_{\text{inorg}} \quad (S5)$$

45 The hygroscopicity parameters κ of BC, organics and inorganics are assumed to be 0, 0.1 and 0.5, respectively (Engelhart et al., 2012). The volume fraction, ε_x , of compound class “x” in the particle can be calculated using:

$$\varepsilon_x = \frac{m_x}{m_{\text{tot}}} \frac{\rho_{\text{tot}}}{\rho_x} \quad (S6)$$

where m_x is the mass of “x” in the particle, m_{tot} is the total particle mass. ρ_x is the material density of “x”, which is assumed to be 1800 kg m^{-3} , 1200 kg m^{-3} and 1700 kg m^{-3} for BC, organics and inorganics, respectively. The mixed particle density, ρ_{tot} , is obtained with:

$$\rho_{\text{tot}} = \frac{1}{\frac{m_{\text{BC}}}{m_{\text{tot}}} \rho_{\text{BC}} + \frac{m_{\text{org}}}{m_{\text{tot}}} \rho_{\text{org}} + \frac{m_{\text{inorg}}}{m_{\text{tot}}} \rho_{\text{inorg}}} \quad (S7)$$

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S5. Dryers and losses

Differences in m_{rBC} and m_{EC} mass concentration can also come from differences in the losses of the respective sampling inlets. Particle losses can be caused by the presence of a dryer in the inlet line. The dryer technology offers different dryer types including diffusion and membrane dryers. Diffusion dryers use chemical adsorbents such as silica gel for the minimization of aerosol losses. When the aerosol passes through the tube, the silica adsorbs water vapor, therefore this chemical needs to be changed and regenerated on a regular basis. Membrane dryers are elastic tubes based on water vapor-permeable polytetrafluoroethylene (PTFE). Commercially available membranes are products such as Nafion®: a sulfonated tetrafluoroethylene working as permeable membrane in which water vapor molecules are transported.

In any case, particle losses by diffusion across the dryer should be accounted for in the data processing. This is done by calculating an equivalent tube length, which is longer than the actual dryer length (Wiedensohler et al., 2012).

In this work we calculate the diffusion losses when dryers were present in the inlet lines. A diffusion dryer was placed in front of the SP2 line during the Bologna campaign; while the EC line had none. The particle transmission efficiency of the diffusion dryer for a flow rate of 2 L min^{-1} was evaluated to be less than 10 % on average on the size range of the SP2 measurements. Nafion dryers (model MD-700, Perma Pure) were set in front of the SP2s in the Melpitz and Cabauw campaigns. The diffusion losses through them were calculated with the hypothesis of laminar flow and were found to be negligible. The particle loss due to the presence of a dryer in the inlet line does not seem to be the main reason of the m_{rBC} and $m_{\text{EC_PM}_{2.5}}$ discrepancies for the campaigns of this study. For this reason we did not correct the rBC mass concentrations for this effect.

Tables SI

Table S1: site, station code, coordinates, altitude and year/season of the field campaigns presented in this work.

Site (country)	Station code	Coordinates	Altitude	Year/season
Palaiseau (FR)	Palaiseau	48.713° N 2.208° E	160	2010/winter
Melpitz (DE)	Melpitz	51°32' N, 12°56' E	86	2017/winter and 2015/summer
Cabauw (NE)	Cabauw	51° 58' N, 4° 55' E	−0.7	2016/autumn
Bologna (IT)	Bologna	44° 31' N, 11° 20' E	39	2017/summer

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Table S2: The first three columns show median, 10th and 90th percentiles of EC, TC and EC/TC filter loading. The last four columns show minima and maxima of EC and TC filter loadings for the field campaigns of this study.

Station code	EC filter loading median (10 th , 90 th) [$\mu\text{g cm}^{-2}$]	TC filter loading median (10 th , 90 th) [$\mu\text{g cm}^{-2}$]	EC/TC filter loading median (10 th , 90 th) [$\mu\text{g cm}^{-2}$]	Min EC filter load [$\mu\text{g cm}^{-2}$]	Max EC filter load [$\mu\text{g cm}^{-2}$]	Min TC filter load [$\mu\text{g cm}^{-2}$]	Max TC filter load [$\mu\text{g cm}^{-2}$]
Palaiseau	1.66 (0.76, 3.19)	9.79 (5.02, 24.03)	0.14 (0.09, 0.30)	0.44	7.53	3.72	37.56
Melpitz winter	4.30 (1.50, 11.08)	44.73 (9.77, 110.72)	0.11 (0.09, 0.17)	0.93	12.53	5.45	115.03
Melpitz summer	0.90 (0.48, 1.43)	12.47 (6.93, 19.08)	0.08 (0.05, 0.10)	0.25	2.30	4.76	24.43
Cabauw	1.47 (0.71, 2.34)	6.87 (3.78, 14.95)	0.18 (0.12, 0.29)	0.48	3.44	2.92	19.10
Bologna	2.49 (1.84, 2.66)	15.50 (13.20, 17.72)	0.16 (0.11, 0.19)	1.63	2.74	12.46	18.42

- 80 Table S3: In this table the aerodynamic particle diameter D_{aero} corresponding to the upper SP2 cut-off, which depends on the BC mass equivalent diameter, D_{ve} (calculating starting from the BC core mass equivalent diameter, D_{rBC} , knowing the particle mixing state), is given for different extreme hypotheses concerning particle shape (χ), mixing state and relative humidity (RH). The calculations are made with the hypothesis of fixed density of BC core $\rho_{rBC} = 1800 \text{ kg m}^{-3}$ and with the hypothesis of coating made by half organic material with $\kappa_{org} = 0.1$ and
- 85 $\rho_{org} = 1200 \text{ kg m}^{-3}$ and half inorganic material with $\kappa_{inorg} = 0.5$ and $\rho_{inorg} = 1700 \text{ kg m}^{-3}$. From these hypothesis and knowing the ratio between the mass of the coating material, m_{coat} and the mass of the BC core, m_{rBC} , the total particle density, ρ_p , can be calculated. The mixing state of the last example particle is constrained with SP2 measurements during the Melpitz winter campaign.

	D_{rBC} [nm]	ρ_p [kg m ⁻³]	κ_{tot}	D_{ve} [nm] Dry	GF (RH = 80 %)	D_{ve} [nm] (RH = 80 %)	GF (RH = 95 %)	D_{ve} [nm] (RH = 95 %)	D_{aero} [nm] Dry	D_{aero} [nm] (RH = 80 %)	D_{aero} [nm] (RH = 95 %)
Fractal-like pure BC: $\chi = 2.4$ (Park et al., 2003)	722.0	1800.0	0.0	722.0	1.0	722.0	1.0	722.0	625.3	625.3	625.3
Spherical pure BC: $\chi =$ 1	722.0	1800.0	0.0	722.0	1.0	722.0	1.0	722.0	968.7	968.7	968.7
Coated BC: $\chi = 1$, $m_{coat} = m_{rBC}$	722.0	1579.4	0.1	909.7	1.2	1063.1	1.6	1423.4	1143.2	1336.0	1788.8
Coated BC: $\chi = 1$, $m_{coat} = 6m_{rBC}$	722.0	1452.2	0.2	1381.1	1.2	1722.4	1.8	2432.5	1664.4	2075.6	2931.3
Coated BC with coating as observed during the Melpitz Winter campaign: $\chi = 1$, $m_{coat} = 2.33$ [0.99–3.17] m_{rBC}	722.0	1505.2	0.2	1078.2	1.2	1310.4	1.7	1815.9	1322.8	1607.7	2227.9

Table S4: AAE (470,950) statistics for the campaigns in this study: median, geometric mean, 10th and 90th percentiles and number of data points.

	Palaiseau	Cabauw	Melpitz summer	Melpitz winter	Bologna
AAE median (10, 90)	1.35 (1.24, 1.53)	1.05 (0.97, 1.12)	1.19 (1.09, 1.26)	1.40 (1.28, 1.50)	1.03 (1.01, 1.07)
AAE geometric mean	1.36	1.04	1.18	1.38	1.04
# points	34	32	49	20	7

95 Table S5: Summary of site name, country, SP2 calibration material, m_{EC} cut-off and TOA thermal protocol, sampling period, site characteristics and geometric mean of the m_{rBC}/m_{EC} ratio for all the data. *One data point

	SP2 Calibration Material	m_{EC} cut-off/ TOA Thermal technique	Site name, country	Season/year and site characteristics	m_{rBC}/m_{EC} median
Palaiseau	Fullerene Soot	PM _{2.5} / EUSAAR-2	Palaiseau, France	Jan/Feb 2010 suburban background	1.20
Cabauw	Fullerene Soot	PM ₁₀ / EUSAAR-2	Cabauw, Netherlands	Oct 2016 rural background	0.53
Bologna	Fullerene Soot	PM _{2.5} / EUSAAR-2	Bologna, Italy	July 2017 urban background	0.65
Melpitz winter	Fullerene Soot	PM _{2.5} / EUSAAR-2	Melpitz, Germany	Feb 2017 rural background	1.29
Melpitz summer	Fullerene Soot	PM _{2.5} / EUSAAR-2	Melpitz, Germany	July 2015 rural background	0.97
Zhang et al. (2016)	Fullerene Soot	PM _{2.5} / IMPROVE	Fresno, California, USA	Jan/Feb 2013 urban background	0.70
Miyakawa et al. (2016)	Fullerene Soot	PM _{2.5} / IMPROVE-like	Yokosuka, Japan	Summer 2014 June 17 – 27 urban	1.07
Sharma et al. (2017)	Aquadag scaled to Fullerene Soot	PM ₁ / EnCan-Total- 900	Nunavut, Canada	From Mar 2011 to Dec 2013 remote site	0.55
Corbin et al. (2019)	Fullerene Soot	PM ₁ / IMPROVE-A (washed)	–	Chamber study – four-stroke ship diesel engine	1.03*
Laborde et al. (2012b)	Fullerene Soot	NIOSH-5040	–	Chamber study – CAST soot	1.10*

Figures SI

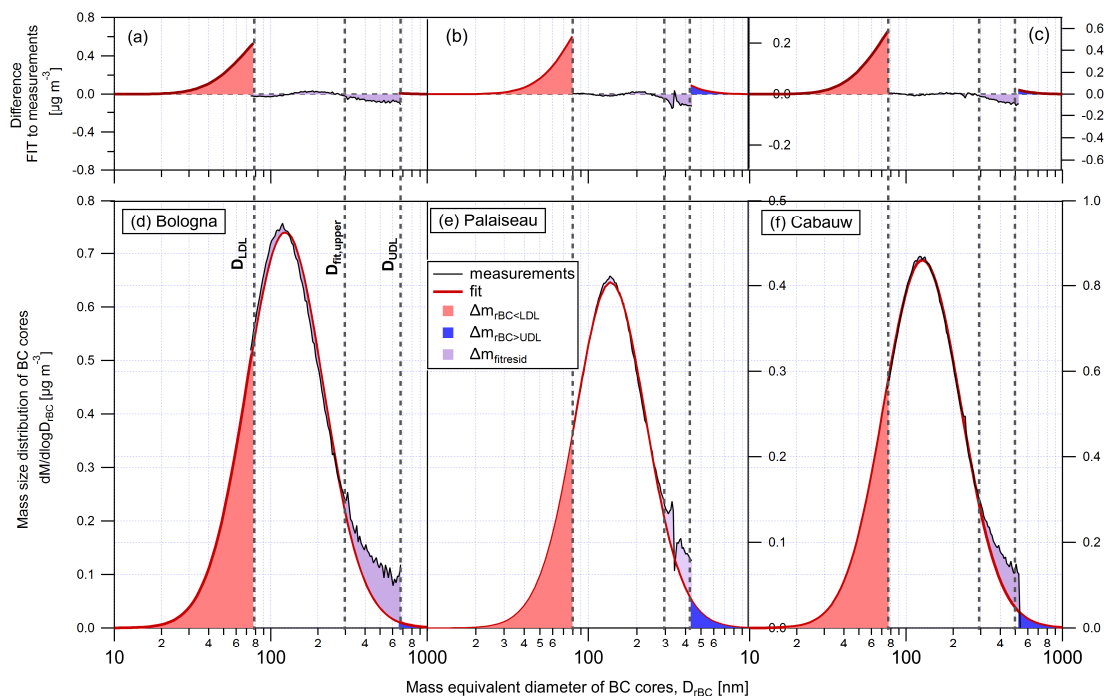
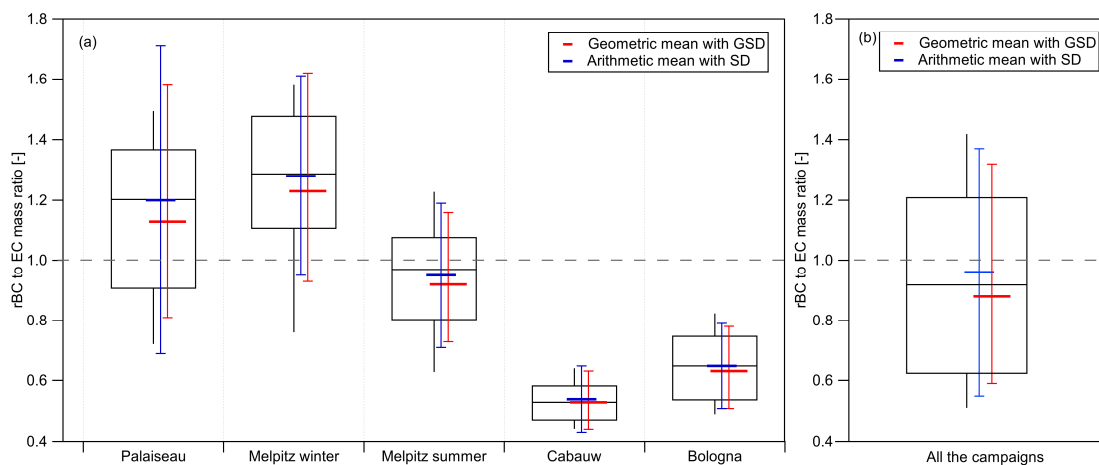


Figure S1: Approach to correct for the rBC mass outside the rBC core size range covered by the SP2 for the Bologna (panels a and d), the Palaiseau (panels b and e) and the Cabauw (panels c and f) campaigns. The bottom three panels show the measured rBC mass size distribution as a function of rBC core mass equivalent diameter, including the SP2 detection limits D_{LDL} and D_{UDL} . The lognormal functions are fitted between D_{LDL} and $D_{fit,upper}$. The integrated area of the red, purple, and blue shadings correspond to $\Delta m_{rBC < LDL}$, $\Delta m_{fitresid}$ and $\Delta m_{rBC > UDL}$, respectively (see Sect. 2.3.5). The top three panels additionally show the same shadings after subtraction of the measured size distribution (and measurement forced to be zero outside the SP2 detection range).



115 Figure S2: Statistics (10th, 25th, 50th, 75th and 90th percentiles, arithmetic and geometric means, SD and GSD) of the rBC to EC mass ratio ($m_{\text{rBC}}/m_{\text{EC}}$) per campaign (panel a) and with all the campaigns of this work (panel b).

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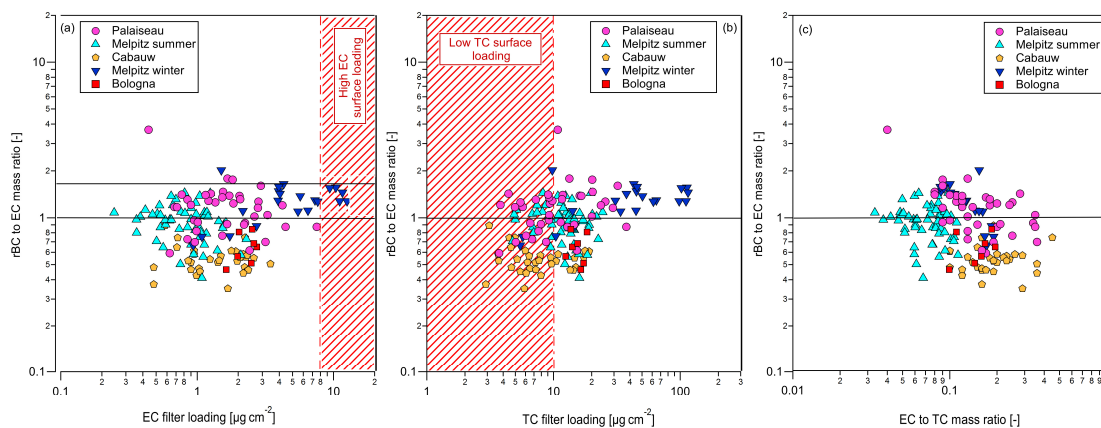


Figure S3: rBC mass concentration versus EC filter loading (panel a), TC filter loading (panel b) and EC/TC mass ratio (panel c). The red shaded areas in panel (a) and (b) indicate the high EC surface loading and the low TC surface loading areas respectively.

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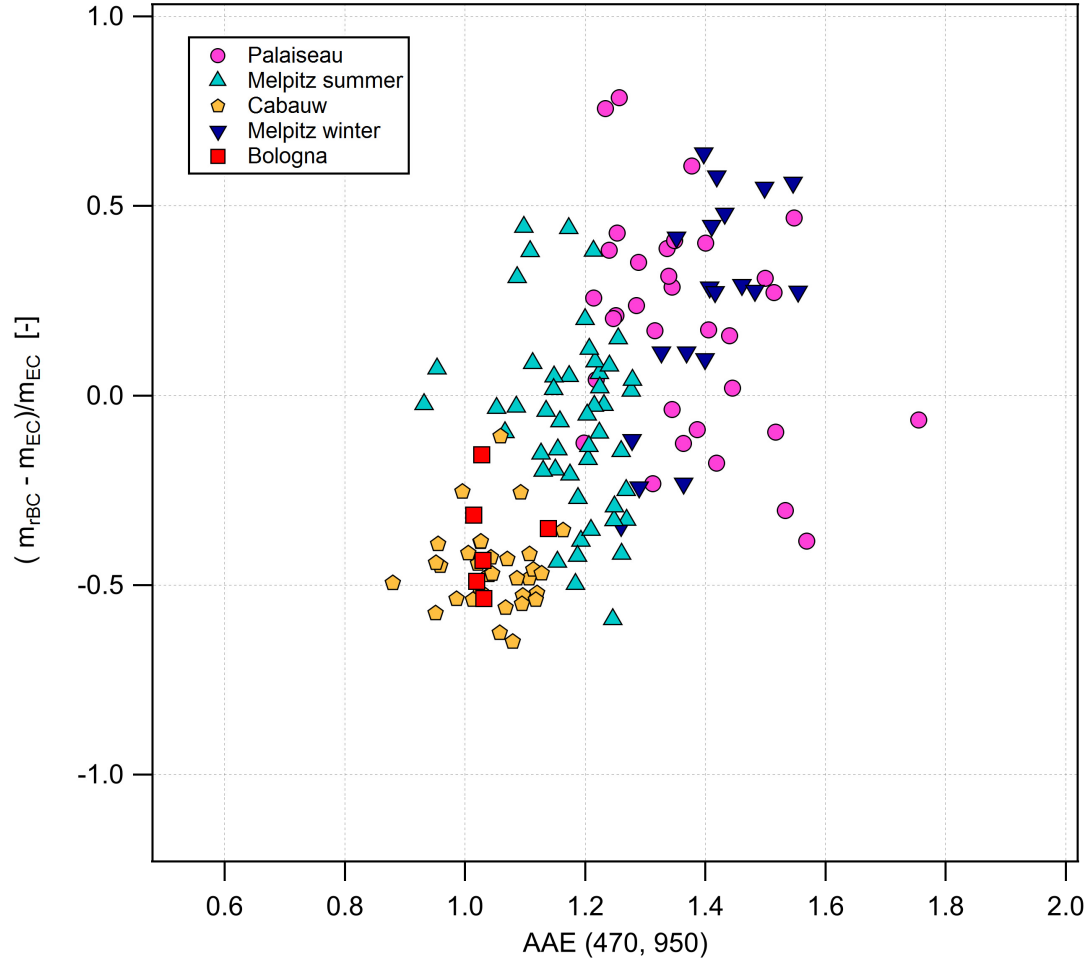


Figure S4: Relative difference between m_{rBC} and m_{EC} versus the AAE(470,950) coloured by campaign.