

J. Brito¹, F. Wurm¹, Y. Liu², S. S. de Sá², S. Carbone¹, L.-V. Rizzo³, G. Cirino⁴, H. Barbosa¹, R. Souza⁵, S. T. Martin² and P. Artaxo¹

¹University of São Paulo, Brazil, ²Harvard University, USA; ³Federal University of São Paulo, Brazil, ⁴National Institute of Amazonian Research, Brazil; ⁵University of the State of Amazonas, Brazil.

Work summary

- **10 month aerosol and trace gases characterization near Manaus**
- **Consistent first-order plume characterization**
- **Strong effect on isoprene mixing ratios due to anthropogenic activities**
- **Significant difference in size distribution between wet/dry seasons**
- **Four factors identified from PMF analysis of organics during dry season: Urban, biomass burning, biogenic and aged OA.**

Sampling site

Results shown here relate to measurements carried out at the T2 site from the GoAmazon 2014/15 experiment, near Manaus.

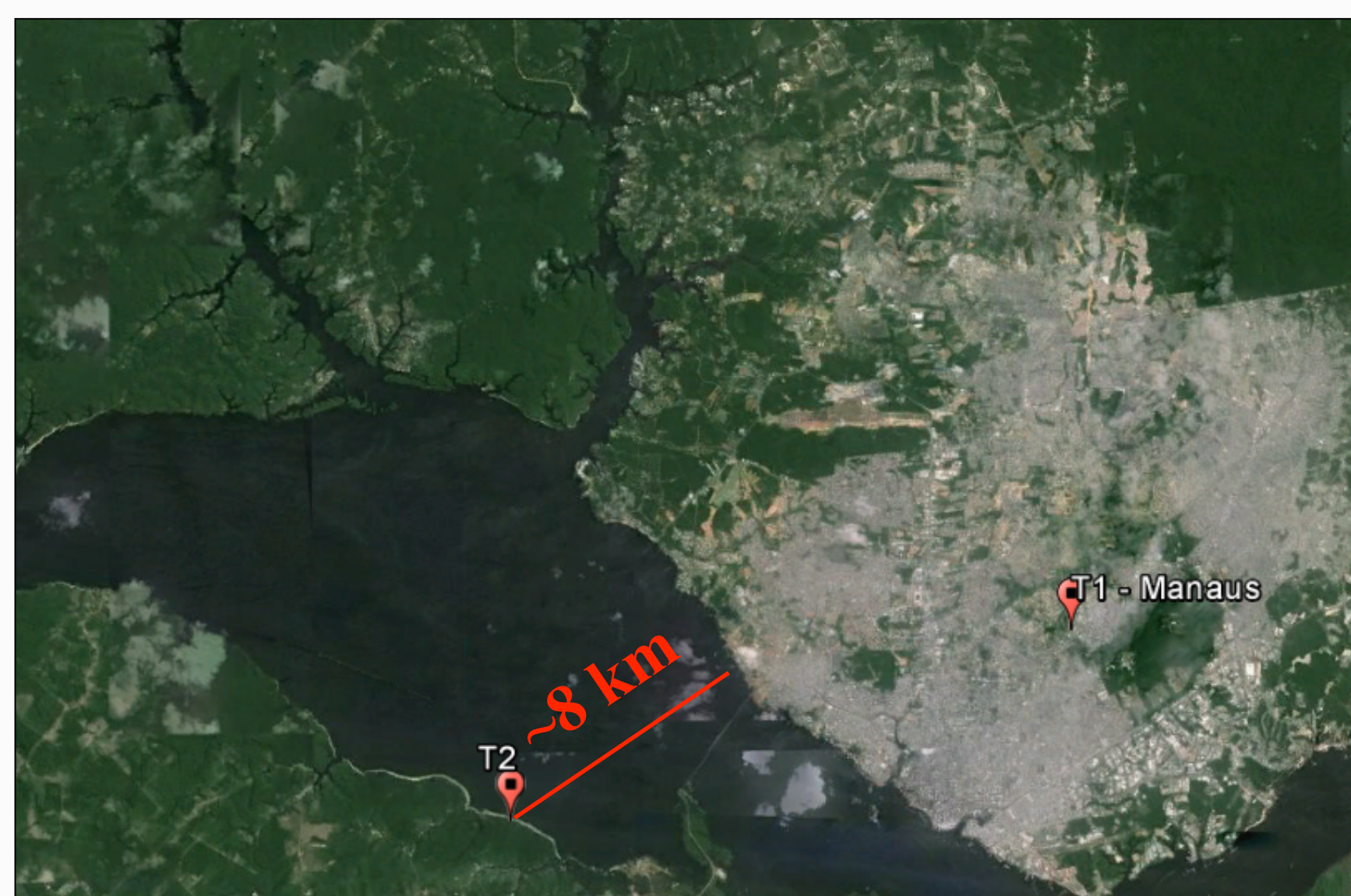


Figure 1 – GoAmazon sampling sites



Table 1 – List of instruments deployed at the GoAmazon T2 site

	Instrument	Analysis	Campaign	
			IOP 1 (Wet)	IOP 2 (Dry)
Gas Phase	PTR-Q-MS (Ionicon)	VOCs	X	X
	49i (Thermo)	Ozone	X	X
	43i (Thermo)	SO ₂	X	X
	CAPS (Aerodyne)	NO ₂		X
	Los Gatos Research	CO, N ₂ O	X	X
Aerosol Phase	Los Gatos Research	CH ₄ , CO ₂	X	
	SMPS (TSI)	Aerosol Size Dist.	X	X
	ACSM (Aerodyne)	PM ₁ NR Composition	X	X
	Filters	Elem. Comp. & EC/OC	X	X
	MAAP (Thermo) & AE33 (Magee)	Black Carbon	X	X
	Nephelometer (Ecotech Aurora)	Light Scattering	X	X
	TEOM (Thermo)	PM _{2.5} & PM ₁₀		X
	CCNC (DMT)	Size-Resolved CCNC		X

Manaus plume characterization

Tracers of anthropogenic activity, BC and CO, have been used to constrain a chemical characterization of the Manaus Plume. Due to the proximity of the Negro River, a significant river breeze – wind pattern according to temperature gradient between water and land – was observed. Fig.1 shows histogram of BC measurements considering easterly winds only according to the hour of the day.

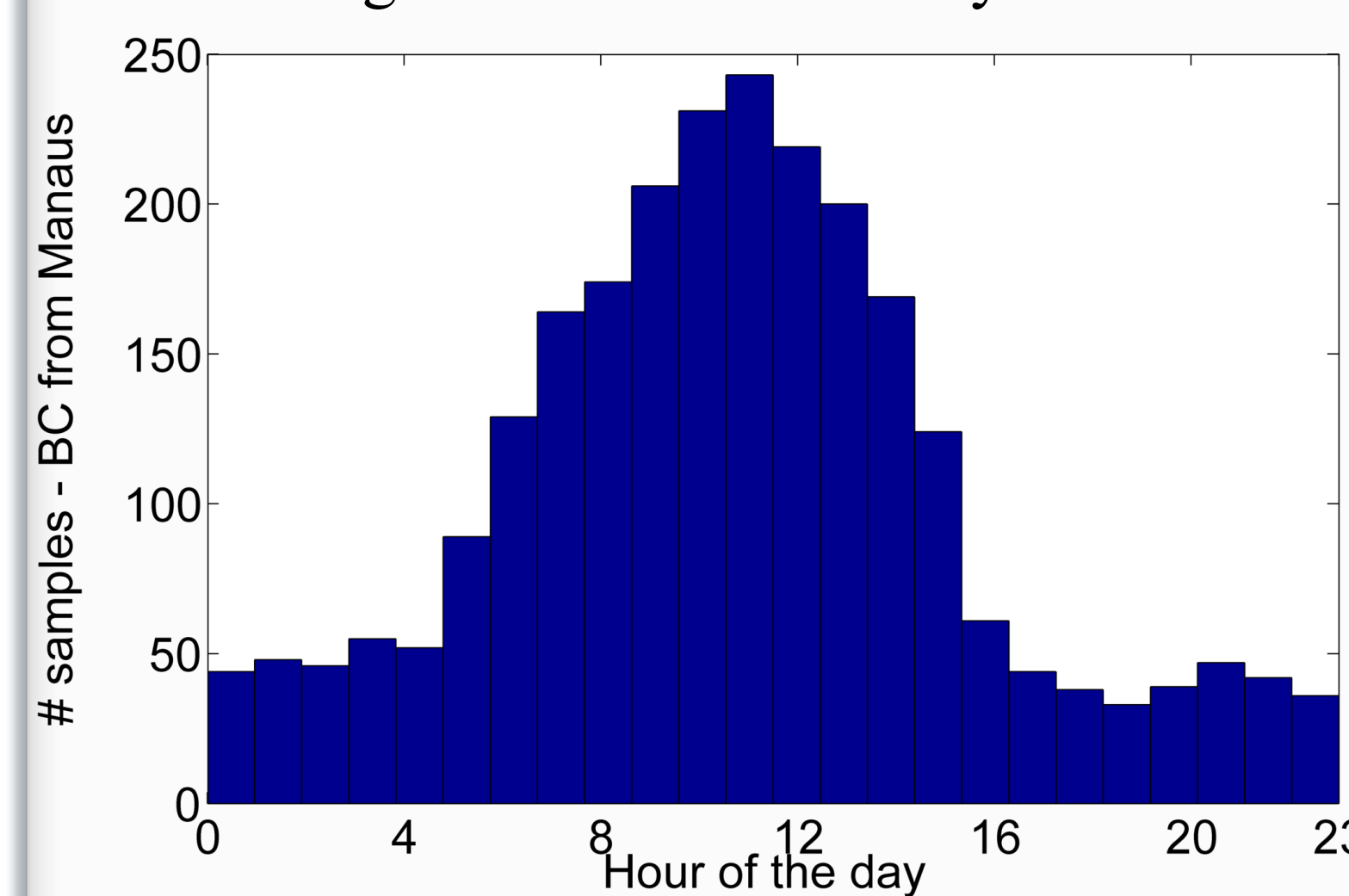


Figure 2 – Histogram of BC measurements considering periods of time of easterly wind only.

By correlating BC and CO data from easterly wind only allows us to constrain a chemical characterization of Manaus plume. Such values shall be used as a guide of Manaus pollution downwind of the site, namely T3.

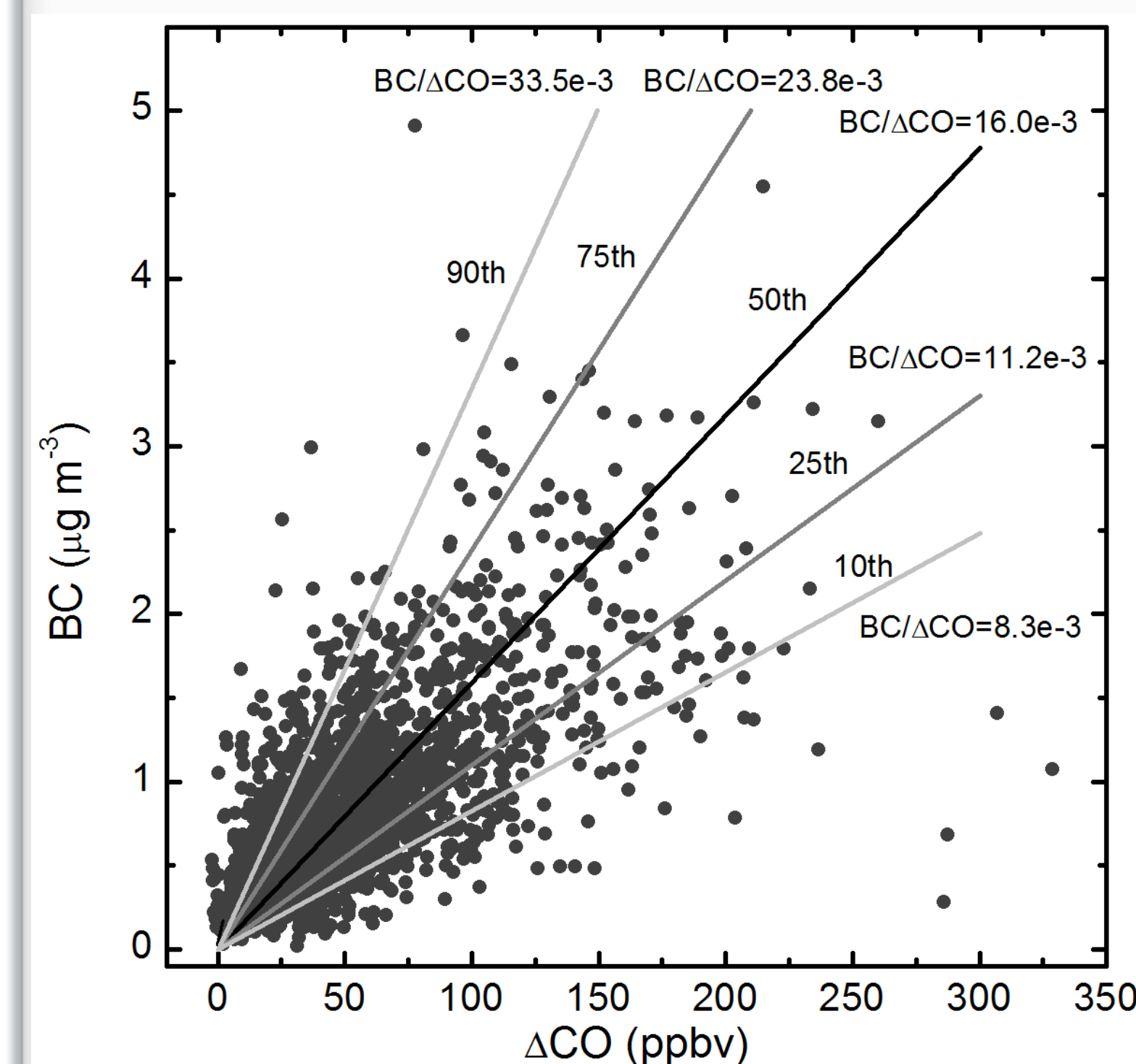


Figure 4 – Correlation of BC and ΔCO from Manaus

Diel variation of isoprene levels have shown a stark difference depending on local winds, northerly being mainly anthropogenic-free and easterly being Manaus direction

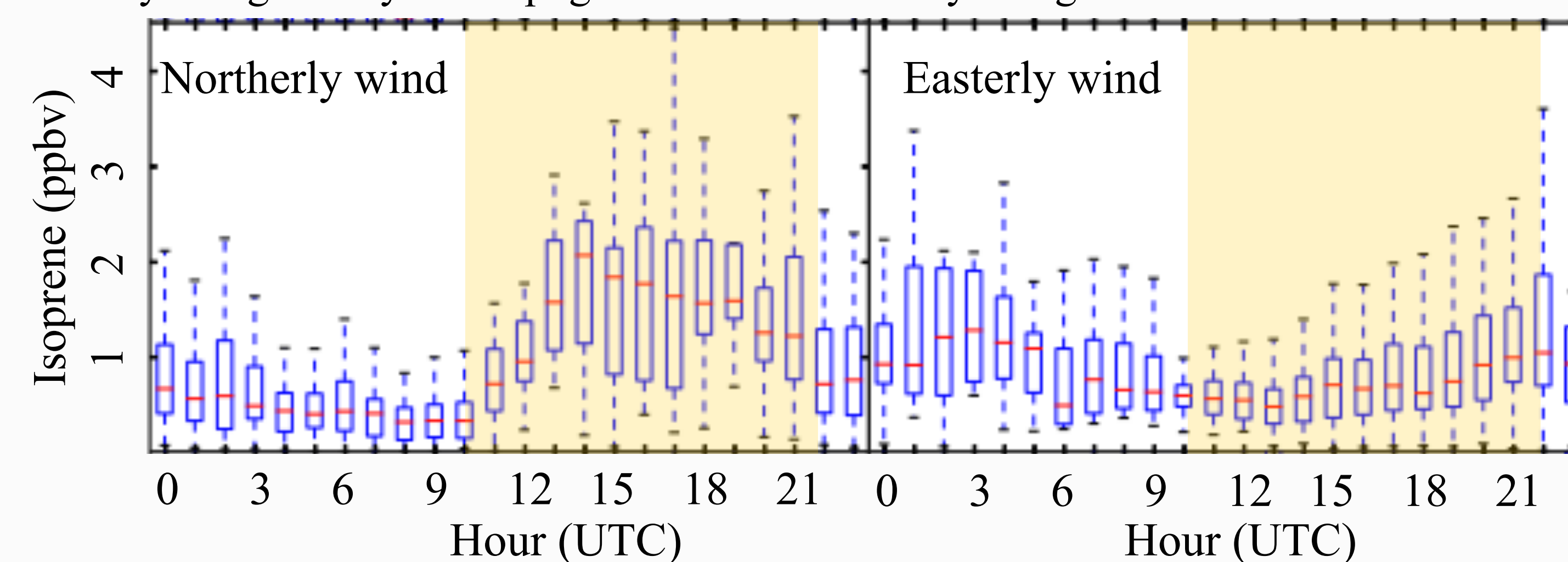


Figure 5 – Diel variation of isoprene mixing ratios depending on local wind direction

Taking in account periods of time of easterly winds (from Manaus), a clear rush hour pattern is observed for both BC and CO (Fig. 3).

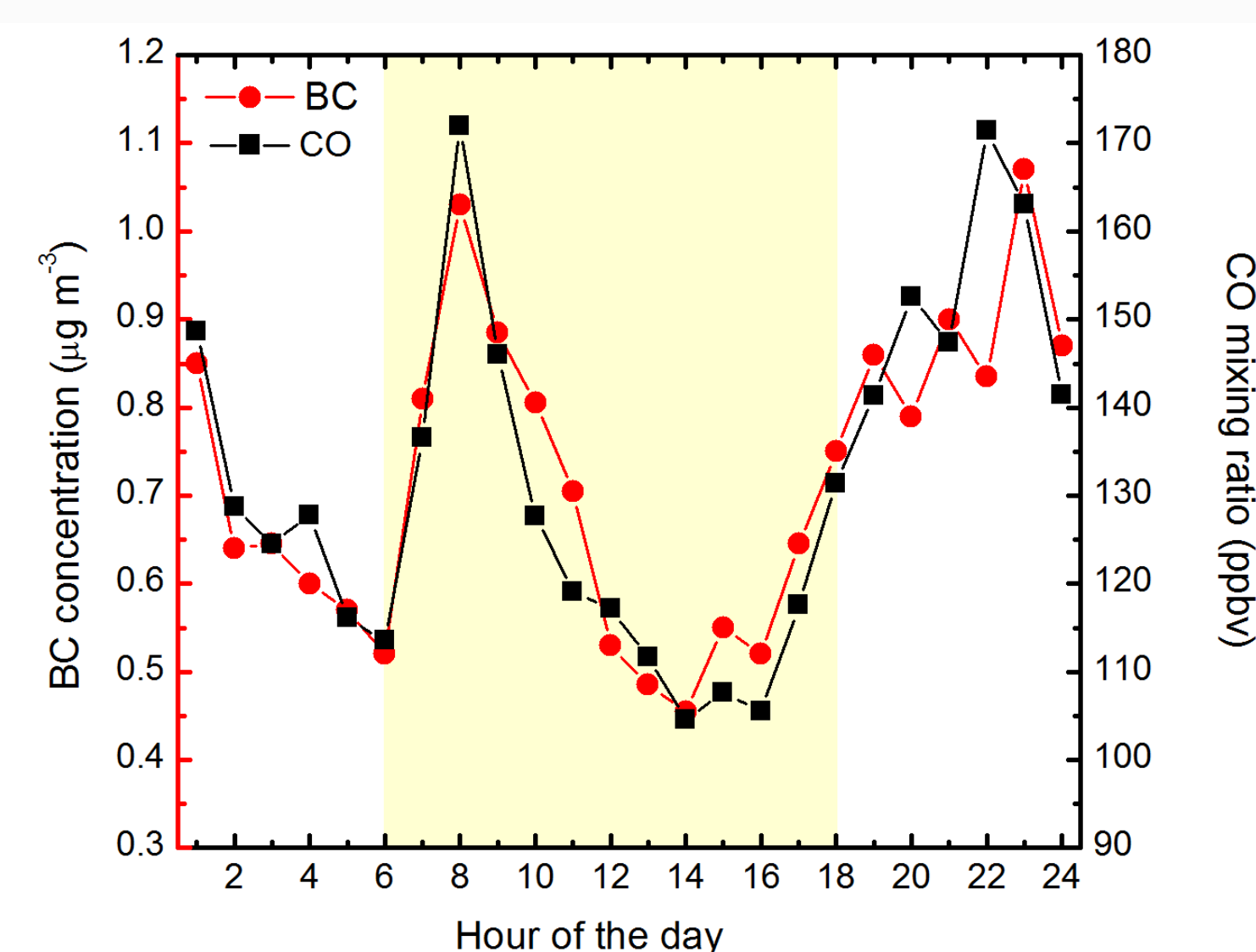


Figure 3 – Diel variation of BC and CO, considering easterly wind only.

In a similar analysis as performed above, SO₂, along with SO₄, do not show a distinct rush hour pattern, indicating the dominance of other sulfur sources on the site, probably from the diesel power plant.

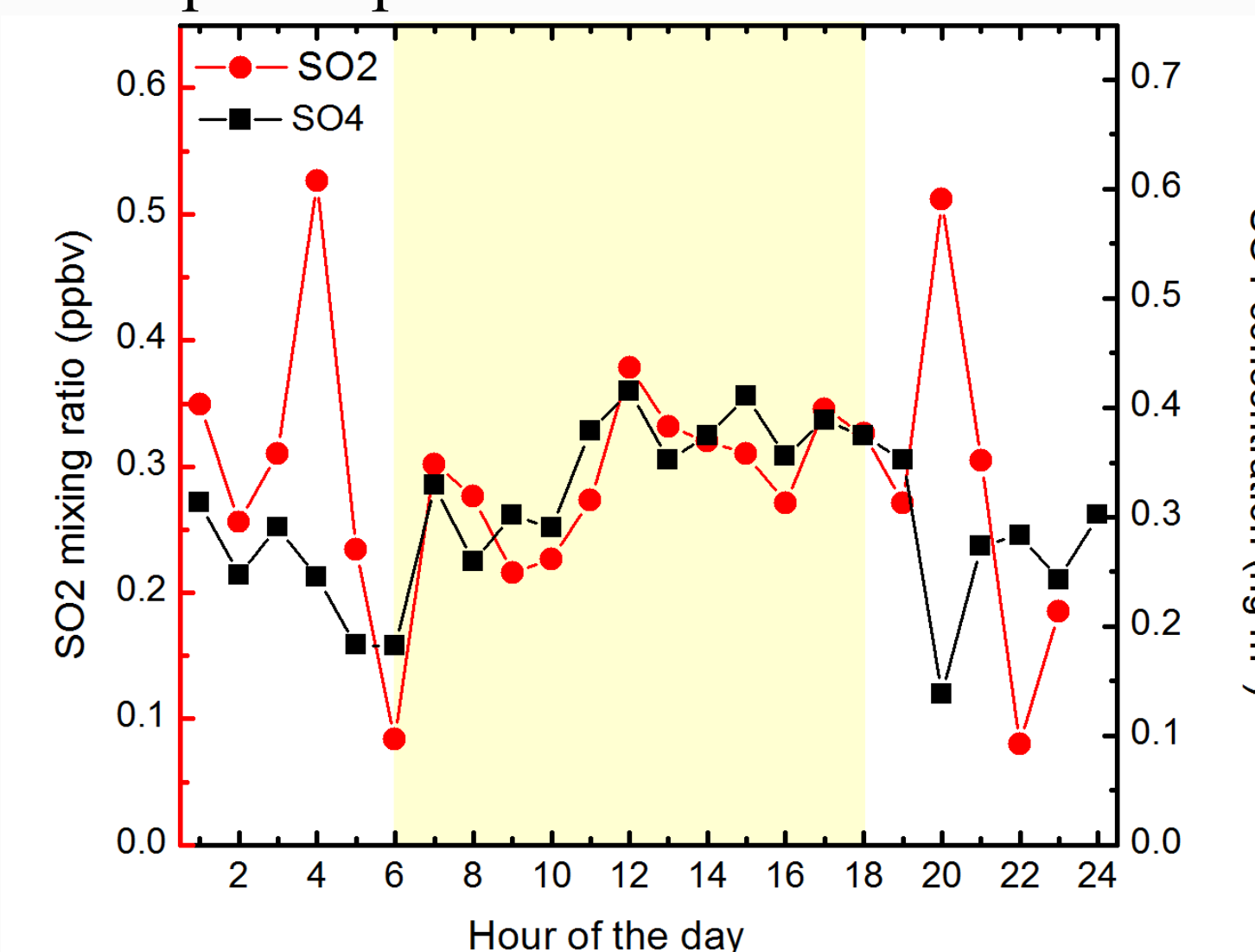


Figure 5 – Diel variation of SO₂ and SO₄, considering easterly wind only.

Aerosol composition and size distribution

Some of the results concerning aerosol chemical composition and size distribution are shown below. A significant shift from nucleation to accumulation mode aerosols is observed from IOP1 (wet season) to IOP2 (dry season). Such shift is reflected in much higher PM₁ mass as well. The metric f₄₄ (linked with O:C ratio of OA) shows relatively highly oxidized aerosols already at T2, with consistent increase in average values during dry season. Both results are consistent with a strong enhance of biomass burning emissions during the dry season.

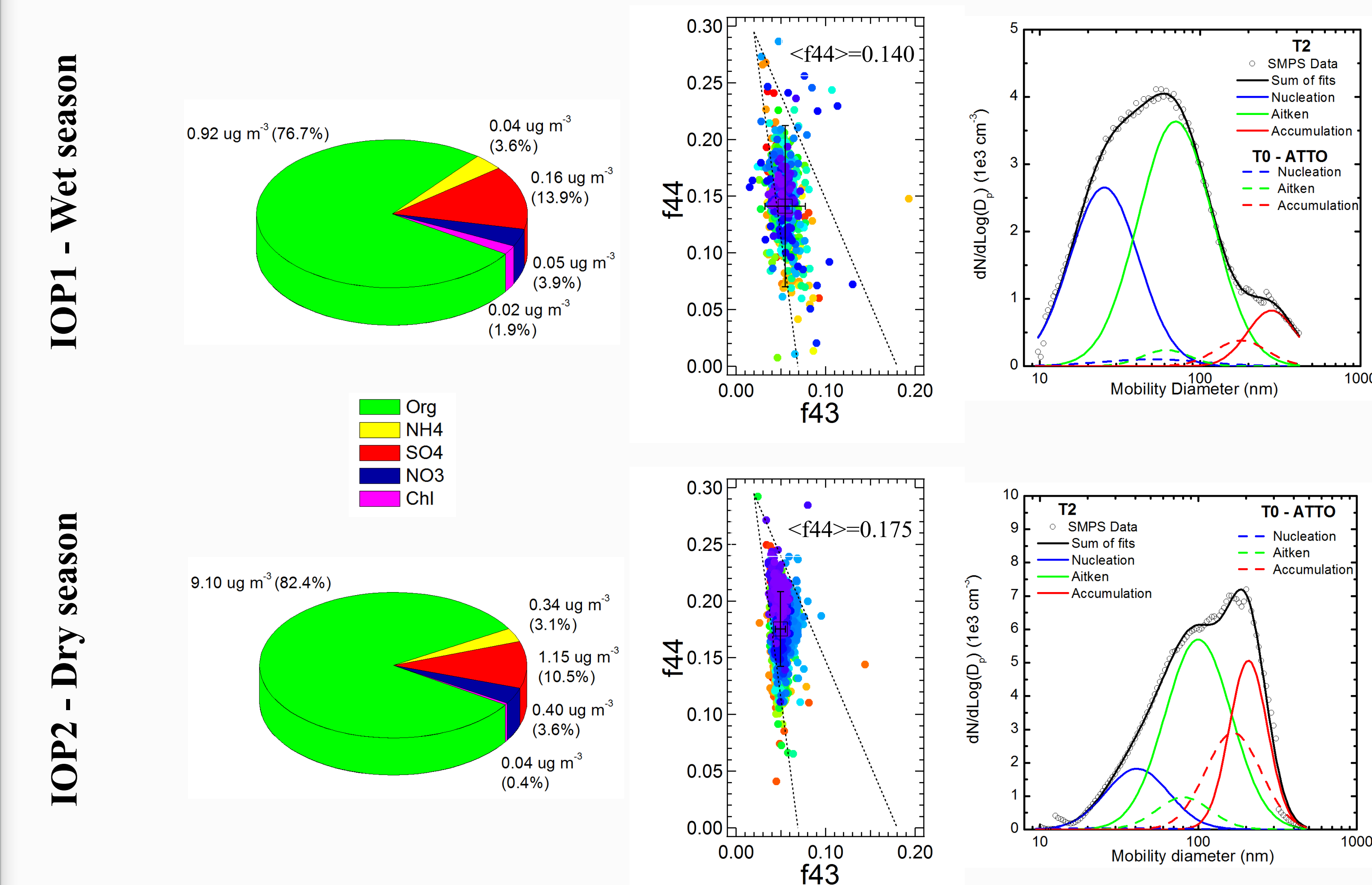


Figure 6 – NR PM₁ chemical composition, triangle plot and average size distribution for: IOP1 – wet season (top) and IOP2 – dry season (bottom).

Organic aerosol source apportionment – IOP2

A Positive Matrix Factorization analysis has been applied to the organic aerosol spectra during the dry season, resulting in four factors: HOA (Urban), BBOA (biomass burning), Fac82 (isoprene SOA under low NO_x condition) and LV-OOA (highly oxidized aerosols). LV-OOA factor dominates the OA mass (40%), whereas the other three factors show similar contribution. Further analysis include running PMF analysis on entire dataset and integration with trace gases measured at the site.

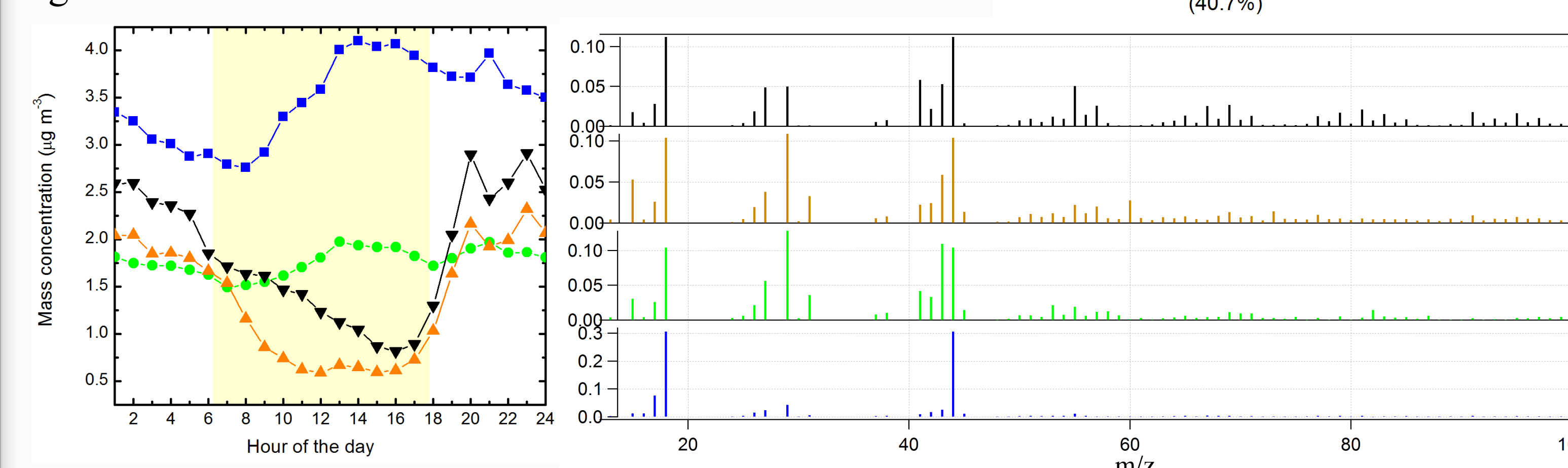


Figure 7 – preliminary results of Positive Matrix Factorization of OA measured during IOP2