



Ultrafine particles and black carbon measurements at an urban background site in Milan

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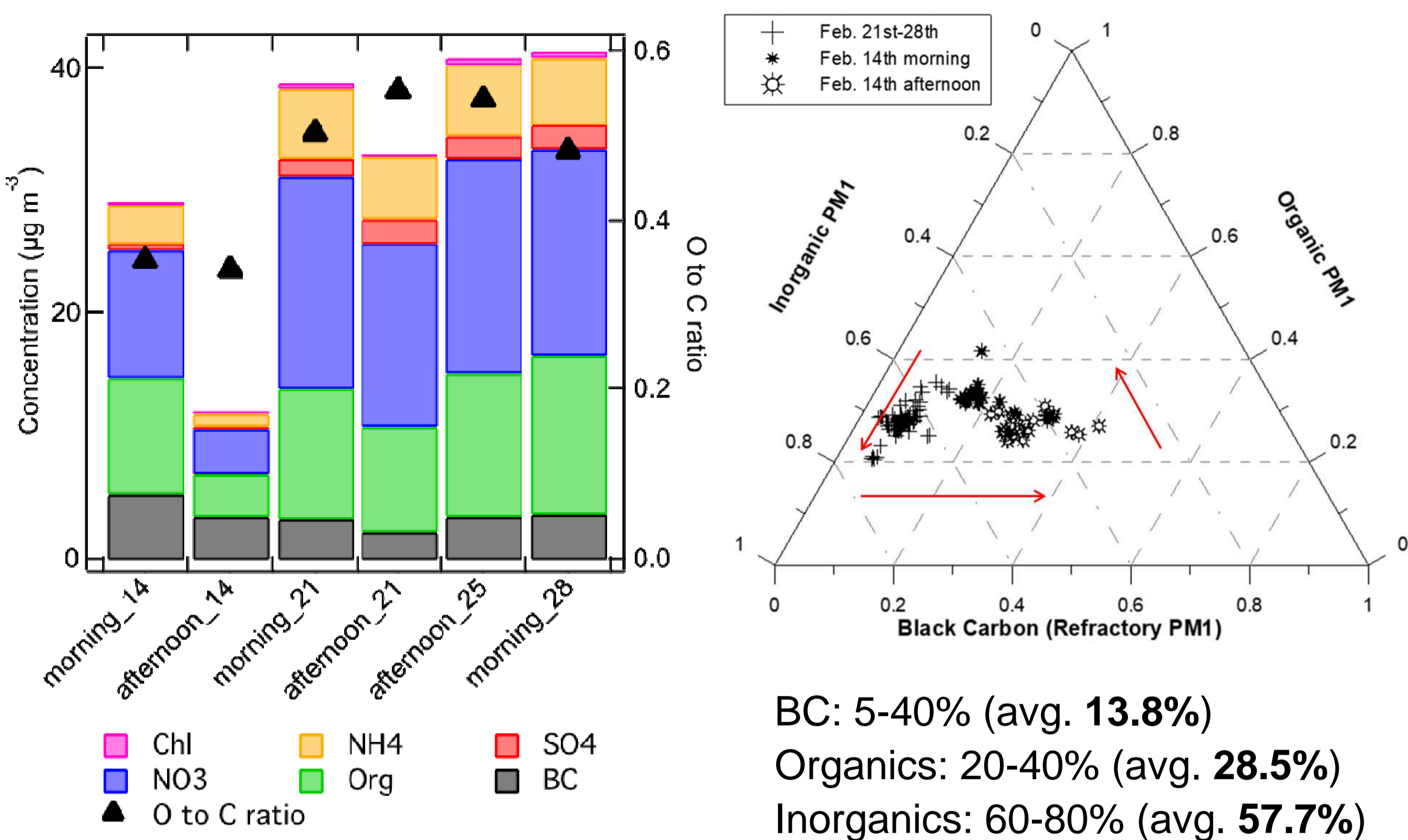
Background

During the PoAir intensive air quality monitoring campaign over Northern Italy, collocated measurements of fine and ultrafine particles were performed at an urban background site in Milan for a few days in February 2014. Concurrent measurements at high time resolution of the different features of airborne particles (optical properties, number size distribution, chemical composition) have been performed in order to investigate the potential for a better understanding of the nature and origin of submicron particles in urban environments.

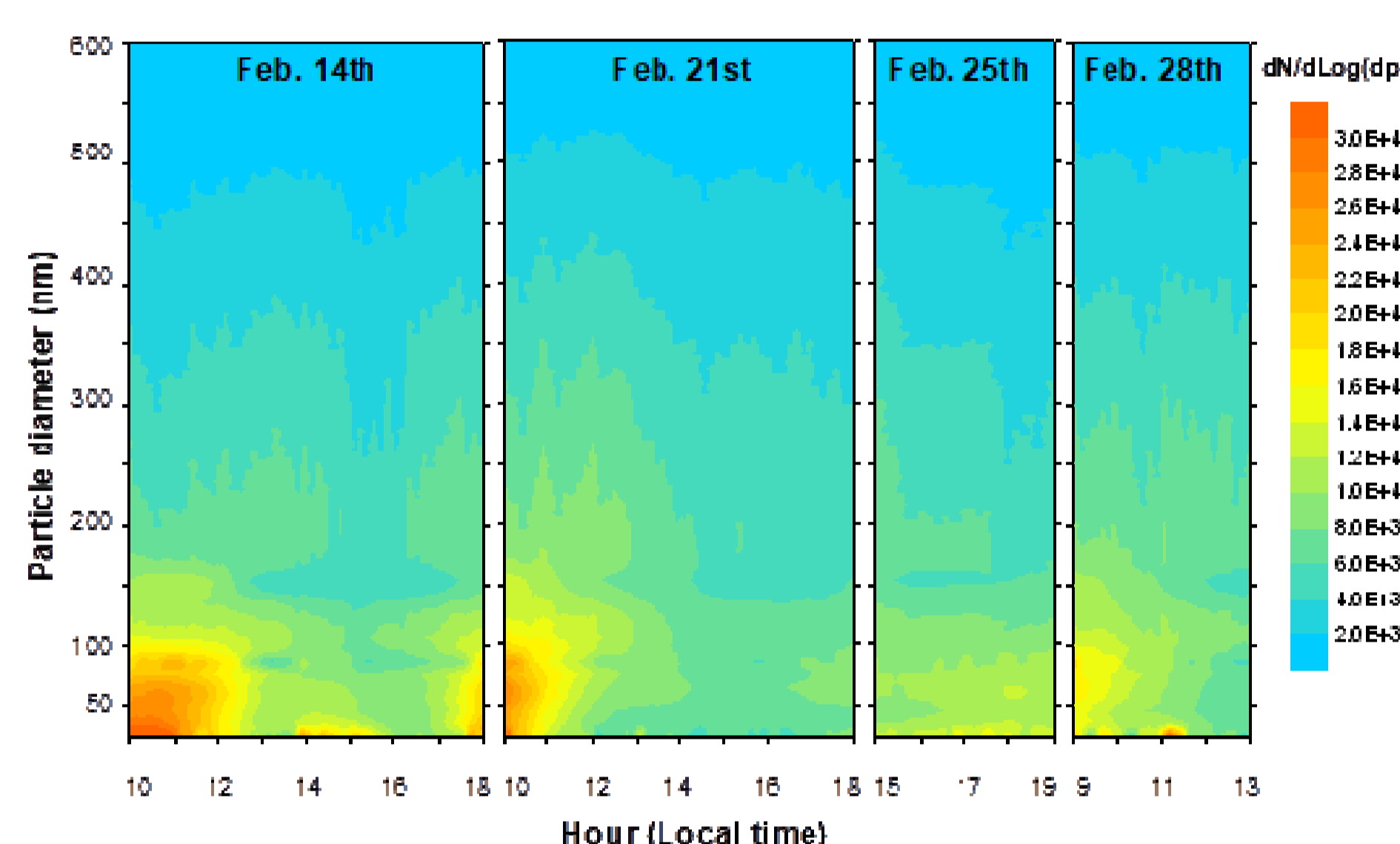
Black carbon (BC) was measured through Laser-Induced Incandescence (LII) technique (Italian Patent ITRM20090617), microaethalometer AE51 (AethLabs), multi-angle absorption photometer (MAAP); size-resolved particle number concentration (PNC) in six size bins (20-30 nm, 30-50 nm, 50-70 nm, 70-100 nm, 100-200 nm, and > 200 nm) through a TSI 3031 UFP monitor; non-refractory PM1 chemical composition through a High Resolution - Time of Flight - Aerosol Mass Spectrometer (HR-TOF-AMS).

All measurements were taken at 10-min time resolution, thus enabling to capture the short-period fluctuations, due to fresh emissions from combustion sources (namely from the traffic source), typical of kerbside locations but that can also affect urban background sites.

PM1 chemical composition



Particle number and size



Concentration levels

PNC₂₀₋₁₀₀₀: 5.7·10³ - 24·10³ cm⁻³
(avg. 10.3·10³ cm⁻³)
PNC₂₀₋₁₀₀: 3.8·10³ - 20·10³ cm⁻³
(avg. 7.8·10³ cm⁻³)

➤ UFP = 70-80% of total PNC

Effect of Relative Humidity

avg. PMD = 73 nm if RH < 60%
avg. PMD = 90 nm if RH > 60%

PM1 average composition and ageing

BC = 3.1 μg m⁻³, Organics = 13.0 μg m⁻³
NO₃⁻ = 13.0 μg m⁻³, SO₄²⁻ = 1.4 μg m⁻³, NH₄⁺ = 4.2 μg m⁻³

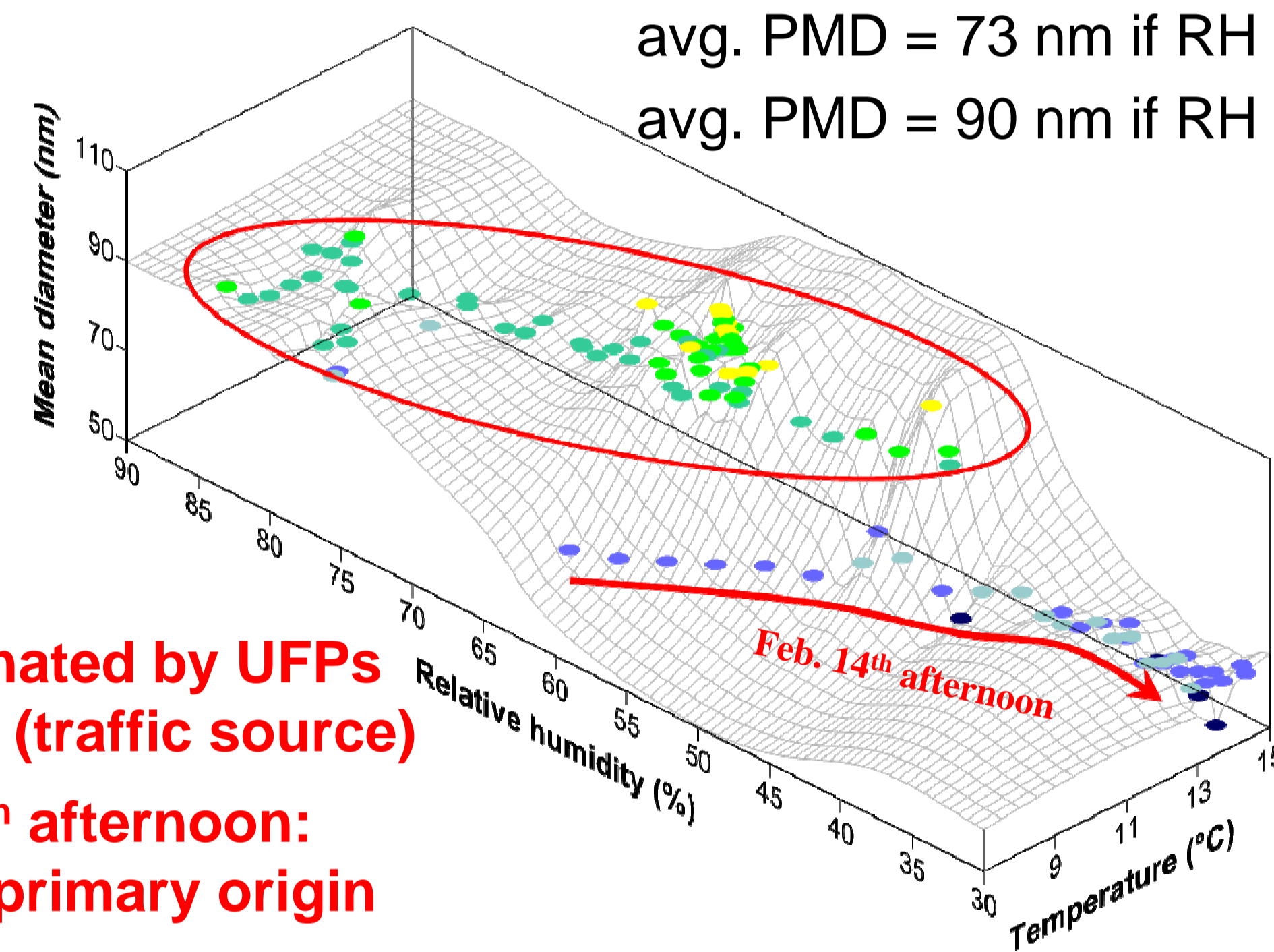
- Ammonium nitrate dominating over ammonium sulfate coherently with the relatively low temperature and with the NO_x (high) and SO₂ (low) concentrations in Milan urban area.

Oxygen-to-carbon ratio of Organic Aerosol: 0.34-0.55 range

- Dominant contribution of processed OA (typically O:C > 0.4) over primary combustion OA (typically O:C < 0.2)

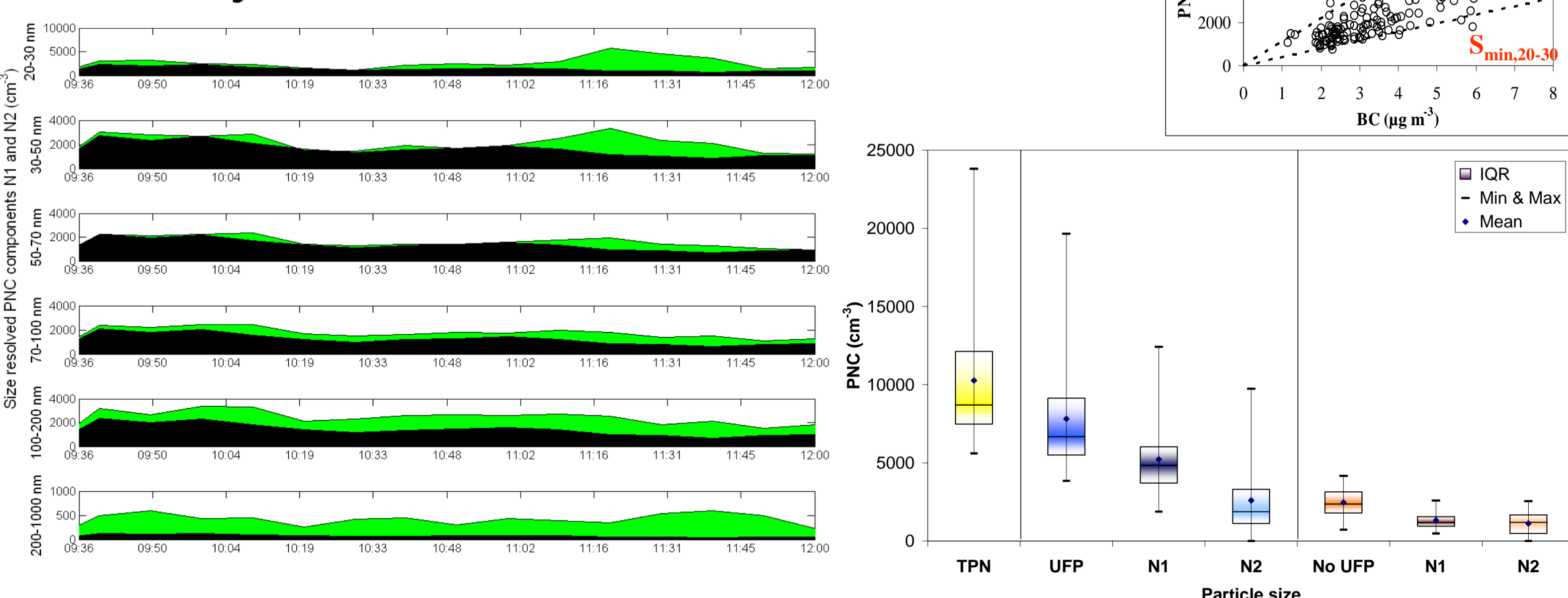
UFP time pattern

- Size distribution dominated by UFPs on mid-morning hours (traffic source)
- "Clean air" on Feb. 14th afternoon: smaller particles of primary origin
- PNC₂₀₋₁₀₀₀ and BC well correlated (r = 0.80 for the entire dataset; r = 0.6-0.9 on single-day basis)



PNC split

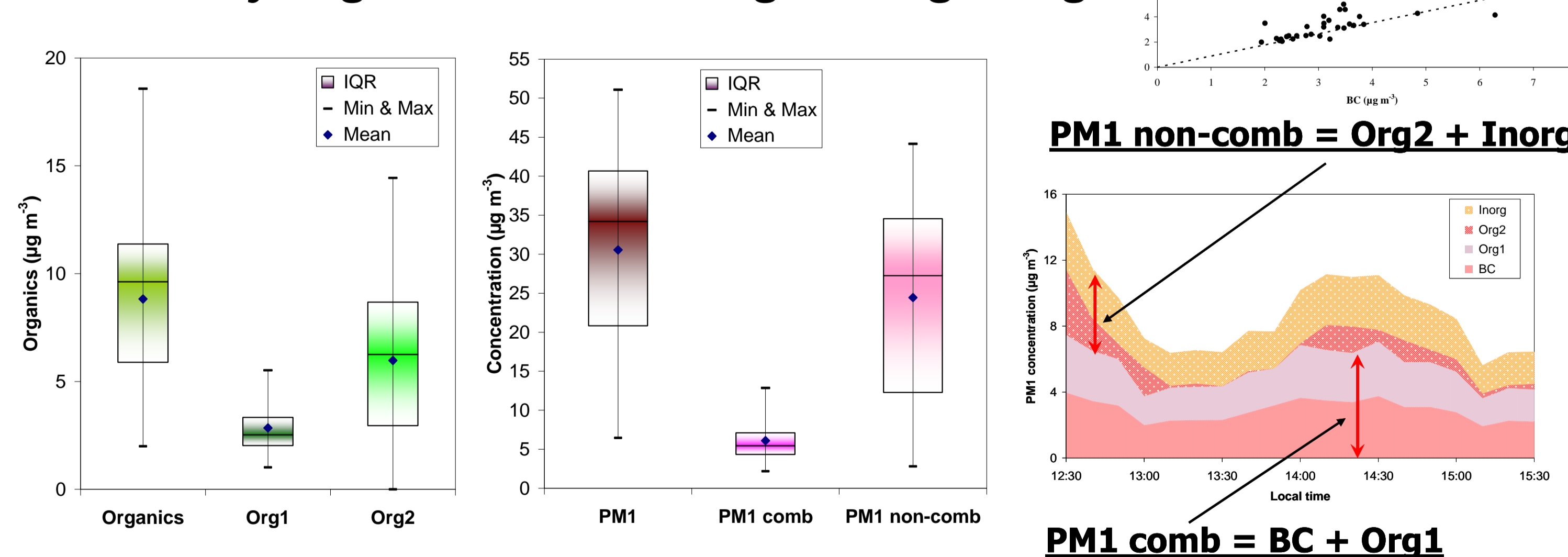
Primary combustion particles: N1 = S_{min,PNC}·BC
Secondary aerosol: N2 = N - N1



- N1: the most part (65%) of PNC₂₀₋₁₀₀₀, with higher contributions (71-77%) for the size bins between 30-100 nm
- N2: only 35% of the total of PNC₂₀₋₁₀₀₀, with the highest contribution (70%) in the PNC₂₀₀₋₁₀₀₀ size bin

PM1 organic matter split

Primary organic particles: Org1 = S_{min,Org}·BC
Secondary organic aerosol: Org2 = Org - Org1



- S_{min,Org} from "clean air" event: 0.88 μg_{Org}/μg_{BC}
- PM1 organic mass mostly secondary (67.8%) giving 19.6% of PM1
- Secondary ("non-combustion generated") organic and inorganic components accounting for about 80% of PM1 mass, but down to 40% on "clean air" event

Conclusions

- Despite the small dataset, highly resolved data show a potential for a better understanding of the nature and origin of ultrafine and submicron particles in urban environments
- BC as a tracer of primary emission from combustion processes allows splitting particle number and organic mass concentration data between primary and secondary components
- Primary, "combustion-generated", aerosol is responsible for the most part of particle number (especially in the UFP range), conversely with limited contribution to PM1 mass, except for the "clean air" event