

Supplement of

In situ formation and spatial variability of particle number concentration in a European megacity

M. Pikridas et al.

Correspondence to: S. N. Pandis (spyros@chemeng.upatras.gr)

The copyright of individual parts of the supplement might differ from the CC-BY 3.0 licence.

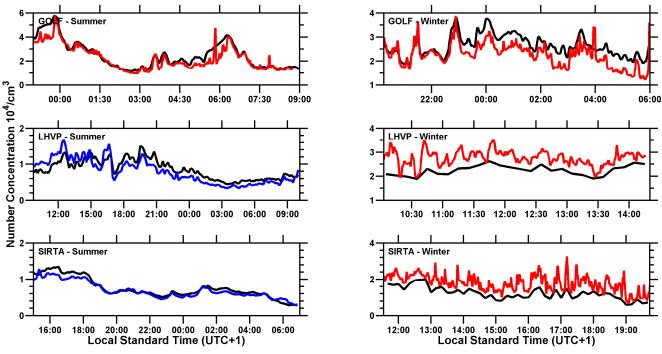


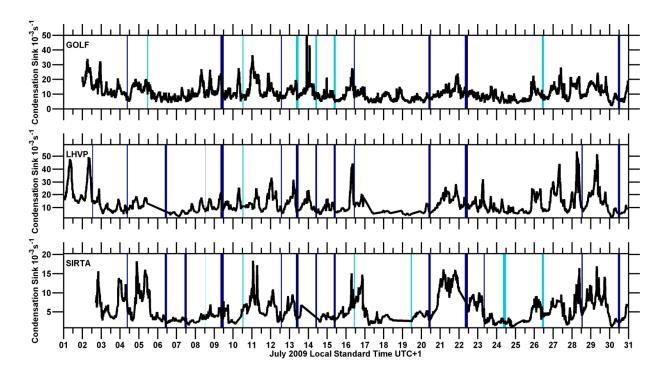


Figure S1. Time series of particle number concentration used for intercomparison between aerosol sizing instrumentation located at the stationary sites (black lines) and the mobile laboratory CPC (MoLa - red line; MOSQUITA - blue line). During summer the difference does not exceed 10% at all sites. The larger discrepancy at SIRTA during winter is attributed to the lower size detection limit of the CPC employed at MoLa (2.5 nm) compared to the 10 nm of the SMPS system at SIRTA.

- 52
- 53

54

55



56 Figure S2. Condensation sink measured at the three ground sites during July 2009. Dark

57 and light blue bars indicate the event and undefined periods, respectively.

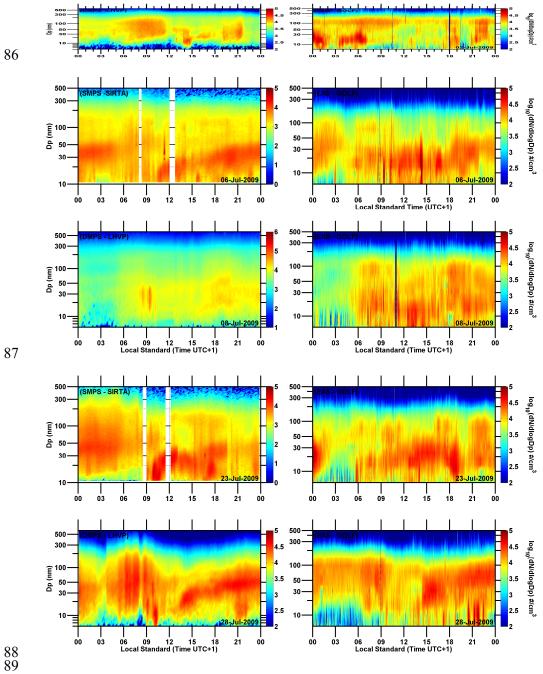


Figure S3. Number size distribution time series when a nucleation event was identified at SIRTA and/or LHVP but not at GOLF. D_p is the particle diameter.

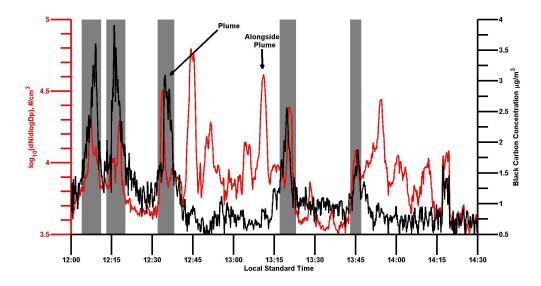
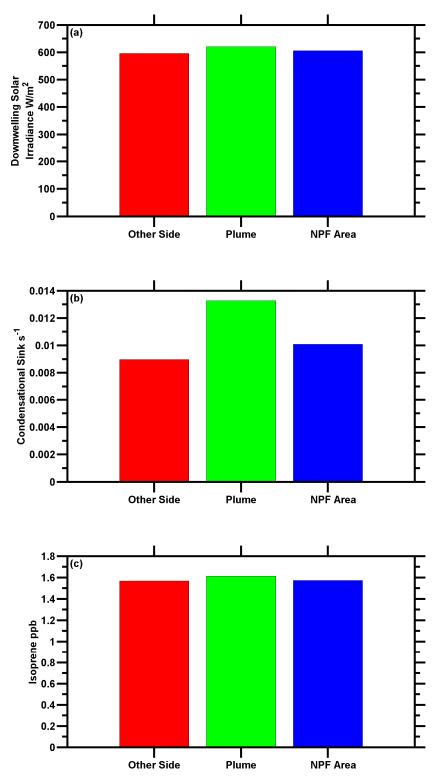


Figure S4. Number (red line) and black carbon (black line) concentrations during airborne measurements on July 1st 2009. Number concentration increases observed simultaneously with increases in black carbon mass concentration (grey areas) were attributed to the Paris plume. Number concentration increases were also observed along the plume.



105 106 Figure S5. Downwelling solar irradiance (top), condensation sink (middle) and isoprene 107 concentration (bottom) comparison of the Paris plume with areas on either side of the 108 plume when high particle concentrations were observed at one side outside of the plume. 109 Significant differences among these areas were not observed with respect to 110 condensation sink, isoprene and solar irradiance.