

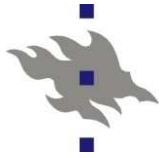


HELSINGIN YLIOPISTO
HELSINGFORS UNIVERSITET
UNIVERSITY OF HELSINKI

Continuous air pollution and surface-atmosphere interaction measurements at the SMEAR III station in Helsinki

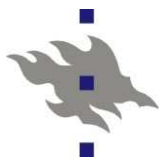
Leena Järvi et al.

Ubcasting workshop 10.9.2008



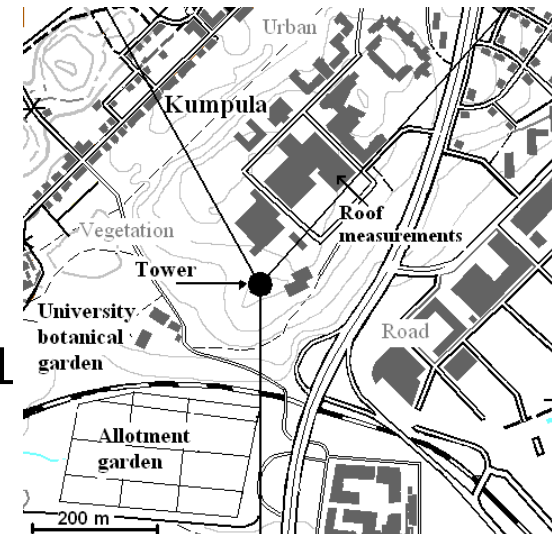
Motivation

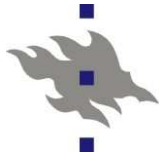
- Urban areas create very different circumstances for the lower level of the atmosphere
 - Many of the pollution sources and majority of people are located in cities
 - Urban areas are characterized by high surface roughness and different thermal conditions which affect turbulent mixing and further pollutant dispersion
- Still lot of information missing about the sources, sinks and mixing of air pollutants
- Measurements at the SMEAR III station try to give answers also these questions



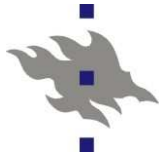
Measurements used in this study

- The aerosol particle number concentration with size range 3 nm-20 μm are measured with twin DMPS and APS
- Particles are divided in three classes due to their different dynamics and sources
 - Ultrafine particles (UFP, $d < 100 \text{ nm}$), accumulation mode particles ($100 \text{ nm} < d < 1 \mu\text{m}$) and coarse particles ($d > 1 \mu\text{m}$)

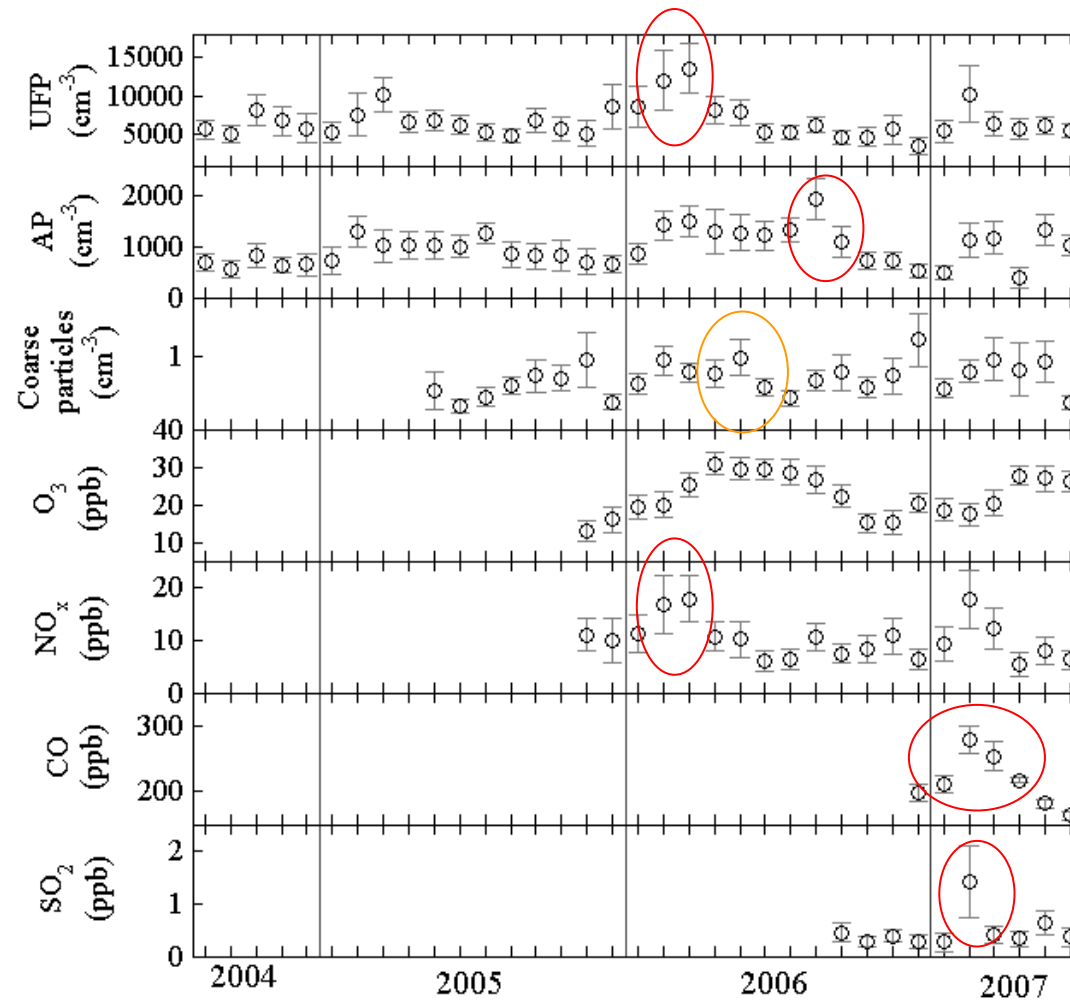


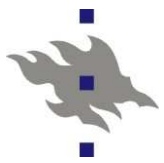


- Pollutant gases: Ozone (O_3), nitrogen oxides (NO_x), carbon monoxide (CO) and sulphur dioxide (SO_2)
- Meteorological variables: Pressure, solar radiation, wind speed and direction, RH and turbulent exchange
- Data between Aug 2004 and Jun 2007 is shown
 - Not all variables available the whole time
- Data was divided into four seasons: winter, spring, summer and fall

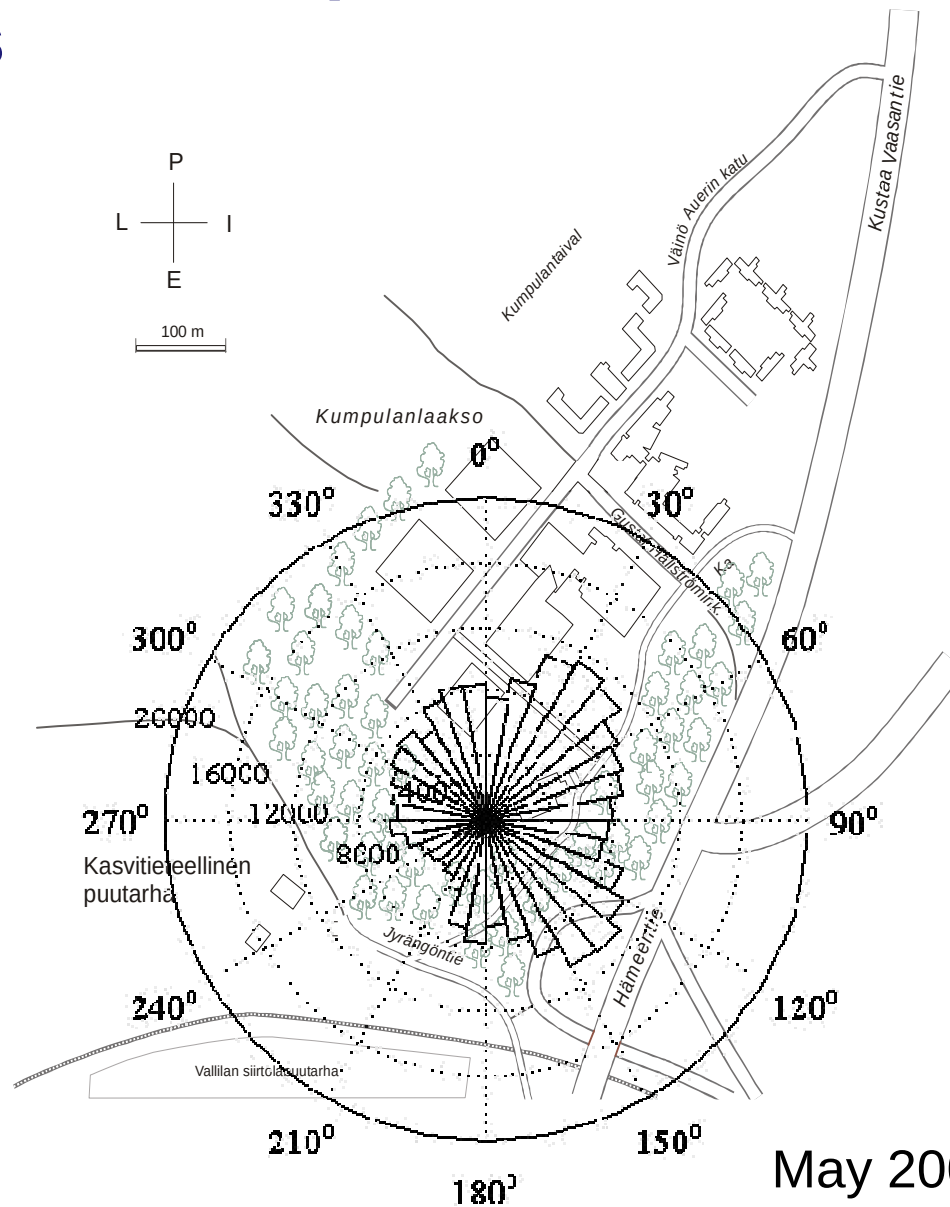


Time series of pollutant concentrations



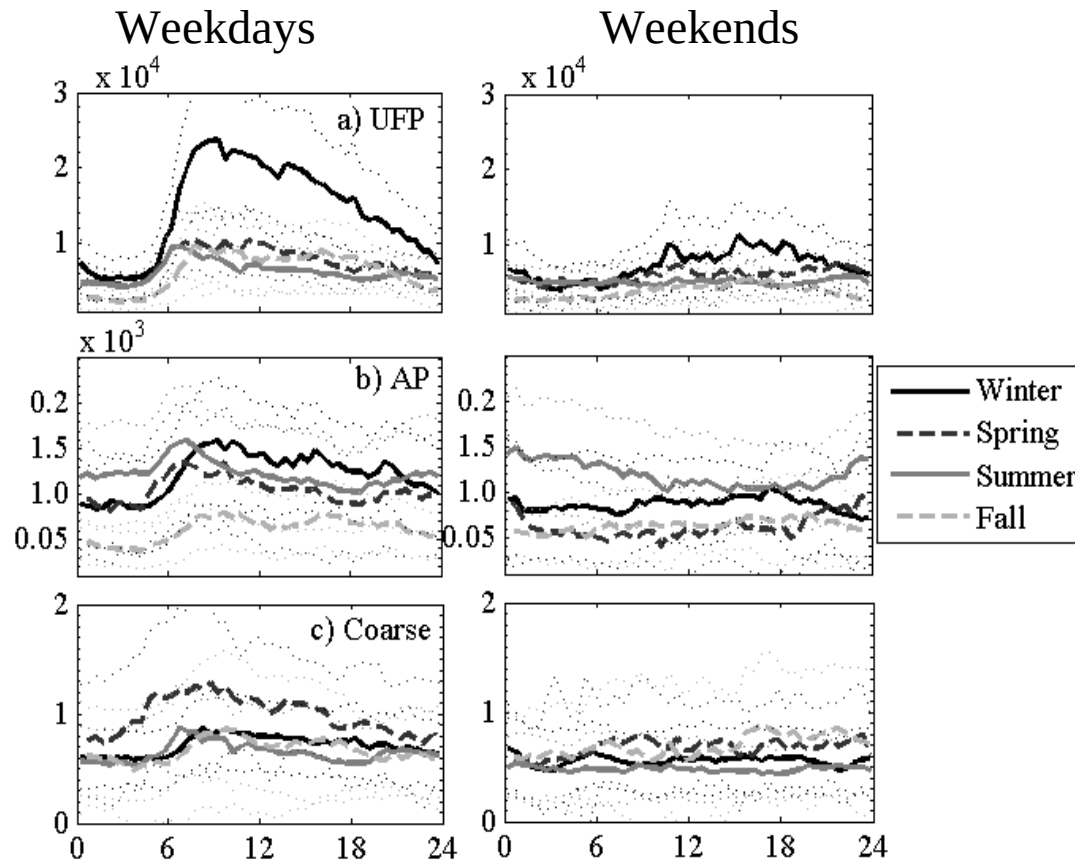


Wind direction dependence of ultrafine particles



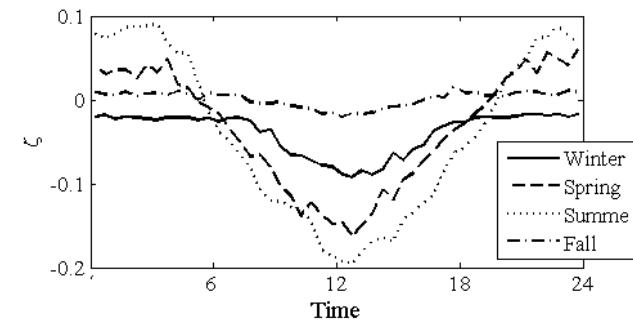


- **Diurnal behavior of aerosol particle concentrations for different seasons and separately for weekdays and weekends**

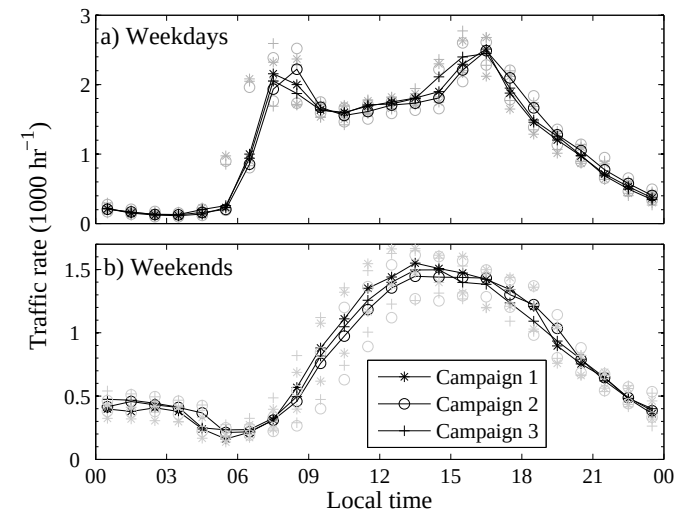


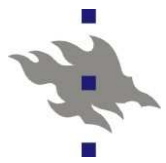
May 2005-Jun 2007

Järvi et al. (2007)



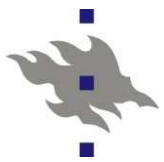
Järvi et al. (2008)



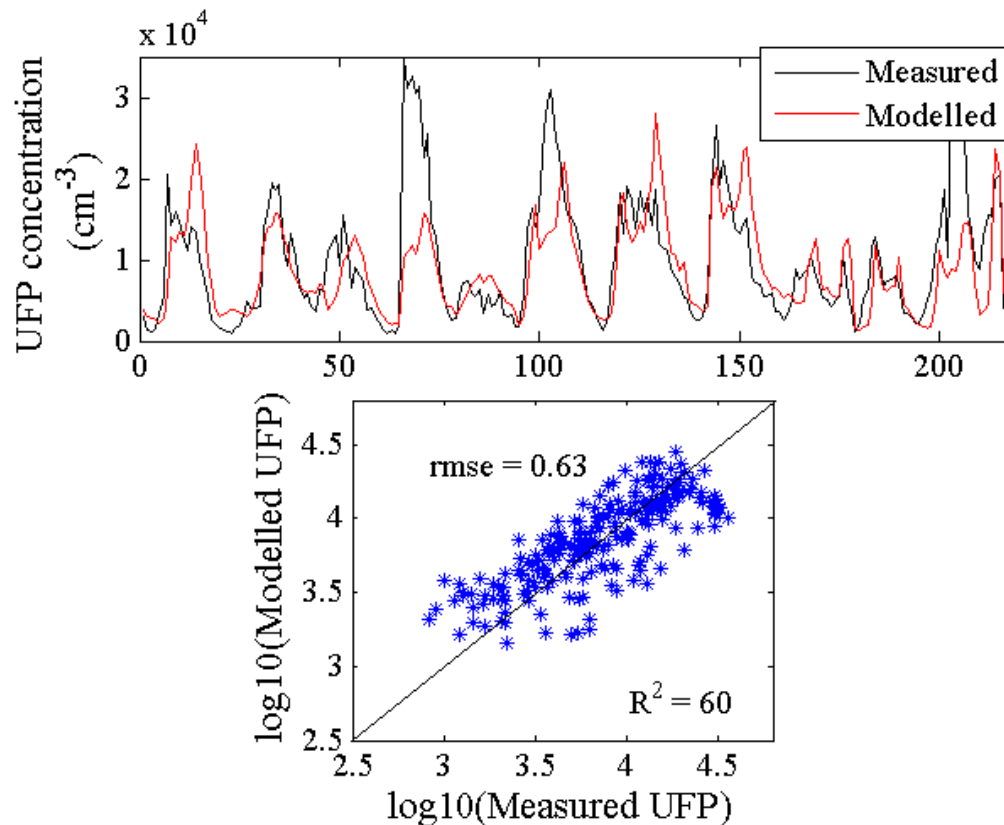


Dependency of particle concentrations on traffic rate and meteorological variables

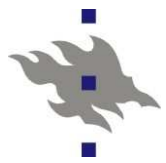
- A multiple linear regression (MLR) models ($Y = b_0 + b_1X_1 + \dots + b_nX_n$) were made for UFP, accumulation mode particle and coarse particle concentrations
- We wanted to find those independent variables which minimize the difference between the measured and modeled concentrations
- We also get the relative importance of each independent variable compared to the other variables in the model
- MLR was made for Dec 2005-Aug 2006.



Example from MLR analysis: Ultrafine particle concentrations in Dec 2005

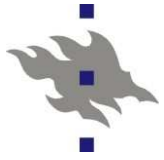


Variable	β -constant
H ₂ O	-0.15
WD ₁	0.04
σ_w	-0.21
Traffic	0.70

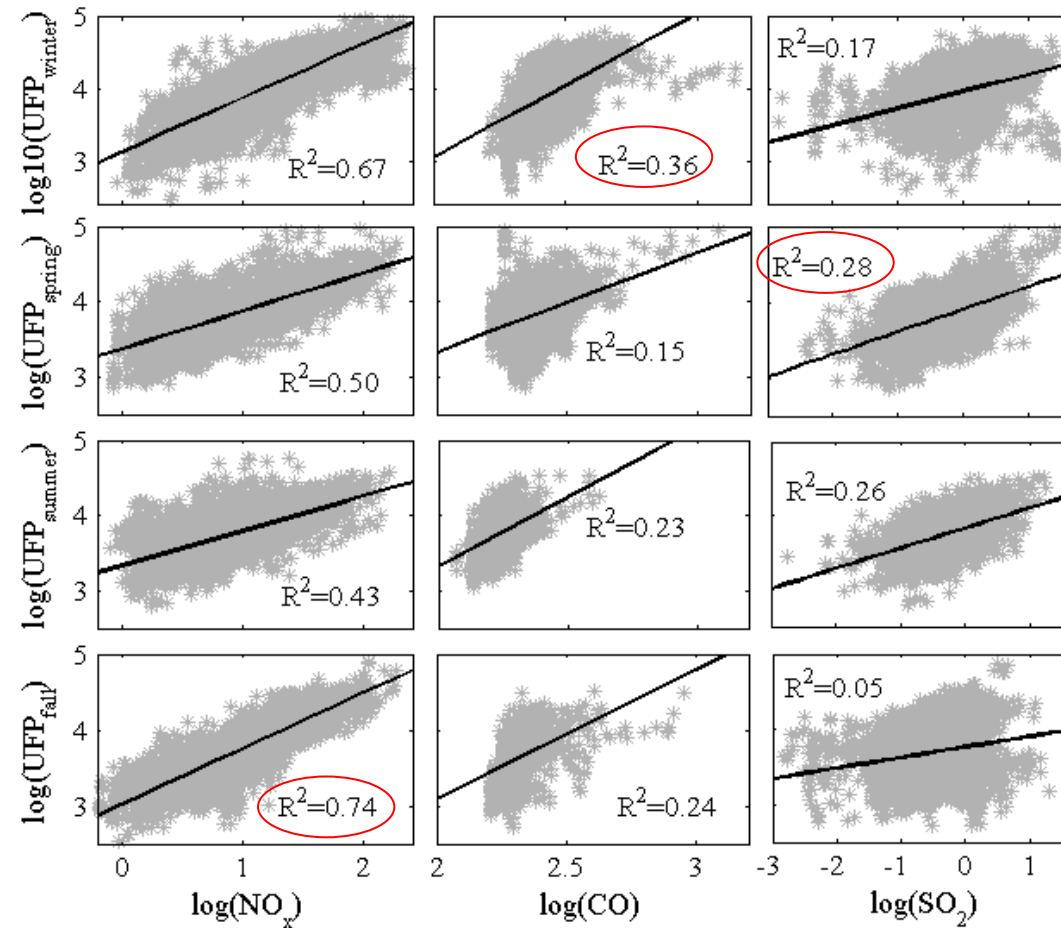


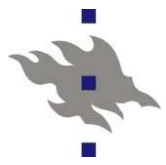
Main findings from the MLR analysis

- UFP concentrations could be explained best with the available variables
 - Most affected by traffic, turbulent mixing and H₂O
- In the case of accumulation mode particles, traffic was equally or less important than meteorological variables
- Coarse particle concentrations could not be explained as well as fine particle concentrations
 - Humidity had an great impact especially in spring
- Turbulent mixing had an inverse effect on different sized of particles
 - Increases the mixing volume and concentrations decrease
 - On the other hand, larger particles are re-suspended more efficiently

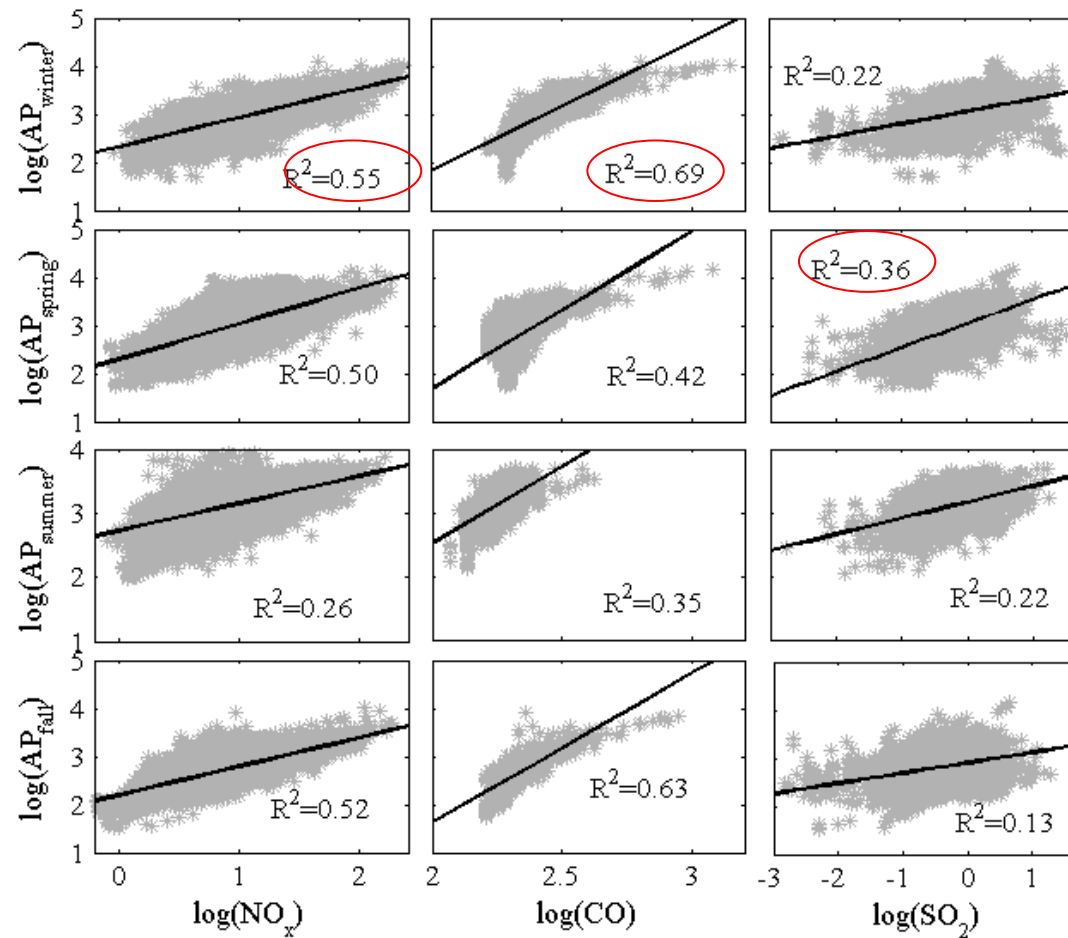


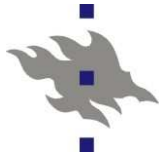
Correlations between ultrafine particles and concentrations of NO_x , CO and SO_2





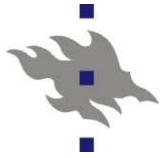
Correlations between accumulation mode particles and concentrations of NO_x , CO and SO_2





Conclusions

- Most the pollutants showed dependence on traffic
 - For particles, the importance decreased with increasing particle size
- Fine particles, NO_x , CO and SO_2 experienced their maxima in winter due to the lowered mixing and enhanced emissions
- The effect of re-suspension could be seen in coarse particle concentrations especially in spring



- Long-range transport had an effect on accumulation mode particles (Aug 2006!)
- Correlation between UFP and SO_2 increased in spring and summer → New particle formation?