

#### Two years air quality study by DOAS within Beijing

Klaus Schäfer<sup>1</sup>, Yuesi Wang<sup>2</sup>, Jinyuan Xin<sup>2</sup>, Hong Ling<sup>2</sup>, Peter Suppan<sup>1</sup>

 <sup>1</sup> Karlsruhe Institute of Technology, Institute for Meteorology and Climate Research, Atmospheric Environmental Research (KIT/IMK-IFU), Garmisch-Partenkirchen, Germany
<sup>2</sup> Chinese Academy of Sciences (CAS), Institute of Atmospheric Physics, State Key Laboratory of Atmospheric Boundary-Layer Physics and Atmospheric Chemistry (LAPC), Beijing, P. R. China

INSTITUTE FOR METEOROLOGY AND CLIMATE RESEARCH, ATMOSPHERIC ENVIRONMENTAL RESEARCH (IMK-IFU)



#### Scientific questions

#### Process studies

Methodology Influences upon air pollution

#### Future tasks

KIT – University of the State of Baden-Württemberg and National Large-scale Research Center of the Helmholtz Association



# Scientific questions for air quality in Beijing

Origin of frequently occurring air pollution events

Origin of pollutants - urban agglomerations one of the most important sources for pollutants

Local and regional wind systems - bring fresh air masses and limit air pollution

Role of mixing layer height (MLH) – mountains West to North



# **Process studies**

# Influences upon air pollution

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# Air quality studies

Continuous determination of mixing layer height (MLH) by ceilometer

Limits the vertical distribution of emitted air pollutants with consequences for dilution and transport

Essential for the determination of speed and range of vertical dispersion

Influenced by future climate change and thus important for quality of living in large cities



MLH influence upon air pollution in urban and sub-urban area Hannover, Munich, Augsburg, Budapest, Zurich Airport, Mexico City International Airport, Athens International Airport, Paris CDG

Correlation with MLH smallest inside street canyons

Correlation with MLH larger in winter than in summer

Influences of MLH upon CO, NO<sub>2</sub> and PM<sub>10</sub> concentrations in the order of 20 %, up to 50 %

therefore better MLH determination necessary
deployment of ceilometers for continuous operation

Schäfer, K., S. Emeis, H. Hoffmann, C. Jahn: Influence of mixing layer height upon air pollution in urban and sub-urban areas. Meteorol. Z., 15, 647-658 (2006).



#### Wind influences upon air pollution

Under strong background flows: reduced concentrations for all pollutants without distinct maxima and minima of diurnal cycle

Under the development of local flows: high concentrations of air pollutants

Influences of wind speed upon CO, NO<sub>2</sub> and PM<sub>10</sub>: in the order of 20 %

Schäfer, K., Emeis, S., Hoffmann, H., Jahn, C., Müller, W., Heits, B., Haase, D., Drunkenmölle, W.-D., Bächlin, W., Schlünzen, H., Leitl, B., Pascheke, F., Schatzmann, M.: Field measurements within a quarter of a city including a street canyon to produce a validation data set. International Journal of Environment and Pollution, 25, 1/2/3/4, 201-216, (2005).

Dandou, A., Tombrou, M., Schäfer, K., Emeis, S., Protonotariou, A.P., Bossioli, E., Soulakellis, N. Suppan, P.: A comparison between modelled and measured mixing layer height over Munich. Boundary-Layer Meteorology 131, 425–440 (2009).

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# Air quality studies in Beijing



Comparison motorway with background concentrations

- diurnal variations
- influences upon gaseous concentrations: emissions, meteorology
- gases of interest for secondary aerosol formation

Path-averaged concentrations of air pollutants NO<sub>2</sub>, SO<sub>2</sub>, O<sub>3</sub>, (benzene, toluene, xylene, NO, NH<sub>3</sub>, HCHO)

- near / across a motorway April 2009 March 2011
- commercial DOAS
- three retro-reflectors
- automatic operation

#### Instrument





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#### **Measurement sites**



#### LAPC tower, ceilometer, DOAS from 13 July 2009 on



Path1- the first floor of the tower 126m(double) Path2- on the roof in our courtyard 266m(double)

Path3- in the middle of the lampstandard 568m(double)

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#### Calibration





**Calibration system** 



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**Light source** 



#### **Comparison - instruments**





- chemiluminescence NO-NO<sub>2</sub>-NO<sub>x</sub> analyzer
- UV photometric O<sub>3</sub> analyzer
- pulsed fluorescence SO<sub>2</sub> analyzer

• GC/MS system (Finnigan Trace 2000/DSQ, ThermoFisher, USA) with fuse-silica capillary column for aromatic hydrocarbons



#### Comparison O<sub>3</sub> (path 2)





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#### Comparison SO<sub>2</sub> (path 2)





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#### Comparison NO<sub>2</sub> (path 2)





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### Comparison NO (path 1)





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#### **Comparison Aromatic Hydrocarbons (path 2)**





Average difference between DOAS and GC/MS:

BEN: 0.94 ppbv (64% higher)

TOL: -0.56 ppbv (20% lower)

PXY: -0.49 ppbv

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#### **Comparison HCHO (path 3)**



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Continuous determination of mixing layer height by ceilometer since February 2009 **DOAS** installation April 2009 until July 2009

4000 m



**Optical remote sensing:** Ceilometer Vaisala LD40 or CL31 wave length: 855 or 910 nm range: Resolution: 10 or 7.5 m Second

Meteorological monitoring at 325 m LAPC tower

#### **Ceilometer measurements**



#### High particulate load and low mixing layer height



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#### Correlations NO<sub>2</sub> – mixing layer height April – May 2009

#### Path 1 across the road nearby



#### Path 2 in the background



#### Path 3 across the motorway



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# Mixing layer height - air quality

If planetary boundary layer > 1000 m: often multiple layering if < 1000 m during daytime: often one layer

Influence of MLH upon NO<sub>2</sub>: relevant – standard error 0.15

Logarithmic regression best correlations: NO<sub>2</sub> well mixed but not at motorway – exponential dependence

Correlations of NO and SO<sub>2</sub> with MLH: not significant

Concentrations of BTX and HCHO: near the detection limit



# **Further tasks**



# Tasks for air quality studies in Beijing

Further influences upon air quality

emissions (in the case of  $NO/NO_2$  and  $SO_2$ )

air chemistry (photochemistry in the case of NO<sub>2</sub>)

by small-scale model studies on the basis of

road traffic data - emission modelling

DOAS together with in situ concentration and meteorological data – transport-chemistry modelling



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