

# Source apportionment of submicron organic aerosol at an urban background site in Zürich

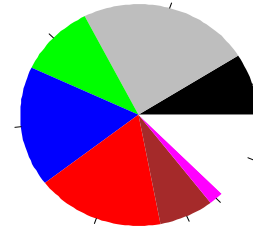
11<sup>th</sup> ETH-Conference on Combustion Generated Nanoparticles

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# Overview

- Introduction: Atmospheric aerosols



- Methods: Receptor modeling

$$\mathbf{x}_i = \mathbf{C}\mathbf{S}_i + \mathbf{E}_i$$

- Results: Zürich-Kaserne



- Conclusions



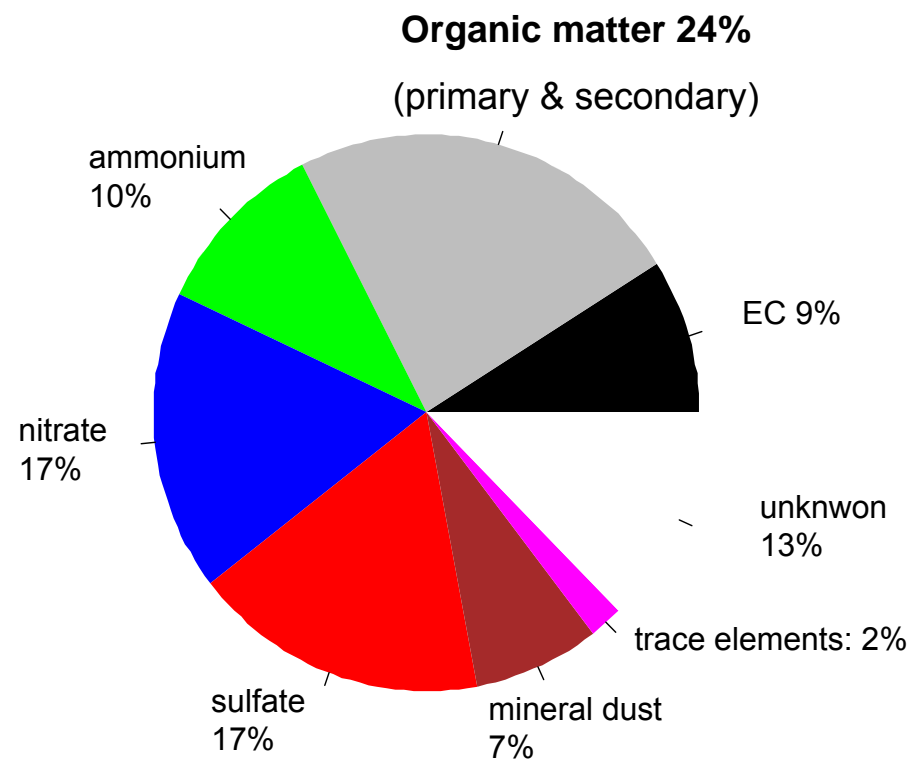
# Atmospheric (organic) aerosols

- Impact on health (climate, ecosystems, visibility, ....)

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- Composition:

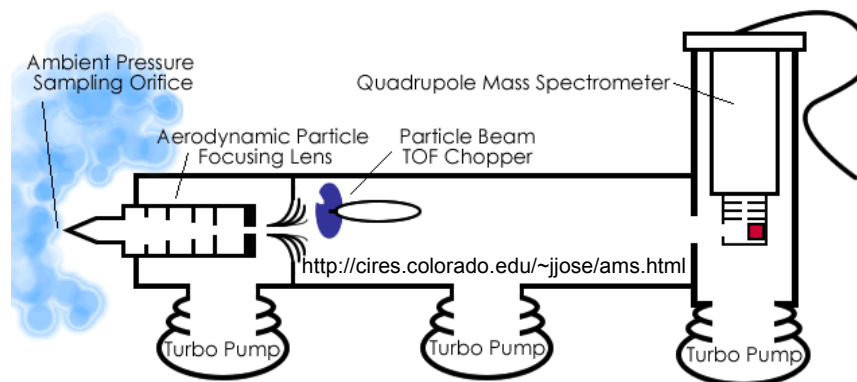


PM<sub>2.5</sub> (Zürich-Kaserne; Hueglin et al., 2005)

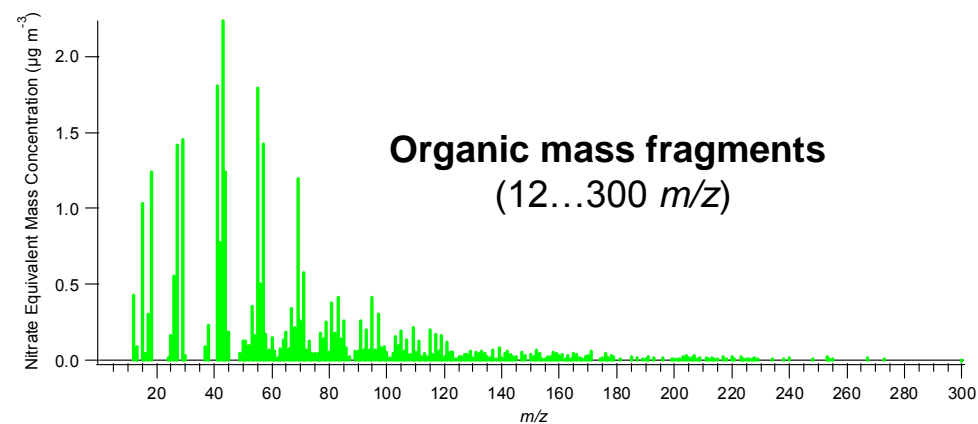


# Aerosol measurements

- Aerosol mass spectrometer (Q-AMS, Aerodyne Inc.)



- Non-refractory, submicron ( $\sim$ PM<sub>1</sub>) aerosol
- Online, (semi-)quantitative
- High time-resolution (sec - minutes)



# Receptor modeling

- **Receptor models** (linear mixing models)

$$\mathbf{X}_i = \mathbf{C}\mathbf{S}_i + \mathbf{E}_i$$

$\mathbf{X}_i$ : multivariate observation (m-vector)

$\mathbf{C}$ : loadings (m x p-matrix),  $C \geq 0$

$\mathbf{S}_i$ : scores (p-vector),  $S \geq 0$

*measured AMS spectrum (270 dimensional)*

*aerosol source profile (270 x p-matrix)*

*p latent aerosol source activities*

- No assumptions about source distributions
- Optional: meteorology/source composition
- Types

CMB - chemical mass balance:

*chemical signature of sources known*

PMF - positive matrix factorization:

“ “ “ “ unknown

# Receptor modeling II

reconstructed data matrix

$$\hat{\mathbf{X}} = \mathbf{CS}$$

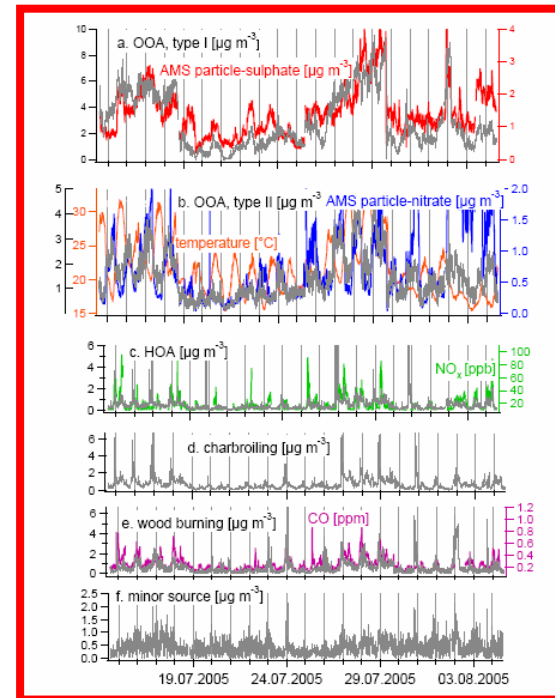
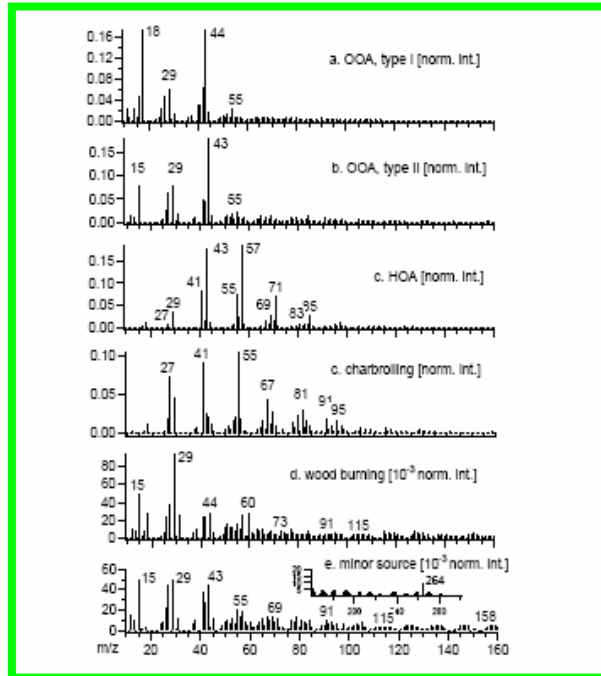
measured data matrix (X)

$$\mathbf{X} - \hat{\mathbf{X}} = \mathbf{E}$$

modeled **source profiles**

modeled time series of **source activities**

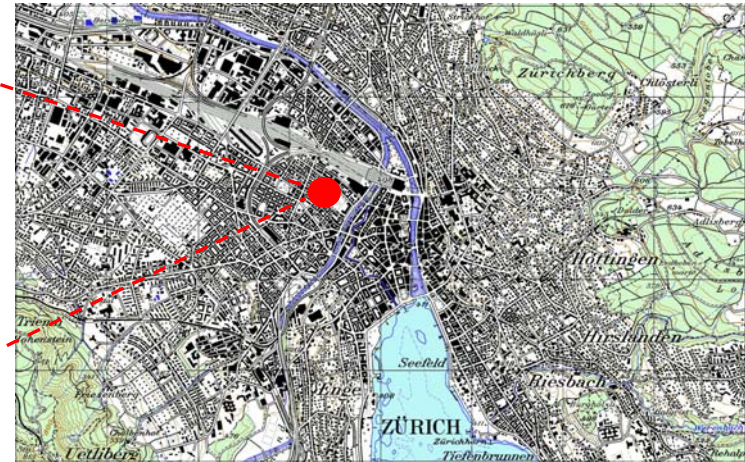
factors/sources



# AMS sampling site: Zürich, urban background



Urban background site (Zürich-Kaerne)  
Public backyard



500 meters from Zurich main station  
1 mile from lakeside

Site of the Swiss National Air Pollution Monitoring Network (NABEL; [www.empa.ch/nabel](http://www.empa.ch/nabel))

## Ancillary data to validate math. model:

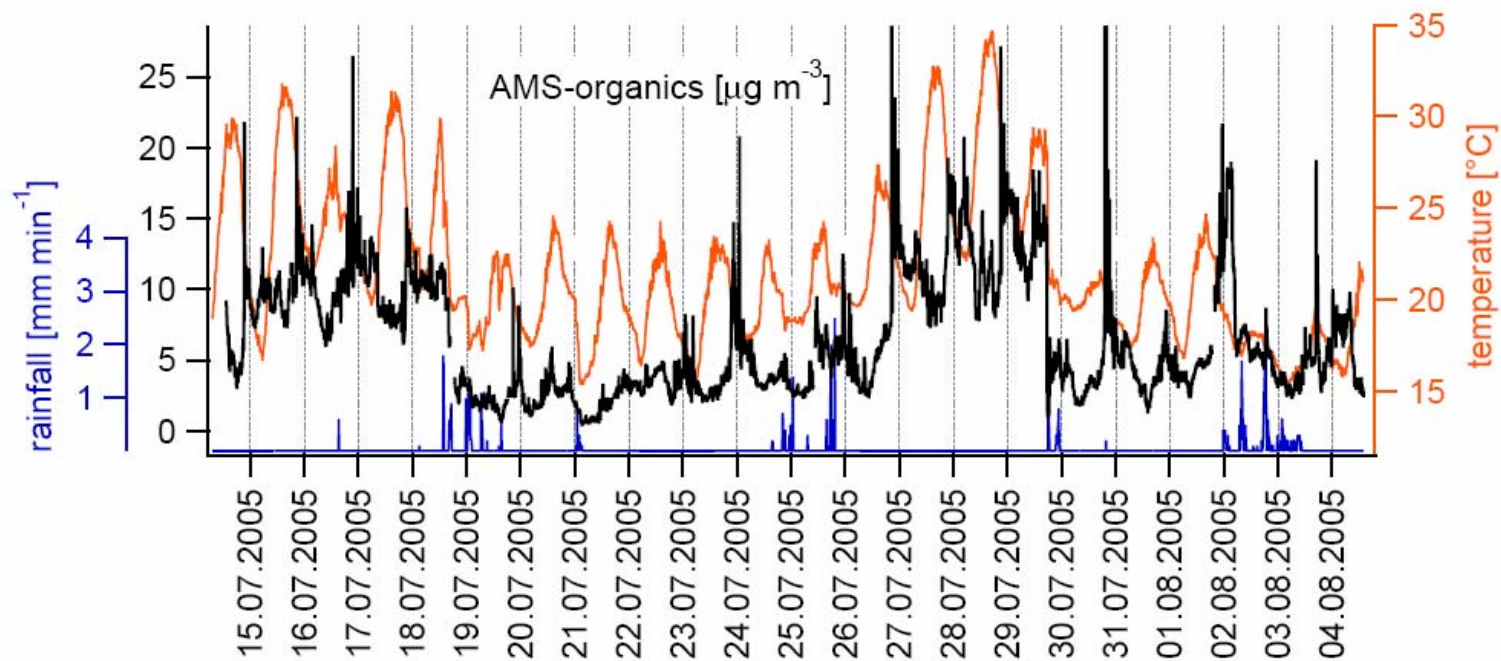
- meteo (radiation, wind direction, temperature...)
- gas-phase (CO, NO<sub>x</sub>, O<sub>3</sub>, SO<sub>2</sub>, VOCs, ...)
- aerosol (PM<sub>10</sub>, OC, EC)





# Zürich, Summer 2005

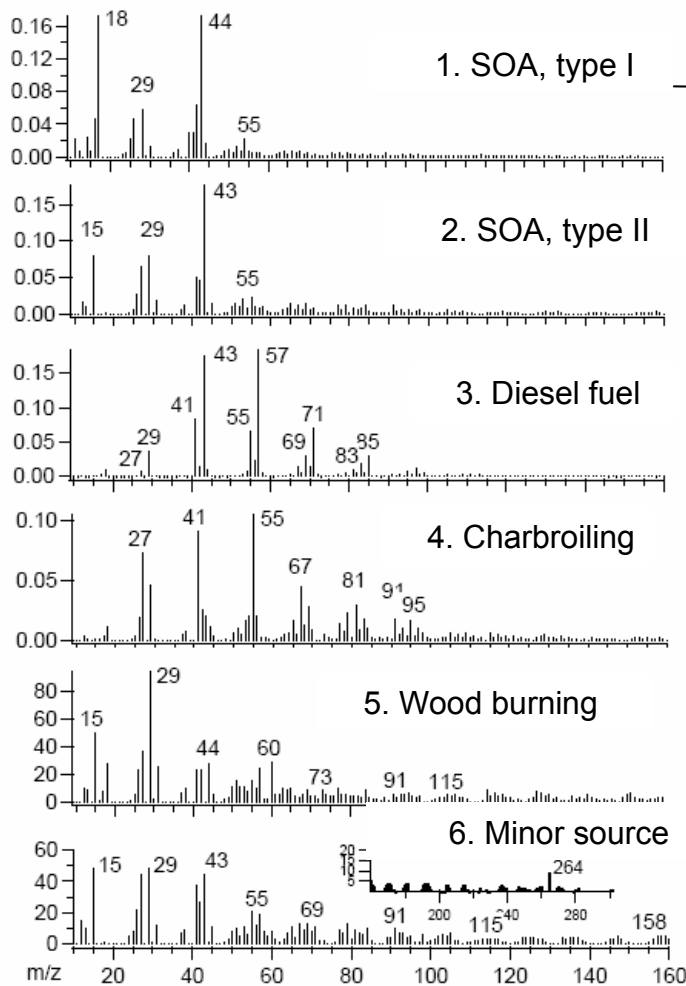
- 14 July – 4 August 2005
- **Photochemical episodes** („summer smog“)
- Positive matrix factorization (PMF)  
(Lanz et al., 2007, Atmos. Chem. Phys.)



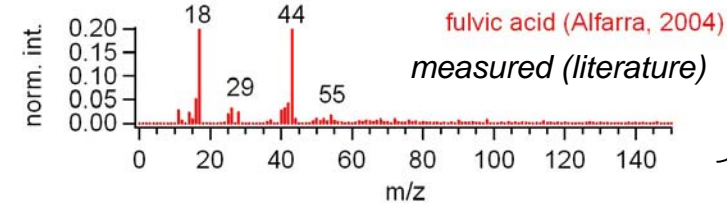
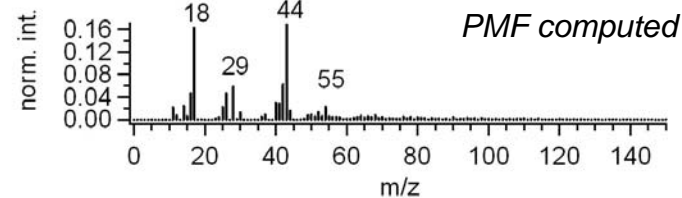
# Zürich, Summer 2005



## Source profiles (C matrix): Verification by independently measured AMS profiles

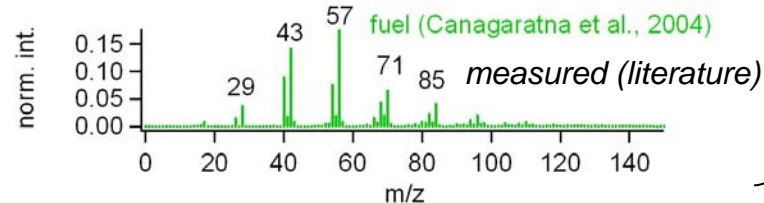
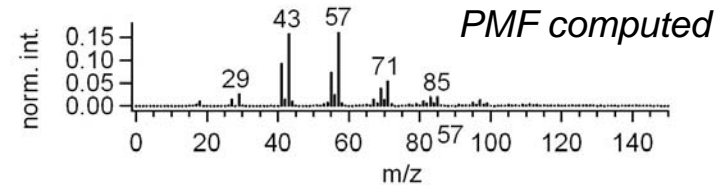


ex. 1



$R^2 = 0.96$

ex. 2



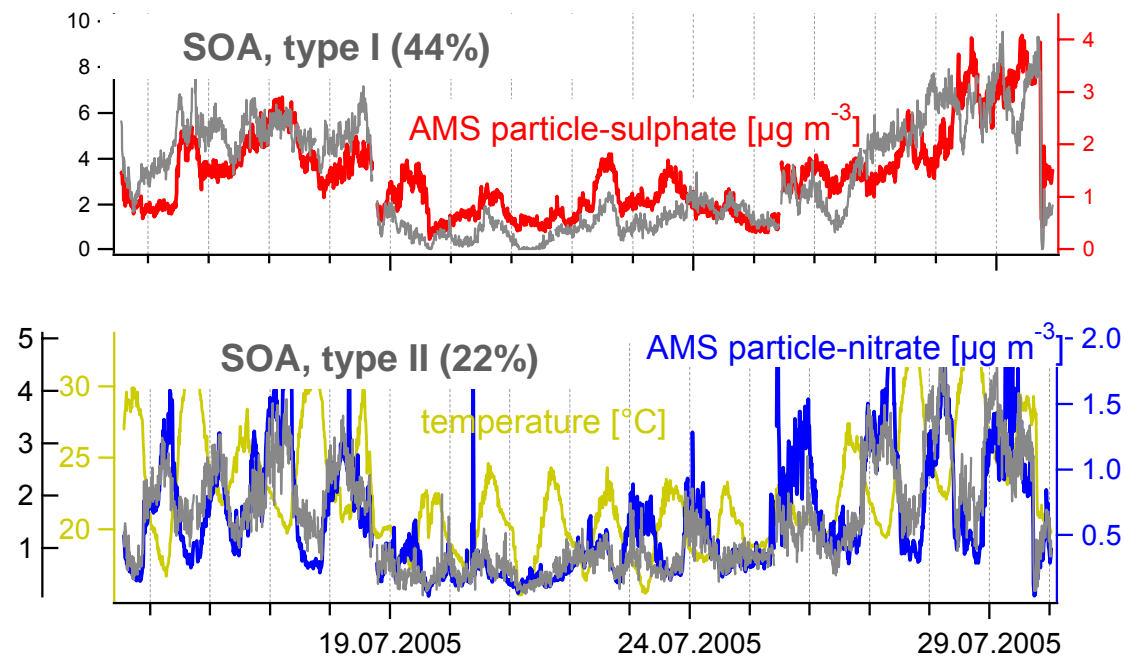
$R^2 = 0.99$



# Zürich, Summer 2005

Source contributions: **66%** of submicron AMS-organics are **SOA**

SOA = secondary organic aerosols (i.e. formed by gas-to-particle reactions in the atmosphere)

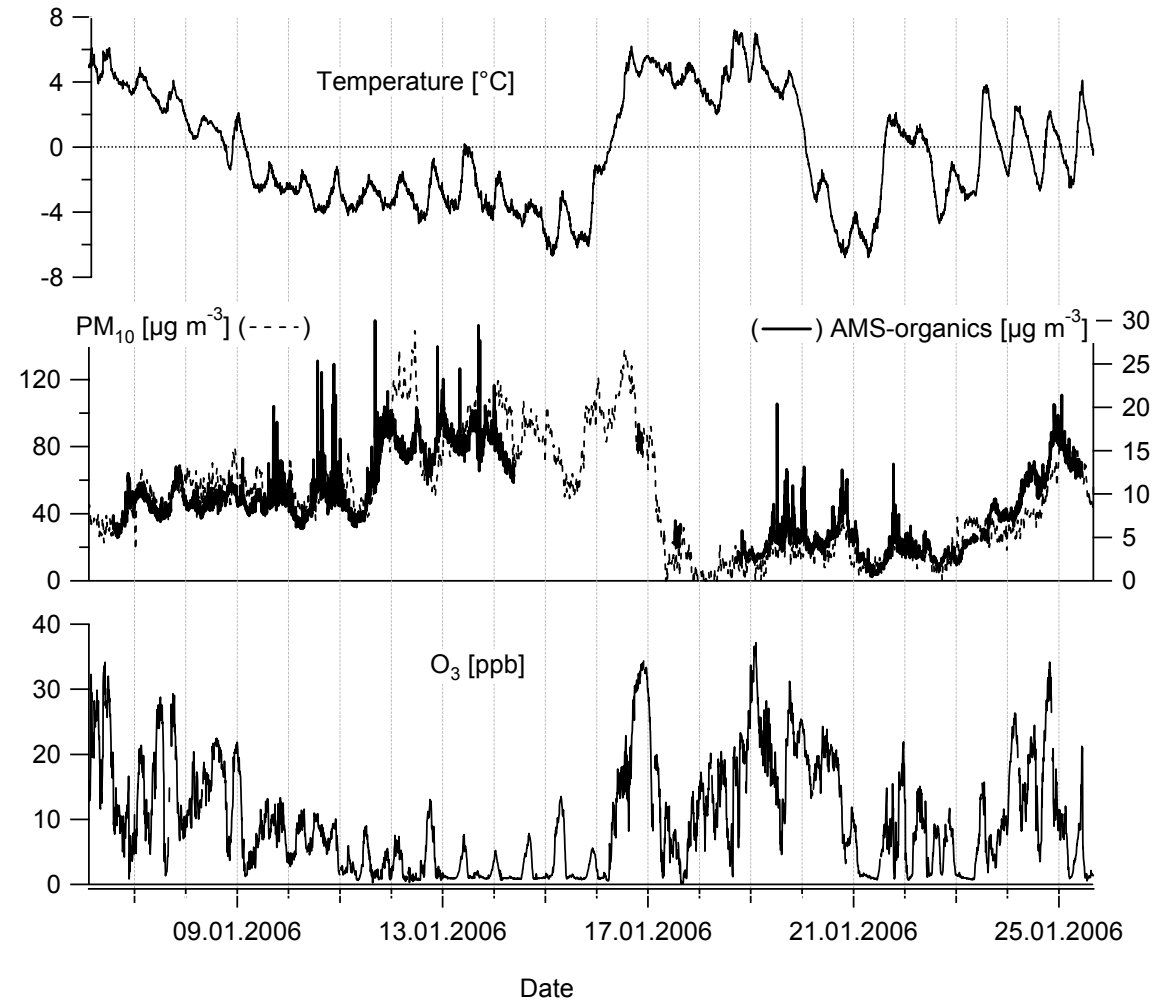


**POA** (=primary organic aerosols) account for **34%** and can directly be attributed to sources (10% wood burning, 7% fuel combustion, 13% charbroiling, <4% food cooking)



# Zürich, Winter 2006

- 6 to 23 January 2006
- **Temperature inversions**  
(„winter smog“)
- Hybrid model: CMB x PMF  
(Lanz et al., ES&T, in review)

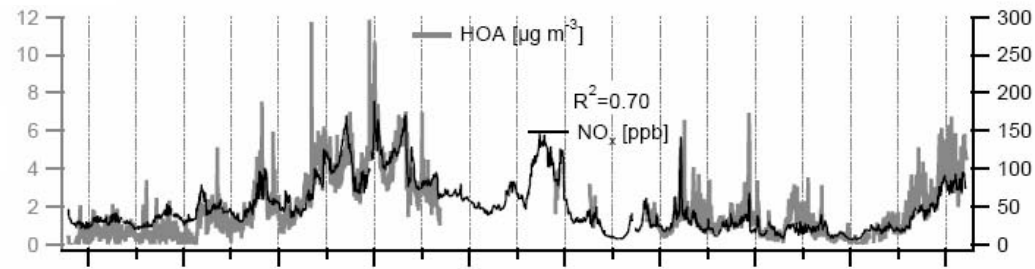


# Zürich, Winter 2006



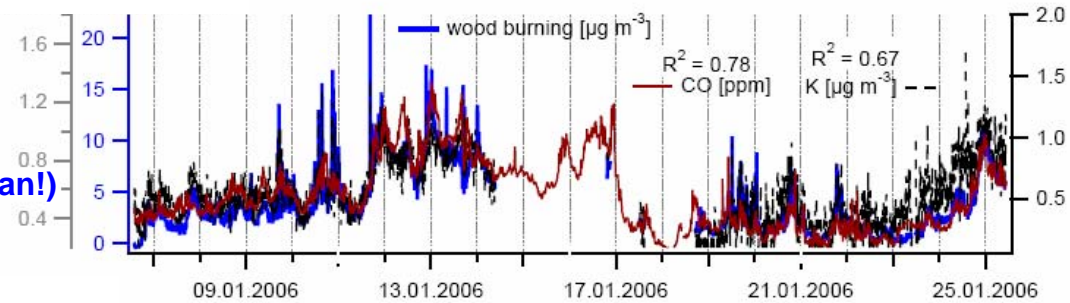
Source contributions:

a. Fuel combustion (7%)

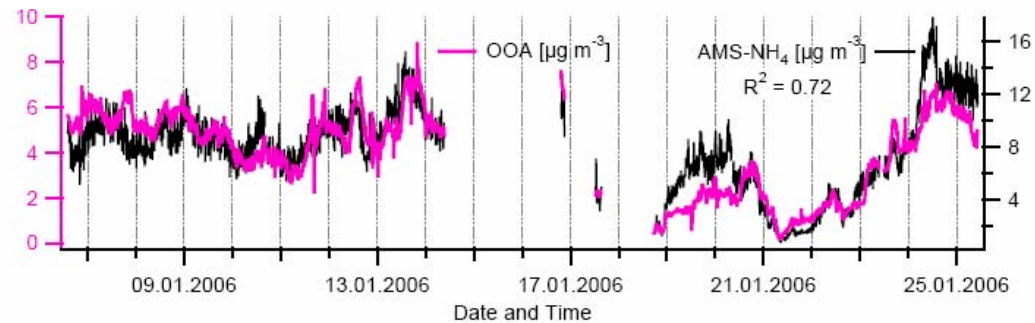


b. Wood burning (38%)

(emission rel. levoglucosan!)

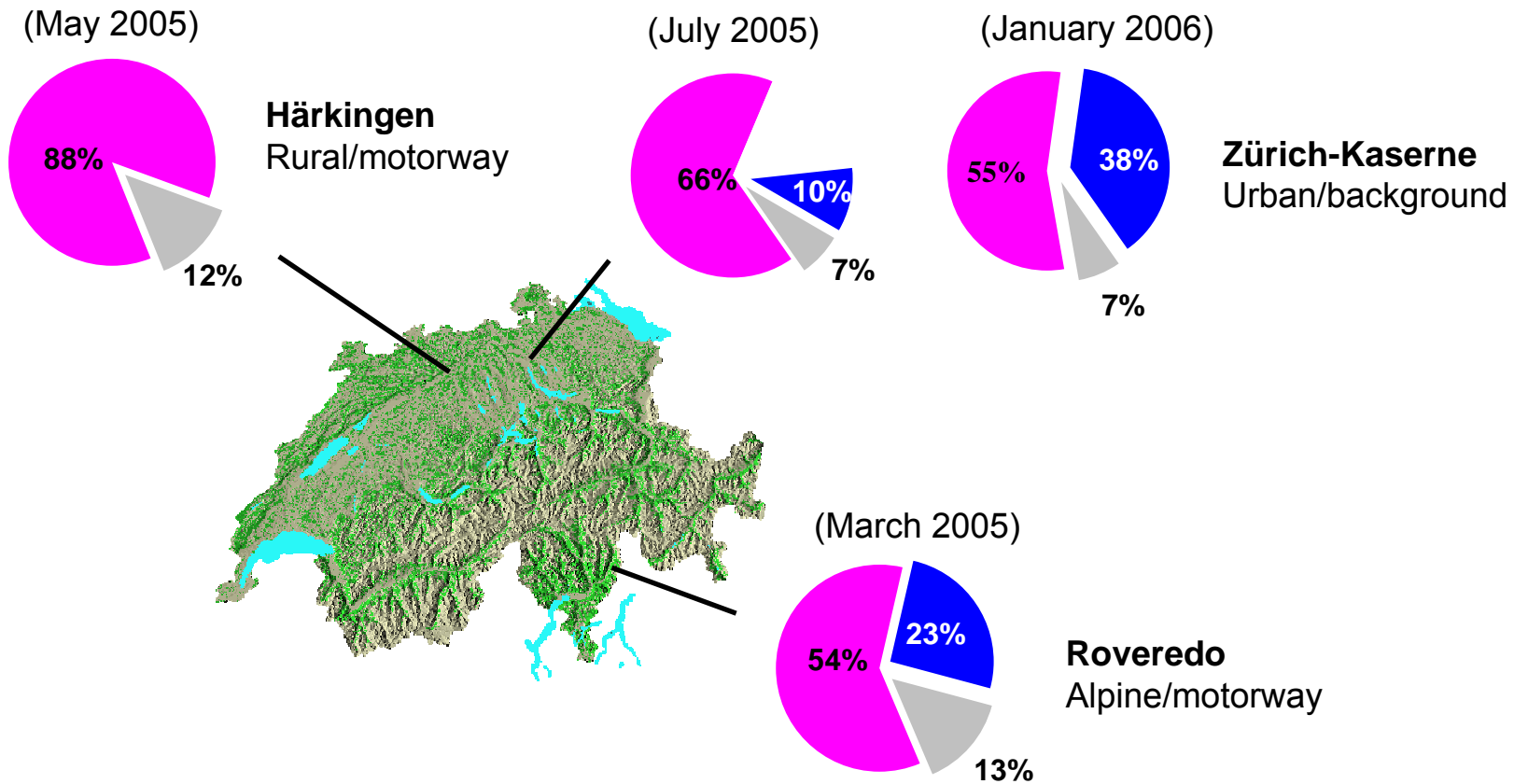


c. SOA (55%)



Primary organic aerosols (POA): 45%

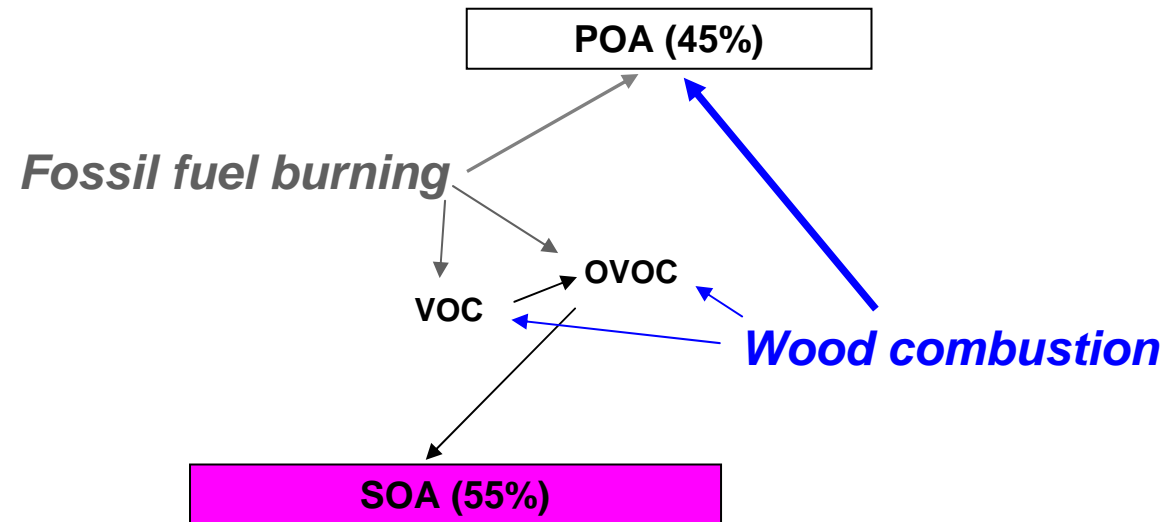
Diesel combustion  
Wood burning aerosol  
Secondary organic aerosol





# Source apportionment of SOA

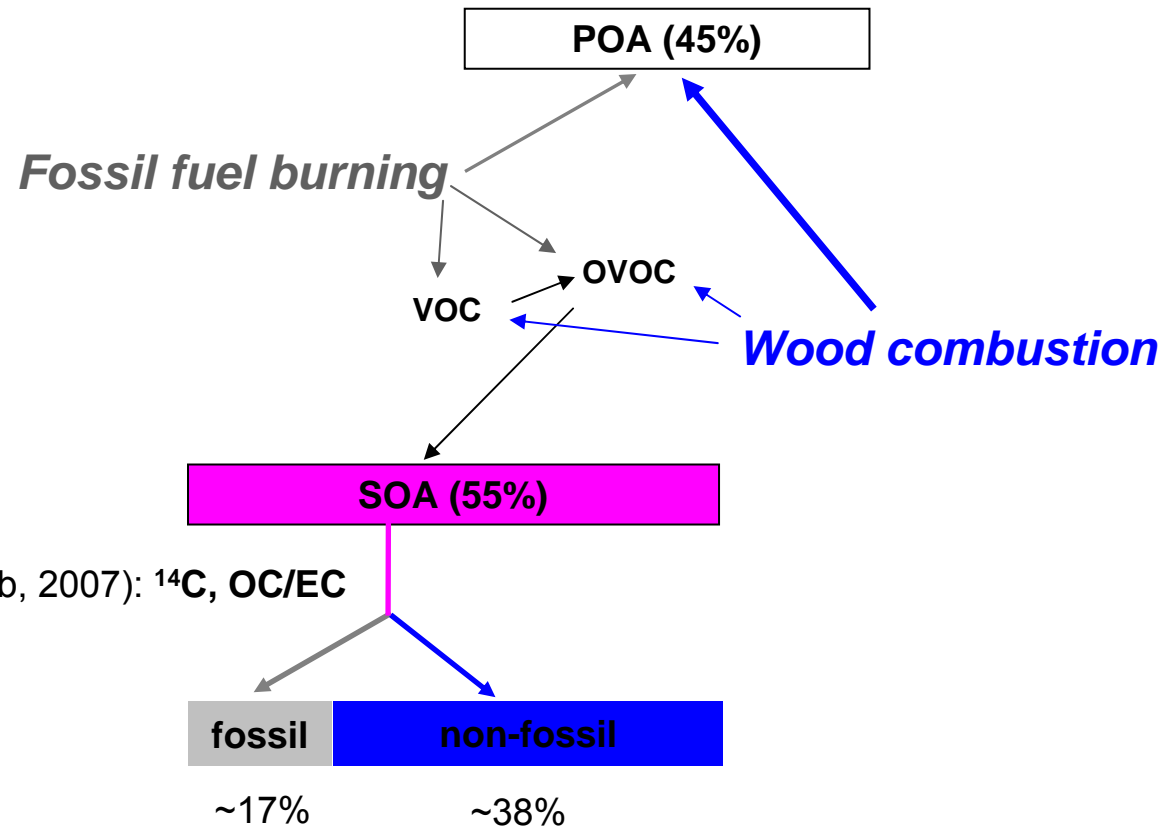
- Apportionment of SOA to emission sources: winter 2006



# Source apportionment of SOA



- Apportionment of SOA to emission sources: winter 2006



Szidat et al. (2006a, 2006b, 2007):  $^{14}\text{C}$ , OC/EC



# Conclusions

- Ambient (AMS) organic aerosols can be attributed to sources by linear mixing models (PMF).

- **SOA** accounted for **50-90%** of the organic aerosol

SOA is dominant both in **summer** (photochemistry) and in **winter** (accumulation, low temperatures)

- **POA** accounted for **10-50%** of organic aerosol

Residential areas/winter: POA dominated by **wood burning**

- $^{14}\text{C}$  measurements give further insights into **SOA source attribution** (fossil/non-fossil) (Winter 2006):

**Fuel combustion** contributes more to SOA (17%) than POA (7%)

**Wood burning** contributes equally to SOA (38%) and POA (38%)

# Thanks to

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