

# **Contribution of natural and anthropogenic factors into near-surface ozone in the Northern Eurasia**

*A.M.Obukhov Institute of Atmospheric Physics RAS*

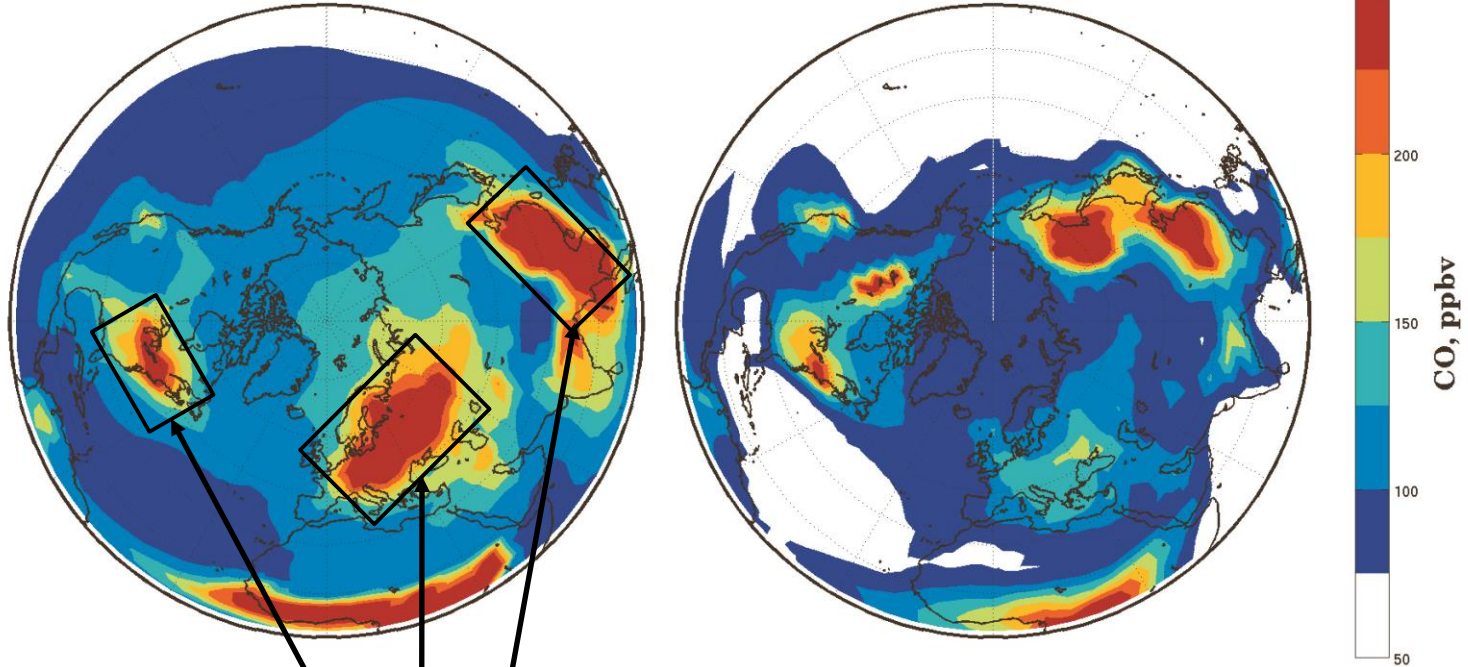
Yu.A. Shtabkin, K.B. Moiseenko

**ENVIROMIS 2016**

# GEOS-Chem near-surface CO field

CO January 2008

CO July 2008



Regions of climatically significant anthropogenic emissions

**ZOTTO measurements: CO<sub>2</sub>, CH<sub>4</sub>, CO, Ozone, NO<sub>x</sub> and aerosols at different heights, meteorology at different heights and on the ground (Temperature, Wind, Humidity), biweekly flask sampling at 301 m height and various irregular ecosystem measurements**



**Background character of the station provide an excellent opportunity to study regional as well as long-range impact of various climatically important sources of pollutants including regional industry and wildfires.**

**[zottoproject.org](http://zottoproject.org)**



# Methods

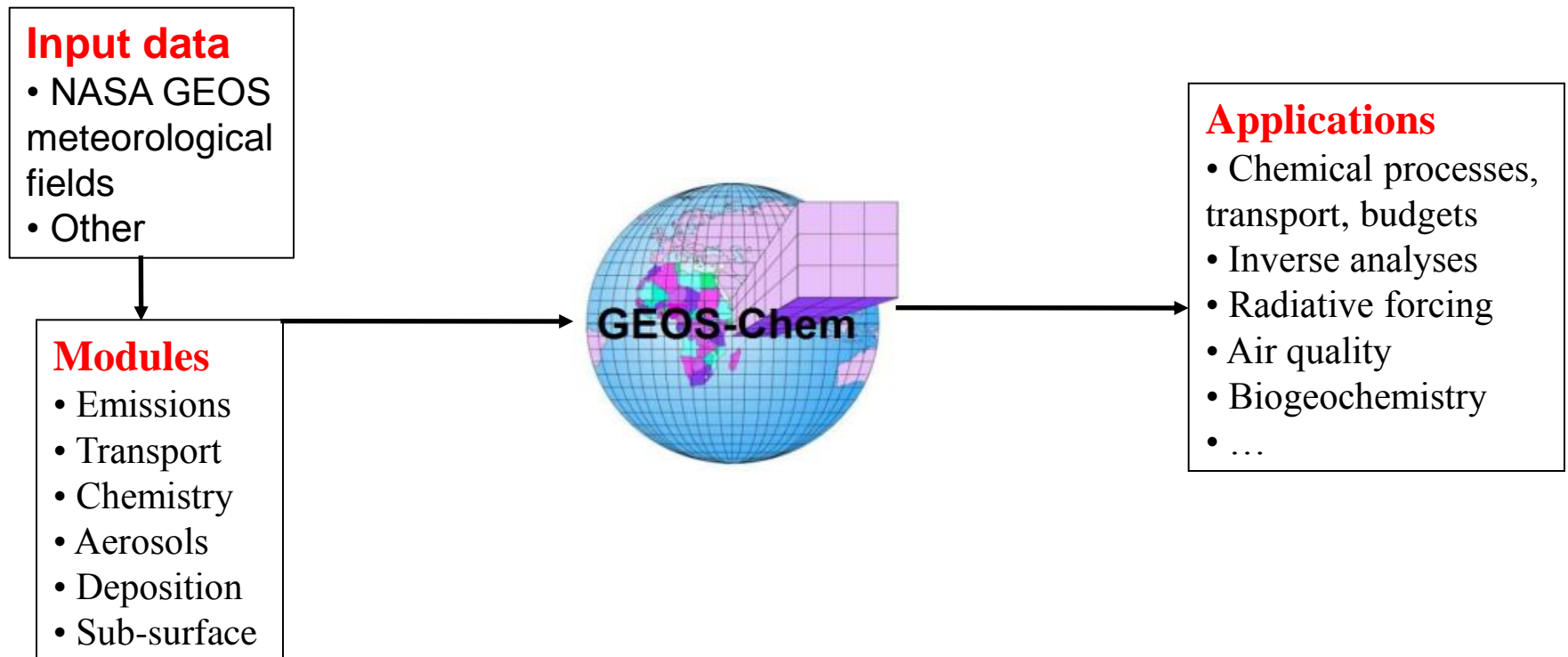
## 1) Emission inventories:

**Anthropogenic** (EDGAR, <http://edgar.jrc.ec.europa.eu>)

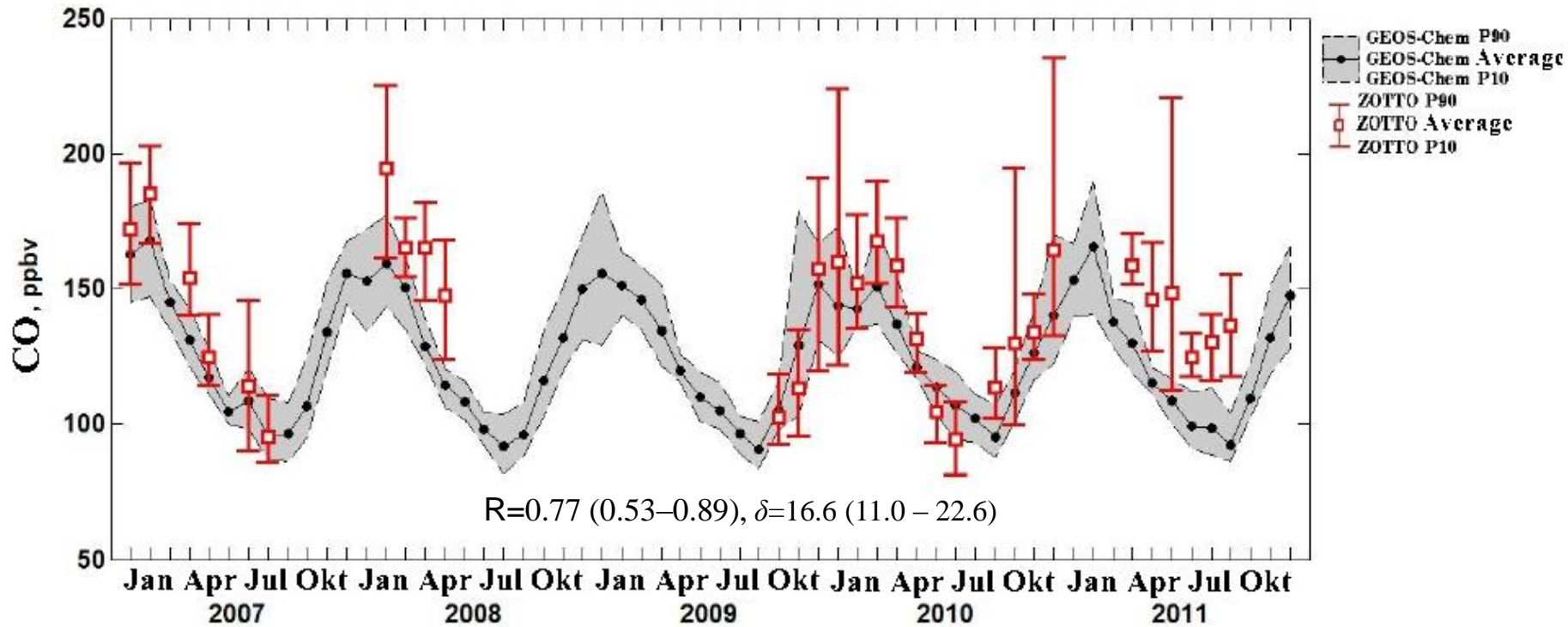
**Biogenic** (VOC oxidation, MEGAN, <http://bai.acd.ucar.edu/MEGAN/>)

**Wildfires** (GFED, <http://www.globalfiredata.org>)

## 2) Global chemical-transport model GEOS-Chem (<http://acmg.seas.harvard.edu/geos>).

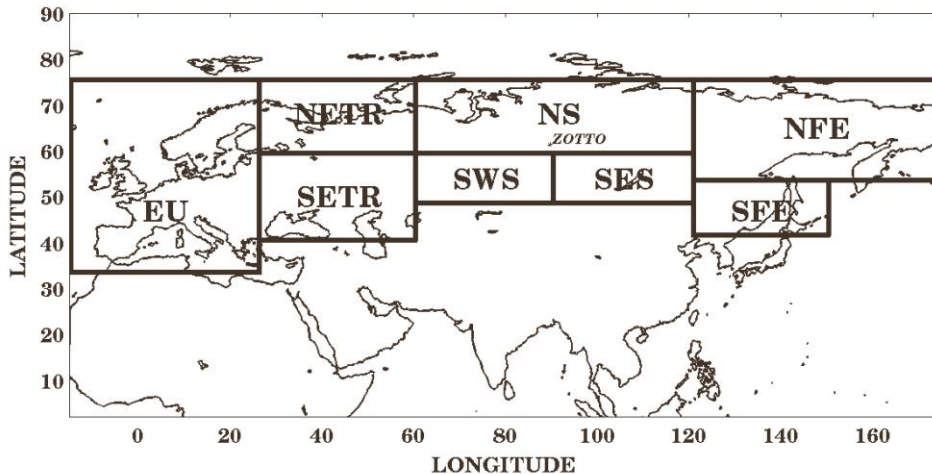


# ZOTTO vs GEOS-Chem



CO concentration at a height of 300 m above the ground observed at ZOTTO in 2007-2011. P10,90 - percentile,  $\square$  - average. The solid and dashed lines - GEOS-Chem model calculation (monthly averaged concentrations at the third model level,  $\sim 320$  m above the ground).

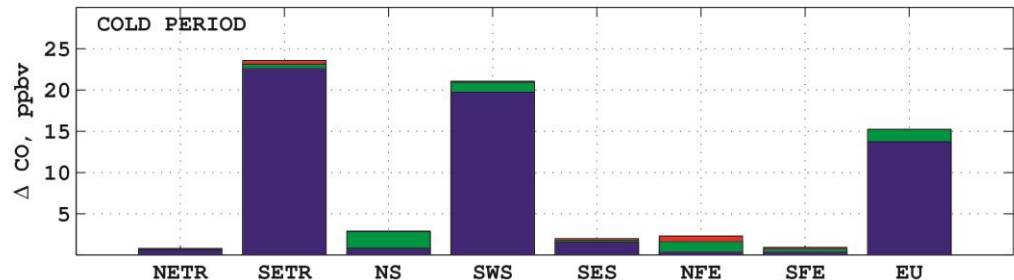
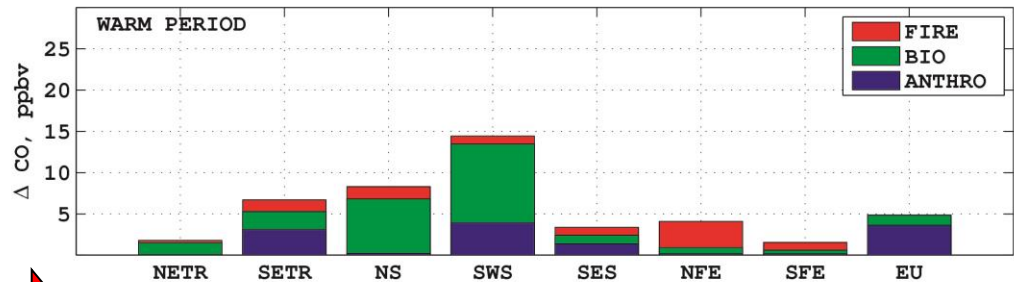
# Model Experiment



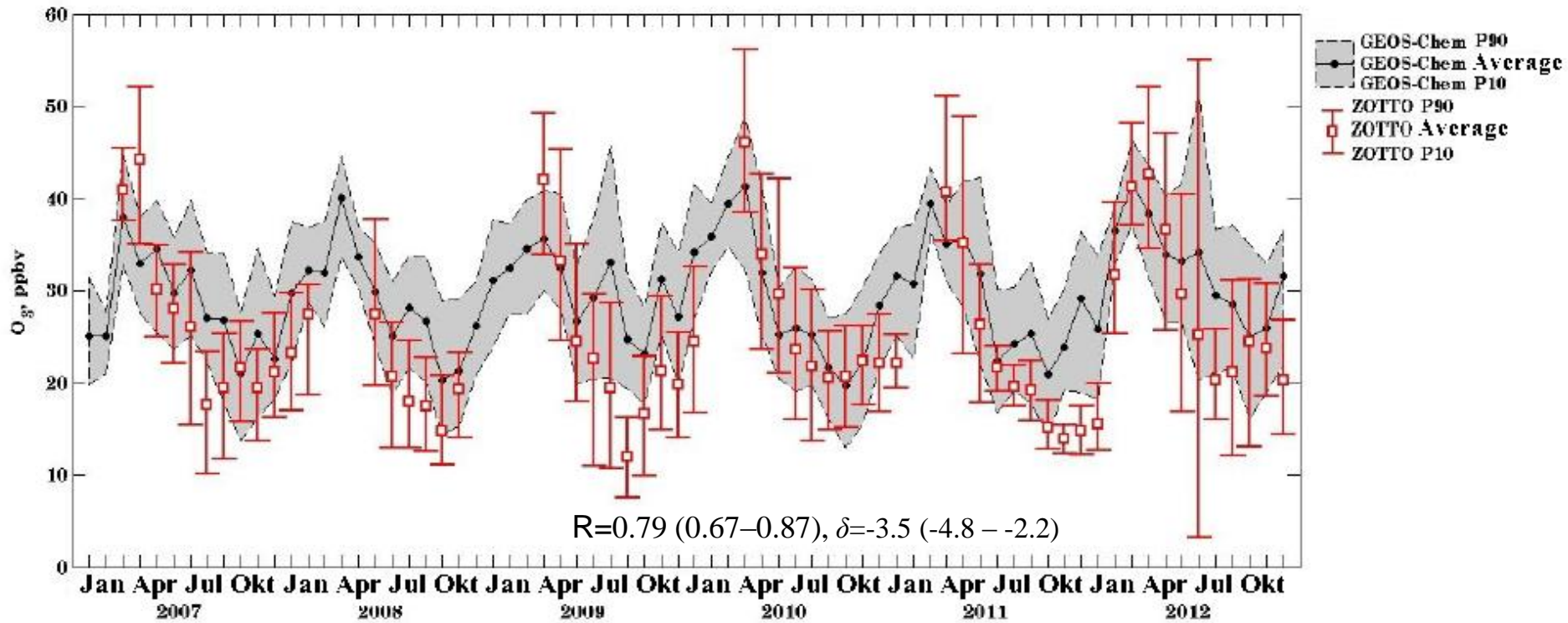
Geographical areas used for CO emission impact evaluation:  
 NETR – north of ETP, SETR – south and midland of ETP, NS – north of Siberia, SWS – south of western Siberia, SES – south of eastern Siberia, NFE – north of Far East, SFE – south of Far East, EU — western Europe.

$$AO_R = \chi(CO)_0 - \chi(CO)_{REG}$$

Summary diagrams of atmospheric response on CO emissions in different regions in warm and cold periods

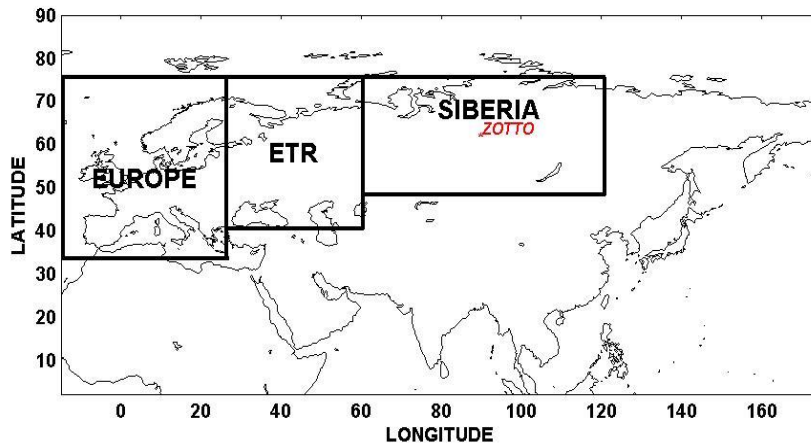


# GEOS-Chem vs ZOTTO



Ozone concentration at a height of 6 m above the ground observed at ZOTTO in 2007-2012. P10,90 - percentile,  $\square$  - average. The solid and dashed lines - GEOS-Chem model calculation (monthly averaged concentrations at the first model level,  $\sim 58$  m above the ground).

# ZOTTO near-surface ozone sensitivity to NO<sub>x</sub> and VOC emissions (I)



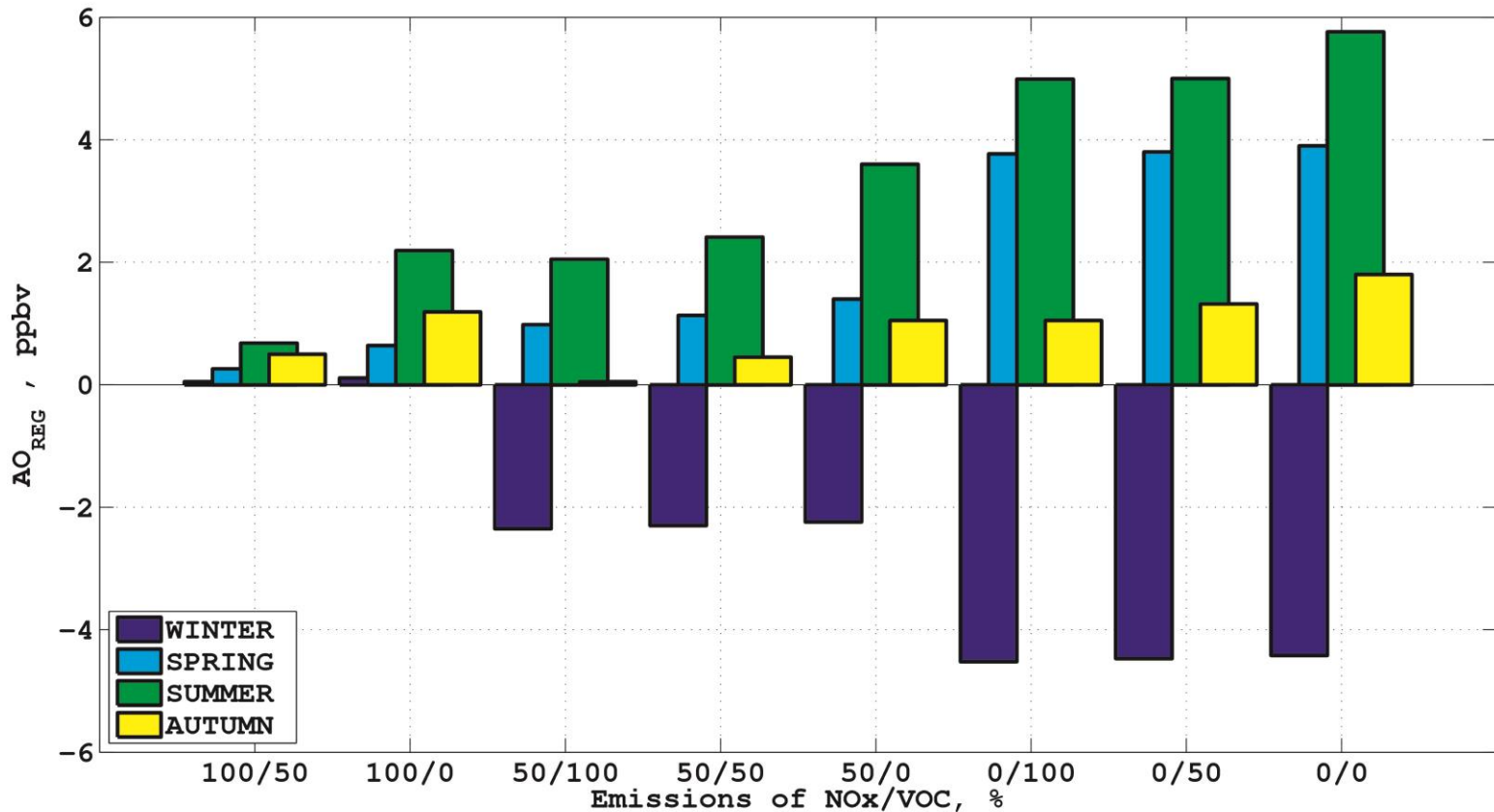
← Geographical areas selected  
for ozone reduction  
calculations

Atmospheric response:

$$AO_R = \chi(O_3)_0 - \chi(O_3)_{REG}$$



# ZOTTO near-surface ozone sensitivity to NO<sub>x</sub> and VOC emissions (II)

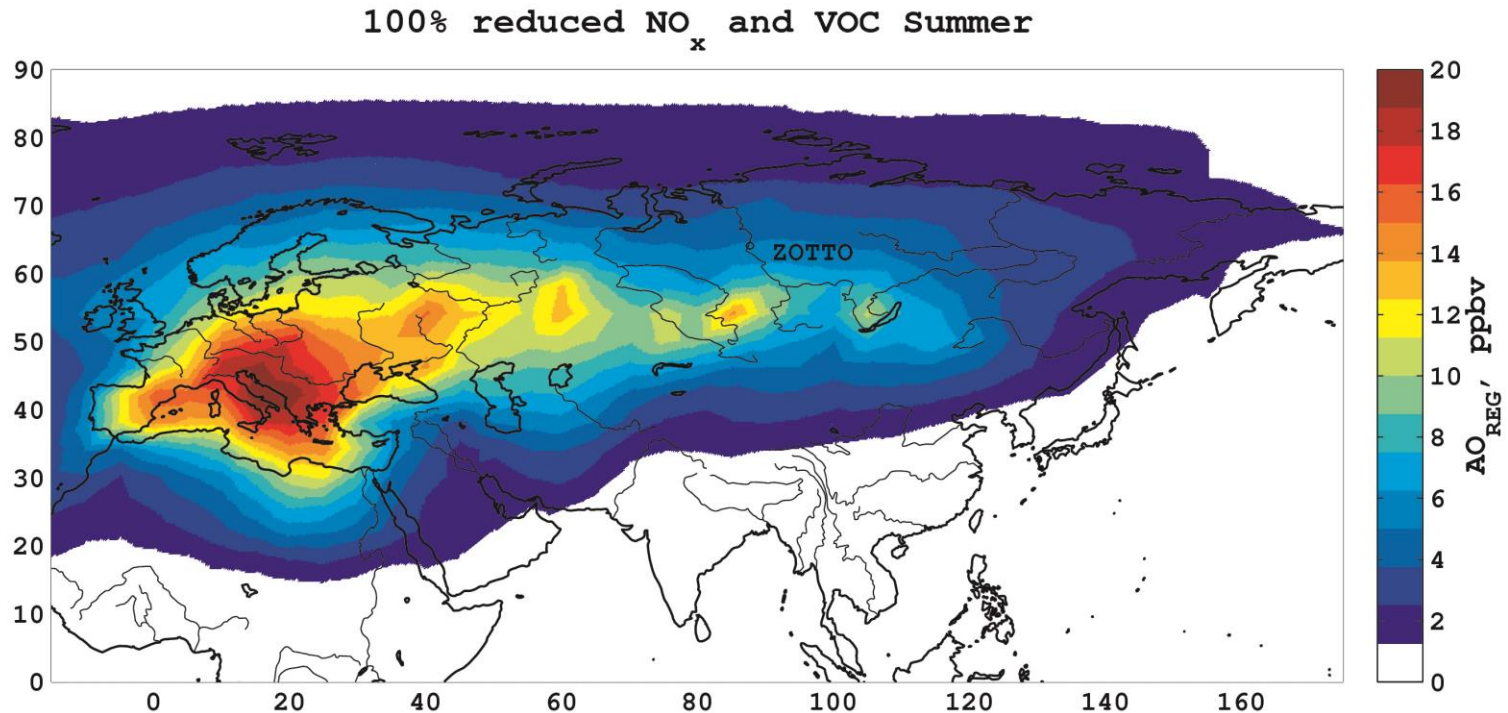


Summer: NO<sub>x</sub> ↓ O<sub>3</sub> ↓      Winter: NO<sub>x</sub> ↓ O<sub>3</sub> ↑

Summary diagram of ozone reduction near ZOTTO station at different biogenic VOCs and anthropogenic NO<sub>x</sub> emission reduced values in Siberia, European Russia and Europe. An averaged values for all 2007 seasons are given.

# ZOTTO near-surface ozone sensitivity to $\text{NO}_x$ and VOC emissions

$$\text{AO}_{\text{REG}} = \chi(\text{O}_3)_0 - \chi(\text{O}_3)_{\text{REG}}$$



Atmospheric response on anthropogenic and biogenic emissions in near-surface ozone field for summer 2007.  $\text{NO}_x$  and VOC emissions are reduced by 100 %