



## **SO<sub>2</sub> retrieved from OMI and OMPS using optimal estimation technique and validation over China**

Congzi Xia (1), Cheng Liu (1), Zhaonan Cai (2), Qihou Hu (3), Fei Zhao (1), Wenjing Su (1), Chengxin Zhang (1), Chengzhi Xing (1), and Wenqiang Zhang (1)

(1) University of Science and Technology of China, University of Science and Technology of China, School of Earth and Space Science, China (czxia17@mail.ustc.edu.cn), (2) Key Laboratory of Middle Atmosphere and Global Environment Observation, Institute of Atmospheric Physics, Chinese Academy of Sciences, Beijing 100029, China, (3) Key Lab of Environmental Optics & Technology, Anhui Institute of Optics and Fine Mechanics, Chinese Academy of Sciences, Hefei, 230031, China

We retrieve sulfur dioxide (SO<sub>2</sub>) vertical columns from the Ozone Monitoring instrument (OMI) and Ozone Mapping and Profiler Suomi National Polar-orbiting Partnership spacecraft (OMPS) using optimal estimation method (OEM) over China from 2013 to 2017. Comparison between OEM retrievals and the principal component analysis (PCA) product shows that a general good agreement between using the different algorithms is obtained with a correlation coefficient of 0.7249 and a slope of 0.8789 over eastern China. Validations with ground-based Multi Axis Differential Optical Absorption Spectroscopy (MAX-DOAS) measurements show that a monthly averaged ground-based SO<sub>2</sub> results and coincident OMI SO<sub>2</sub> results using OEM agree very well. The seasonal cycle of SO<sub>2</sub> is consistent in both data sets with a maximum in winter, on average,  $6 \times 10^{16}$  molecules\*cm<sup>-2</sup> in Xianghe (a key pollution area), and a minimum in summer, which has a mean value of  $2 \times 10^{16}$  molecules\*cm<sup>-2</sup> there. Winter is the domestic heating season. Both show that SO<sub>2</sub> originates mainly from human sources rather than natural ones. The spatio-temporal distribution over China shows that the pollution is mainly concentrated in Beijing-Tianjin-Hebei area and Sichuan Basin. And the yearly averaged SO<sub>2</sub> results show that the SO<sub>2</sub> vertical columns are decreasing from 2013,  $4.57 \times 10^{16}$  molecules\*cm<sup>-2</sup>, to 2017,  $0.89 \times 10^{16}$  molecules\*cm<sup>-2</sup> in those key pollution areas. SO<sub>2</sub> and NO<sub>2</sub> are major aerosol precursors, and SO<sub>2</sub> and NO<sub>2</sub> respectively are sources of pollution mainly from coal-fired power plants and motor vehicle emissions. We also investigate the relationship between SO<sub>2</sub> emission and aerosol production in Beijing. A stronger correlation between the SO<sub>2</sub> concentrations and aerosol optical depths (AODs) measured by the MODIS satellite instrument than NO<sub>2</sub> concentrations with AODs obtained in winter suggests that anthropogenic SO<sub>2</sub> is the major contributor to the aerosol content during the period of the year.