

# **Photoinduced reactions of Anthraquinone-2-sulfonate towards aerosol constituents in tropospheric aqueous solution**

T. Schaefer, Leipzig/D, H. Herrmann, Leipzig/D

Thomas Schaefer, Leibniz Institute for Tropospheric Research (TROPOS),  
Atmospheric Chemistry Department (ACD), Permoserstraße 15, 04318 Leipzig

The formation of secondary organic aerosol (SOA) is centered on the traditional models on the absorption of semi-volatile organic into preexisting aerosols. However, the occurring aerosol aging is associated with chemical transformation by photooxidation chemistry within the particle. [1-4] Unfortunately, the radical sources driving this chemistry remain highly uncertain due to an incomplete understanding of interacting organics in the aerosol particles.

This study presents the results from Anthraquinone-2-sulfonate (AQS) acting as photosensitizer, including time-resolved absorbance spectra ( $\lambda = 300 - 700$  nm) of the excited states, the reaction rate constant with molecular oxygen ( $k_{298\text{ K}} = (5.1 \pm 1.1) \times 10^8 \text{ L mol}^{-1} \text{ s}^{-1}$ ) as well as other aerosol constituents. These results were obtained by using a laser flash photolysis-laser long path absorption (LFP-LLPA) setup. The product analysis of the photo-induced oxidation reaction of AQS, were done by several analytical techniques like UPLC-HRMS and GC-MS.

## Literature:

[1] M. E. Monge, PNAS 2012, 109, 6840-6844. [2] K. Z. Aregahegn, Faraday Discuss. 2013, 165, 123-134. [3] S. Rossignol, Environ. Sci. Technol. 2014, 48, 3218-3227. [4] C. George, Chem. Rev. 2015, 115, 4218-4258.