# Atmospheric Chemistry and Physics An Interactive Open Access Journal of the European Geosciences Union

| EGU.eu | | EGU Journals | Contact

## Online Library ACP

- Recent Final Revised **Papers**
- Volumes and Issues
- Special Issues
- Library Search
- Title and Author Search

Online Library ACPD

Alerts & RSS Feeds

General Information

**Submission** 

Production

Subscription

### Comment on a Paper



lindexed



■ Volumes and Issues
■ Contents of Issue 4

Atmos. Chem. Phys., 10, 1545-1554, 2010 www.atmos-chem-phys.net/10/1545/2010/ © Author(s) 2010. This work is distributed under the Creative Commons Attribution 3.0 License.

# Heterogeneous ozonation kinetics of 4phenoxyphenol in the presence of photosensitizer

S. Net, L. Nieto-Gligorovski, S. Gligorovski, and H. Wortham Universités d'Aix-Marseille I, II, III - CNRS UMR 6264: Laboratoire Chimie Provence Equipe Instrumentation et Réactivité Atmosphérique Case courrier 29, 3 place Victor Hugo, 13331 Marseille Cedex 03, France

Abstract. In this work we have quantitatively measured the degradation of 4-phenoxyphenol adsorbed on silica particles following oxidative processing by gas-phase ozone. This was performed under dark conditions and in the presence of 4-carboxybenzophenone under simulated sunlight irradiation of the particles surface.

At the mixing ratio of 60 ppb which corresponds to strongly polluted ozone areas, the first order of decay of 4-phenoxyphenol is  $k_1 = 9.95 \times 10^{-6} \text{ s}^{-1}$ . At a very high ozone mixing ratio of 6 ppm the first order rate constants for 4phenoxyphenol degradation were the following:  $k_1 = 2.86 \times 10^{-5} \text{ s}^{-1}$  under dark conditions and  $k_1 = 5.58 \times 10^{-5} \text{ s}^{-1}$  in the presence of photosensitizer (4-carboxybenzophenone) under light illumination of the particles surface. In both cases, the experimental data follow the modified Langmuir-Hinshelwood equation for surface reactions. The Langmuir-Hinshelwood and Langmuir-Rideal mechanisms for bimolecular surface reactions are also discussed along with the experimental results.

Most importantly, the quantities of the oligomers such as 2-(4-Phenoxyphenoxy)-4-phenoxyphenol and 4-[4-(4-Phenoxyphenoxy) phenoxy]phenol formed during the heterogeneous ozonolysis of adsorbed 4-phenoxyphenol were much higher under solar light irradiation of the surface in comparison to the dark conditions.

■ Final Revised Paper (PDF, 373 KB) ■ Supplement (57 KB) **Discussion** Paper (ACPD)

Citation: Net, S., Nieto-Gligorovski, L., Gligorovski, S., and Wortham, H.: Heterogeneous ozonation kinetics of 4-phenoxyphenol in the presence of photosensitizer, Atmos. Chem. Phys., 10, 1545-1554, 2010. ■ Bibtex ■ EndNote ■ Reference Manager



Library Search Author Search

- Bringing Down Geoscientific **Barriers**
- New Tax Regulation for Service Charges
- Sister Journals AMT & GMD
- Public Relations & Background Information

### Recent Papers

01 | ACP, 19 Feb 2010: Tropospheric photooxidation of CF<sub>3</sub>CH<sub>2</sub>CHO and CF<sub>3</sub>(CH<sub>2</sub>) CHO initiated by CI atoms and OH radicals

02 | ACP, 19 Feb 2010: Estimations of climate sensitivity based on top-ofatmosphere radiation imbalance

03 | ACP, 19 Feb 2010: Numerical simulations of contrail-to-cirrus transition -Part 2: Impact of initial ice crystal number, radiation, stratification, secondary nucleation and layer depth