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▶ To cite this version:

Marius Duncianu, Alexandre Tomas, Nadine Locoge, Sébastien Dusanter. Development of a mobile photochemical reactor to investigate gas-phase chemistry and aerosol formation in the field. 25th International Symposium on Gas Kinetics and Related Phenomena, Jul 2018, Lille, France. hal-04250522

HAL Id: hal-04250522 https://hal.science/hal-04250522

Submitted on 20 Oct 2023

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Gas-Phase Chemistry and Aerosol Formation in the Field

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Context

The development of innovative tools is important to tackle atmospheric complexity, i.e. its composition and future changes. Investigating chemical transformations during intensive field campaigns is not straightforward due to the impact of emissions, vertical dilution, and other meteorological parameters on ambient concentrations of trace gases and aerosols. Investigations of these transformations in the laboratory using smog chambers provide a much more controlled environment but are limited to simple chemical systems, which are not representative of complex atmospheres.

Preliminary Results from the CERVOLAND Field Campaign (July 2018)

- > Flushing time: 45 min / 90 min to reach equilibrium b/w the GP species and reactor surfaces
- \blacktriangleright Mixing time < 10 min
- Good stability of primary BVOCs and their oxidation products in the chamber



Objectives

The objective of the current study is to combine both concepts through the construction of a mobile photochemical reactor and its deployment in the field during intensive measurement campaigns. This new tool will allow trapping real air masses, directly on site, to characterize the fate of primary emitted trace gases using all the benefits of an environmental chamber, i.e. decoupling the chemistry from other processes (emission, transport...).

Strategy

A 9 m³ photo-reactor was constructed to trap ambient air at atmospheric pressure into a FEP foil bag maintained by an aluminium frame.

The reactor allows both solar and artificial irradiations, whose irradiance is monitored by a spectroradiometer.

The chemical composition, including volatile organic compounds, oxidants and aerosols, is probed in real time during an oxidation experiment using state-of-the-art instrumentation (PTR-ToFMS and offline sampling on cartridges for VOCs, CRM for OH reactivity, PERCA for peroxy radicals, O_3 and NO_x analysers, hygrometer, CPC for particles). **Filling line**







✓ Field experiments will be reproduced in the laboratory inside the same simulation chamber on the basis of a "simplified" air mass. Comparison between field observations and laboratory results will indicate whether any minor species not included in the mixture or unmeasured ambient species are important players for the oxidation chemistry occurring in the chamber. ✓ Simulations from 0D box modeling will be compared with the field results to test our understanding of the BVOC chemistry.

