



## **Chemical conversion of sulphur dioxide on Eyjafjallajökull's volcanic ash from the 2010 eruption**

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Volcanic eruptions induce important climatic and weather modifications. When volcanic ashes are emitted into the atmosphere they can travel for several weeks according to their size distribution and altitude of the emission. Eyjafjallajökull eruption, between April 14th and May 23th, is considered as a medium-size eruption. The upper level winds advected ashes over the UK and continental Europe.

During volcanic eruptions high amounts of SO<sub>2</sub> were injected into the atmosphere (from 50 to 200 ppbv)[1]. Previous works showed that SO<sub>2</sub> could be convert into sulfate on mineral dust surfaces under dark conditions[2]. However, no conversion has been studied with real volcanic ashes and under day conditions (light exposure). For this study, real Eyjafjallajökull's ashes samples, collected on the 2010.04.18 at Seljavellir, were used. The ashes were deposited on a horizontal cylindrical coated-wall flow tube reactor surrounded by 5 fluorescent lamps (340-420 nm). The kinetic studies revealed that the presence of UV-A irradiation enhanced the conversion of SO<sub>2</sub> on ashes samples.

Moreover chemical analyses as XPS, Ion Chromatography and SEM were performed on volcanic ashes before and after exposition to SO<sub>2</sub>. XPS and ion chromatography analyzes showed that the presence of light increase the SO<sub>2</sub> uptake on ashes surfaces and convert it into ions sulphate. Beside SEM analyses disclosed that the conversion takes place systematically on an iron oxide site . By combining kinetics and chemical analysis we are able to propose a new mechanism for the SO<sub>2</sub> conversion on mineral surfaces under light conditions.

1. Self, S., et al., Volatile fluxes during flood basalt eruptions and potential effects on the global environment: A Deccan perspective. *Earth and Planetary Science Letters*, 2006. 248(1-2): p. 518-532.
2. Zhang et al., Heterogeneous Reactions of Sulfur Dioxide on Typical Mineral Particles, *J. Phys. Chem. B*, 2006