

Name of research institute or organization:

**Empa - Materials Science and Technology**

Title of project:

Hydrogen observations at Jungfraujoch, Switzerland

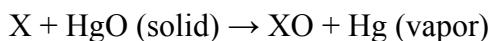
Project leader and team:

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Project description:

Molecular hydrogen ( $H_2$ ) is one of the most abundant atmospheric trace gases. Yet despite its classification as a trace constituent of the atmosphere,  $H_2$  is widely regarded as a potential key component in the future energy chain. For this reason, measurements of molecular  $H_2$  have recently gained broader scientific appeal, as a more complete understanding of atmospheric  $H_2$  is essential as we move towards an increasingly hydrogen-intensive economy. To better ascertain the impacts of enhanced  $H_2$  emissions to the atmosphere, however, the current  $H_2$  budget must be better understood. In order to ameliorate our understanding of  $H_2$  in the atmosphere, along with its various sources and sinks, continuous  $H_2$  measurements have been conducted at Jungfraujoch since 2005. The goal of these observations is to evaluate various trends (diurnal, seasonal, annual), in addition to identifying European sources.

The instrument used to measure  $H_2$  is a modified reduction gas analyzer (RGA3, Trace Analytical), which uses a technique based on chromatographic separation followed by the reduction of mercuric oxide ( $HgO$ ):



where X represents an appropriate reducing gas. The resultant mercury vapor is quantitatively determined through ultraviolet light absorption detection. Mercury vapor is easily measured in minute concentrations, which provides this method significant sensitivity and very low detection limits.

$H_2$  is not a direct greenhouse gas, although its behavior in the atmosphere has the ability to indirectly influence weather and climate patterns. An increase in atmospheric  $H_2$  concentrations could have effects on chemistry in both the troposphere and stratosphere. An increase in tropospheric  $H_2$ , for example, could lead to a reduction in the hydroxyl radical ( $OH$ ), which could consequently push methane ( $CH_4$ ) concentrations upwards ( $H_2$  and  $CH_4$  similarly compete for the oxidizing  $OH$  radical), increasing the radiative forcing of  $CH_4$ , a potent greenhouse gas. An increase in tropospheric  $H_2$  could also lead to an augmentation in stratospheric  $H_2$ . Greater concentrations of  $H_2$  in the stratosphere could lead to a rise in stratospheric water vapor and the formation of polar stratospheric clouds, which in turn could accelerate ozone ( $O_3$ ) depletion and exacerbate the development of the  $O_3$  hole, particularly in the Antarctic due to the delayed breakdown of the polar vortex.

Source identification is an important function of the continuous measurements at Jungfraujoch. Figure 1 illustrates an example of the influence imposed by the southern hemisphere on northern mid-latitudes. On October 1, 2005, advection of clean maritime air masses from low northern latitudes led to depressed  $CH_4$ , carbon

monoxide (CO), and chlorofrom ( $\text{CHCl}_3$ ) concentrations at Jungfraujoch. On the other hand, molecular  $\text{H}_2$  concentrations were simultaneously elevated. Figure 2 below depicts the 10-day backward trajectories of the maritime air masses that produced the results observed in Figure 1. The trajectories are based on 3-D wind analysis from the European Centre for Medium-Range Weather Forecasts (ECMWF).

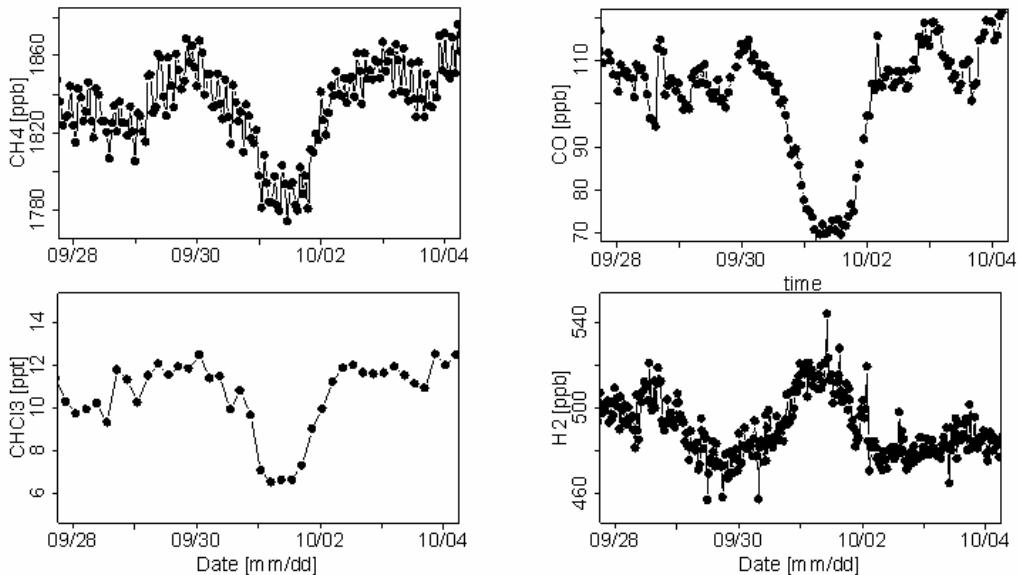


Figure 1. Concentrations of various substances depicting influence of southern hemisphere on northern mid-latitudes.

Anthropogenic activities are major sources of  $\text{CH}_4$ , CO, and  $\text{CHCl}_3$ , and are a significant factor in the global budget of each of these substances. As a result of

human activities, concentrations of these pollutants are higher in the northern hemisphere. As air masses from low northern latitudes (which are influenced by the lower concentrations of these compounds in the southern hemisphere) are carried north, measured concentrations of these pollutants are reduced at the Jungfraujoch station. Alternatively, the soil sink is the major factor in the global  $\text{H}_2$  budget. With the greater soil surface area in the northern hemisphere,  $\text{H}_2$  concentrations are lower north of the equator, despite correspondingly

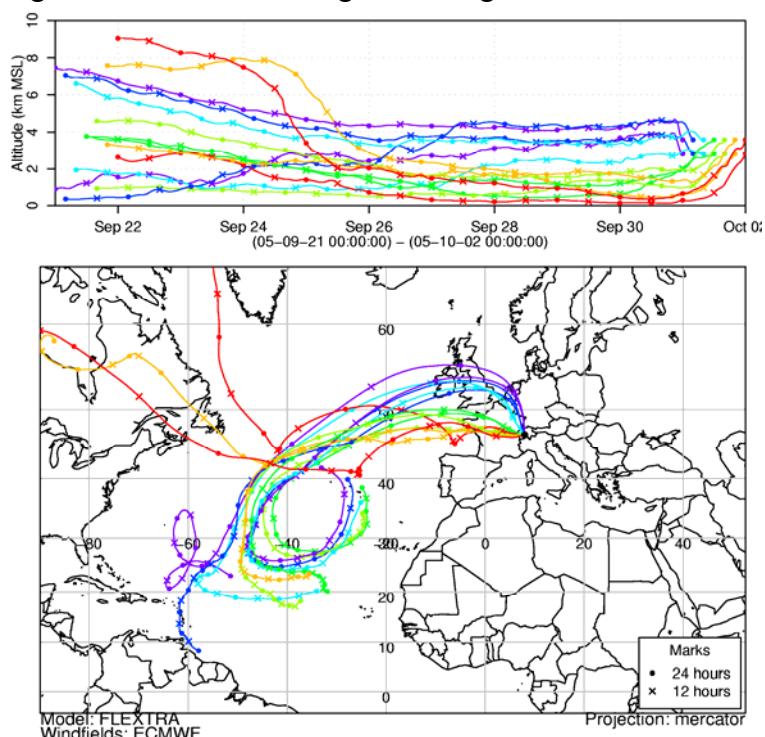


Figure 2. 10-day backward air mass trajectories leading to measurements at Jungfraujoch on October 1, 2005.

higher anthropogenic emissions. As such, as air from low northern latitudes (with high H<sub>2</sub> concentrations from the influence of the southern hemisphere, e.g. from biomass burning) are carried north, measurements at Jungfraujoch reveal elevated H<sub>2</sub> concentrations, as ambient alpine air is displaced by air from the south.

The continuously measured H<sub>2</sub> at Jungfraujoch will be further analysed for trends and for emissions from European sources. This activity is part of the EUROHYDROS project, where data from several European measurement stations are integrated. Furthermore, measurements are also used within the Competence Centre for Energy and Mobility (CCEM-CH), an activity within the ETH domain.

**Key words:**

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Hydrogen, H<sub>2</sub>, Jungfraujoch, Atmosphere

**Collaborating partners/networks:**

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- EUROHYDROS – A European Network for Atmospheric Hydrogen Observations and Studies
  - Competence Centre for Energy and Mobility (CCEM-CH)

**Scientific publications and public outreach 2007:**

**Refereed journal articles**

Steinbacher, M., A. Fischer, M.K. Vollmer, B. Buchmann, S. Reimann, C. Hueglin, 2007. Perennial observations of molecular hydrogen (H<sub>2</sub>) at a suburban site in Switzerland. *Atmospheric Environment*, 41, 2111-2124.

Vollmer, M. K., N. Juergens, M. Steinbacher, S. Reimann, M. Weilenmann, B. Buchmann, 2007. Road vehicle emissions of molecular hydrogen (H<sub>2</sub>) from a tunnel study. *Atmospheric Environment*, 41, 8355-8369, doi:10.1016/j.atmosenv.2007.06.037.

**Conference contributions**

Vollmer, M. K., Steinbacher, M., Reimann, S., Buchmann, B., Weilenmann, M., Fischer, A., Hill, M., Juergens, N. Atmospheric molecular hydrogen (H<sub>2</sub>): sources, sinks, and Empa's links, TECAT seminar, Empa Dubendorf, May 5, 2007.

Steinbacher, M., Vollmer, M.K., Henne, S., Brunner, D., Buchmann, B., Reimann, S. Non-CO<sub>2</sub> Greenhouse Gas Mixing Ratios at Jungfraujoch, Switzerland - Influence of Air Mass Origin, 14th WMO/IAEA Meeting of Experts on Carbon Dioxide, Other Greenhouse Gases, and Related Tracer Measurement Techniques, Helsinki, Finland, September 10-13, 2007.

Steinbacher, M., Continuous H<sub>2</sub> observations and H<sub>2</sub> road tunnel studies in Switzerland, EUROHYDROS 1<sup>st</sup> annual meeting, Norwich, September 18-19, 2007.

Bond, S., Vollmer, M. K., Sources and Sinks of Atmospheric H<sub>2</sub> during the Transition to Hydrogen-based Transportation, CONCAWE, Brussels, November 5-6, 2007.

**Poster presentations**

Bond, S., Reimann, S., Vollmer, M. K., Steinbacher, Hill, M., Buchmann, B., Weilenmann, M., Sources and Sinks of Atmospheric H<sub>2</sub> during the Transition to Hydrogen-based Transportation, Empa PhD Symposium 2007, Empa Akademie, Dübendorf, November 21, 2007.

*International Foundation HFSJG*  
*Activity Report 2007*

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