

Fachbereich Biologie, Chemie, Pharmazie  
Institut für Chemie und Biochemie

Freie Universität Berlin, Institut für Chemie und Biochemie  
Takustrasse 3, D-14195 Berlin

**Prof. Dr. E. Illenberger**  
Physikalische und Theoretische Chemie  
Takustrasse 3, D-14195 Berlin

Telefon: +49 30 838 55350 (52096)  
Fax: +49 30 838 56612 (52717)  
E-Mail: [iln@chemie.fu-berlin.de](mailto:iln@chemie.fu-berlin.de)  
Internet: [www.fu-berlin.de](http://www.fu-berlin.de)

Berlin, 01 March 2006

## Host Report

Reference code: **COST-STSM-P9-01803**

During his visit to our laboratory Richard Balog studied electron stimulated desorption (ESD) of fragment anions and thermal desorption spectroscopy (TDS) in condensed phase  $\text{CF}_3\text{COOH}$  following low energy electron impact (0-20 eV). In ESD, the fragment ions  $\text{H}^-$  and  $\text{F}^-$  are observed via resonant features in the energy range 5 – 12 eV similar to the dissociative electron attachment (DEA) resonances present in gas phase  $\text{CF}_3\text{COOH}$ . Extended electron irradiation of the film leads to the formation of  $\text{CO}_2$  and  $\text{CHF}_3/\text{CF}_4$  as revealed by a combination of TDS and ESD experiments. The experiments concerning reactions induced by sub-excitation electrons ( $\approx 1\text{eV}$ ) were so far not conclusive. Richard Balog will be the co-author of a publication entitled

*Reactions in nanofilms of trifluoroacetic acid ( $\text{CF}_3\text{COOH}$ )  
driven by low energy electrons*

to be submitted to Phys. Chem. Chem. Phys. this month.

Eugen Illenberger  
Professor of Physical Chemistry

Richard Balog  
Department of Physics and Astronomy  
University of Aarhus  
Ny Munkegade, Bld. 520  
DK-8000 Aarhus C

Telefon: +45 8942 3774  
Fax: +45 8612 0740  
E-Mail: balog@phys.au.dk

## Scientific Report

Reference code: COST-STSM-P9-01803

The aim of my visit at the host institute in Berlin was to study the reactivity of  $\text{CF}_3\text{COOD}$  under the electron attack in the energy range 0-20eV. The products of reactions were investigated by means of electron stimulated desorption (ESD) and temperature programmed desorption (TPD).

We have observed that  $\text{CF}_3\text{COOD}$  efficiently transforms into  $\text{CF}_3\text{COOH}$  via D/H exchange inside the gas inlet system. The ratio of deuterated and hydrogenated samples deposited onto platinum substrate was monitored using TPD measurements, where amount of fragment ions  $\text{COOD}^+$  (mass 46) and  $\text{COOH}^+$  (mass 45) have been recorded. During the course of measurements after several fill and pump cycles we have obtained mass ratio 46/45 to be 2/1.

In ESD measurements two fragment ions have been observed, namely  $\text{H}^-$  (measured previously from  $\text{CF}_3\text{COOH}$  sample) and  $\text{F}^-$ . Both yields show a resonant structure in the energy range 5-14eV, which does not reveal any dramatic change in their shape and/or intensity with number of repetitive scans (irradiation). Obtained results were directly compared with results from the gas phase and cluster measurements performed in the same laboratory.


We have applied the TPD technique to analyze products generated by bombardment of  $\text{CF}_3\text{COOD}$  samples with electrons of fixed energy. One observation is that irradiation leads to the formation of  $\text{CO}_2$  and most probably  $\text{CF}_3\text{D}$ . Extended irradiations at different incident energies show that  $\text{CO}_2$  is efficiently formed above some threshold energy and continuously increases with increasing electron energy.

In the cluster measurements, electron attachment to  $\text{CF}_3\text{COOH}$  leads to the formation of  $\text{H}_2\text{O}\cdot\text{CF}_3\text{COO}^-$  complex at 0.4eV. The question is whether such reaction takes place also in condensed phase. One can then check if negative species like  $\text{H}_2\text{O}\cdot\text{CF}_3\text{COO}^-$  desorb from the surface after the heating. So far we did not observe negative ions using TPD technique. It is expected that anions formed at the top of the

molecular film diffuse to the substrate during the heating and then charge transfer to the metal substrate takes place prior to desorption. Another possibility is to check if H<sub>2</sub>O or equivalently D<sub>2</sub>O desorbs from the surface by monitoring masses 18 and 20. We have observed that, both masses do desorb even from nonirradiated CF<sub>3</sub>COOD (CF<sub>3</sub>COOH) film. Such signal may originate from water impurity in the sample and/or from the dimers. However, mass analysis of the evaporated film does not indicate any presence of the dimers.

Because of the unexpected shut down of the apparatus (change of the gold substrate by platinum substrate and consecutive backing of the chamber) at the beginning of my visit, there was no time left to finish the measurements proposed and search for water production from the irradiated CF<sub>3</sub>COOD sample was still in progress.

Date: 03.03.2006 .....

Signature:  .....

