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Study of Copper Corrosion in Pure Water

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**SKB:s referensgrupp möte för kopparkorrosion i syrefri miljö.
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The Ångström Laboratory



Materials Chemistry

Other Departments at the Ångström Laboratory:
Technology, Physics, Physical Chemistry,
Photochemistry & Molecular Science, Mathematics,



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Materials Chemistry

Syntheses of new compounds and materials

Thin films

Nano-sized particles

Bulk

Characterization

X-ray diffraction

electron microscopy

Electrochemical methods

ESCA

Raman spectroscopy





Copper corrosion in water

- The process of metal corrosion in pure water accompanies by hydrogen gas evolution: $\text{Me} + \text{H}_2\text{O} \leftrightarrow \text{Me}^+ + \text{H}_2\uparrow$
- In the experimental study (*Hultquist, Corr.Sci.1986*) it was assumed that copper corroded to Cu_2O according to: $2\text{Cu} + \text{H}_2\text{O} \rightarrow \text{Cu}_2\text{O} + \text{H}_2\uparrow$
- The reaction was further modified (*Szkalos et.al. Electrochem Solid State Lett. 2007*) to:
$$\text{Cu} + y\text{H}_2\text{O} \rightarrow \text{H}_x\text{CuO}_y + (2y-x)\text{H}_{\text{ads}}$$



Calculation of Cu corrosion rate from hydrogen evolution

- *Hultquist, Corrosion Science* **26**, 173–176 (1986).
0.98 $\mu\text{m}/\text{year}$ or **9.8 cm**/100 000 years
- *Szakálos et.al, Electrochemical and Solid-State Letters*
10, C63–C67 (2007)
 $10^{-5}\text{cm}/\text{year}$ or **1 cm**/100 000years
- *Hultquist, Catal. Lett.* **132**, 311-316 (2009).
3.7 nm/year or **0.037 cm**/100 000 years



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Experimental

Sample preparation: Cu 99.95% purity

1. Mechanically polished
2. Electrochemically polished

Three experimental set-ups:

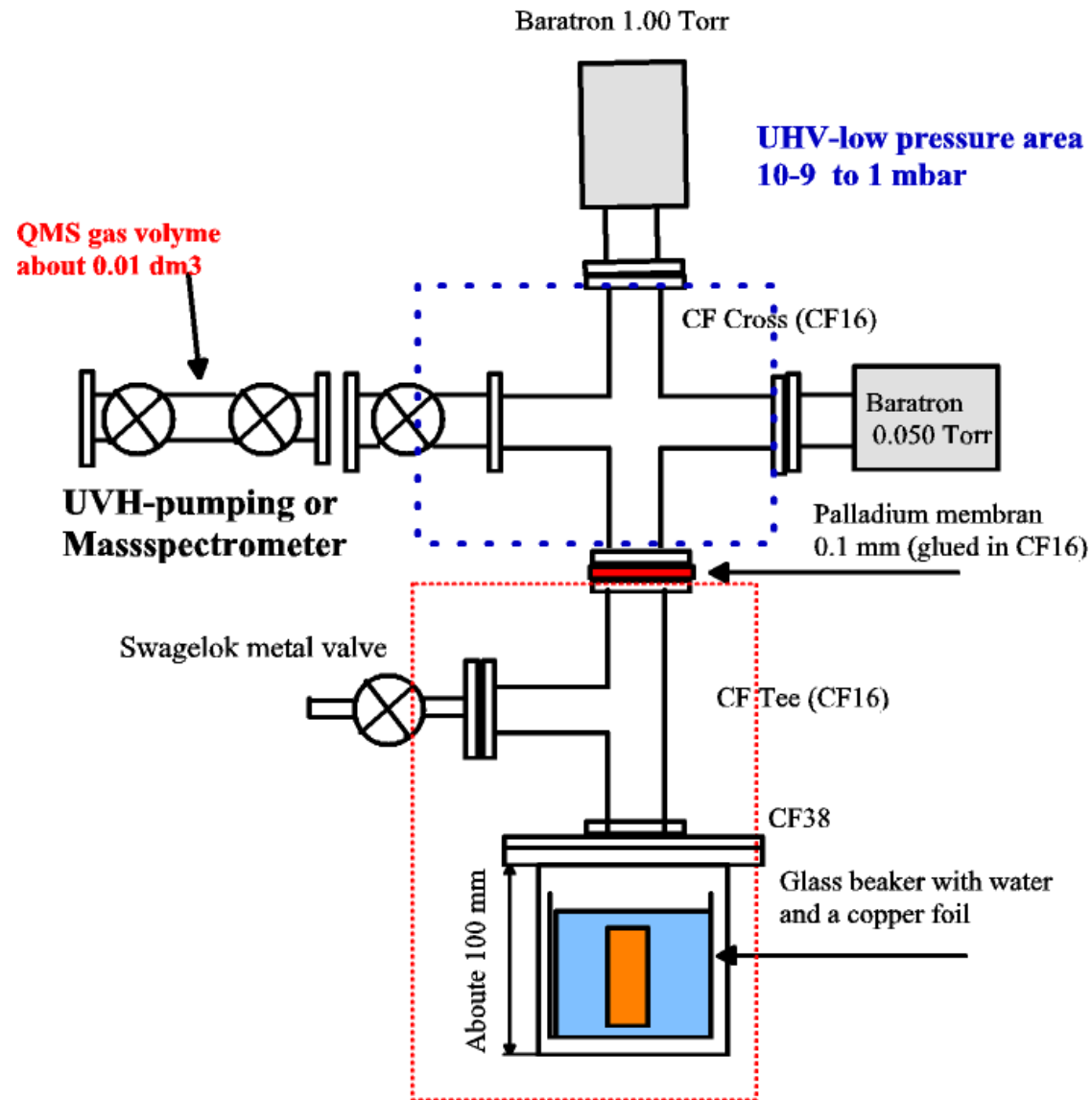
1. Milli-Q water
2. Cu in water
3. Cu in diluted acid





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Vacuum system for measurements of hydrogen production during Copper corrosion.



High pressure area (0.5-1 Atm)
Volyme about 0.2 dm³ heated to 70 C



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Factors, which may influence the detection limit of H₂

- UHV system leaks (external and interior)
- Outgassing rate of UHV components (chambers, glue)
- Palladium membrane
- Cu metal surface can catalyze hydrogen development



Experimental methods to study Cu corrosion products

- The thickness of the surface layer as well as its composition can be studied by **XPS (ESCA)** analysis.
- **ICP-MS** to measure trace amounts of elements in the water. The detection limit for Cu in solution is 2×10^{-11} g/mL.
- **GD-OES** (Glow discharge optical emission spectrometry) to study hydrogen concentration gradients in Cu.
- **FTIR** (Fourier Transform Infrared Spectroscopy) and **SERS** (Surface Enhanced Raman spectroscopy) to detect the O-H, Cu (I)-O, Cu(II)-O etc. vibrations.
- **GI-XRD** (X-ray diffraction and grazing incidence X-ray studies)
- **pH** of the water solution will be controlled after immersing the Cu foil into the water
- Soft X-ray analysis (**XAS** and **XES**).