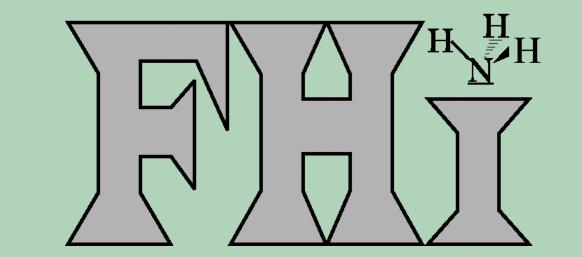


Investigation of the ammonia oxidation over copper with in situ NEXAFS spectroscopy: Influence of the copper oxides on the reaction products

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1. Introduction

Ammonia containing exhaust gas is still a problem in many chemical processes, e.g. in the ammonia slipstream treatment after deNO_x-SCR [1] or in the purification of reformats for fuel-cell applications [2]. Besides the decomposition of ammonia to nitrogen and hydrogen the selective catalytic oxidation to nitrogen and water

$$4NH_3 + 3O_2 \rightarrow 2N_2 + 6H_2O$$
 (1)

is consistent with a wide range of applications [3] and the subject of this investigation. In addition to reaction (1) the total oxidation

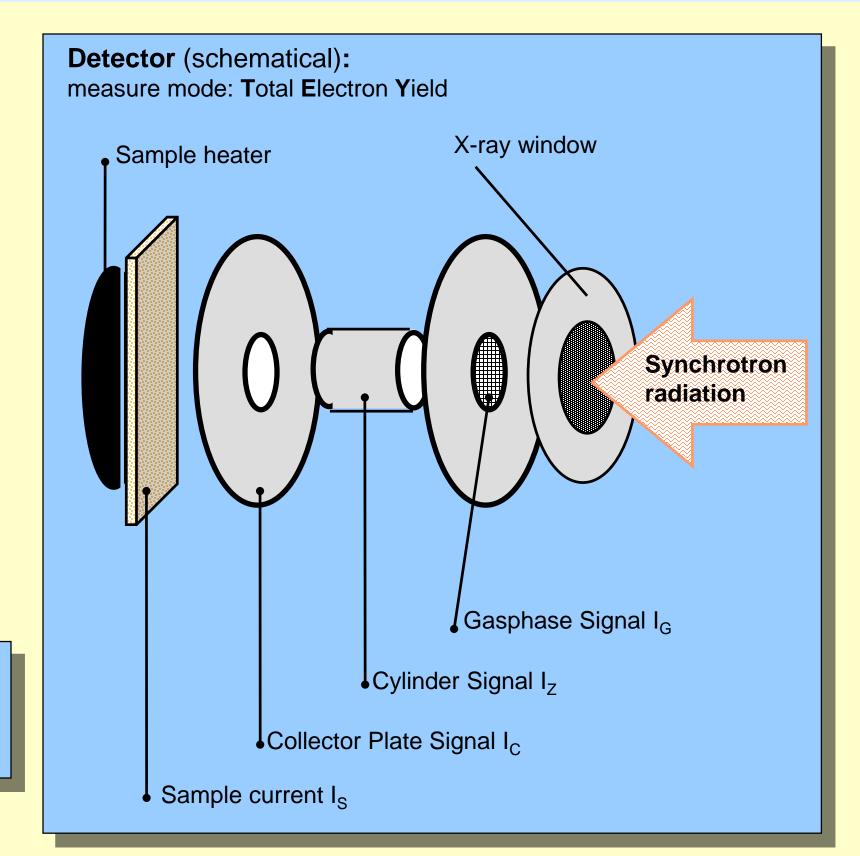
$$4NH_3 + 5O_2 \rightarrow 4NO + 6H_2O$$
 (2)

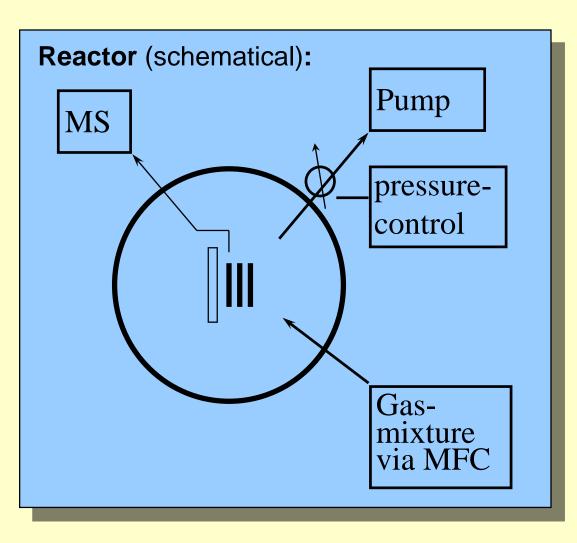
is possible in oxygen containing feed gas. Because of the production of hazardous nitric oxide this reaction path has to be totally suppressed. This is the most important selection criteria for a suitable catalyst.

2. Method

- Near Edge X-Ray Absorption Fine Structure (NEXAFS) Spectroscopy in soft energy range (hv = 350 - 550eV)
- provides **electronic structure** of involved species and by comparison with reference spectra chemical state of the catalyst surface
- in situ: reaction is observed via mass spectrometry
- high pressure: up to several mbars

> aim: detect the catalytically active surface structure of the catalyst





Beamlines at BESSY II:

 U49/1 and UE56/2 (Undulator) PM1 (bending magnet)

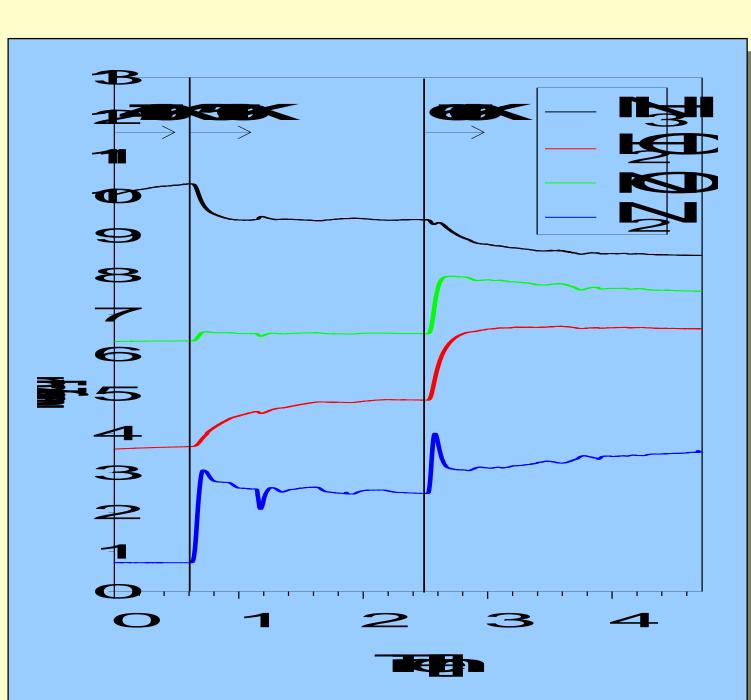
3a. Results @ p=0.4mbar and CuO

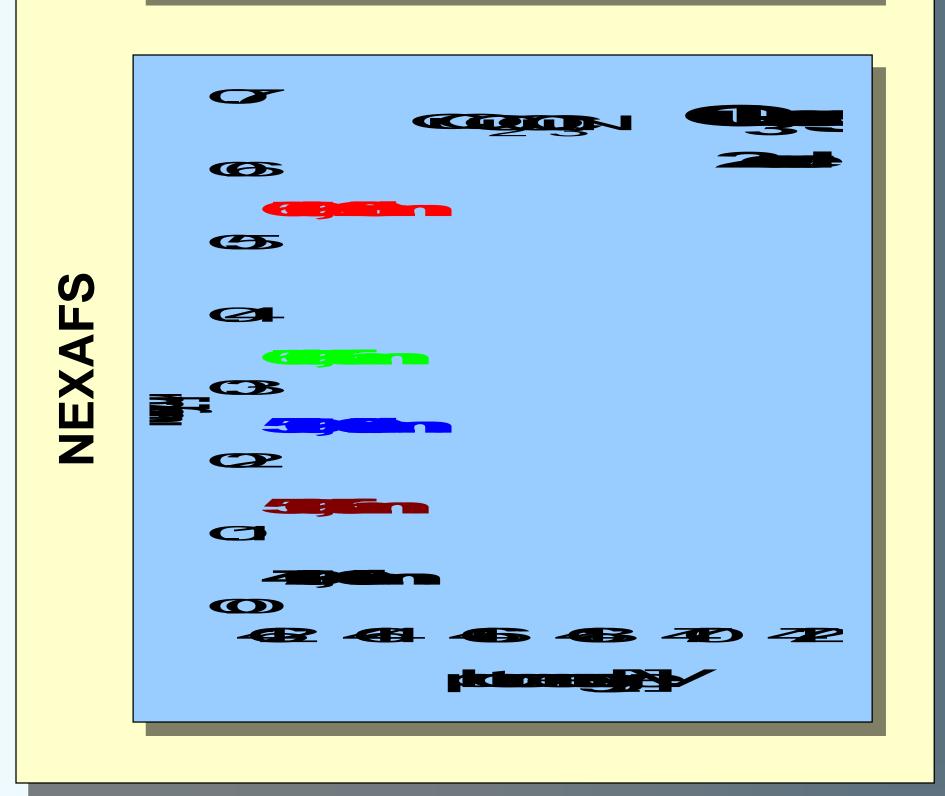
with $NH_3:O_2 = 1:12$ and increasing temperature

Fast Deactivation due to formation of copper(I) nitride (Ref. [4])

3b. Results @ p=1.2mbar and CuO

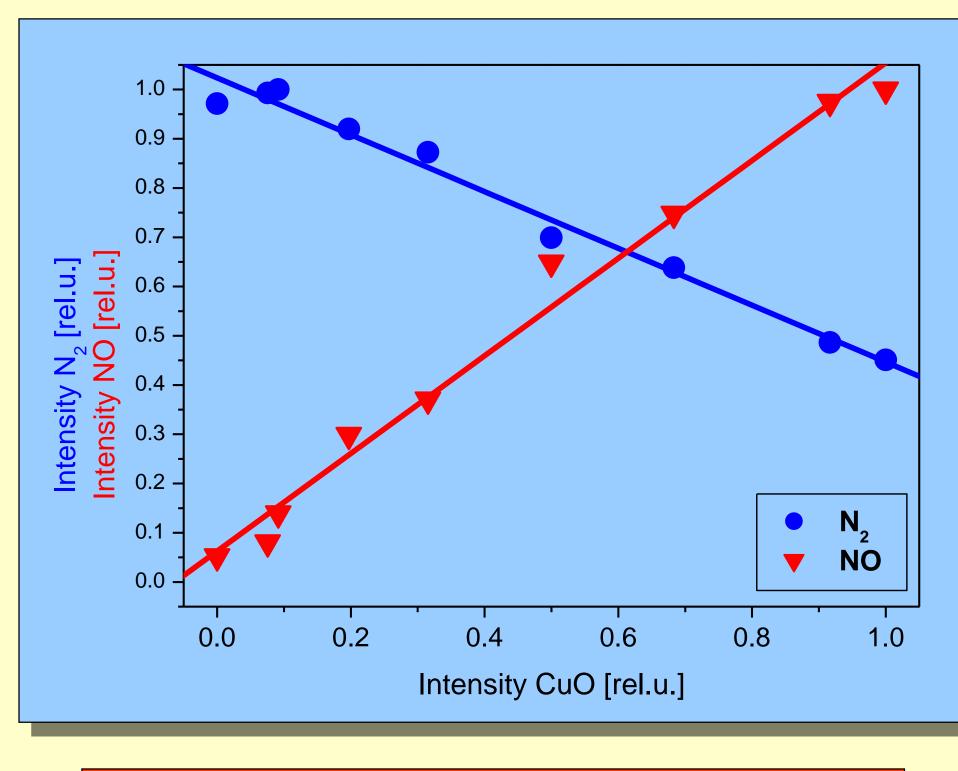
with $NH_3:O_2 = 1:12$ and increasing temperature





4. Conclusions

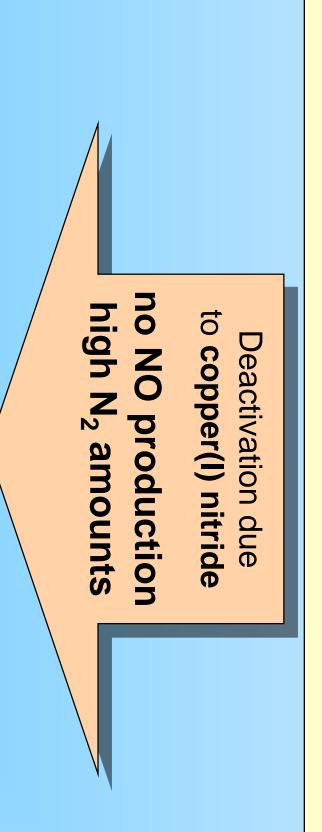
Correlation of the products with CuO surface intensity:

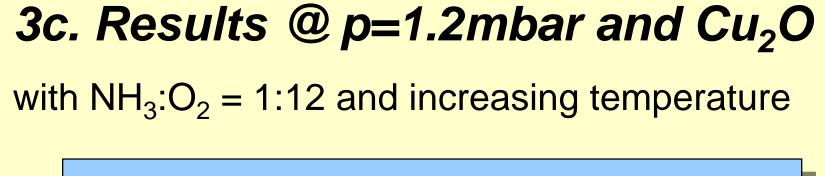


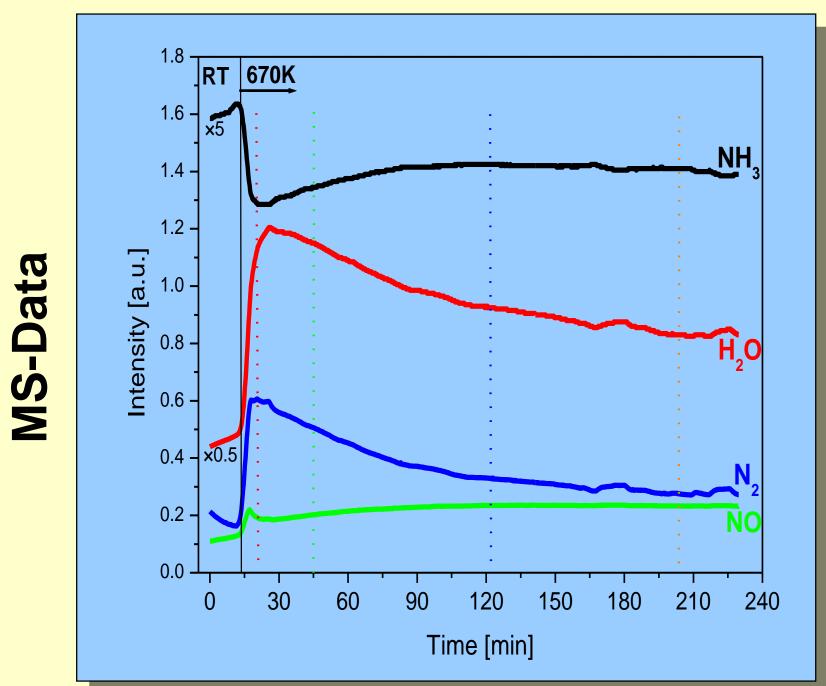
copper(II)oxide catalyzes the total oxidation of ammonia to NO

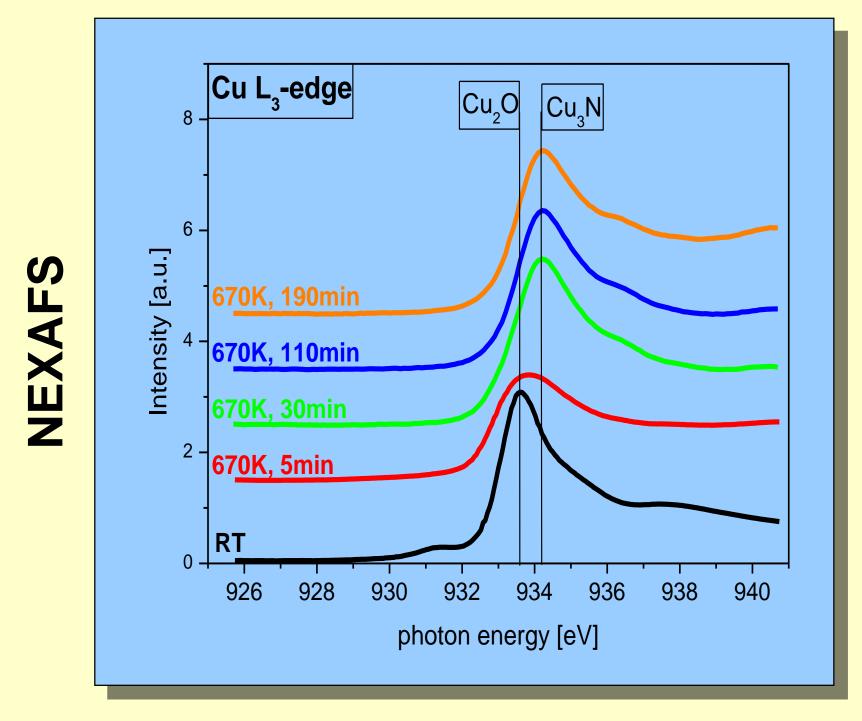
partial oxidation to nitrogen is catalyzed by copper(I)oxide

> but: fast deactivation of Cu₂O due to copper(I)nitride formation





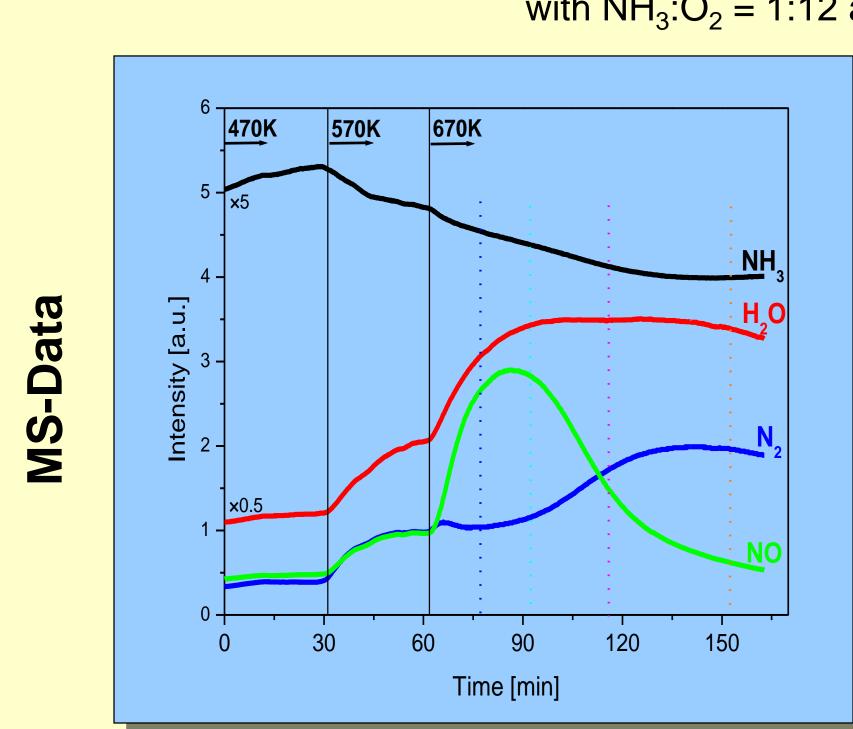




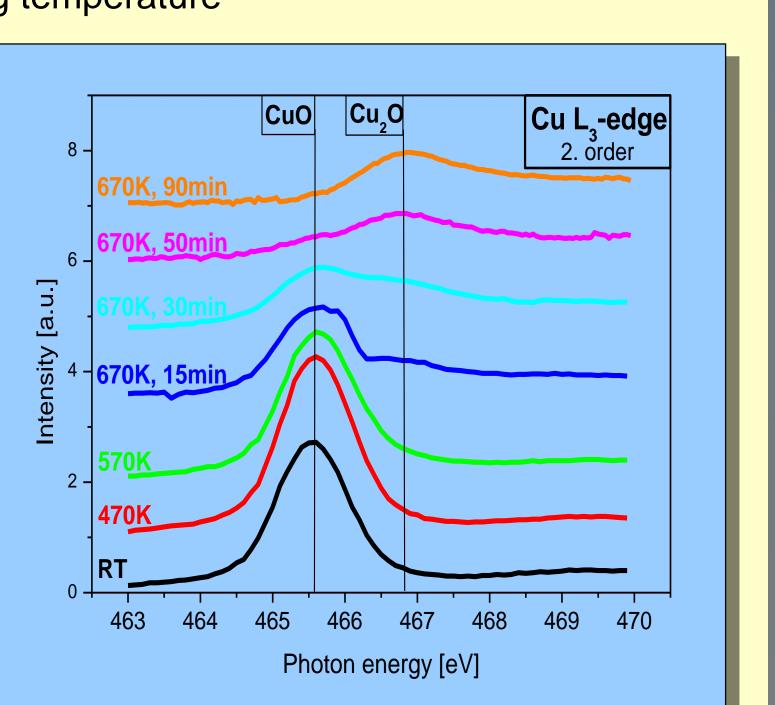
High **NO** production with CuO High N₂ production with Cu₂O

3d. Results @ p=0.8mbar and CuO

with $NH_3:O_2 = 1:12$ and increasing temperature



NEXAFS



References

- [1] H.Bosch and F.J.J.G. Janssen, Catal. Today 2 (1988) 369
- [2] Lu Gang, J. van Grondelle, B.G. Anderson, and R.A. van Santen, J.Catal. 186 (1999) 100
- [3] T. Curtin, F. O'Regan, C. Deconinck, N. Knüttle, and B.K. Hodnett, Catal. Today **55** (2000) 189
- [4] R.W. Mayer, M.Hävecker, A. Knop-Gericke, F.Senf, and R. Schlögl, submitted to Catal. Lett.

Acknowledgement

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