Paired copper monomers in zeolite omega: the active site for methane-to-methanol conversion


Abstract: The direct conversion of methane to methanol using oxygen is a challenging but potentially rewarding pathway towards utilizing methane. By using a stepwise chemical looping approach, copper-exchanged zeolites can convert methane to methanol, but productivity is still too low for viable implementation. However, if the nature of the active site could be elucidated, that information could be used to design more effective catalysts. By employing anomalous X-ray powder diffraction with support from theory and other X-ray techniques, we have derived a quantitative and spatial description of the highly selective, active copper sites in zeolite omega (Cu-omega). This is the first comprehensive description of the structure of non-copper-oxo active species and will provide a pivotal model for future development for materials for methane to methanol conversion.

The development of chemical systems that can convert methane to methanol directly are of high interest, because methane is a cheap and readily available feedstock that continues to be flared globally on a massive scale (143 billion m$^3$/a).[1]–[3] However, this conversion comes with significant challenges, in particular, overcoming the facile over-oxidation of methanol compared to that of methane. Therefore, strategies and systems that can minimize over-oxidation and optimize productivity are required.[4]–[7] One of the most promising and highly investigated approaches involves a chemical looping process over copper-cation-exchanged zeolites, where the oxidative activation of the copper is decoupled from the reaction and product evolution.[8]

Successful partial oxidation of methanol using oxygen relies not only on the storing of oxygen at the active extra-framework copper site, but also on the ability of the zeolite framework to stabilize that active site while simultaneously preventing over-oxidization of the product methanol. This can occur very selectively in copper-zeolites, but productivity remains insufficient for industrial implementation.[9] Therefore, a detailed understanding of the active site upon activation in oxygen and of its structure-property relationships are critical to the development of more active and productive materials. Numerous active-site configurations have been proposed,[8,10–18] but determining the nature of the active site unambiguously or quantifying it directly has not been possible. Instead, proposed active site configurations have come from a combination of several techniques, largely spectroscopic, which are used to infer, rather than determine, the nature and location of the site(s).[19] Despite the debate, all sites that have been identified are copper-oxo species, which form only at high temperatures. [8,10,13–15,18] A pair of monomeric coppers, a precursor to these copper-oxo species, was proposed recently as a potential active site in low-aluminum copper mordenite [19] and in non-microporous systems derived from the grafting of copper onto alumina.[20] Such an active site would have favorable implications regarding minimizing the activation temperature during pre-treatment, and therefore opens up possibilities for the use of new supports. However, the nature of these monomers is unknown, and more importantly, the local environment which is needed to stabilize these monomers is unknown. By exploiting anomalous X-ray diffraction (AXPD) to highlight the copper species and their relation to the crystal structure, we have achieved the most unequivocal assignment of the active sites in methane-to-methanol conversion to date. We identify, both directly and quantitatively, a pair of copper hydroxide monomers, i.e. a non-copper-oxo species, as the active site in zeolite omega, thus giving unprecedented and holistic structure-property information that can serve as a guide for future development of materials for the methane-to-methanol reaction.

In methane-to-methanol conversion, copper-cation exchanged zeolite omega (framework type code MAZ, Figure 1a)[21] can achieve high selectivity (no detectable products of over-oxidation observed by FTIR) and methanol yields for both high (723K) and low (473K) activation temperatures with methanol yields of 200 µmol/gram-zeolite, and 141 µmol/gram-zeolite respectively.[22,23] The reason behind this superior...
The occupancies of the three copper positions refined to 0.56, 0.78 and 0.80 Cu\(^{2+}\) per unit cell and that of O(10) to 1.62 O atoms per unit cell. The electron density of the O(10) position was somewhat elongated, so it was split into O(10) and O(11), each with half the previous occupancy, and these positions and occupancies proved to be stable in refinement. All equivalent positions of the extra-framework atoms are shown in Figure S1. There are 6 Cu(1), 12 Cu(2), 12 Cu(3), 12 O(10) and 12 O(11) symmetry equivalent positions per unit cell, but only a small fraction of them (less than one) are occupied in any given unit cell (approximately 0.5 Cu(1), 0.75 Cu(2), 0.75 Cu(3), 0.75 O(10) and 0.75 O(11) per unit cell). A selection of positions reflecting a possible arrangement of the Cu\(^{2+}\) ions at Cu(1), Cu(2) and Cu(3), is shown in Figure 2. The Cu\(^{2+}\) ions at Cu(1) in the 6-ring are off-center and coordinate to four framework oxygen atoms (two at O(61) and two at O(5)) in the plane of the 6-ring. Only about one 6-ring in four is occupied. The Cu\(^{2+}\) ions at Cu(2) in the 8-ring bond to framework oxygen atoms at O(6) at 1.90 Å and approach two more at O(2) at 2.22 Å. The non-framework oxygen atom at O(11) completes a distorted square planar coordination around the Cu\(^{2+}\). The Cu\(^{2+}\) ions at Cu(3) bond to framework oxygen atoms at O(4) at 1.88 Å and to the non-framework oxygen atom at O(10) at 1.89 Å. The latter has two possible positions. It could act as a bridge to the Cu\(^{2+}\) ion at Cu(2) at 1.89 Å, making the Cu\(^{2+}\) coordination at Cu(2) square pyramidal (as shown in Figure 2), or it could point in the other direction down the narrow 8-ring channel, where it is still 1.89 Å from Cu(3) but not associated with Cu(2). With either arrangement, the Cu\(^{2+}\) ions at Cu(2) and Cu(3) are 3.45 Å apart, and these copper pairs appear to exist as two proximal Cu[OH]\(^{+}\) monomers.

The Rietveld refinement shows two copper sites in the 8-ring, but the occupancy per unit cell is low and their distribution over the available symmetry-equivalent sites cannot be determined directly. To establish whether these coppers are paired across the narrow 8-ring channel, and work in tandem, ex-situ EXAFS from the activated sample and in-situ PDF were analyzed (see supplementary information). The EXAFS fits confirm the presence of a copper-copper distance at 3.4 Å. A PDF derived from in-situ experiments at 723K in oxygen results in a consistent viewpoint with a copper-copper pair distance at 3.3 Å. Thus confirming that these two coppers identified by AXPD do indeed exist in 8-rings of adjacent gme cavities, where they can interact with one another across the narrow 8-ring channel, and thereby facilitate the two-electron redox mechanism for the conversion of methane to methanol.\[26\]
Further confirmation is evident from the refinement of the structure of the reacted sample (Cu-CH$_4$-473K). Refinement was initiated by using just the 0.7105 Å data and the framework structure for the activated material. Cu(1) was unaltered by the reaction step, as expected. However, changes were observed in the coppers in the 8-ring (Figure 2, bottom right). Specifically, Cu(3) moved from its position on the horizontal mirror plane to one where it can interact with two rather than just one framework oxygen and is further away from Cu(2). Only one extra-framework oxygen position was apparent (O(10)), with one of its equivalent positions located 1.99 Å from Cu(2) and a different one 1.90 Å from Cu(3). Finally, a peak close to O(10) appeared in a difference electron density map, and it was added as C(1). These extra-framework oxygen and carbon atoms probably constitute the adsorbed methane conversion intermediate, the methoxy [Cu$^{1+}$-OCH$_3$] species. This intermediate species goes on to form methanol in the extraction step, and corroborates the hypothesis that the reaction is taking place between the coppers on either side of the narrow 8-ring channel where methane has to diffuse through the convoluted 3-dimensional 8-ring channel system to reach the active sites.

As with the activated sample, the other two Cu-CH$_4$-473K data sets were then included in further refinement. The copper ions at Cu(1) were assumed to correspond to the unreactive ones, and were refined as Cu$^{2+}$, while those at Cu(2) and Cu(3) were refined as Cu$^{1+}$, as it has been previously shown that two copper reductions are correlated with each methanol molecule formed. Occupancy parameters for the three copper positions were similar to those found in the Cu-O$_2$-723K structure. The occupancy of O(10) refined consistently to values close to the sum of the occupancies of Cu(2) and Cu(3), and that of carbon C(1) somewhat lower. In the final structural model then, the occupancies of Cu(2) and Cu(3) were constrained to be equal.

Figure 2: Structures of Cu-O$_2$-723K and Cu-CH$_4$-473K. (top) Coordination of the Cu$^{2+}$ ions at Cu(1) in a 6-ring of a zeolite cavity showing the difference between an occupied (front) and an unoccupied (back) 6-ring, and (bottom) coordination geometries of Cu(2) and Cu(3) before and after the introduction of CH$_4$. 

These methoxy species goes on to form methanol in the extraction step, and corroborates the hypothesis that the reaction is taking place between the coppers on either side of the narrow 8-ring channel where methane has to diffuse through the convoluted 3-dimensional 8-ring channel system to reach the active sites.
that of O(10) to be twice that of Cu(2), and that of C(1) to be 1.5 times (to account for the three electrons from H in CH$_3$) that of Cu(2). All equivalent positions of the extra-framework atoms are shown in Figure S1, and a selection of these positions reflecting a possible arrangement is shown in Figure 2.

From the results of the refinement of the occupancies and positions of these extra-framework species, we can now specify the nature and geometry of the active copper site, how the zeolite framework affects the nature and geometry of this site, and how these arrangements lead to the selective conversion of methane. It appears that maintaining a stable distance between the monomers is critical, and this proximity is defined by the zeolite’s framework structure—in particular, the geometry of the narrow 8-ring channels. The proximity of the copper ions to one another may result in a weak interaction between the two [Cu-OH]$^+$ species, causing Cu(3) to move in the gme 8-ring as suggested by the refinement. DFT modelling shows that if these monomers are too close together, they will form a dimer with two bridging hydroxyl groups, while in the 8-ring channels of Cu-omega, proximal monomers can form stable associated pairs. Molecular dynamics simulation of these proximal monomers at 723K provides a glimpse of how these monomers interact. The copper hydroxide maintains its monomeric character, but the distance between the Cu(2) and the Cu(3) may oscillate, and these two monomers can loosely interact (Figure S6). The average Cu(2)-Cu(3) distances derived from the different characterization methods are provided in Figure 3.

Cu-omega is one of the most selective and highest methanol-yielding zeolite systems for the stepwise conversion of methane to methanol. From the measurements reported here, it is clear that this results directly from a framework topology that can stabilize paired [CuOH]$^+$ monomers at specific sites with a relatively high density. By refining the occupancies and locations of all extra-framework species using X-ray powder diffraction techniques, one of the first holistic views of an active copper speciation and its location in the zeolite structure has been achieved. Furthermore, the unprecedented and precise information that we have derived provides a blueprint for a local arrangement of proximal monomers that can be applied beyond zeolites to other non-microporous supports. Just as the development of copper-exchanged Zeolites for methane-to-methanol conversion has been bio-inspired from copper-containing enzymes, the elucidation of the active site and relevant structural properties in zeolite omega, presented here, can lead to the next generation of zeolite-inspired materials for methane-to-methanol conversion.

Acknowledgements

DP gratefully acknowledges the use of computational facilities provided by the Swiss National Supercomputing Centre. We thank Prof. Coperet for use of the UV-Vis spectrometer, Dr. Nachtegaal for assistance at the SuperXAS beamline, Dr. Sushkevich for capillary sealing, Dr. Rzepka for discussions of copper in zeolites, and Dr. Calby-Muzyka for English editing. We would also like to thank SLS for giving us access to the material science and the SuperXAS beamlines.

Keywords: copper zeolites • methane • zeolite omega • methanol • anomalous X-ray diffraction


By employing anomalous X-ray powder diffraction along with support from theory and other X-ray techniques, the active copper site for methane-to-methanol conversion in zeolite omega has been determined spatially and quantitatively.