Spectroscopic Definition of a Highly Reactive Site in Cu-CHA for Selective Methane Oxidation: Tuning a mono- μ -oxo dicopper(II) Active Site for Reactivity

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Abstract

Using UV-Vis and resonance Raman spectroscopy, we identify a $[\text{Cu}_2\text{O}]^{2+}$ active site in O_2 and N_2O activated Cu-CHA that reacts with methane to form methanol at low temperature. The Cu-O-Cu angle (120°) is smaller than for the $[\text{Cu}_2\text{O}]^{2+}$ core on Cu-MFI (140°) and its coordination geometry to the zeolite lattice is different. Site-selective kinetics obtained by operando UV-Vis show that the $[\text{Cu}_2\text{O}]^{2+}$ core on Cu-CHA is more reactive than the $[\text{Cu}_2\text{O}]^{2+}$ site in Cu-MFI. From DFT calculations we find that the increased reactivity of Cu-CHA is a direct reflection of the strong $[\text{Cu}_2\text{OH}]^{2+}$ bond formed along the H-atom abstraction reaction coordinate. A systematic evaluation of these $[\text{Cu}_2\text{O}]^{2+}$ cores reveals that the higher O-H bond strength in Cu-CHA is due to the relative orientation of the two planes of the coordinating bidentate O-Al-O T-sites that connect the $[\text{Cu}_2\text{O}]^{2+}$ core to the zeolite lattice. This work along with our earlier study (*J. Am. Chem. Soc.* 2018, 140, 9236–9243.) elucidates how zeolite lattice constraints can influence active site reactivity.

1. Introduction

The direct conversion of methane into methanol would have large advantages in industry as current processes to produce methanol require harsh and expensive conditions. This one-step oxidation is difficult due to the strong C-H bond of CH₄ and the

Scheme 1: A) Structure of alpha-O site located in a 6MR B) Structure of [Cu₂O]²⁺ site with bidentate O-Al-O T-Site coordination.

tendency for further oxidation of the methanol product.^{1,2} In nature, the Cu- and Fe- enzymes particulate and soluble methane monooxygenase (pMMO and sMMO) catalyze this

reaction at ambient conditions, but the nature of the active sites in these enzymes remains an open issue.^{3–7} Cu- and Fe- doped zeolites also convert methane to methanol in the gas phase at relatively low temperatures.^{8–12} The iron active site, alpha-O, has been well defined as a high-spin Fe(IV)=O site located in a 6-membered zeolite ring (Scheme 1, A) and can react with methane at room temperature.^{9,13} This alpha-O site is formed when the alpha-Fe precursor reacts with N₂O at 160°C.¹³ Whether alpha-O can be formed by O₂ and in what topologies is an open issue.¹⁴ In Cu zeolites, a variety of active sites has been proposed to hydroxylate methane^{12,15–18} but only the [Cu₂O]²⁺ active sites in Cu-MFI and Cu-MOR (Scheme 1, B) have been well-defined in terms of structure and reactivity.^{8,10} These [Cu₂O]²⁺ sites have the advantage of being generated with O₂ in addition to N₂O, however they are less reactive with CH₄ than the alpha-O site in Fe-zeolites that can react at room temperature. Identifying a Cu/O active site that can react at a lower temperature would be advantageous for potential applications in CH₄ conversion and abatement.¹⁹

Copper-exchanged CHA zeolites have received significant research interest recently, both for their outstanding DeNOx capabilities, being commercialized for selective catalytic reduction (SCR) of NOx gasses in heavy vehicles,^{20–22} and for their partial methane oxidation performance.²³ Cu-CHA zeolites activated with either O₂ or N₂O can react with methane to form methanol at 200°C.^{15,16,24} Building on past studies on Cu-MFI¹⁰ and Cu-MOR,^{8,25} O₂ activated Cu-CHA has been studied by several groups using diffuse reflectance (DR-UV-Vis-NIR) and resonance Raman (rR) spectroscopies.^{15,16,26,27} This has led to various active site proposals and combinations thereof, including a trans-μ-1,2-peroxo dicopper(II) species ([CuOOCu]²⁺), a mono-μ-oxo dicopper(II) species ([Cu₂O]²⁺) and Cu(II) superoxo species (both end-on and side-on).^{15,16,26}

Due to the simple structure with a unique T-site and its relevant applications, Cu-CHA is a model-system for understanding structure-activity relationships in zeolite catalysis, which makes a correct assignment of its active

Scheme 2. A) Cage structure in CHA lattice with 8MR windows B) 10MR channel in MFI lattice

Cu site an important addition to the field.²³ To unambiguously determine the active site(s) responsible for methane activation in O₂ activated Cu-CHA, we performed an in-depth spectroscopic study. By collecting the rR excitation profile, extending the rR spectrum out to a higher energy region, identifying active site formation with N₂O, and using an ¹⁸O₂ perturbation of the rR vibrations, we elucidated the presence of a [Cu₂O]²⁺ site that is different from the past assignments in Cu-CHA literature.^{15,16} This [Cu₂O]²⁺

also differs structurally from the one found in Cu-MFI, leading to higher methane activation reactivity ($[\text{Cu}_2\text{O}]^{2+}$ in CHA is ~3 times more reactive than MFI at 25 °C). Using DFT, two contributions to reactivity were evaluated: the different contributions of physisorption in a Cu-CHA cage versus a 10MR channel in Cu-MFI (Scheme 2), and the differences in the $[\text{Cu}_2\text{O}]^{2+}$ active site structure. Complementing our earlier study on Cu-MOR where confinement increased physisorption and influenced reactivity,²⁵ the results here define a second means through which second sphere effects of the zeolite lattice contribute to reactivity. These structure-reactivity insights can help guide the engineering process towards materials with Cu/O active sites that exhibit higher reactivity, similar to alpha-O sites in Fe-zeolites. Combined with the advantage of using O_2 as an oxidant in Cu-zeolites, this will broaden their potential applications.

2. Results and Analysis

2.1 Physical properties

Cu-CHA samples were prepared with $\text{Si}/\text{Al} = 15.7$ and $\text{Cu}/\text{Al} = 0.29$ (1.8 wt% Cu) according to the procedure in the Experimental Sections 1 and 2 in the SI. PXRD patterns of the calcined H-CHA materials as well as the Cu-CHA samples after Cu ion exchange (Experimental Section 2 in SI) and after a reaction cycle correspond to the standard pattern of pure, crystalline H-CHA zeolites with no large copper oxide clusters (Figure S1). The microporosity was determined by N_2 physisorption on calcined H-CHA ($0.30 \text{ cm}^3/\text{g}$) and Cu-CHA ($0.28 \text{ cm}^3/\text{g}$), in good agreement with literature (Figure S2).¹⁵

2.2 DR-UV-Vis-NIR Spectroscopy of the Active Copper Site

DR-UV-Vis-NIR spectra were measured after each sequential step in the activation procedure (Experimental Section 3 in SI). The DR-UV-Vis-NIR spectra of Cu-CHA after calcination in O₂ flow at 450°C, autoreduction in He flow at 700°C, N₂O activation at 550°C and after CH₄ reaction at 200°C (1 atm, 60 minutes) are shown in Figure 1A. Figure 1B shows the analogous cycle replacing N₂O with O₂ in the activation step.

Figure 1. DR-UV-Vis NIR spectra after each step of a typical Cu-CHA activation cycle with A) N₂O as the oxidant and B) O₂ as the oxidant.

After the calcination step, four Cu²⁺ d-d absorption bands can be distinguished at 11,000, 13,600, 16,600 and 20,000 cm⁻¹. These bands have often been observed in literature,^{15,26–31} but an unequivocal assignment in terms of location or coordination of the Cu²⁺ has not yet been made.

After the autoreduction step in He at 700°C, the intensity of the Cu²⁺ d-d bands has reduced, indicating that part of the Cu²⁺ has been converted to Cu⁺. A similar autoreduction occurs in He at 500°C (Figure S3).

Upon treatment with N₂O or O₂ at 550°C, an absorption band appears around 22,400 cm⁻¹, accompanied by a weaker absorption band around 6,200 cm⁻¹. Both bands are more intense after N₂O than after O₂. Similar 6,200 and ~ 22,000 cm⁻¹ absorption bands in Cu-MOR⁸ and Cu-MFI¹⁰ were assigned using rR spectroscopy and ¹⁸O

perturbation to $[\text{Cu}_2\text{O}]^{2+}$ species that react with methane to form methanol. A linear relationship between the intensity of this absorption band and methanol production was previously obtained for Cu-MFI.³²

The $22,400\text{ cm}^{-1}$ band disappears after 60 minutes of exposure to methane at 200°C (Figure 1), but not in Helium (Figure S4), indicating a reaction with methane. Liquid extraction (see Experimental Section 4 in the SI) recovers $82.2\text{ }\mu\text{mol methanol/g}$ for N_2O activated Cu-CHA and $63.6\text{ }\mu\text{mol methanol/g}$ for O_2 activated Cu-CHA. The molar methanol/Cu ratios are 0.29 and 0.22 respectively for N_2O and O_2 activation, comparable to the largest values reported in literature.¹⁶ Absorption bands around $28,100\text{ cm}^{-1}$ and $35,300\text{ cm}^{-1}$ were also lost in the CH_4 reaction step (See UV-Vis difference spectra in Figure S5). Both have been linked to methanol production in literature,^{15,27,28} however their decay does not correlate with that of the $22,400\text{ cm}^{-1}$ band and they were already present after He 700°C , suggesting they represent different sites. Methane reaction immediately after He 700°C (no O_2 or N_2O activation) recovers $25.0\text{ }\mu\text{mol methanol/g}$, which we attribute to the copper species related to the $28,100\text{ cm}^{-1}$ and $35,300\text{ cm}^{-1}$ absorption bands (Figure 1). Similar methanol production of autoreduced Cu-zeolites after reaction with CH_4 has been observed in other systems.^{15,33,34}

2.3 Spectroscopic Definition of the Active Copper Site

Laser excitation into the 22,400 cm^{-1} absorption feature resonantly enhances similar vibrations in both the N_2O and O_2 activated Cu-CHA that are not observed after the autoreduction step (Figure 2). Like the 22,400 cm^{-1} band, these are not present before activation and are eliminated upon reaction with methane (Figure S6). The vibrations resonantly enhanced in the N_2O activated Cu-CHA sample (Figure 2, middle) are identical to those enhanced in the O_2 activated Cu-CHA sample (Figure 2, bottom). The vibrations are more intense in the N_2O activated sample consistent with the more intense

Figure 2. Comparison of the resonance Raman spectra ($\lambda_{\text{ex}} = 458 \text{ nm}$) of autoreduced (top black), N_2O activated (middle blue) and O_2 activated (bottom red) Cu-CHA. The features that are resonantly enhanced are identified by the dotted lines (* indicates a feature from the glass sample holder, ∇ indicates a lattice feature in the autoreduced sample). The rR spectrum of H-CHA can be found in SI Figure 7 for reference.

22,400 cm^{-1} band in the N_2O activated Cu-CHA in Figure 1A. Thus, the same species is formed upon activation of the autoreduced zeolite with either O_2 or N_2O . Scanning the rR laser lines along the 22,400 cm^{-1} chromophore demonstrates that all these vibrations profile with the 22,400 cm^{-1} absorbance indicating that all the vibrations are associated with the same species (Figure 3).

The most intense Raman feature in Figure 2 at 581 cm^{-1} is found to be $^{18}\text{O}_2$ isotope sensitive ($\Delta(^{18}\text{O}_2) = 21 \text{ cm}^{-1}$, see Figure S8) as well as the features at 837 cm^{-1} ($\Delta(^{18}\text{O}_2) = 43 \text{ cm}^{-1}$) and 1161 cm^{-1} ($\Delta(^{18}\text{O}_2) = 32 \text{ cm}^{-1}$). No other isotope shifts were observed above noise due to the lower Raman intensity of O_2 activated Cu-CHA and an incomplete labeling with $^{18}\text{O}_2$.

Cu-MFI and Cu-MOR stabilize $[\text{Cu}_2\text{O}]^{2+}$ cores that can be formed with both O_2 and N_2O and exhibit similar rR features that are associated with their $\sim 22,000 \text{ cm}^{-1}$

Figure 3: A) Zoom in of the 22,400 cm^{-1} DR UV-Vis absorbance feature. B) Full rR profile of the 22,400 cm^{-1} absorption feature formed upon activation with N_2O . Bars on the 22,400 cm^{-1} absorbance spectrum in A are the energies of the rR lines used in (B). Dotted lines on the rR spectrum indicate features in resonance with the absorbance feature. (*) indicates features that are not resonantly enhanced by the 22,400 cm^{-1} feature but are enhanced by other bands.

absorbance
bands.^{8,10}
Based on this
similarity, the
most intense
feature at 581
 cm^{-1} ($\Delta(^{18}\text{O}_2) =$
21 cm^{-1}) is
assigned as the
symmetric

stretch of an oxo group bridging two Cu centers and the vibration at 837 cm^{-1} is assigned as the oxo antisymmetric stretch. This assignment is validated by the observation of progressions in the region between 1,000 cm^{-1} and 2,000 cm^{-1} (Figure 2). The second and third quanta of the 581 cm^{-1} symmetric stretch are observed at 1,161 and 1,743 cm^{-1} , respectively. The second member of a progression in the 837 cm^{-1} antisymmetric vibration is observed at 1,659 cm^{-1} . The 360 cm^{-1} vibration is assigned as

a Cu-O-Cu bend vibration. Finally, the tetrahedral site (T-site) ligand shows a resonance enhanced vibration at 510 cm^{-1} (no isotope perturbation), and this forms a combination band with the symmetric stretch (581 cm^{-1}) at 1091 cm^{-1} .

Our assignment of the rR features of the Cu-CHA active site is different from those previously proposed in the literature. The features at 581 cm^{-1} and 837 cm^{-1} were previously assigned as the Cu-O stretch and the O-O stretch of a trans- μ -1,2-peroxo species, respectively.^{15,16} Trans- μ -1,2-peroxo species have been extensively defined in the literature and do indeed have Cu-O stretches between $550\text{--}600\text{ cm}^{-1}$ and O-O stretches between $800\text{--}850\text{ cm}^{-1}$.^{35–38} However, due to the nature of the charge transfer (CT) transition responsible for the resonance enhancement in trans- μ -1,2-peroxo bridged Cu(II)₂ sites, the O-O stretch is more intense than the Cu-O stretch; this intensity pattern is opposite to that experimentally observed in Cu-CHA. The 837 cm^{-1} feature is weak and now assigned as an antisymmetric stretch which is not rR allowed in idealized Cu₂O C_{2v} symmetry and the 581 cm^{-1} feature is intense as it is the rR allowed symmetric stretch of the Cu-O-Cu core. As shown in Figure 3, all the rR enhanced vibrations of Cu-CHA profile its absorbance feature, which is also inconsistent with a trans- μ -1,2-peroxo species where the CT absorbance feature is made up of three distinct transitions that give different resonance enhancement profiles.³⁵ In another study, the rR features at $1,091\text{ cm}^{-1}$ and $1,161\text{ cm}^{-1}$ were proposed to be O-O stretches of copper superoxide species,¹⁶ which are commonly observed in this energy region.³⁹ However, if these rR vibrations originated from a different Cu/O species they would not have the same resonance enhancement profile as the symmetric and antisymmetric stretches of the [Cu₂O]²⁺ site and they would also not be

formed with N_2O . Additionally, copper superoxide species with O-O vibrations in this region generally have $\Delta^{18}\text{O}_2$ shifts around 60 cm^{-1} . Experimentally we see a smaller value ($\Delta^{18}\text{O}_2 = 32\text{ cm}^{-1}$) for the $\nu_{2\text{sym}}$ vibration at $1,161\text{ cm}^{-1}$. This is consistent with its assignment as the 2x symmetric stretch which has an experimental value of 581 cm^{-1} ($\Delta^{18}\text{O}_2 = 21\text{ cm}^{-1}$) and a DFT calculated frequency of 543 ($\Delta^{18}\text{O}_2 = 10\text{ cm}^{-1}$) (*vide infra*).

Three $[\text{Cu}_2\text{O}]^{2+}$ cores have been previously identified in other zeolite lattices, one in Cu-MFI¹⁰ and two in Cu-MOR.⁸ The rR spectra with 458 nm excitation into the $\sim 22,000\text{ cm}^{-1}$ absorbance

feature of Cu-MFI and Cu-CHA

are compared in Figure 4. The

rR vibrations of Cu-CHA are

shifted compared to those of

Cu-MFI. The red highlights in

Figure 4 compare the

symmetric (ν_{sym}) and 2x

symmetric ($\nu_{2\text{sym}}$) stretches of

the $[\text{Cu}_2\text{O}]^{2+}$ cores in Cu-MFI

(top) and Cu-CHA (bottom). The

ν_{sym} at 581 cm^{-1} and the $\nu_{2\text{sym}}$ at

1161 cm^{-1} in CHA are both higher in energy than in Cu-MFI (ν_{sym} at 456 cm^{-1} and $\nu_{2\text{sym}}$

at 908 cm^{-1}). In contrast, the weaker antisymmetric (ν_{anti}) and the 2x antisymmetric

features ($\nu_{2\text{anti}}$) (blue highlights in Figure 4) are shifted to lower energy in Cu-CHA

Figure 4. Comparison of the vibrations associated with the $[\text{Cu}_2\text{O}]^{2+}$ core in Cu-MFI (top) and Cu-CHA (bottom). The symmetric stretches are shown in the red highlights. The antisymmetric stretches are shown in the blue highlights. The T-sites are shown in the gray highlights and the combination band of the T-site + symmetric stretch is shown in the green highlights.

compared to the previously assigned vibrations in Cu-MFI. Interestingly, the T-site resonance enhancement (ν_{Tsite} , gray highlight) in Cu-CHA at 510 cm^{-1} is located at almost the same energy as the T-site at 514 cm^{-1} in Cu-MFI. The $\nu_{\text{Tsite}} + \nu_{\text{sym}}$ combination band also increases in energy from 974 to 1091 cm^{-1} (green highlight) reflecting the $\sim 120 \text{ cm}^{-1}$ increase in ν_{sym} for Cu-CHA. A normal coordinate analysis (NCA) of a metal-oxygen-metal core⁴⁰ was used to correlate the symmetric and antisymmetric vibrations (see Explanation S1) and determined that the Cu-O-Cu angle in Cu-CHA is 120°, compared to $\sim 140^\circ$ in Cu-MFI and Cu-MOR.

2.4 Kinetic analysis of the active site reaction with CH₄:

Figure 5. Eyring plot for the reaction of CH₄ with [Cu₂O]²⁺ in Cu-CHA (left) and Cu-MFI (right).

The decay of the 22,400 cm^{-1} band of activated Cu-CHA and Cu-MFI was monitored during reaction with methane at a series of temperatures with

operando UV-Vis spectroscopy to obtain site-selective kinetics of [Cu₂O]²⁺ (See

Table 1: Comparison of kinetic parameters of the [Cu₂O]²⁺ species in Cu-CHA and Cu-MFI.

	$\Delta H^\ddagger$ (kcal/mol)	$\Delta S^\ddagger$ (cal/(mol*K))
CHA	6.2	-51.7
MFI	8.7	-45.5

Experimental Section 8 in SI). An Eyring plot was obtained (Figure 5, and

Explanation S2), and from the fitted slope

and intercept an enthalpy of activation $\Delta H^\ddagger$ and entropy of activation $\Delta S^\ddagger$ were obtained for the [Cu₂O]²⁺ species in Cu-CHA and Cu-MFI (Table 1). The enthalpy barrier is 2.5 kcal/mol lower in Cu-CHA.

2.5 Active Site Models: Comparison to Experiment

Previous studies determined that the $[\text{Cu}_2\text{O}]^{2+}$ active site in Cu-MFI is stabilized in a 10MR channel, each Cu(II) with bidentate ligation to the oxygens of the Al T-site and a Cu-O-Cu angle of 140° .¹⁰ The CHA lattice does not contain long channels but is made up of cages connected through adjoining 8MR windows (Scheme 2, A). Large lattice models (~150 atoms) of both the MFI channel and the CHA cage were optimized using Density Functional Theory (DFT). The model of MFI stabilizes a $[\text{Cu}_2\text{O}]^{2+}$ core on one side of a 10MR in an O-Al-O-(Si-O)₂-Al-O sequence with the oxo of the $[\text{Cu}_2\text{O}]^{2+}$ site pointed towards the middle of the ring (Figure 6, A). Due to the smaller size of the CHA 8MR, a $[\text{Cu}_2\text{O}]^{2+}$ site with an angle of 120° can only be stabilized across the 8MR and is in an O-Al-O-(Si-O)₃-Al-O sequence with the bridging O-atom of the $[\text{Cu}_2\text{O}]^{2+}$ pointing out of the 8MR into a CHA cage (Figure 6, B).

Figure 6. $[\text{Cu}_2\text{O}]^{2+}$ cores in the MFI (A, green box) and CHA (B, purple box) lattices and schematic of bidentate oxygen ligation of $[\text{Cu}_2\text{O}]^{2+}$ cores in MFI (C) and CHA (D) assigning out of plane (OOP) and in plane (IP) ligation with respect to the Cu-O-Cu plane.

Note that the $[\text{Cu}_2\text{O}]^{2+}$ active sites identified in CHA and MFI also differ in their bidentate coordination to the lattice oxygens of the Al T-site. In CHA, the oxygen atoms of both bidentate T-site ligands are oriented out of the Cu-O-Cu plane by 93° and 96° (Figure 6, D). In contrast,

in MFI one set of bidentate lattice oxygens is more in the Cu-O-Cu plane (IP) (18°) and one set is out of plane (OOP) (100°) (Figure 6, C).

A comparison of the experimental and DFT predicted $[\text{Cu}_2\text{O}]^{2+}$ structures and spectroscopic features for both lattices are presented in Table 2. The calculated angles

of the $[\text{Cu}_2\text{O}]^{2+}$ cores
are in good

Table 2. Comparison of experimental and predicted spectroscopic and structural features of MFI and CHA and their corresponding large models.

agreement with the

experimental values

determined by rR

and NCA

	MFI (Exp)	MFI Channel	CHA (Exp)	CHA Cage
$\angle \text{Cu-O-Cu}$	140	140	120	126
$d(\text{Cu-O}_{\text{lattice}}) / \text{\AA}$	—	1.98-2.04	—	1.93-2.04
$d(\text{Cu-O}_{\text{oxo}}) / \text{\AA}$	—	1.75	—	1.73-1.75
Abs / cm^{-1}	22700	24930	22000	23246/26099
$\nu_{\text{sym}} (\Delta^{18}\text{O}_2) / \text{cm}^{-1}$	456 (8)	454 (5)	580 (21)	543 (10)
$\nu_{\text{asym}} (\Delta^{18}\text{O}_2) / \text{cm}^{-1}$	870 (40)	847 (41)	837 (43)	842 (40.5)

(Explanation S1) (139.5° vs. 140° for MFI; 126° vs. 120° for CHA). The calculated Cu-O-Cu symmetric and antisymmetric vibrations as well as their $\Delta^{18}\text{O}_2$ shifts reinforce the experimental assignments.

2.6 Reaction Coordinate Calculations

To elucidate the origin of the higher experimental reactivity of methane with the active site in Cu-CHA compared to Cu-MFI (Table 1), the H atom abstraction (HAA) reaction coordinates of CH_4 to form $[\text{Cu}_2\text{OH}]^{2+}$ and a $\cdot\text{CH}_3$ radical were calculated for the large lattice models, both with dispersion (solid lines) and with the dispersion removed (dotted lines) (Figure 7). The corresponding enthalpy values are given in Table 3. The difference between the calculations with and without dispersion gives an estimate of the physisorption contribution to the energetics of each step. The sites were optimized with dispersion, then the structure was fixed, and the dispersion was eliminated. Starting from the left of the reaction coordinate in Figure 7, gas phase

methane is first physisorbed to the Cu-O-Cu active site (Reactant Complex, RC). The RC then goes through a transition state (TS) to form a product complex (PC) which is a Cu^I-OH-Cu^{II} core with a physisorbed ·CH₃ radical. The ·CH₃ radical was then removed from the zeolite lattice to calculate the physisorption contribution to the energetics of the product complex.

Physisorption of CH₄ into the zeolite lattice was evaluated to determine its effect on the apparent activation barrier. Relative to the calculations without dispersion, methane physisorption from the gas phase to the active site is favorable by ~ 10 kcal/mol (Figure 7, left to RC). There is not much difference in the physisorption energetics between the CHA and MFI models, suggesting that this does not contribute to the increased reactivity in CHA. This is in contrast to the results in MOR where the physisorption energy difference was the dominating contribution to the difference in reactivity between a more and a less confined [Cu₂O]²⁺ species.²⁵ On the right side of Figure 7, physisorption of the methyl radical to the Cu-OH-Cu product (Fig 7, right to

PC) is calculated to be favorable by ~10 kcal/mol. There is an additional interaction of the new H-bond formed between the methyl radical and the Cu-OH-Cu core that results from the HAA reaction that is worth ~3 kcal/mol (based on the stabilization of the $\cdot\text{CH}_3$ radical by the $[\text{Cu}_2\text{OH}]^{2+}$ site without dispersion). Overall, the models exhibit similar physisorption into the zeolite for both the CH_4 substrate and the $\cdot\text{CH}_3$ radical.

Now focusing on the TS energies, CHA has a 4.4 kcal/mol lower TS energy compared to MFI (CHA lower by 3.3 kcal/mol in the calculations without dispersion). If we look at apparent values where we compare the $\text{CH}_4 + [\text{Cu}_2\text{O}]^{2+}$ to the HAA TS, the difference is 2.2 kcal/mol with dispersion. This value is in good agreement with the experimental enthalpy difference with CHA lower than MFI by 2.5 kcal/mol (Table 1). In order to determine the effect of

Figure 7. CH_4 reaction coordinates for large channel/cage models for MFI (green) and CHA (purple). Dotted lines are calculations without dispersion. Values for each step in the reaction coordinate for each structure are given in Table 3. Reactant Complex energies are all set to zero.

Table 3. Enthalpy values (kcal/mol) for the different complexes in Figure 7.

ΔH (kcal/mol)	no dispersion		dispersion	
	CHA Cage	MFI Channel	CHA Cage	MFI Channel
$\text{CH}_4 + [\text{Cu}_2\text{O}]^{2+}$	0.0	0.4	9.5	11.7
Reactant Complex	0.0	0.0	0.0	0.0
HAA Transition State	16.6	19.9	16.9	21.3
Product Complex	12.0	15.6	10.3	15.0
$\text{CH}_3 + [\text{Cu}_2\text{OH}]^{2+}$	15.4	19.8	23.0	29.2

thermodynamics on the TS, Marcus theory⁴¹ was used to extract the intrinsic barriers (i.e. with no thermodynamic driving force, see Explanation S3). These values are given in Table 4. The calculated intrinsic barriers for the Cu-CHA and Cu-MFI reaction

coordinates in the large lattice models are similar (11.2 vs. 12.7 kcal/mol) indicating that most of the 4.4 kcal/mol of TS difference is due to the thermodynamic difference between the reactant and the product.

The difference in energy of the two extremes on the reaction coordinate in Figure 7 (where the CH₄ and •CH₃ are in the gas phase removed from the active site cores) gives direct insight into the thermodynamics for the reaction coordinate, reflecting the strength of the O-H bond formed along the HAA reaction coordinate (values are given in

Table 4). The O-H bond formed in Cu-CHA is calculated to be 4 kcal/mol stronger than MFI.

Table 4: HAA Transition State energies (in kcal/mol), Marcus corrected intrinsic barriers and calculated O-H bond strengths for the large lattice models (with respect to the 103.3 kcal/mol calculated C-H bond of CH₄).

ΔH (kcal/mol)	HAA Transition State	Intrinsic Barrier	OH Bond Strength
CHA CAGE	16.9	11.2	89.8
MFI Channel	21.3	12.7	85.8

Therefore, the higher reactivity

exhibited in Cu-CHA is due to the stronger O-H bond that forms in Cu-CHA. We also calculated HAA reaction coordinates for small ring models (SRMs) that do not include the walls of the zeolite lattice. These SRMs reproduce the trends observed in the large lattice models (Explanation S4).

To determine the geometric and electronic structure contributions to this OH bond strength difference between CHA and MFI, smaller models were developed by extracting the Cu-O-Cu reactant and Cu-OH-Cu product cores along with their coordinating O-Al-O T-sites from the large lattice models (Figure 8). The lattice Si atoms capping the Al T-sites were replaced with hydrogen atoms set along the bond direction to a bond length of 0.96 Å. To ensure that the small core model had the same coordinates (angles, bond lengths, bidentate O angles, etc.) as the larger lattice models,

no additional optimization was performed after extracting the core and capping the Al T-sites with hydrogens. A single point calculation was performed to obtain the energy of each structure. These extracted core models of MFI and CHA have the same number of atoms and electrons and can be directly compared (Figure 8).

Figure 8. Relative stability of extracted Cu-O-Cu reactant and Cu-OH-Cu product cores for MFI (green) and CHA (purple).

The reactant Cu-O-Cu core in Cu-CHA was calculated to be 4.1 kcal/mol higher in energy than in Cu-MFI while the Cu-OH-Cu product was calculated to be only 0.3 kcal/mol higher in Cu-CHA than in Cu-MFI. This indicates that the higher OH bond strength in CHA derives from a destabilization of the reactant Cu-O-Cu core imposed by the CHA lattice.

As mentioned earlier in the study, the main structural differences between the reactant cores of CHA and MFI are the angle of the Cu-O-Cu core (120° in MFI and 140° in CHA) and the rotation of the bidentate oxygen ligands with respect to the Cu-O-Cu plane (close to 90°/0° in MFI and 90°/90° in CHA). The contributions of these differences to the relative destabilization of 4.1 kcal/mol of the reactant core in CHA were evaluated using a series of small models (Experimental Section 10 in SI). A series of fixed ligand rotations was evaluated (0°/0°, 90°/0°, +45°/-45° and 90°/90°) with respect to the Cu-O-Cu plane. These ligand rotation sets were analyzed for three Cu-O-Cu angles (161° from unconstrained optimization, 140° to reflect MFI and 120° to reflect

CHA). The relative energies for these models are given in Table 5 and their structure

and LUMOs in Figure S14. Table 5 shows that for models with the same bidentate oxygen ligand rotations, the Cu-O-Cu angle does not significantly affect the energy. The models with the two bidentate oxygen pairs in the same plane as each other (rotated 0°/0° and 90°/90° with respect to the Cu-O-Cu plane) are calculated to be ~ 6 kcal/mol higher in energy than those with ligand sets rotated by 90° from each other (rotated +45°/-45° and 90°/0° with respect to the Cu-O-Cu plane) across all evaluated Cu-O-Cu angles. The actual rotation of the bidentate ligands with respect to the Cu-O-Cu plane does not significantly impact the energy, only their rotations with respect to each other.

Table 5: Comparison of energies (kcal/mol) of models with various Cu-O-Cu angles and bidentate oxygen ligand rotation.

Energy (kcal/mol) Bidentate Oxygen Ligand Rotations (with respect to the Cu-O-Cu plane)	Cu-O-Cu angle		
	161°	140°	120°
+45° / - 45°	0.4	1.4	0.2
0° / 0°	6.4	7.2	14.5*
90° / 0°	0.0	2.5	0.3
90° / 90°	6.7	5.5	5.9

(*) larger energy difference due to steric interaction between two in plane Al T ligands at the acute 120° angle.

The bidentate O-Al-O T-site ligation on each Cu results in a highest energy singly occupied d x^2-y^2 orbital on each metal. The $[\text{Cu}_2\text{O}]^{2+}$ models with the bidentate oxygen ligands in the same plane as each other (Figure 9, right) have two frontier molecular orbitals (FMOs) (the β LUMO and β LUMO+1) that are the (–) and (+) combinations, respectively, of the d x^2-y^2 orbitals on each Cu(II), delocalized over the model. The (+) combination (Figure 9 top right) of the two Cu d x^2-y^2 orbitals interacts with an OOP oxo 2p orbital leading to 29.8% oxo character while the (–) combination (Figure 9, bottom right) has only a limited interaction with the IP oxo 2p orbital and thus a small amount (8.3%) of oxo character. In the models with ligand sets rotated 90° from each other, the

two FMOs are again (+) and (−) combinations of $d x^2-y^2$ orbitals but now each is localized on a different Cu(II) which interacts with a different oxo p orbital, resulting in ~24% oxo character for both combinations (Figure 9, left).

Figure 9. Comparison of FMOs for bidentate oxygen ligands rotated 90° from each other (left) and in the same plane (right) (triplet ground state; see SI Figure S13 for the full series).

Thus, for the ligand sets with 90° relative rotation as in MFI, each Cu $d x^2-y^2$ orbital has a strong bonding interaction with a different oxo p orbital resulting in 10% more oxo

character distributed over the two LUMOs relative to the 0° ligand rotation on Figure 9, right (47.7% vs 38.1% total oxo character). This results in the stabilization of this active site relative to the 0° rotation as in CHA, where each Cu $d x^2-y^2$ orbital competes with the same oxo 2p orbital for bonding. Thus, the copper atoms in CHA bind less strongly to the oxo than in MFI destabilizing the $[\text{Cu}_2\text{O}]^{2+}$ active site and priming it for reactivity. This conclusion holds throughout the series of small models with varying Cu-O-Cu angles in Figure S14.

3. Discussion

A $[\text{Cu}_2\text{O}]^{2+}$ core with an angle of 120° is identified as the active site in Cu-CHA that converts methane to methanol at low temperature. This $[\text{Cu}_2\text{O}]^{2+}$ active site assignment in Cu-CHA is different from previous proposals of multiple Cu/O species

contributing to the features observed in the Cu-CHA rR spectrum.^{15,16} These previous assignments were largely based on empirical correlations to Cu/O structures present in model and enzyme literature.^{10,35,42,43} However, in the present study additional new data were collected including the rR spectrum of Cu-CHA activated with N₂O, the ¹⁸O₂ perturbation on the rR spectrum, the full resonance enhancement profile over the associated absorption feature and an extension of the rR spectrum to the higher energy overtone region. These additional experiments enabled an unambiguous assignment of the rR features of Cu-CHA, highlighting the importance of rigorous spectroscopy and the possibility of inaccurate assignments relying on empirical correlations to define an intermediate.

From rR, the [Cu₂O]²⁺ site defined in Cu-CHA has a more acute Cu-O-Cu angle than the one defined in Cu-MFI and from kinetic experiments is found to have a lower enthalpic barrier by 2.5 kcal/mol. DFT modeling reproduces the experimental results and indicates that this difference in reactivity is due to the thermodynamic driving force associated with the stronger O-H bond formed in the [Cu₂OH]²⁺ CHA intermediate generated in the HAA reaction. While the [Cu₂OH]²⁺ intermediates in the two lattices are similar in energy, the reactant [Cu₂O]²⁺ site in Cu-CHA is destabilized compared to that in Cu-MFI and this lowers the activation barrier in Cu-CHA. This destabilization is not due to the change in the Cu-O-Cu angle but derives from the lattice-imposed relative orientations of the two bidentate oxygen ligand sets.

The results from the present study on Cu-CHA and Cu-MFI are different from what was found in our prior studies on Cu-MOR where two [Cu₂O]²⁺ sites are present with different reactivity but similar Cu-O-Cu structures.²⁵ In MOR, the two [Cu₂O]²⁺ sites

are located in different parts of the zeolite lattice, one in the constrained 8MR side pocket and one in a large 12MR channel. Their difference in the apparent activation is due to the large difference in physisorption enthalpy for the two sites ($\Delta\Delta H=7.7$ kcal/mol) reflecting stabilization by the constrained pocket. In the present study, physisorption was calculated to be similar in both CHA and MFI (Figure 7, left side) and it is not a dominating contribution to the difference in apparent activation barriers between CHA and MFI. The lattice constraints associated with the bidentate T-site ligation in CHA impose a destabilizing geometry on the $[\text{Cu-O-Cu}]^{2+}$ core, reminiscent of the entatic state in metalloproteins where protein constraints enforce an unstable geometry that is associated with high reactivity.¹ It is interesting to note that the approximate perpendicular orientation of the bidentate O-Al-O T-Site ligand planes in MFI and their parallel orientation in CHA would reflect the lower and upper limits on this role of the zeolite lattice in activating a $[\text{Cu}_2\text{O}]^{2+}$ core. Improving upon the reactivity of $[\text{Cu}_2\text{O}]^{2+}$ in Cu-CHA could occur via topologies with a more confined cage which still maintains coplanar geometry of the four lattice oxygen ligands. The latter not only depends on the zeolite topology, but also on the location of Al substitution which can, in some cases, be guided during the zeolite synthesis.^{44–46}

Together, the previous study on MOR and the present study demonstrate two means through which a carefully chosen zeolite support can enhance the reactivity of a $[\text{Cu}_2\text{O}]^{2+}$ species, providing a direction for optimization towards room temperature reactivity. This would open up new applications for these materials, such as the ambient capturing of CH_4 from dilute emission streams or from the atmosphere as a negative emission technology.¹⁹

Notes:

The authors declare no competing financial interest.

Associated Content:

The following information is available free of charge on the ACS Publications Website: Experimental Section, PXRD patterns, nitrogen sorption isotherms, additional DR UV-Vis spectra and rR spectra of Cu-CHA, rR of $^{18}\text{O}_2$ activated Cu-CHA, normal coordinate analysis of a Cu-O-Cu angle, Arrhenius plot of the reaction of CH_4 with Cu-CHA and Cu-MFI, DFT of small ring Cu-CHA and Cu-MFI models, DFT cluster model coordinates

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