

Electronic Structure of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ Cluster in Mordenite Zeolite and Its Effects on the Methane to Methanol Oxidation

Vogiatzis, Konstantinos D.; Li, Guanna; Hensen, Emiel J M; Gagliardi, Laura; Pidko, Evgeny A.

DOI

[10.1021/acs.jpcc.7b08714](https://doi.org/10.1021/acs.jpcc.7b08714)

Publication date

2017

Document Version

Final published version

Published in

The Journal of Physical Chemistry C

Citation (APA)

Vogiatzis, K. D., Li, G., Hensen, E. J. M., Gagliardi, L., & Pidko, E. A. (2017). Electronic Structure of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ Cluster in Mordenite Zeolite and Its Effects on the Methane to Methanol Oxidation. *The Journal of Physical Chemistry C*, 121(40), 22295-22302. <https://doi.org/10.1021/acs.jpcc.7b08714>

Important note

To cite this publication, please use the final published version (if applicable).
Please check the document version above.

Copyright

Other than for strictly personal use, it is not permitted to download, forward or distribute the text or part of it, without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license such as Creative Commons.

Takedown policy

Please contact us and provide details if you believe this document breaches copyrights.
We will remove access to the work immediately and investigate your claim.

Electronic Structure of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ Cluster in Mordenite Zeolite and Its Effects on the Methane to Methanol Oxidation

Konstantinos D. Vogiatzis,^{*,†,id} Guanna Li,^{‡,§,id} Emiel J. M. Hensen,^{‡,||,id} Laura Gagliardi,^{⊥,id}
and Evgeny A. Pidko^{*,‡,||,#,▽,id}

[†]Department of Chemistry, University of Tennessee, Knoxville, Tennessee 37996, United States

[‡]Inorganic Materials Chemistry Group, Eindhoven University of Technology, PO Box 513, Eindhoven 5600 MB, The Netherlands

[§]Catalysis Engineering, Department of Chemical Engineering, Delft University of Technology, Van Oord Massag 9, 2629 HZ Delft, The Netherlands

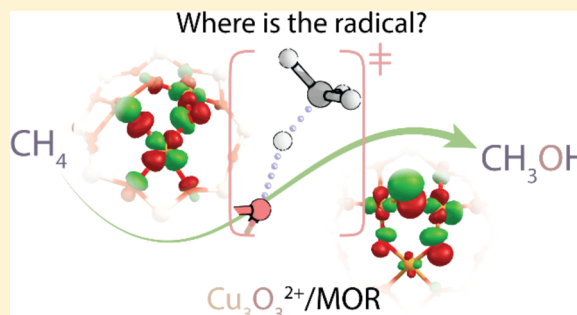
^{||}Institute for Complex Molecular Systems, Eindhoven University of Technology, Eindhoven 5600 MB, The Netherlands

¹Department of Chemistry, and Minnesota Supercomputing Institute, University of Minnesota, Minneapolis, Minnesota 55455, United States

[#]Theoretical Chemistry Group, ITMO University, Kronverkskiy pr., 49, St. Petersburg 197101, Russia

S Supporting Information

ABSTRACT: Identifying Cu-exchanged zeolites able to activate C–H bonds and selectively convert methane to methanol is a challenge in the field of biomimetic heterogeneous catalysis. Recent experiments point to the importance of trinuclear $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ complexes inside the micropores of mordenite (MOR) zeolite for selective oxo-functionalization of methane. The electronic structures of these species, namely, the oxidation state of Cu ions and the reactive character of the oxygen centers, are not yet fully understood. In this study, we performed a detailed analysis of the electronic structure of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ site using multiconfigurational wavefunction-based methods and density functional theory. The calculations reveal that all Cu sites in the cluster are predominantly present in the Cu(II) formal oxidation state with a minor contribution from Cu(III), whereas two out of three oxygen anions possess a radical character. These electronic properties, along with the high accessibility of the out-of-plane oxygen center, make this oxygen the preferred site for the homolytic C–H activation of methane by $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$. These new insights aid in the construction of a theoretical framework for the design of novel catalysts for oxyfunctionalization of natural gas and suggest further spectroscopic examination.



■ INTRODUCTION

Natural gas is considered the main source of conventional energy for the future due to recent technological advances in the excavation techniques and the lower environmental impact of its utilization compared to other fossil fuels.¹ The primary component of natural gas is methane. Its low price and large availability have motivated scientists and engineers to explore new routes for the efficient conversion of methane to liquid fuels and chemicals.² The selective catalytic oxyfunctionalization of methane to methanol is considered one of the most viable conversion routes.^{1,3,4} Methanol is a reactive and easily condensable chemical that can serve both as the energy carrier and as a versatile platform chemical, and it thus plays a central role in future chemical and energy technologies.⁵ An important advantage of methanol over other chemicals is the ease of integration into the existing chemistry infrastructure and the possibility to derive it from a wide range of potential feedstocks. The selective activation of methane and its direct conversion to methanol represent an important challenge for catalysis and

chemical engineering. The established technology to obtain methanol is only profitable at a very large scale. It involves first the steam reforming of methane over a nickel catalyst to produce syngas, which is then converted to methanol over a Cu/ZnO/Al₂O₃ catalyst in a separate process. The overall conversion scheme is optimized for the large-scale operation under high temperature and pressure conditions.⁶ The availability of a cheaper and more flexible process at smaller scale not only would allow developing relatively small remote natural gas fields such as shale gas and clathrate reserves but would also constitute the game changer in the chemical industry as it would enable the production of a wide range of chemicals and fuels.

Methane monooxygenases (MMOs) are a family of enzymes that nature employs for selective oxidation of the inert C–H bond of methane.⁷ The MMOs are sorted in two different

Received: September 1, 2017

Published: September 8, 2017

categories based on their position in the cell. The cytoplasmic, soluble MMOs (sMMOs) contain a bis(μ -oxo)diiron core,⁸ while the membrane-bound MMOs (particulate MMOs or pMMOs) oxidize the C–H bond over a Cu-based active site. Different handling conditions in the purification of the pMMOs resulted in different Cu concentrations giving rise to a substantial uncertainty about the nature of the Cu-containing active site. Whereas dimeric Cu complexes have been postulated as the active centers by Rosenzweig et al.,^{9–11} an alternative hypothesis of the trinuclear speciation of the reactive Cu sites is advocated by Chan et al.¹²

Biomimetic catalysis is a promising approach to the design of functional materials, but it also provides new insights into how enzymes work. Nanoporous inorganic matrices such as zeolites and metal–organic frameworks (MOFs) provide well-defined confinement space and coordination environments suitable for the stabilization of highly reactive intermediates often regarded as enzymatic analogues. Cu-containing reactive species have been the synthetic target for homogeneous^{13,14} and heterogeneous^{15–25} catalysis. Cationic Cu complexes stabilized in zeolite micropores have been shown to be efficient catalysts for the selective oxyfunctionalization of methane.¹ However, the exact structure of the catalytic sites in such materials is still not fully understood.^{15,26–35} Originally, a binuclear bis-oxo $[\text{Cu}(\mu\text{-O})_2\text{Cu}]^{2+}$ ion was proposed as the active site in Cu-ZSM-5 zeolite.¹⁶ In later studies, Woertink et al. employed Raman spectroscopy to characterize the working Cu-ZSM-5 catalyst and suggested the mono-(μ -O) dicopper site $[\text{Cu}(\mu\text{-O})\text{Cu}]^{2+}$ to be the active complex, which does not have analogues in biological or organometallic systems.³⁵ Similar dicopper sites were proposed for Cu-SSZ-13 and Cu-SSZ-39 zeolites based on synchrotron X-ray diffraction, in situ UV–vis, Raman spectroscopy, and theory.³⁶ Recent theoretical studies addressed the stability and catalytic performance of mono-CuOH,³⁷ dicopper,³⁸ and polynuclear Cu_xO_y ($x = 1\text{--}5$) sites³⁹ as possible catalytic sites responsible for methane oxidation. An alternative structural proposal suggesting the trinuclear active site formulation $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ (Figure 1) has been put forward

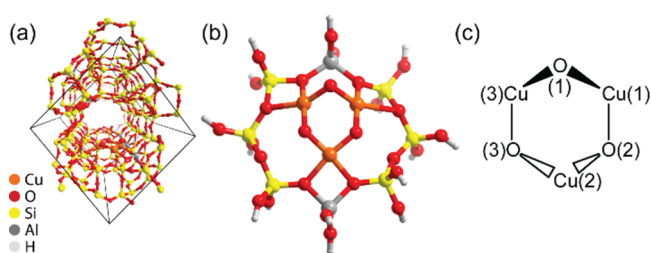


Figure 1. (a) Optimized periodic structure of Cu/MOR containing a $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cationic cluster in the $S = 1/2$ spin state and (b) the respective cluster with (c) the schematic representation and atom numbering within the trinuclear copper-oxo core.

by Grudner et al. on the basis of a combined experimental and computational study on well-defined Cu-exchanged mordenite catalysts.^{21,40,41} The active site is the ring-type $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ extraframework cluster, in which the positive charge is balanced by two Al sites of the zeolite framework. A similar active site has been proposed for pMMO,¹³ and it has also been isolated in a molecular complex.^{42,43} The importance of the trinuclear structure for the activity of Cu-ZSM-5 catalysts has also been recently discussed.⁴⁴

Understanding the relationships between the electronic structure, geometry, and reactivity of such multinuclear clusters is essential not only to explain the mechanism of selective oxidation of methane by zeolites but also to design new and improved catalyst systems. The key challenge here is that the electronic configuration of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster as well as other potential multinuclear Cu-oxo complexes cannot be readily rationalized based on two-electron Lewis models. The presence of at least three paramagnetic centers and of covalent bonds may give rise to a multiconfigurational system, whose description needs more than one (“multiple”) electronic configuration. Most computational studies on Cu–zeolite systems reported up to date have been performed using single-determinant density functional theory (DFT) that has intrinsic limitations for multiconfigurational systems.⁴⁵ The multiconfigurational character of the Cu-oxo clusters that may be responsible for metal-to-ligand charge transfer was claimed to be crucial for their ability to selectively activate methane.²¹ The evidence of such effects further comes from X-ray absorption spectroscopy, indicating the predominance of Cu(II) in the active materials, whereas the formal oxidation state of Cu in the trimer is Cu(III).

In this study, we employed a high-level ab initio methodology based on multiconfigurational wave function theory to obtain detailed information about the ground spin state of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster, the partial oxidation state of each Cu center, and the oxo/oxy character of the three (μ -O) atoms. High-level quantum chemical calculations have previously been employed to investigate Cu-exchanged zeolites⁴⁶ and Fe-oxo sites in zeolitic materials^{4,47} and elucidate their complex electronic structure. The results of the multiconfigurational calculations are compared with those obtained with a range of popular density functionals providing thus a basis for calibrating the choice of suitable methods for applied computational catalysis studies on related systems. An important finding is that the substantial anisotropy activity of the (μ -O) centers correlates with both their accessibility and their radical character.

■ COMPUTATIONAL DETAILS

Periodic DFT Calculations. All spin-polarized periodic DFT (spin-polarized pDFT) calculations were performed using the VASP software with a generalized gradient-approximated PBE exchange-correlation functional.^{48,49} The projected-augmented wave method and a plane-wave basis set with a cutoff of 400 eV were employed.⁵⁰ Brillouin zone sampling was restricted to the Γ point. A supercell of all-silica MOR, constructed by doubling of the monoclinic primitive cell along the c axis with lattice parameters of $a = b = 13.648$, $c = 15.015$ Å, and $\gamma = 97.2^\circ$ as optimized by DFT, was used as the initial model (Figure 1(a)).^{51,52} To compensate for the positive charge of the extra-framework cationic Cu complexes, two framework Si^{4+} ions in the MOR supercell were substituted by two Al^{3+} at the side-pocket positions $\text{Al}^{\text{I}}_{\text{sp}}$. The other two $[\text{AlO}_2]^-$ units at the remote side-pocket positions ($\text{Al}^{\text{II}}_{\text{sp}}$) were charge compensated by two Brønsted acid sites.⁵³ The resulting MOR model had a Si/Al ratio of 11. The climbing image nudged elastic band method⁵⁴ was used to determine the minimum energy path and locate the transition state for the homolytic C–H activation step resulting in a hydroxylated Cu site and a CH_3 radical species.^{40,44} The maximum energy geometry along the reaction path obtained by the climbing image nudged elastic band method was further optimized using

Table 1. Relative Energy Differences (in kJ/mol) from the CASSCF/CASPT2(11,11) and RASSCF/RASPT2(19,21) Calculations

	CASSCF(11,11) ^a	RASSCF(19,21) ^a	CASPT2(11,11)	RASPT2(19,21)
sextet	22.6	57.1	78.1	108.6
quartet	6.2	13.1	20.7	22.1
doublet	0	0	0	0

^aThe weights of the dominant configurations demonstrate the multiconfigurational character of the doublet state, and they are 46%, 25%, 15%, and 8% (CASSCF(11,11)) and 53%, 18%, 14%, and 5% (RASSCF(19,21)).

a quasi-Newton algorithm, as implemented in VASP. In this step, only the extra-framework atoms were relaxed. Spin-polarized calculations were performed throughout this study. Vibrational frequencies for optimized structures of reaction intermediates and transition states were calculated using the finite-difference method, as implemented in VASP. Small displacements (0.02 Å) were used to estimate the numerical Hessian matrix. The transition states were confirmed by the presence of a single imaginary frequency corresponding to the reaction path.

Multiconfigurational Calculations. For all multiconfigurational calculations a cluster model of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ core stabilized inside the mordenite zeolite micropores was constructed from the periodic DFT optimized structure with spin $S = 1/2$. The cluster was saturated with hydrogen atoms (Figure 1b). Cu(1), O(2), Cu(3), and O(3) (Figure 1c) are located on the same plane; Cu(2) is out-of-plane and closer to the framework of the MOR zeolite; and O(1) is located also out-of-plane and points to the main channel of MOR. All calculations were performed without any symmetry restrictions (point group C_1). The structure reported in Figure 1b was used in the multiconfigurational calculations to elucidate the electronic effects of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ core on the methane functionalization, while it was not employed to model the oxidation reaction.

The complete active space self-consistent field (CASSCF) and restricted active space self-consistent field (RASSCF) methods^{55,56} and their extension through second-order perturbation theory (CASPT2⁵⁷ and RASPT2,⁵⁸ respectively) were employed for the cluster model calculations. In CASSCF, a multiconfigurational wave function is considered by performing a full configuration interaction (full CI) expansion within a limited orbital space. The notation CAS(n,m) is followed for the description of the active spaces, where n is the number of electrons and m the number of orbitals considered for the CI expansion. In the RASSCF method, three individual orbital subspaces are considered, RAS1, RAS2, and RAS3, and a CI expansion is formed by applying restrictions between the three RAS spaces. A full CI is performed in RAS2, similarly to CAS. RAS1 includes doubly occupied orbitals, and a maximum number of holes is allowed; RAS3 includes unoccupied orbitals, and a maximum number of particles (electrons) is allowed. Since only two holes and two particles are considered in this work for RAS1 and RAS3, respectively, a simple RAS(n,m) notation is used, similar to the CAS notation.

Apart from state-specific (SS) CAS/RAS calculations, multiple states of the same spin multiplicity were examined by the state-averaged (SA) CASSCF/RASSCF methods. For the SA variants, the multistate (MS)-CASPT2/RASPT2 extensions were applied to include dynamical electron correlation. The nomenclature followed is SA(n)-CASSCF/RASSCF and MS(n)-CASPT2/RASPT2, where n is the number of the optimized states.

For the choice of a proper active space, a formal charge of 2+ for each Cu was initially assumed, corresponding to a formal $3d^9$ electronic configuration per Cu. Based on this assumption, and in order to have a total charge of 2+ for the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ core, one O^{2-} and two $\text{O}^{\bullet-}$ atoms are expected. The energetically most stable states were first explored using a minimal active space of size (5,5), namely, the bonding (σ)/antibonding (σ^*) combinations of the $3d_{x^2-y^2}$ orbitals of two Cu atoms with the 2p of two neighboring O atoms, respectively, and a singly occupied $3d_{x^2-y^2}$ atomic orbital of the third Cu atom.

After a systematic examination of active spaces of various sizes (see Supporting Information), we concluded that a CAS(11,11) and a RAS(19,21) are adequate to describe the low-lying $S = 1/2$, $3/2$, and $5/2$ states of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ model system. CAS(11,11) includes the five key orbitals of CAS(5,5), the three 2p orbitals of O(3), and the three 3p orbitals of O(3). RAS(19,21) includes all 2p/3p orbitals of the three O atoms and the $3d_{x^2-y^2}$ orbitals of the three Cu atoms. The correlated orbital space of RASSCF(19,21) was organized as such: RAS2 includes the (5,5) minimal space described above. RAS1 includes the remaining seven 2p orbitals of the three O atoms, which do not participate in the bonding/antibonding combinations with the $3d_{x^2-y^2}$ orbitals of Cu atoms. RAS3 includes the nine unoccupied 3p orbitals of the three O atoms. All configurations that involve up to two electron excitations from RAS1 to RAS2 and RAS3, and from RAS1 and RAS2 to RAS3 are considered in the CI expansion. It was found that the doubly occupied 3d atomic orbitals of the three Cu atoms do not participate in the bonding between the Cu and the ($\mu\text{-O}$) atoms. Therefore, they were not further considered in this study.

The triple- ζ quality atomic natural orbital basis set^{59,60} (ANO-RCC-VTZP) was used for the three Cu and three O atoms of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ core, the double- ζ basis set analogue (ANO-RCC-VDZP) for the six O atoms of the zeolite framework coordinated to the three Cu atoms, and a minimal basis (ANO-RCC-MB) for all remaining atoms (Si, Al, O, and H). In all calculations, scalar relativistic effects were included using the all-electron relativistic basis sets of ANO-RCC type and a second-order Douglas–Kroll–Hess Hamiltonian.^{61,62} A shifted zeroth-order Hamiltonian⁶³ (IPEA shift) with the default value of 0.25 au and an imaginary shift⁶⁴ of 0.2 au were applied to all CASPT2/RASPT2 calculations. The two-electron integral evaluation was simplified by using the Cholesky decomposition technique.⁶⁵ All CAS/RAS calculations were performed using the MOLCAS 8.0 program package.⁶⁶

RESULTS AND DISCUSSION

Electronic Structure of $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$. Table 1 reports the energy differences among the three lowest states, namely, the doublet ($S = 1/2$), quartet ($S = 3/2$), and sextet ($S = 5/2$)

states, calculated at the CASSCF/CASPT2(11,11) and RASSCF/RASPT2(19,21) levels of theory. The multiconfigurational SCF methods (CASSCF and RASSCF) are used to generate a multideterminant zeroth-order wave function and provide only qualitative energy differences between different spin states since they lack dynamical correlation. This type of correlation energy is recovered by second-order perturbation theory (PT2). Therefore, in this section we analyze the nature of the CASSCF/RASSCF wave function, but we discuss only the energy differences obtained at the CASPT2/RASPT2 levels.

The doublet state is the lowest in energy, followed by the quartet, 20.7 kJ/mol higher, and by the sextet, 78.1 kJ/mol higher, at the CASPT2(11,11) level of theory. These energy differences are in good agreement with the values obtained from PBE (21.1 and 67.8 kJ/mol, respectively) and BLYP^{67,68} functionals (22.7 and 71.2 kJ/mol, respectively). The results obtained with different popular density functionals are summarized in the [Supporting Information](#). The RASPT2-(19,21) doublet/quartet energy difference (22.1 kJ/mol) is in excellent agreement with the CASPT2(11,11) result (20.7 kJ/mol), while the doublet/sextet difference is predicted to be slightly higher by RASPT2(19,21), 108.6 kJ/mol, than by CASPT2(11,11), 78.1 kJ/mol.

The five molecular orbitals (MOs) of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster with occupation numbers between 0.04 and 1.96 are shown in [Figure 2](#), as they were obtained from the

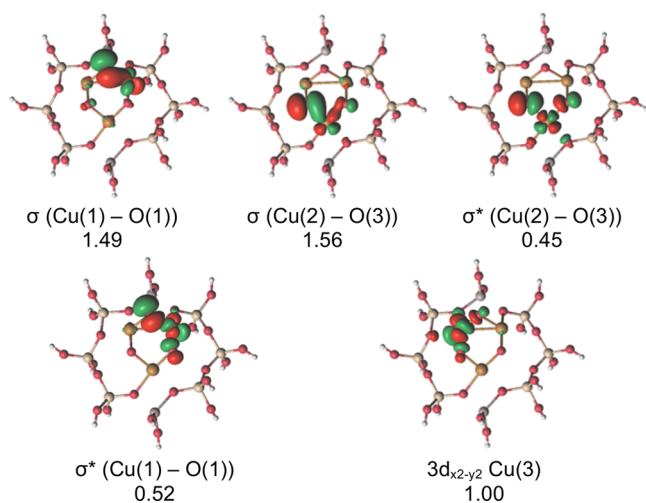


Figure 2. Five most relevant MOs of the doublet state of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster, as calculated from the RASSCF(19,21) level of theory. A description and their corresponding occupation numbers are given below each MO. The atom labels are as in [Figure 1c](#).

RASSCF(19,21) calculation. While they are singly occupied in the sextet state, they are responsible for the multiconfigurational character of the doublet and quartet states. RASSCF(19,21) shows that Cu(3) has a pure d^9 electronic configuration, while Cu(1) and Cu(2) form bonding (σ)/antibonding (σ^*) combinations with the 2p orbitals of O(1) and O(3), respectively. The O(1) and O(3) atoms have a partially occupied 2p atomic orbital and some $\text{O}^{\bullet-}$ (oxyl) character. On the other hand, O(2) has the three 2p orbitals doubly occupied (O^{2-} character). Similar orbitals were obtained in the CASSCF(11,11) calculation (see [Supporting Information](#)).

The first five low-lying doublet states were examined using SA(5)-RASSCF(19,21)/MS(5)-RASPT2(19,21) level of theory

(for the choice of these five states, see [Supporting Information](#)). For the first three states, one MO is characterized by a single 3d orbital of only one Cu center: Cu(3) for state 1, Cu(1) for state 2, and Cu(2) for state 3, respectively. The other four MOs are bonding (σ)/antibonding (σ^*) combinations between the remaining two Cu centers and the O(1) and O(3) atoms (Cu(1)-O(1)/Cu(2)-O(3) for state 1, Cu(3)-O(1)/Cu(2)-O(3) for state 2, and Cu(1)-O(1)/Cu(3)-O(3) for state 3). The corresponding five MOs of states 4 and 5 are combinations of one 3d orbital of the three Cu centers and a 2p atomic orbital of O(1) and O(3). The occupation numbers of these MOs are close to one.

In the three most stable doublet states, all three Cu centers are equivalent and have an overall $3d^9$ atomic configuration characteristic for the +2 (Cu(II)) oxidation state. The O(2) has a closed-shell ($2p^6$) electronic configuration. In other words, O(2) has an O^{2-} character, whereas the O(1) and O(3) centers “share” an electron and a neighboring Cu resulting in σ/σ^* MO combinations. We can assign to these O atoms a $2p^5$ open-shell electronic configuration corresponding to an oxyl ($\text{O}^{\bullet-}$) radical. A similar interpretation can be given by considering the lowest two states (states 1 and 2) as near-degenerate states, since they differ by less than 18 kJ/mol (0.2 eV, [Table S2](#)). The configuration of the two states differs by one σ/σ^* pair (Cu(1)-O(1) for state 1, Cu(3)-O(1) for state 2). A transition between these two states would involve the spontaneous formation of O^{2-} for O(1) and the oxidation of either Cu(1) or Cu(3) to Cu^{3+} , as shown in [Figure 3](#). On the

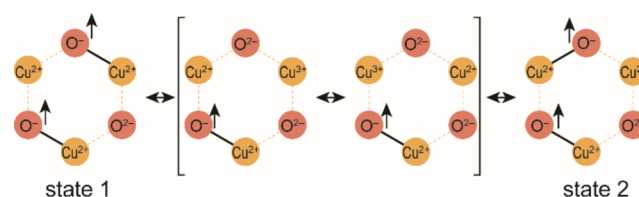


Figure 3. Proposed resonance mechanism between states 1 and 2. The arrows represent the radical character of the $\text{O}^{\bullet-}$ atoms.

other hand, Cu(2) is accessible in this “resonance” mechanism only through O(3) since O(2) was found to have constantly an oxo character (O^{2-}). Therefore, state 3 is higher in energy (35.1 kJ/mol or 0.4 eV at the MS(5)-RASPT2 level). The nature of the oxygen centers is crucial for understanding the fundamental factors that control the reactivity of the intra-zeolite Cu sites. Given the importance of the electrophilicity of the reactive oxygen centers for C-H activation,^{69–71} one expects that these two O atoms would provide low-energy paths for C-H activation. However, if the reactivity is dominated by the basicity of the proton-accepting site, the C-H cleavage should be more facile over the electron-rich closed-shell O(2) center.

The above analysis is based only on one conformer of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster with an asymmetric geometry, in which only Cu(3) is closer to the walls of the MOR zeolite: Cu(3)-O(MOR-1) and Cu(3)-O(MOR-2) bond distances are 2.105 and 2.044 Å, respectively ([Table 2](#)). (The labels MOR-1 and MOR-2 correspond to the two different oxygen atoms of the zeolite coordinated with the Cu centers.) Similarly, the out-of-plane Cu(2) points to the zeolitic framework and is located 1.955 and 2.016 Å from the two oxygen atoms of the framework. Cu(1) lies significantly further from the zeolite since the bond distance between Cu(1) and one of the oxygen

Table 2. Bond Distances (in Å) between the Three Cu Centers and the Oxygen Atoms of the Mordenite Zeolite from the Structure Optimized by Periodic DFT (Doublet Spin State) and Used for the Multiconfigurational Calculations

	Cu(1)	Cu(2)	Cu(3)
O(MOR-1)	1.987	1.955	2.105
O(MOR-2)	2.628	2.016	2.044

atoms of the framework is 2.628 Å, about 0.5 Å longer than the equivalent distance for Cu(3). On the other hand, we expect an equal probability of the cluster conformers that are located closer to MOR, from either the Cu(1) or the Cu(3) site due to the symmetric nature of the MOR pore and the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ ring cluster. In other words, there should be an isostructural and isoenergetic conformer of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ site where Cu(1) is closer to the walls of the zeolite, while Cu(3) is further from the oxygen atoms of the zeolite (about 2.6 Å). For this conformer the electronic configurations of O(2) and O(3) are interchanged, with O(2) having a radical, oxyl-character, and O(3) a closed-shell O^{2-} character. The two in-plane oxygens (O(2) and O(3)) are thus equivalent, and they resonate between $\text{O}^{\bullet-}$ and O^{2-} . Therefore, their reactivity toward homolytic C–H cleavage should be at most comparable but most likely less significant than O(1), which provides the optimal accessibility for the reacting alkane molecules.

Active Site Anisotropy for Homolytic C–H Cleavage.

The above considerations suggest that the oxidation of methane is more facile over the more electronegative O(1) center which forms a stronger O–H bond. In order to verify this hypothesis, we calculated the C–H activation barrier (electronic energies) over three different oxygen sites of the Cu trimer stabilized in the MOR model by means of periodic DFT calculations (Figure 4) to capture all effects from the full pore of the zeolite. An activation barrier of 37 kJ/mol was obtained for the hydrogen transfer from methane to O(1), resulting in the formation of a hydroxo species and a methyl radical confined in the main channel of mordenite. The same process is energetically less favorable for O(2) and O(3); the corresponding barriers from periodic DFT are 78 and 74 kJ/mol, respectively. These results indicate that O(1) is catalytically more active for methane C–H bond activation than the other two O sites, which is consistent with our conclusions drawn by multiconfigurational analysis. Notice that the anisotropy will be

substantially altered after the activation of the first methane molecule and consumption of the O(1) site as this would result in the formation of a partially reduced $[\text{Cu}_3\text{O}_2]^{2+}$ with electronic structure substantially different from the original active site.

We propose that this reactivity difference can only manifest itself within a continuous catalytic process. The catalyst will be exposed to methane feed at elevated temperature commonly employed within the sequential process schemes^{21,72} with separate methane activation/zeolite regeneration. This exposure will level the intrinsic reactivity differences of the various O sites within the trinuclear clusters and other potentially reactive intrazeolite Cu complexes.

Our calculations provide a convincing evidence of the importance of the radical anion character of the reactive O sites in the zeolite-stabilized extraframework Cu-oxo clusters for the activation of the C–H bonds in methane. Although we have explored only one type of active sites,⁷³ which can be selectively generated in MOR zeolite,²¹ the theoretical insights emerging from this study allow rationalizing reactivity trends predicted for other types of Cu-oxo complexes. Moreover, they provide guidelines for active site design to enable a low-temperature catalytic process for methane oxofunctionalization. Previous DFT studies^{37,39} showed that the activity of Cu complexes in C–H bond activation of methane follows the sequence $\text{CuOH}^+ < [\text{CuO}_2\text{Cu}]^{2+} < [\text{CuOCu}]^{2+} < [\text{Cu}_3\text{O}_3]^{2+} \approx [\text{Cu}_4\text{O}_x]^{2+} \approx [\text{Cu}_5\text{O}_x]^{2+}$. This trend confirms our finding that the initial reactivity increase with larger nuclearity is associated with the number of spin domains in the cluster. This is directly connected to the radical character of the ligand oxygens. The effect of spin delocalization and, accordingly, reactivity enhancement level off for clusters containing more than three copper sites. We expect that further increase of the cluster size would result in a complete localization of the spin density on the Cu centers, as observed in bulk CuO.

CONCLUSIONS AND OUTLOOK

The electronic structure of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ complex deposited inside the pores of the MOR zeolite was studied by means of multiconfigurational wave function methods. The $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ site can activate strong C–H bonds and oxidize methane to methanol, a catalytic process with great industrial interest due to the large deposits of methane. Understanding the underlying details that drive this reaction will allow us to

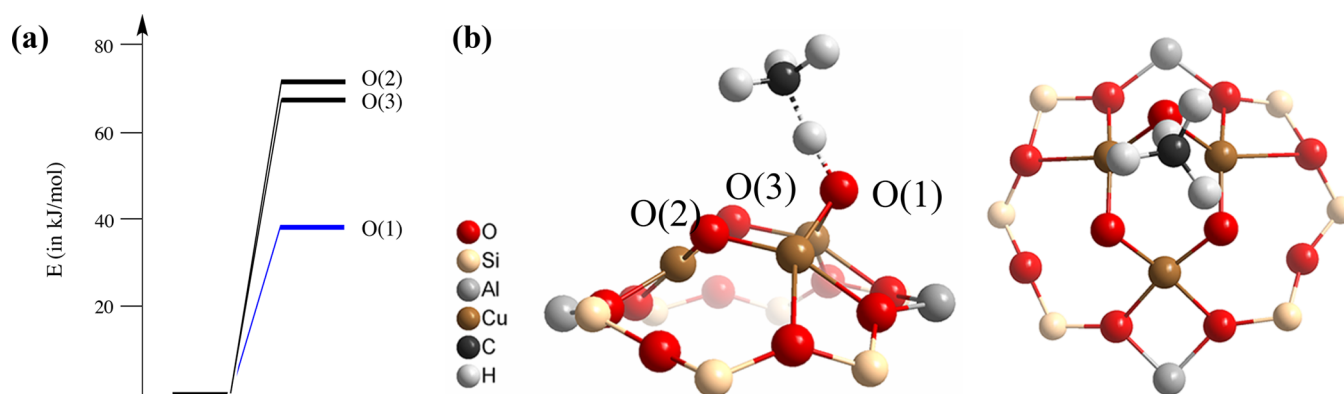


Figure 4. (a) Relative electronic energies of the C–H activation barriers (in kJ/mol, $S = 1/2$) from the three oxygen atoms of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ cluster confined in MOR zeolite. (b) Optimized local geometry of the transition state of the C–H activation for O(1).

optimize catalytic systems with enhanced performance for the functionalization of C–H bonds.

The analysis of the low-lying energy states confirms the presence of mixed valence Cu(II)/Cu(III) oxidation states of the three Cu cations. For the geometrical conformer that was studied here, Cu(2) has a more pure Cu(II) character, while Cu(1) and Cu(3) have a mixed Cu(II/III) composition. The two oxygen atoms with an oxyl character, O(1) and O(3), are expected to activate the C–H bond of methane. Despite the fact that two oxygen atoms have both radical character, O(1) is expected to be catalytically more active than O(3) because of the lower symmetry of the structural conformer used in our high-level calculations. Cu(2) and Cu(3) are closer to the zeolitic walls than Cu(1), an effect that perturbs the electronic structure of the cluster. Due to the higher symmetry of the $[\text{Cu}_3(\mu\text{-O})_3]^{2+}$ site, the in-plane O(2) and O(3) are expected to be equivalent, and only O(1) is expected to be more catalytically active. These conclusions were verified by periodic DFT. The optimized transition state geometries adopt the proper symmetry and equalize the two in-plane O atoms. This suggestion was reflected on the calculated C–H activation barriers: the out-of-plane O(1) was found about 30 kJ/mol lower than the respective barriers from the in-plane oxygen atoms. These conclusions can be further examined by EPR spectroscopy on the activated catalyst. This technique has been already successfully applied in mono-Cu sites deposited on zeolites^{46,74,75} and can provide further insight on the electronic structure of the copper cluster. Future experimental examination of the reaction with operando spectroscopy can provide further insight into the evolution of the catalyst and help us to accurately model the next reaction steps and better understand the activation of the second methane molecule.

■ ASSOCIATED CONTENT

■ Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jpcc.7b08714.

Evaluation of the computational details, comparison of different density functionals, Cartesian coordinates, and cif files from periodic DFT (PDF)

■ AUTHOR INFORMATION

Corresponding Authors

*E-mail: kvogiatz@utk.edu (K. D. V.).

*E-mail: e.a.pidko@tudelft.nl (E. A. P.).

ORCID

Konstantinos D. Vogiatzis: 0000-0002-7439-3850

Guanna Li: 0000-0003-3031-8119

Emiel J. M. Hensen: 0000-0002-9754-2417

Laura Gagliardi: 0000-0001-5227-1396

Evgeny A. Pidko: 0000-0001-9242-9901

Present Address

[†](E.A.P.) Inorganic Systems Engineering group, Department of Chemical Engineering, Faculty of Applied Sciences, Delft University of Technology, Van der Maasweg 9, 2629 HZ Delft, The Netherlands

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

K.D.V. would like to acknowledge the University of Tennessee for financial support of this work (start-up grant). G.L. acknowledges financial support from NWO for her personal VENI grant (no. 016.Veni.172.034) and NWO-SurfSARA for providing access to supercomputer resources. E.J.M.H. and E.A.P. acknowledge MCEC, a Gravitation programme of The Netherlands Organization for Scientific Research (NWO) funded by the Ministry of Education, Culture and Science of the government of The Netherlands. E.A.P. thanks the Ministry of Education and Science of Russian Federation (GosZadanie no. 11.1706.2017PP). K.D.V. and L.G. thank the Inorganometallic Catalyst Design Center, an EFRC funded by the DOE, Office of Basic Energy Sciences (DE-SC0012702). The authors acknowledge the Minnesota Supercomputing Institute (MSI) at the University of Minnesota and the Newton High-Performance Computing program at the University of Tennessee for providing resources that contributed to the research results reported within this paper.

■ REFERENCES

- (1) Saha, D.; Grappe, H. A.; Chakraborty, A.; Orkoulas, G. Postextraction Separation, On-Board Storage, and Catalytic Conversion of Methane in Natural Gas: A Review. *Chem. Rev.* **2016**, *116*, 11436–11499.
- (2) Lin, R.; Amrute, A. P.; Pérez-Ramírez, J. Halogen-Mediated Conversion of Hydrocarbons to Commodities. *Chem. Rev.* **2017**, *117*, 4182–4247.
- (3) Olivos-Suarez, A. I.; Szécsényi, À.; Hensen, E. J. M.; Ruiz-Martinez, J.; Pidko, E. A.; Gascon, J. Strategies for the Direct Catalytic Valorization of Methane Using Heterogeneous Catalysis: Challenges and Opportunities. *ACS Catal.* **2016**, *6*, 2965–2981.
- (4) Snyder, B. E. R.; Vanelderen, P.; Bols, M. L.; Hallaert, S. D.; Böttger, L. H.; Ungur, L.; Pierloot, K.; Schoonheydt, R. A.; Sels, B. F.; Solomon, E. I. The Active Site of Low-Temperature Methane Hydroxylation in Iron-Containing Zeolites. *Nature* **2016**, *536*, 317–321.
- (5) Olah, G. A. Beyond Oil and Gas: The Methanol Economy. *Angew. Chem., Int. Ed.* **2005**, *44*, 2636–2639.
- (6) Guo, Z.; Liu, B.; Zhang, Q.; Deng, W.; Wang, Y.; Yang, Y. Recent Advances in Heterogeneous Selective Oxidation Catalysis for Sustainable Chemistry. *Chem. Soc. Rev.* **2014**, *43*, 3480–3524.
- (7) Solomon, E. I.; Heppner, D. E.; Johnston, E. M.; Ginsbach, J. W.; Cirera, J.; Qayyum, M.; Kieber-Emmons, M. T.; Kjaergaard, C. H.; Hadt, R. G.; Tian, L. Copper Active Sites in Biology. *Chem. Rev.* **2014**, *114*, 3659–3853.
- (8) Rosenzweig, A. C.; Frederick, C. A.; Lippard, S. J.; Nordlund, P. Crystal Structure of a Bacterial Non-Haem Iron Hydroxylase that Catalyses the Biological Oxidation of Methane. *Nature* **1993**, *366*, 537–543.
- (9) Lieberman, R. L.; Rosenzweig, A. C. Crystal Structure of a Membrane-Bound Metalloenzyme that Catalyses the Biological Oxidation of Methane. *Nature* **2005**, *434*, 177–182.
- (10) Balasubramanian, R.; Smith, S. M.; Rawat, S.; Yatsunyk, L. A.; Stemmler, T. L.; Rosenzweig, A. C. Oxidation of Methane by a Biological Dicomplex. *Nature* **2010**, *465*, 115–119.
- (11) Himes, R. A.; Barnese, K.; Karlin, K. D. One is Lonely and Three is a Crowd: Two Coppers Are for Methane Oxidation. *Angew. Chem., Int. Ed.* **2010**, *49*, 6714–6716.
- (12) Chan, S. I.; Chen, K. H.-C.; Yu, S. S.-F.; Chen, C.-L.; Kuo, S. S.-J. Toward Delineating the Structure and Function of the Particulate Methane Monooxygenase from Methanotrophic Bacteria. *Biochemistry* **2004**, *43*, 4421–4430.
- (13) Chan, S. I.; Wang, V. C.-C.; Lai, J. C.-H.; Yu, S. S.-F.; Chen, P. P.-Y.; Chen, K. H.-C.; Chen, C.-L.; Chan, M. K. Redox Potentiometry Studies of Particulate Methane Monooxygenase: Support for a

Trinuclear Copper Cluster Active Site. *Angew. Chem., Int. Ed.* **2007**, *46*, 1992–1994.

(14) Nagababu, P.; Maji, S.; Kumar, M. P.; Chen, P. P.-Y.; Yu, S. S.-F.; Chan, S. I. Efficient Room-Temperature Oxidation of Hydrocarbons Mediated by Tricopper Cluster Complexes with Different Ligands. *Adv. Synth. Catal.* **2012**, *354*, 3275–3282.

(15) Smeets, P. J.; Groothaert, M. H.; Schoonheydt, R. A. Cu Based Zeolites: A UV–vis Study of the Active Site in the Selective Methane Oxidation at Low Temperatures. *Catal. Today* **2005**, *110*, 303–309.

(16) Groothaert, M. H.; Smeets, P. J.; Sels, B. F.; Jacobs, P. A.; Schoonheydt, R. A. Selective Oxidation of Methane by the Bis(μ -oxo)dicopper Core Stabilized on ZSM-5 and Mordenite Zeolites. *J. Am. Chem. Soc.* **2005**, *127*, 1394–1395.

(17) Vanelderen, P.; Hadt, R. G.; Smeets, P. J.; Solomon, E. I.; Schoonheydt, R. A.; Sels, B. F. Cu-ZSM-5: A Biomimetic Inorganic Model for Methane Oxidation. *J. Catal.* **2011**, *284*, 157–164.

(18) Alayon, E. M.; Nachtegaal, M.; Ranocchiari, M.; van Bokhoven, J. A. Catalytic Conversion of Methane to Methanol over Cu–Mordenite. *Chem. Commun.* **2012**, *48*, 404–406.

(19) Alayon, E. M. C.; Nachtegaal, M.; Kleyenov, E.; van Bokhoven, J. A. Determination of the Electronic and Geometric Structure of Cu Sites During Methane Conversion over Cu-MOR with X-ray Absorption Spectroscopy. *Microporous Mesoporous Mater.* **2013**, *166*, 131–136.

(20) Vanelderen, P.; Vancauwenbergh, J.; Tsai, M.-L.; Hadt, R. G.; Solomon, E. I.; Schoonheydt, R. A.; Sels, B. F. Spectroscopy and Redox Chemistry of Copper in Mordenite. *ChemPhysChem* **2014**, *15*, 91–99.

(21) Grundner, S.; Markovits, M. A. C.; Li, G.; Tromp, M.; Pidko, E. A.; Hensen, E. J. M.; Jentys, A.; Sanchez-Sanchez, M.; Lercher, J. A. Single-Site Trinuclear Copper Oxygen Clusters in Mordenite for Selective Conversion of Methane to Methanol. *Nat. Commun.* **2015**, *6*, 7546.

(22) Narsimhan, K.; Iyoki, K.; Dinh, K.; Román-Leshkov, Y. Catalytic Oxidation of Methane into Methanol over Copper-Exchanged Zeolites with Oxygen at Low Temperature. *ACS Cent. Sci.* **2016**, *2*, 424–429.

(23) Le, H. V.; Parishan, S.; Sagaltchik, A.; Göbel, C.; Schlesiger, C.; Malzer, W.; Trunschke, A.; Schomäcker, R.; Thomas, A. Solid-State Ion-Exchanged Cu/Mordenite Catalysts for the Direct Conversion of Methane to Methanol. *ACS Catal.* **2017**, *7*, 1403–1412.

(24) Tomkins, P.; Ranocchiari, M.; van Bokhoven, J. A. Direct Conversion of Methane to Methanol under Mild Conditions over Cu-Zeolites and beyond. *Acc. Chem. Res.* **2017**, *50*, 418–425.

(25) Latimer, A. A.; Kulkarni, A. R.; Aljama, H.; Montoya, J. H.; Yoo, J. S.; Tsai, C.; Abild-Pedersen, F.; Studt, F.; Nørskov, J. K. Understanding Trends in C–H Bond Activation in Heterogeneous Catalysis. *Nat. Mater.* **2017**, *16*, 225–229.

(26) Groothaert, M. H.; van Bokhoven, J. A.; Battiston, A. A.; Weckhuysen, B. M.; Schoonheydt, R. A. Bis(μ -oxo)dicopper in Cu-ZSM-5 and Its Role in the Decomposition of NO: A Combined in Situ XAFS, UV–Vis–Near-IR, and Kinetic Study. *J. Am. Chem. Soc.* **2003**, *125*, 7629–7640.

(27) Groothaert, M. H.; Pierloot, K.; Delabie, A.; Schoonheydt, R. A. Identification of Cu(II) Coordination Structures in Cu-ZSM-5, Based on a DFT/ab initio Assignment of the EPR Spectra. *Phys. Chem. Chem. Phys.* **2003**, *5*, 2135–2144.

(28) Beznis, N. V.; Weckhuysen, B. M.; Bitter, J. H. Cu-ZSM-5 Zeolites for the Formation of Methanol from Methane and Oxygen: Probing the Active Sites and Spectator Species. *Catal. Lett.* **2010**, *128*, 14–22.

(29) Vanelderen, P.; Vancauwenbergh, J.; Sels, B. F.; Schoonheydt, R. A. Coordination Chemistry and Reactivity of Copper in Zeolites. *Coord. Chem. Rev.* **2013**, *257*, 483–494.

(30) Vanelderen, P.; Snyder, B. E. R.; Tsai, M.-L.; Hadt, R. G.; Vancauwenbergh, J.; Coussens, O.; Schoonheydt, R. A.; Sels, B. F.; Solomon, E. I. Spectroscopic Definition of the Copper Active Sites in Mordenite: Selective Methane Oxidation. *J. Am. Chem. Soc.* **2015**, *137*, 6383–6392.

(31) Itadani, A.; Sogawa, Y.; Oda, A.; Ohkubo, T.; Yumura, T.; Kobayashi, H.; Sato, M.; Kuroda, Y. Possibility of Copper-Ion-Exchanged MFI-Type Zeolite as C–H Bond Activation Material for Propane and the Driving Force for Activation. *J. Phys. Chem. C* **2015**, *119*, 21483–21496.

(32) Itadani, A.; Yumura, T.; Ohkubo, T.; Kobayashi, H.; Kuroda, Y. Existence of Dual Species Composed of Cu⁺ in CuMFI Being Bridged by C₂H₂. *Phys. Chem. Chem. Phys.* **2010**, *12*, 6455–6465.

(33) Alayon, E. M. C.; Nachtegaal, M.; Bodi, A.; van Bokhoven, J. A. Reaction Conditions of Methane-to-Methanol Conversion Affect the Structure of Active Copper Sites. *ACS Catal.* **2014**, *4*, 16–22.

(34) Hammond, C.; Forde, M. M.; Rahim, M. H. A.; Thetford, A.; He, Q.; Jenkins, R. L.; Dimitratos, N.; Lopez-Sanchez, J. A.; Dummer, N. F.; Murphy, D. M.; et al. Direct Catalytic Conversion of Methane to Methanol in an Aqueous Medium by using Copper-Promoted Fe-ZSM-5. *Angew. Chem., Int. Ed.* **2012**, *51*, 5129–5133.

(35) Woertink, J. S.; Smeets, P. J.; Groothaert, M. H.; Vance, M. A.; Sels, B. F.; Schoonheydt, R. A.; Solomon, E. I. A [Cu₂O]²⁺ Core in Cu-ZSM-5, the Active Site in the Oxidation of Methane to Methanol. *Proc. Natl. Acad. Sci. U. S. A.* **2009**, *106*, 18908–18913.

(36) Ipek, B.; Wulfers, M. J.; Kim, H.; Göltl, F.; Hermans, I.; Smith, J. P.; Booksh, K. S.; Brown, C. M.; Lobo, R. F. Formation of [Cu₂O]²⁺ and [Cu₂O]²⁺ toward C–H Bond Activation in Cu-SSZ-13 and Cu-SSZ-39. *ACS Catal.* **2017**, *7*, 4291–4303.

(37) Kulkarni, A. R.; Zhao, Z.-J.; Siahrostami, S.; Nørskov, J. K.; Studt, F. Monocopper Active Site for Partial Methane Oxidation in Cu-Exchanged 8MR Zeolites. *ACS Catal.* **2016**, *6*, 6531–6536.

(38) Mahyuddin, M. H.; Staykov, A.; Shiota, Y.; Miyaniishi, M.; Yoshizawa, K. Roles of Zeolite Confinement and Cu–O–Cu Angle on the Direct Conversion of Methane to Methanol by [Cu₂(μ -O)]²⁺-Exchanged AEI, CHA, AFX, and MFI Zeolites. *ACS Catal.* **2017**, *7*, 3741–3751.

(39) Palagin, D.; Knorpp, A. J.; Pinar, A. B.; Ranocchiari, M.; van Bokhoven, J. A. Assessing the Relative Stability of Copper Oxide Clusters as Active Sites of a CuMOR Zeolite for Methane to Methanol Conversion: Size Matters? *Nanoscale* **2017**, *9*, 1144–1153.

(40) Grundner, S.; Luo, W.; Sanchez-Sanchez, M.; Lercher, J. A. Synthesis of Single-Site Copper Catalysts for Methane Partial Oxidation. *Chem. Commun.* **2016**, *52*, 2553–2556.

(41) Kim, Y.; Kim, T. Y.; Lee, H.; Yi, J. Distinct Activation of Cu-MOR for Direct Oxidation of Methane to Methanol. *Chem. Commun.* **2017**, *53*, 4116–4119.

(42) Chen, P. P.-Y.; Yang, R. B.-G.; Lee, J. C.-M.; Chan, S. I. Facile O-atom Insertion into C–C and C–H Bonds by a Trinuclear Copper Complex Designed to Harness a Singlet Oxene. *Proc. Natl. Acad. Sci. U. S. A.* **2007**, *104*, 14570–14575.

(43) Chen, P. P.-Y.; Nagababu, P.; Yu, S. S.-F.; Chan, S. I. Development of the Tricopper Cluster as a Catalyst for the Efficient Conversion of Methane into MeOH. *ChemCatChem* **2014**, *6*, 429–437.

(44) Li, G.; Vassilev, P.; Sanchez-Sanchez, M.; Lercher, J. A.; Hensen, E. J. M.; Pidko, E. A. Stability and Reactivity of Copper Oxo-Clusters in ZSM-5 Zeolite for Selective Methane Oxidation to Methanol. *J. Catal.* **2016**, *388*, 305–312.

(45) Ghosh, A.; Taylor, P. R. High-Level ab initio Calculations on the Energetics of Low-Lying Spin States of Biologically Relevant Transition Metal Complexes: A First Progress Report. *Curr. Opin. Chem. Biol.* **2003**, *7*, 113–124.

(46) Pierloot, K.; Delabie, A.; Groothaert, M. H.; Schoonheydt, R. A. A Reinterpretation of the EPR Spectra of Cu(II) in Zeolites A, Y and ZK4, Based on ab initio Cluster Model Calculations. *Phys. Chem. Chem. Phys.* **2001**, *3*, 2174–2183.

(47) Göltl, F.; Michel, C.; Andrikopoulos, P. C.; Love, A. M.; Hafner, J.; Hermans, I.; Sautet, P. Computationally Exploring Confinement Effects in the Methane-to-Methanol Conversion Over Iron-Oxo Centers in Zeolites. *ACS Catal.* **2016**, *6*, 8404–8409.

(48) Kresse, G.; Hafner, J. Ab initio Molecular Dynamics for Open-Shell Transition Metals. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1993**, *48*, 13115–13118.

- (49) Perdew, J. P.; Burke, K.; Ernzerhof, M. Generalized Gradient Approximation Made Simple. *Phys. Rev. Lett.* **1996**, *77*, 3865–3868.
- (50) Blöchl, P. E. Projector Augmented-Wave Method. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1994**, *50*, 17953–17978.
- (51) van Koningsveld, H.; Jansen, J. C.; van Bekkum, H. The Monoclinic Framework Structure of Zeolite H-ZSM-5. Comparison with the Orthorhombic Framework of as-Synthesized ZSM-5. *Zeolites* **1990**, *10*, 235–242.
- (52) Pidko, E. A.; van Santen, R. A.; Hensen, E. J. M. Multinuclear Gallium-Oxide Cations in High-Silica Zeolites. *Phys. Chem. Chem. Phys.* **2009**, *11*, 2893–2902.
- (53) Pidko, E. A.; van Santen, R. A. Structure–Reactivity Relationship for Catalytic Activity of Gallium Oxide and Sulfide Clusters in Zeolite. *J. Phys. Chem. C* **2009**, *113*, 4246–4249.
- (54) Henkelman, G.; Uberuaga, B. P.; Jónsson, H. A Climbing Image Nudged Elastic Band Method for Finding Saddle Points and Minimum Energy Paths. *J. Chem. Phys.* **2000**, *113*, 9901–9904.
- (55) Olsen, J.; Roos, B. O.; Jørgensen, P.; Jensen, H. J. A. Determinant Based Configuration Interaction Algorithms for Complete and Restricted Configuration Interaction Spaces. *J. Chem. Phys.* **1988**, *89*, 2185.
- (56) Malmqvist, P. Å.; Rendell, A.; Roos, B. O. The Restricted Active Space Self-Consistent-Field Method, Implemented with a Split Graph Unitary Group Approach. *J. Phys. Chem.* **1990**, *94*, 5477–5482.
- (57) Andersson, K.; Malmqvist, P.-Å.; Roos, B. O. Second-Order Perturbation Theory with a Complete Active Space Self-Consistent Field Reference Function. *J. Chem. Phys.* **1992**, *96*, 1218.
- (58) Malmqvist, P. Å.; Pierloot, K.; Shahi, A. R. M.; Cramer, C. J.; Gagliardi, L. The Restricted Active Space Followed by Second-Order Perturbation Theory Method: Theory and Application to the Study of CuO₂ and Cu₂O₂ systems. *J. Chem. Phys.* **2008**, *128*, 204109.
- (59) Roos, B. O.; Lindh, R.; Malmqvist, P.-Å.; Veryazov, V.; Widmark, P.-O. Main Group Atoms and Dimers Studied with a New Relativistic ANO Basis Set. *J. Phys. Chem. A* **2004**, *108*, 2851–2858.
- (60) Roos, B. O.; Lindh, R.; Malmqvist, P.-Å.; Veryazov, V.; Widmark, P.-O. New Relativistic ANO Basis Sets for Transition Metal Atoms. *J. Phys. Chem. A* **2005**, *109*, 6575–6579.
- (61) Douglas, M.; Kroll, N. M. Quantum Electrodynamical Corrections to the Fine Structure of Helium. *Ann. Phys.* **1974**, *82*, 89–155.
- (62) Hess, B. A. Relativistic Electronic-Structure Calculations Employing a Two-Component No-Pair Formalism with External-Field Projection Operators. *Phys. Rev. A: At., Mol., Opt. Phys.* **1986**, *33*, 3742.
- (63) Ghigo, G.; Roos, B. O.; Malmqvist, P.-Å. A Modified Definition of the Zeroth-Order Hamiltonian in Multiconfigurational Perturbation Theory (CASPT2). *Chem. Phys. Lett.* **2004**, *396*, 142–149.
- (64) Forsberg, N.; Malmqvist, P. Å. Multiconfiguration Perturbation Theory with Imaginary Level Shift. *Chem. Phys. Lett.* **1997**, *274*, 196–204.
- (65) Aquilante, F.; Lindh, R.; Pedersen, T. B. Unbiased Auxiliary Basis Sets for Accurate Two-Electron Integral Approximations. *J. Chem. Phys.* **2007**, *127*, 114107.
- (66) Aquilante, F.; Autschbach, J.; Carlson, R. K.; Chibotaru, L. F.; Delcey, M. G.; Vico, L. D.; Galván, I. F.; Ferré, N.; Frutos, L. M.; Gagliardi, L.; et al. Molcas 8: New Capabilities for Multiconfigurational Quantum Chemical Calculations Across the Periodic Table. *J. Comput. Chem.* **2016**, *37*, 506–541.
- (67) Becke, A. D. Density-Functional Exchange-Energy Approximation with Correct Asymptotic Behavior. *Phys. Rev. A: At., Mol., Opt. Phys.* **1988**, *38*, 3098.
- (68) Lee, C.; Yang, W.; Parr, R. G. Development of the Colle-Salvetti Correlation-Energy Formula into a Functional of the Electron Density. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1988**, *37*, 785.
- (69) Rosa, A.; Ricciardi, G.; Baerends, E. J. Is [FeO]²⁺ the Active Center Also in Iron Containing Zeolites? A Density Functional Theory Study of Methane Hydroxylation Catalysis by Fe-ZSM-5 Zeolite. *Inorg. Chem.* **2010**, *49*, 3866–3880.
- (70) Kazaryan, A.; Baerends, E. J. Ligand Field Effects and the High Spin–High Reactivity Correlation in the H Abstraction by Non-Heme Iron(IV)–Oxo Complexes: A DFT Frontier Orbital Perspective. *ACS Catal.* **2015**, *5*, 1475–1488.
- (71) Verma, P.; Vogiatzis, K. D.; Planas, N.; Borycz, J.; Xiao, D. J.; Long, J. R.; Gagliardi, L.; Truhlar, D. G. Mechanism of Oxidation of Ethane to Ethanol at Iron(IV)–Oxo Sites in Magnesium-Diluted Fe₂(dobdc). *J. Am. Chem. Soc.* **2015**, *137*, 5770–5781.
- (72) Sushkevich, V. L.; Palagin, D.; Ranocchiari, M.; Bokhoven, J. A. v. Selective Anaerobic Oxidation of Methane Enables Direct Synthesis of Methanol. *Science* **2017**, *356*, 523–527.
- (73) Pidko, E. A. Toward the Balance between the Reductionist and Systems Approaches in Computational Catalysis: Model versus Method Accuracy for the Description of Catalytic Systems. *ACS Catal.* **2017**, *7*, 4230–4234.
- (74) Baute, D.; Arieli, D.; Neese, F.; Zimmermann, H.; Weckhuysen, B. M.; Goldfarb, D. Carboxylate Binding in Copper Histidine Complexes in Solution and in Zeolite Y: X- and W-band Pulsed EPR/ENDOR Combined with DFT Calculations. *J. Am. Chem. Soc.* **2004**, *126*, 11733–11745.
- (75) Moreno-González, M.; Blasco, T.; Góra-Marek, K.; Palomares, A. E.; Corma, A. Study of Propane Oxidation on Cu-Zeolite Catalysts by in-situ EPR and IR Spectroscopies. *Catal. Today* **2014**, *227*, 123–129.