



Showcasing research from Professors Tonner-Zech, Heine, and Asmis, members of the Research Training Group on Hydrogen Isotopes centred at Leipzig University, Saxony, Germany.

Direct evidence for ligand-enhanced activity of Cu(I) sites

In a combined experimental and computational effort, we study the strength and isotopologue selectivity of hydrogen adsorption on the undercoordinated copper (I) site. Exploiting gas-phase complexes as model systems, we show that the high isotopologue selectivity in dihydrogen binding results from a large difference in the adsorption zero-point energies. We discuss the role of the environment and the coordination geometry of the adsorption site in achieving a high selectivity and the ramifications for identifying and designing future materials for adsorptive dihydrogen isotopologue separation.

As featured in:



See Jiaye Jin, Ralf Tonner-Zech, Thomas Heine, Knut R. Asmis *et al.*, *Chem. Sci.*, 2024, **15**, 14635.

Cite this: *Chem. Sci.*, 2024, 15, 14635

All publication charges for this article have been paid for by the Royal Society of Chemistry

Direct evidence for ligand-enhanced activity of Cu(I) sites†

Elvira Gouatieu Dongmo, ^{‡ab} Shabnam Haque, ^{‡a} Florian Kreuter, ^a Toshiki Wulf, ^{ac} Jiaye Jin, ^{*a} Ralf Tonner-Zech, ^{*a} Thomas Heine ^{*bcd} and Knut R. Asmis ^{*a}

Little is known about the strong mediating effect of the ligand sphere and the coordination geometry on the strength and isotopologue selectivity of hydrogen adsorption on the undercoordinated copper(I) site. Here, we explore this effect using gas-phase complexes $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_n$ (with $n \leq 3$) as model systems. $\text{Cu}^+(\text{H}_2\text{O})$ attracts dihydrogen (82 kJ mol^{-1}) more strongly than bare Cu^+ (64 kJ mol^{-1}) does. Combining experimental and computational methods, we demonstrate a high isotopologue selectivity in dihydrogen binding to $\text{Cu}^+(\text{H}_2\text{O})$, which results from a large difference in the adsorption zero-point energies (2.8 kJ mol^{-1} between D_2 and H_2 , including an anharmonic contribution of 0.4 kJ mol^{-1}). We investigate its origins and the bond strengthening between Cu^+ and H_2 upon addition of a single H_2O ligand. We discuss the role of the environment and the coordination geometry of the adsorption site in achieving a high selectivity and the ramifications for identifying and designing future materials for adsorptive dihydrogen isotopologue separation.

Received 10th July 2024
Accepted 9th August 2024

DOI: 10.1039/d4sc04582c

rsc.li/chemical-science

Introduction

Porous materials containing under-coordinated Cu(I) centres with a strong tendency to adsorb dihydrogen are promising candidates for efficient H_2/D_2 separation.^{1–5} A benchmark material with such centres is the metal–organic framework (MOF) Cu(I)-MFU-4l, containing trigonal pyramidal Cu^+ sites linked by BTDD (bis-(1*H*-1,2,3-triazolo-[4,5-*b*],[4',5'-*i*])dibenzo-[1,4]-dioxin).^{2,4,5} Its high D_2/H_2 adsorption selectivity of 11 at the relatively high temperature of 100 K is due to the combination of high enthalpies of adsorption for H_2 (31 kJ mol^{-1}) and D_2 (34 kJ mol^{-1}) and the relatively high difference of these values between the two isotopologues. The latter is governed by the zero-point energy (ZPE) difference, and leads to a strong preference towards adsorption of the higher mass isotopologue.^{6–8}

The rational design of high-performance separation materials requires a detailed understanding of the binding of

dihydrogen isotopologues to metal centres, particularly Cu(I), which has shown a strong dihydrogen affinity in many cases.^{2–4} However, spectroscopic studies of dihydrogen coordinated to Cu(I) centres in bulk materials typically suffer from structural heterogeneity, complicating an in-depth understanding. This can be avoided by performing experiments on well-defined and isolated gas-phase complexes using vibrational action spectroscopy in combination with electronic structure calculations.⁹ This approach allows characterization of the geometric structure of relevant M– H_2 motifs,¹⁰ and also provides a deeper understanding of their binding nature as well as the isotopologue selectivity.

The adsorption thermodynamics of H_2 isotopologues at binding sites in porous materials are governed by the ZPE effects associated with the conversion of the three translational and two rotational degrees of freedom of free H_2 into five molecule-adsorption-site vibrations.¹¹ These low-frequency vibrational modes usually display strongly anharmonic behaviour and pronounced nuclear quantum effects,¹² raising a research interest in model complexes that show the influence of these vibrational modes on dihydrogen isotopologue separation.

A recent spectroscopic and theoretical study of the $\text{Cu}^+(\text{H}_2)_4$ complex and its isotopologues revealed a substantial red-shift of the dihydrogen stretching frequency ($\nu_{\text{HH}} = 3729 \text{ cm}^{-1}$ and $\nu_{\text{DD}} = 2678 \text{ cm}^{-1}$) upon complexation ($\nu_{\text{HH}} = 4162 \text{ cm}^{-1}$ and $\nu_{\text{DD}} = 2994 \text{ cm}^{-1}$).¹³ The exceptionally high sequential bond dissociation energy (BDE) for $\text{Cu}^+(\text{H}_2)_n \rightarrow \text{Cu}^+(\text{H}_2)_{n-1} + \text{H}_2$ with $n = 1$ of 64 kJ mol^{-1} reduces to only 21 kJ mol^{-1} with $n = 4$.^{13,14} This

^aWilhelm-Ostwald-Institut für Physikalische und Theoretische Chemie, Universität Leipzig, Linnéstr. 2, 04103 Leipzig, Germany. E-mail: jiaye.jin@uni-leipzig.de; ralf.tonner@uni-leipzig.de; knut.asmis@uni-leipzig.de

^bInstitute of Resource Ecology, Research Site Leipzig, Helmholtz-Zentrum Dresden-Rossendorf, Permoserstr. 15, 04318 Leipzig, Germany

^cFaculty of Chemistry and Food Chemistry, School of Science, TU Dresden, 01062 Dresden, Germany. E-mail: thomas.heine@tu-dresden.de

^dDepartment of Chemistry and ibs for Nanomedicine, Yonsei University, Seodaemun-gu, Seoul 120-749, Republic of Korea

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4sc04582c>

‡ These authors contributed equally.

study motivated the investigation of H_2 -affine $Cu(I)$ complexes with oxygen-donor ligands that can significantly increase the Cu^+-H_2 BDE.^{8,14,15} The aqua complex $Cu^+(H_2O)$, arguably the simplest such system, represents a useful model for an under-coordinated adsorption site in a zeolite or a MOF. The experimental BDE of the Cu^+-H_2 bond in $Cu^+(H_2O)(H_2)$ is 82 kJ mol^{-1} ,¹⁴ which is 18 kJ mol^{-1} larger than that in $Cu^+(H_2)$. Such high BDEs are desirable for adsorptive separation of dihydrogen isotopologues, both because they allow higher operating temperatures and also because they typically result in larger ZPE differences, which lead to a higher selectivity.

In this paper, we disentangle why a single H_2O ligand already markedly strengthens the $Cu(I)-H_2$ interaction. We first investigate the interaction of dihydrogen and its isotopologues with $Cu^+(H_2O)$ by ion-trap mass spectrometry at variable temperature. We then combine cryogenic ion-trap vibrational action spectroscopy^{16,17} of $Cu^+(H_2O)(H_2)_2$ isotopologues with quantum-chemical calculations to characterize their structures and vibrational properties. Finally, we make use of bond analysis methods to rationalize the strong chemical bonding of H_2 to the $Cu(I)$ centre as well as the large ZPE difference of the dihydrogen isotopologues in order to identify the relevant factors that govern the isotopologue selectivity of dihydrogen adsorption.

Results and analysis

Ion-trap mass spectrometry

We begin our investigation by characterizing the adsorption yield of H_2 and D_2 on mass-selected $Cu^+(H_2O)$ cations confined (on average) for 50 ms in an ion-trap reactor under multiple-collision conditions at ion-trap temperatures ranging from 295 K to 15 K. Under these conditions, $Cu^+(H_2O)(H_2/D_2)_n$ complexes with $n \leq 4$ can be formed *via* three-body collisions. Their relative abundances are detected using time-of-flight (TOF) mass spectrometry. Integration over the corresponding normalized TOF-MS peaks yields the relative ion yields (see Fig. S1† for all TOF mass spectra and relative ion yields).



Fig. 1 Total relative ion yield of $Cu^+(H_2O)(H_2)_{n>0}$ and $Cu^+(H_2O)(D_2)_{n>0}$ for ion trap temperatures ranging from 15 K to 200 K, obtained from the TOF mass spectra shown in Fig. S1†. The TOF mass spectra were recorded after storing $Cu^+(H_2O)$ cations on average 50 ms in a temperature-controlled ion trap filled with roughly 1 mbar of H_2 or D_2 .

Summation over all ion yields with $n > 0$ represents the total relative yields of (H_2/D_2) -containing complexes, which is plotted as a function of ion-trap temperature between 15 K and 200 K in Fig. 1. The formation of dihydrogen complexes is already observed at room temperature, confirming the relatively strong interaction between the $Cu(I)$ cation and dihydrogen.¹⁴ Ion yields larger than $\sim 4\%$ are observed for temperatures of 200 K and below. Dihydrogen adsorption increases with decreasing temperature. Moreover, at sufficiently low temperatures (< 80 K) the adsorption of additional dihydrogen molecules is possible leading to the formation of $n > 2$ complexes (see Fig. S1†). The ion yields in Fig. 1 show that D_2 -containing complexes are more likely formed than the H_2 -containing complexes over the complete temperature range, revealing a substantial isotope effect in dihydrogen adsorption at $Cu^+(H_2O)$. This isotopologue selectivity increases with decreasing temperature, reaching a maximum at 120 K. At lower temperatures the contribution of $n > 2$ complexes and eventually saturation of the dihydrogen binding sites needs to be considered.

Infrared photodissociation spectra

In order to characterize the nature of H_2/D_2 adsorption on $Cu^+(H_2O)$ in a more detailed way, we measured vibrational action spectra of the complexes using infrared



Fig. 2 Vibrational spectra of $Cu^+(D_2O)(H_2)_2$ and $Cu^+(H_2O)(D_2)_2$. (A) and (B): Experimental IRPD spectra. (C) and (D): Predicted infrared spectra considering anharmonic contributions (VPT2/MP2/def2-TZVPP) as well as a 10 cm^{-1} wide Gaussian lineshape function. Bands related to the excitation of the dihydrogen stretching fundamentals ν_{HH} (a_1) and ν_{DD} (b_1), are color-coded in blue and the corresponding combination bands in orange (see Table 1 for the band assignments).

photodissociation (IRPD) spectroscopy. For a comparison of the experimental IRPD spectra to computed IR spectra it proves helpful to measure the IRPD spectra in the linear absorption regime in the range of interest (2500–4500 cm⁻¹), which covers the excitations of the O–H/O–D and H–H/D–D fundamental transitions. To avoid spectral overlap of the ligand-specific excitations, *e.g.*, O–H/O–D stretching transitions, we focus on the Cu⁺(D₂O)(H₂) and Cu⁺(H₂O)(D₂) complexes here.

The BDE for Cu⁺(D₂O)(H₂) → Cu⁺(D₂O) + H₂ is too high to allow for efficient single-photon dissociation. Therefore the IRPD spectra of the *n* = 2 complexes, Cu⁺(D₂O)(H₂)₂ and Cu⁺(H₂O)(D₂)₂, were measured and these are compared to the corresponding calculated IR spectra in Fig. 2 (see Fig. S2† for the IRPD spectra of Cu⁺(H₂O)(H₂)₂ and Cu⁺(D₂O)(D₂)₂). The most prominent IRPD bands (see Table 1 for band positions) are assigned to the partly-resolved rovibrational transitions of the symmetric (ν^s_{OH/D}) and the antisymmetric O–H/O–D stretching fundamentals (ν^{as}_{OH/D}). The corresponding bands in the IRPD spectrum of Cu⁺(H₂O)(D₂)₂ are observed at 3628 cm⁻¹ (ν^s_{OH}) and 3700 cm⁻¹ (ν^{as}_{OH}), which is 4 cm⁻¹ higher than the previously reported values for Cu⁺(H₂O)Ar₂.¹⁸ Likewise, ν^s_{OD} and ν^s_{OD} for Cu⁺(D₂O)(H₂)₂ are at 2651 cm⁻¹ and 2751 cm⁻¹, only 2 cm⁻¹ different from the bands for Cu⁺(D₂O)Ar₂. This minor effect of dihydrogen adsorption on the O–H/O–D stretching frequencies in Cu⁺(H₂O)/D₂O is in line with H₂/D₂ binding directly to the Cu⁺ cation.

The bands related to the excitation of H–H stretching mode (ν_{HH}) are best observed in the IRPD spectrum of Cu⁺(D₂O)(H₂)₂, shown in Fig. 2A, and labelled a_{1–4}. The ν_{HH} fundamental (a₁) appears at 3568 cm⁻¹, substantially lower than the ν_{HH} fundamentals at 3729 cm⁻¹ in the previously reported IRPD spectrum of Cu⁺(H₂)₄, indicating a stronger Cu⁺–H₂ interaction in Cu⁺(D₂O)(H₂)₂ than in Cu⁺(H₂)₄.¹³ Features with excitation energies above a₁ are observed at 3693 cm⁻¹ (a₂), 3733 cm⁻¹ (a₃) and 4350 cm⁻¹ (a₄). These are tentatively assigned to combination bands (orange-coloured bands in Fig. 2B). Corresponding features are also observed in the IRPD spectrum of Cu⁺(H₂O)(D₂)₂ (Fig. 2B). They are assigned to the ν_{DD} fundamental (b₁, 2567 cm⁻¹) as well as the combination bands at 2675 cm⁻¹ (b₂), 2783 cm⁻¹ (b₃), 3125 cm⁻¹ (b₄) and 3414 cm⁻¹ (b₅).

Computational results

Structure and energetics. MP2/def2-TZVPP minimum-energy structures and BDEs of Cu⁺(H₂O)_{0,1}(H₂/D₂)_{0–2} are shown in Fig. 3. The global minimum-energy structures of Cu⁺(H₂O)(H₂)_{1,2} are of C_{2v} symmetry. Both hydrogen molecules bind directly to the Cu⁺ ion. A single H₂ ligand binds to Cu⁺(H₂O) remarkably strongly with a BDE (D₀) of 76.8 kJ mol⁻¹, which is considerably higher than the 55.5 kJ mol⁻¹ found for one H₂ bound to the bare Cu⁺ ion. This underscores the role of the *trans*-positioned water molecule in increasing the bond

Table 1 Band positions (in cm⁻¹) observed in the IRPD spectra of Cu⁺(D₂O)(H₂)₂ and Cu⁺(H₂O)(D₂)₂ shown in Fig. 1, VPT2/MP2/def2-TZVPP vibrational frequencies and intensities (in km mol⁻¹ in parenthesis) and band assignments^a

Label	IRPD	Calculated frequency (intensity)	Assignment ^a
Cu⁺(D₂O)(H₂)₂			
	2651 ^b	2650 (112)	ν ^s _{OD}
	2751 ^b	2754 (165)	ν ^a _{OD}
	2840	2857 (7)	ν ^{as} _{OD} + δ ^{oop} _{OD}
	3177	3190 (2)	ν ^{as} _{OD} + δ ^{ip} _{OD}
a ₁	3568	3564 (0.7), 3563 (0.2)	ν _{HH}
a ₂	3693	3698 (0.1)	ν _{HH} + τ ^{oop} _{OD}
a ₃	3733	3733 (17), 3777 (18)	ν _{HH} + δ ^{ip} _{CuH₂}
	3944	3945 (2), 4005 (0.3)	ν ^{as} _{OD} + δ ^{oop} _{DD}
a ₄	4350	4287 (4), 4362 (5), 4316 (2), 4338 (0.7)	ν _{HH} + ν _{CuH₂}
Cu⁺(H₂O)(D₂)₂			
b ₁	2567	2566 (0.4), 2568 (0.1)	ν _{DD}
b ₂	2675	2692 (0.2), 2695 (0.1)	ν _{DD} + δ ^{ip} _{CuD₂}
b ₃	2783	2800 (0.2), 2810 (0.1)	ν _{DD} + δ ^{ip} _{CuD₂} , ν _{DD} + δ _{D₂CuD₂}
b ₄	3125	3151 (0.3), 3151 (2), 3158 (1), 3165 (1)	ν _{DD} + ν _{CuD₂}
	3244	3222 (3)	2 × δ _{DD}
b ₅	3414	3441 (0.5), 3442 (0.3), 3450 (0.1)	ν _{DD} + ν _{DCuD}
	3628 ^b	3622 (198)	ν ^s _{OH}
	3700 ^b	3703 (120)	ν ^a _{OH}
	3819	3826 (18)	ν ^a _{OH} + τ ^{oop} _{OH}
	3889, 3900	3860 (4)	ν ^s _{OH} + δ ^{oop} _{CuD₂}
	4207	4170 (0.1)	ν ^s _{OH} + ν ^{as} _{CuD₂}
	4283	4250 (3), 4286 (1)	

^a Labeling of vibrational modes: ν (stretching), ν^s (symmetric stretching), ν^{as} (antisymmetric stretching), δ (bending), δ^{ip} (in-plane bending), δ^{oop} (out-of-plane bending), τ^{oop} (out-of-plane hindered rotation, or libration). Only modes with IR intensities larger than 0.1 km mol⁻¹ are listed.

^b Values are obtained from the fitting of the rovibrational profiles (see Fig. S2).



strength between Cu^+ and H_2 . However, only 17.9 kJ mol^{-1} is gained when binding a second H_2 , which is in line with a higher Cu^+-H_2 distance of $1.64 \text{ \AA}$ for the two H_2 molecules in $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$ (compared to $1.58 \text{ \AA}$ in the complex with one H_2). We will revisit the effect of H_2 adsorption on the ion-ligand bond lengths in the bonding analysis section. The addition of a third H_2 molecule yields only 2.37 kJ mol^{-1} .

When compared to the bond length in the free hydrogen molecule, the H-H distance in the $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$ and $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$ complexes is elongated by 5.3 pm and 4.7 pm , respectively (see Table S6†). This activation of the H-H (D-D) bond is substantially larger than the $2.3\text{--}3.1 \text{ pm}$ reported for $\text{Cu}(\text{H}_2)_4^+$ and is in line with the much stronger redshift of the H-H (D-D) stretching fundamental observed for the present system.

Vibrational spectra and band assignments. In order to assign the bands observed in the IRPD spectra, we calculated IR spectra including anharmonic contributions using vibrational perturbation theory (see computational section for details of the VPT2/MP2/def2-TZVPP calculations). The IR spectra for $\text{Cu}^+(\text{D}_2\text{O})(\text{H}_2)_2$ and $\text{Cu}^+(\text{H}_2\text{O})(\text{D}_2)_2$ (shown in Fig. 2C and D) are determined from VPT2 frequencies and intensities (listed in Table 1) using a 10 cm^{-1} wide Gaussian lineshape function.

Only the VPT2 IR bands involving excitation of H_2 or D_2 stretching modes, either as a fundamental (blue bands) or as part of a combination transition (orange bands), are color-coded. Based on the agreement between the experimental and computed vibrational spectra shown in Fig. 2, we can unambiguously assign all the observed bands (see Table 2). The excitation of the H-H (D-D) stretching fundamental a_1 (b_1) is observed at $\nu_{\text{HH}} = 3568 \text{ cm}^{-1}$ ($\nu_{\text{DD}} = 2567 \text{ cm}^{-1}$), resulting in a frequency ratio $\nu_{\text{HH}}/\nu_{\text{DD}}$ of 1.390 . This ratio is close to the value of 1.388 obtained from the VPT2 IR spectra, confirming the high quality of the MP2/def2-TZVPP potential along the vibrational

Table 2 EDA-NOCV result for $\text{Cu}-\text{H}_2$ bond of $\text{Cu}^+(\text{H}_2)_2(\text{H}_2\text{O})$, $\text{Cu}^+(\text{H}_2)(\text{H}_2\text{O})$ and $\text{Cu}^+(\text{H}_2)^a$

	$\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)-(\text{H}_2)$	$\text{Cu}^+(\text{H}_2\text{O})-(\text{H}_2)$	$\text{Cu}^+-(\text{H}_2)$
ΔE_{int}	−71	−112	−95
$\Delta E_{\text{int}}(\text{disp})^b$	−3(4%)	−2(2%)	−2(2%)
$\Delta E_{\text{int}}(\text{elec})^b$	−68(96%)	−110(98%)	−93(98%)
ΔE_{Pauli}	193	188	189
$\Delta E_{\text{elstat}}^c$	−151(58%)	−172(58%)	−155(55%)
ΔE_{orb}^c	−110(42%)	−126(42%)	−127(45%)
$\Delta E_1(\sigma(\text{H}_2) \rightarrow \text{Cu}(\text{d}))^d$	−58(53%)	−79(63%)	−69(54%)
$\Delta E_2(\text{Cu}(\text{d}) \rightarrow \sigma^*(\text{H}_2))^d$	−24(22%)	−19(15%)	−38(30%)
$\Delta E_3(\text{Cu}(\text{d}) \rightarrow \sigma^*(\text{H}_2))^d$	−26(24%)	−25(20%)	−18(14%)
ΔE_{prep}	44	5	4
$\Delta E_{\text{prep}}(\text{H}_2)$	4	5	4
$\Delta E_{\text{prep}}(\text{Cu}^+\text{L}_{0-3})$	40	0	0
ΔE_{bond}	−27	−107	−91
$r(\text{Cu}-\text{H}_2)^e$	1.72	1.64	1.70

^a All energies in kJ mol^{-1} , bond distances in $\text{\AA}$ computed with B3LYP-D4/TZ2P. Fragments are closed-shell species $\text{Cu}^+\text{L}_{0-3}$ and H_2 ($\text{L} = \text{H}_2\text{O}, \text{H}_2$). ^b Percentage values give the relative contributions of dispersion and electronic effects to ΔE_{int} . ^c Percentage values give the relative contributions to the attractive pEDA terms ΔE_{elstat} and ΔE_{orb} . ^d Percentage values give the relative contributions of the NOCV to ΔE_{orb} . ^e Distance between Cu and the centre of the H_2 fragment.

mode. The combination bands a_2 to a_4 , observed at 3693 cm^{-1} , 3733 cm^{-1} , and 4350 cm^{-1} , respectively, are assigned to combined excitations of the H-H stretching modes (ν_{HH}) with either of the following low-frequency vibrational modes: out-of-plane OD hindered rotation ($\tau_{\text{OD}}^{\text{OOP}}$), in-plane Cu^+-H_2 bending ($\delta_{\text{CuH}_2}^{\text{IP}}$), and Cu^+-H_2 stretching modes (ν_{CuH_2}). For the latter two modes, the present assignment is similar to the one made previously for the IRPD spectrum of $\text{Cu}^+(\text{H}_2)_4$.¹³ Analogously, the combination bands b_2 to b_5 , observed at 2675 cm^{-1} , 2783 cm^{-1} , 3125 cm^{-1} and 3414 cm^{-1} , respectively, are assigned to combination excitations of one of the two D-D stretching modes (ν_{DD}) with Cu^+-D_2 bending ($\delta_{\text{CuD}_2}^{\text{IP}}$ and $\delta_{\text{CuD}_2}^{\text{OOP}}$) or the Cu^+-D_2 stretching excitations ($\nu_{\text{CuD}_2}^s$ and ν_{DCuD}).

The remarkably good agreement between the IRPD and VPT2 spectra in Fig. 2, which is discussed in detail in the ESI,† confirms that the chosen computational method provides an adequate description of also the low-frequency $\text{M}^+-\text{H}_2/\text{D}_2$ vibrational modes, which play an important role in modelling the isotopologue selectivity. However, this appears to be in part due to a favourable error compensation between, *i.e.*, the neglect of substantial relativistic effects, basis set insufficiencies, and the overestimation of London dispersion interaction inherent in the MP2 method (see ESI†).

Bonding analysis of Cu^+-H_2 and $\text{Cu}^+-\text{H}_2\text{O}$ bonds. The nature of the Cu^+-H_2 and $\text{Cu}^+-\text{H}_2\text{O}$ interactions in the complexes investigated was analysed with an Energy Decomposition Analysis^{19–23} together with the Natural Orbital for Chemical Valence approach (EDA-NOCV)^{20–24} method. EDA allows the decomposition of the bonding energy ΔE_{bond} between fragments, which correspond to negative dissociation energy, into chemically meaningful contributions. The bonding energy

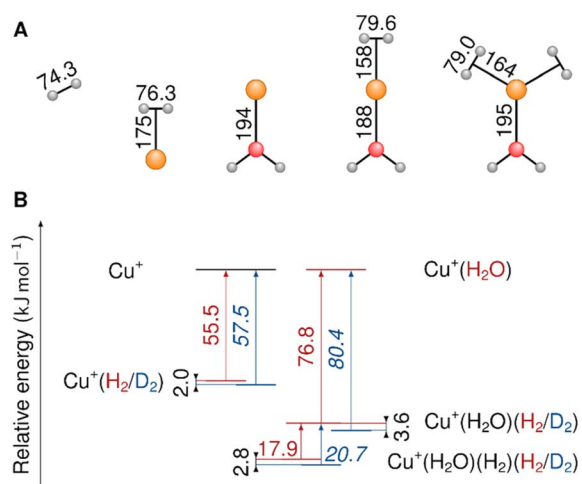


Fig. 3 (A) MP2/def2-TZVPP bond lengths (pm) in $\text{Cu}^+(\text{H}_2)$, $\text{Cu}^+(\text{H}_2\text{O})$ and $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_{1,2}$ (colours: Cu orange, O red, H grey). (B) MP2/def2-TZVPP dissociation energies (kJ mol^{-1}) with anharmonic (VPT2) ZPE correction. The red values (blue values in italic font) represent the dissociation energies of H_2 (D_2). Dissociation ZPE differences are highlighted.



ΔE_{bond} is represented as a sum of interaction energy ΔE_{int} and preparation energy ΔE_{prep} where the latter describes the deformation of the fragments during the bond formation. The interaction energy ΔE_{int} can be further split into dispersion $\Delta E_{\text{int}}(\text{disp})$ and electronic effects $\Delta E_{\text{int}}(\text{elec})$. The electronic effects can then be divided into quasi-electrostatic contribution ΔE_{elec} , Pauli repulsion ΔE_{Pauli} and orbital interaction ΔE_{orb} . The NOCV extension helps to find the most important orbital interactions to the bond as further decomposition of ΔE_{orb} .

The EDA-NOCV results for the complexes investigated in this work are shown in Table 2. They confirm a donor–acceptor-type interaction in the $\text{Cu}^+\text{--H}_2$ bond in all investigated complexes ($\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$, $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$, and $\text{Cu}^+(\text{H}_2)$). At first glance, the stronger bonding in $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$ compared to $\text{Cu}^+(\text{H}_2)$ is due to an increase in the electrostatic attraction (difference $\Delta\Delta E_{\text{elstat}} = -17 \text{ kJ mol}^{-1}$) while the orbital interaction (ΔE_{orb}) and Pauli repulsion (ΔE_{Pauli}) do not change significantly. However, a closer look reveals a more intriguing trend. The bond in $\text{Cu}^+(\text{H}_2)$ is significantly longer compared to $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$. If we analyse the main deformation densities (Fig. 4), we find that upon coordination of H_2O , the σ -donation from $\sigma(\text{H}_2)$ into the d-orbitals at Cu (ΔE_1) increases by 10 kJ mol^{-1} . But the in-plane back-bonding (ΔE_2) decreases by a larger amount (-19 kJ mol^{-1}), while the out-of-plane back-bonding (ΔE_3) is slightly increased ($+7 \text{ kJ mol}^{-1}$). This reduced in-plane back-bonding is due to the involvement of the Cu(d)-orbitals in the bonding to the H_2O ligand, which can be seen in the deformation densities $\Delta\rho_1$ and $\Delta\rho_2$ in Fig. 4. This leads to a lower Pauli repulsion and allows the H_2 molecule to bind closer to the Cu centre. While orbital attraction and Pauli repulsion thus balance at the same values compared to $\text{Cu}^+(\text{H}_2)$, the electrostatic attraction is increased. The main reason is thus lowering of Pauli repulsion leading to a shorter Cu– H_2 distance in $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$.

The weakening of the $\text{Cu}^+\text{--H}_2$ bond in $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$ compared to $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$ is attributed mainly to the large preparation energy resulting from the transition from a linear to

Table 3 Calculated CCSD(T)/aug-cc-pVTZ(-PP) BDEs (D_0) including the desorption ZPE ($\Delta_{\text{des}}\text{ZPE}$) calculated using VPT2 on the MP2/def2-TZVPP potential energy surface; anharmonic contribution (VPT2 level) for the dissociation of H_2 isotopologues, *i.e.* $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2 \rightarrow \text{Cu}^+(\text{H}_2\text{O})(\text{H}_2) + \text{H}_2$. All values reported in kJ mol^{-1}

Species	D_0	$\Delta_{\text{des}}\text{ZPE}$	Anharmonic contribution
$\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$	16.9	−10.8	0.8
$\text{Cu}^+(\text{H}_2\text{O})(\text{D}_2)_2$	19.7	−8.0	0.4
$\text{Cu}^+(\text{H}_2\text{O})(\text{T}_2)_2$	21.0	−6.7	0.3
$\text{Cu}^+(\text{H}_2\text{O})(\text{HD})_2$	18.4	−9.3	0.6
$\text{Cu}^+(\text{H}_2\text{O})(\text{HT})_2$	19.2	−8.5	0.5
$\text{Cu}^+(\text{H}_2\text{O})(\text{DT})_2$	20.4	−7.3	0.3

a trigonal structure. Additionally, the larger bond lengths are due to the stronger Pauli repulsion between the higher number of ligands and contribute to this phenomenon because they weaken the attractive interactions compared to $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$.

Note that the $\text{Cu}^+\text{--H}_2\text{O}$ bond is stronger in $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$ compared to $\text{Cu}^+(\text{H}_2\text{O})$, which is mainly due to reduced Pauli repulsion as an effect of the back-bonding contributions from Cu^+ to H_2 (see NOCV in Fig. S9 in ESI†). Adding another H_2 molecule, *i.e.* the formation of $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$, leads to a weaker $\text{Cu}^+\text{--H}_2\text{O}$ bond due to competition of the three ligands for the electrons at the central atom.

Anharmonic contribution to the BDEs for different isotopologues

As shown in Table 3, going from $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2$ to $\text{Cu}^+(\text{H}_2\text{O})(\text{T}_2)_2$, the BDE of the H_2 isotopologue increases due to the increasing mass of the isotopologues and the consequently lower ZPE. The difference in the adsorption ZPE between the D_2 and the H_2 complexes is 2.8 kJ mol^{-1} (1.0 kJ mol^{-1} between T_2 and D_2), which is slightly larger than the one observed in Cu(I)-MFU-4l (2.5 kJ mol^{-1}) by I. Weinrauch *et al.*⁴ and about one-third greater than the ZPE change for the exchange reaction $\text{Cu}^+(\text{D}_2)_3(\text{H}_2) + \text{D}_2 \rightarrow \text{Cu}^+(\text{D}_2)_4 + \text{H}_2$ (2.1 kJ mol^{-1}) observed in the previous study conducted by some of us.¹³

Our calculations also show that the anharmonic contribution to the H_2 desorption ZPE for the $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)_2 \rightarrow \text{Cu}^+(\text{H}_2\text{O})(\text{H}_2) + \text{H}_2$ reaction is more than three times as big as the value obtained for free H_2 (0.2 kJ mol^{-1}). Similar factors are observed for D_2 (3.1) and T_2 (3.1). This demonstrates the substantial influence of anharmonic contributions on the thermodynamic equilibrium of the isotopologue exchange reaction, reducing, *e.g.*, the energy difference between H_2 and D_2 adsorption by 12%. As expected, the anharmonic contributions to the vibrational frequencies and ZPEs are inversely proportional to the reduced mass of the dihydrogen isotopologue.

H_2 isotopologue selectivity

As shown in Fig. 5, the predicted isotopologue selectivity of H_2 adsorption is much higher at $\text{Cu}^+(\text{H}_2\text{O})$ (B) than at Cu^+ (A) and decreases again for a second H_2 adsorbing at $\text{Cu}^+(\text{H}_2\text{O})(\text{H}_2)$. We



Fig. 4 Selected deformation densities ($\Delta\rho_i$) from EDA-NOCVs for Cu– H_2 bond with energy contribution (ΔE_i) to ΔE_{orb} in kJ mol^{-1} and eigenvalues (v_i) Charge depletion (red) and charge accumulation (blue) with isosurface value 0.0025 for $\text{Cu}(\text{H}_2)^+$, 0.0020 for $\text{Cu}(\text{H}_2)(\text{H}_2\text{O})^+$ and 0.0012 for $\text{Cu}(\text{H}_2)_2(\text{H}_2\text{O})^+$.





Fig. 5 Predicted selectivities for dihydrogen isotopologue adsorption at Cu⁺, Cu⁺(H₂O), and Cu⁺(H₂O)(H₂) as a function of temperature. For the latter, two models are compared (see text). Literature data of Cu(I)-MFU-4l are shown for comparison.⁴

study two scenarios for the resulting Cu⁺(H₂O)(H₂)₂ complex: in the first one, we assume that the second H₂ molecule adsorbs at the free space of a hypothetical Cu⁺(H₂O)(H₂) complex where H₂O and the first H₂ are already positioned as in Cu⁺(H₂O)(H₂)₂, thus mimicking a scenario as it could occur on an undercoordinated adsorption site in a porous solid (C). Here, only the six degrees of freedom of the considered H₂ molecule are taken into account to calculate the isotopologue selectivity of dihydrogen adsorption, an approach that has been used in previous studies of MOFs.⁴

In the second one (D), all complexes are fully optimized and the degrees of freedom of all involved H₂ moieties are considered, which can be compared directly with the IRPD experiments. As a consequence of the weaker binding energy per H₂ molecule, the ZPE per H₂ and finally the overall selectivity is reduced significantly.

We conclude that a model with a rigid ligand structure is insufficient when dealing with structurally flexible adsorption sites such as undercoordinated metal ions, as it will lead to spurious results due to ligand reorientation. Conversely, as

evidenced by prior studies,^{4,8} rigid adsorption sites with structurally constraining ligands like in the hypothetical starting structure in Fig. 5C may be essential for achieving a high adsorption energy and isotopologue selectivity.

Summary and conclusions

We elucidated the strong hydrogen isotopologue selectivity of undercoordinated Cu(I) sites by examining the model complexes Cu⁺(H₂O)(H₂)_n (with *n* = 0, 1, 2) using experiment and theory. Both Cu⁺(H₂O) and Cu⁺(H₂O)(H₂/D₂) show an obvious isotope effect in the adsorption of H₂ and D₂. The observed vibrational spectra of Cu⁺(H₂O)(H₂)₂ and its isotopologues in the range from 2500–4500 cm^{−1} match the calculated frequencies of the fundamentals and related combination bands of H₂/D₂ when taking anharmonicity into account. This suggests that these systems are ideal model complexes for gas-phase studies of the chemistry at individual active sites as they occur in framework materials. The stronger H₂ affinity of Cu(I) coordinated by a single oxygen-donor ligand, in comparison to the bare cation, is explained by EDA and traced back to lowering of Pauli repulsion which allows the H₂ molecule to bind closer to the Cu centre. Finally, we used the computational data to predict the dihydrogen isotopologue selectivity of the formation of Cu⁺(H₂O)(H₂)_n complexes, observing a significant selectivity decrease for *n* = 2 vs. *n* = 1. The present study underlines that the structural environment of undercoordinated metal centres in framework materials play a crucial role in the local chemistry and demonstrate particular promise for rigid frameworks with highly exposed metal sites.

Methods

Experimental methods

The experiments were performed on the Leipzig 5 K ring-electrode ion-trap triple mass spectrometer described previously.¹⁶ Cu⁺(H₂O) cations are transferred to the gas phase from a 5 mmol CuSO₄ (Sigma Aldrich: CuSO₄·5H₂O 99% in CH₃OH 99.9% and distilled water) solution using a nanospray ion source under open atmospheric conditions, while the Cu⁺(D₂O) cations are produced *via* H/D exchange in moist air conditions saturated with heavy water vapour (99.9% D₂O). The beam of cations is skimmed, thermalized to room temperature in a Helium-filled radio-frequency ion guide, and then mass-selected in a quadrupole mass filter. Mass-selected Cu⁺(H₂O/D₂O) ions are continuously trapped in a radio-frequency ring-electrode ion trap, held at a temperature in-between 15 K and 295 K using a closed-cycle helium cryostat and filled with ≈ 1 mbar H₂(D₂). Cu⁺(H₂O/D₂O)(H₂/D₂)_n adducts are formed by three-body collisions and thermalized to the ambient temperature by many collisions with other gas molecules.

All cations are extracted from the ion trap every 100 ms and weakly focused both temporally and spatially into the centre of the extraction region of an orthogonally mounted double-focusing reflectron time-of-flight (TOF) mass spectrometer. For the determination of the ion yields, the ions are extracted and accelerated without using the reflectron. Furthermore, TOF



Hoja and A. Daniel Boese at University of Graz for fruitful scientific discussions and computational resources. JJ and KA thank André Fielicke (Fritz-Haber-Institut der Max-Planck-Gesellschaft) for helpful discussions of low-frequency M^+-H_2 modes. SH, JJ and KA thank Alexandra Giermann for the preparation of sample solutions.

References

- Hoja and A. Daniel Boese at University of Graz for fruitful scientific discussions and computational resources. JJ and KA thank André Fielicke (Fritz-Haber-Institut der Max-Planck-Gesellschaft) for helpful discussions of low-frequency M^+-H_2 modes. SH, JJ and KA thank Alexandra Giermann for the preparation of sample solutions.
- ## References
- (a) D. Sengupta, P. Melix, S. Bose, J. Duncan, X. Wang, M. R. Mian, K. O. Kirlikovali, F. Joodaki, T. Islamoglu, T. Yildirim, R. Q. Snurr and O. K. Farha, *J. Am. Chem. Soc.*, 2023, **145**, 20492; (b) B. R. Barnett, H. A. Evans, G. M. Su, H. Z. H. Jiang, R. Chakraborty, D. Banyeretse, T. J. Hartman, M. B. Martinez, B. A. Trump and J. D. Tarver, others, *J. Am. Chem. Soc.*, 2021, **143**, 14884.
 - R. Xiong, L. Zhang, P. Li, W. Luo, T. Tang, B. Ao, G. Sang, C. Chen, X. Yan, J. Chen and M. Hirscher, *Chem. Eng. J.*, 2020, **391**, 123485.
 - B. Ipek, R. A. Pollock, C. M. Brown, D. Uner and R. F. Lobo, *J. Phys. Chem. C*, 2018, **122**, 540.
 - I. Weinrauch, I. Savchenko, D. Denysenko, S. M. Souliou, H.-H. Kim, M. Le Tacon, L. L. Daemen, Y. Cheng, A. Mavrandonakis, A. J. Ramirez-Cuesta, D. Volkmer, G. Schütz, M. Hirscher and T. Heine, *Nat. Commun.*, 2017, **8**, 14496.
 - J. Teufel, H. Oh, M. Hirscher, M. Wahiduzzaman, L. Zhechkov, A. Kuc, T. Heine, D. Denysenko and D. Volkmer, *Adv. Mater.*, 2013, **25**, 635.
 - H. Oh and M. Hirscher, *Eur. J. Inorg. Chem.*, 2016, **2016**, 4278.
 - T. Wulf and T. Heine, *Int. J. Quantum Chem.*, 2018, **118**, e25545.
 - T. Wulf and T. Heine, *J. Phys. Chem. C*, 2020, **124**, 9409.
 - (a) K. R. Asmis, A. Fielicke, G. von Helden and G. Meijer, in *Atomic clusters*, ed. D. Woodruff, Elsevier, Amsterdam and Boston, 2007, pp. 327–375; (b) E. J. Bieske and O. Dopfer, *Chem. Rev.*, 2000, **100**, 3963; (c) J. M. Lisy, *Int. Rev. Phys. Chem.*, 1997, **16**, 267; (d) M. A. Duncan, *Int. Rev. Phys. Chem.*, 2003, **22**, 407; (e) A. Fielicke, *Chem. Soc. Rev.*, 2023, **52**, 3778.
 - (a) V. Dryza and E. J. Bieske, *Int. Rev. Phys. Chem.*, 2013, **32**, 559; (b) V. Dryza, B. L. J. Poad and E. J. Bieske, *Phys. Chem. Chem. Phys.*, 2012, **14**, 14954.
 - (a) G. Piccini and J. Sauer, *J. Chem. Theory Comput.*, 2014, **10**, 2479; (b) K. Sillar, A. Hofmann and J. Sauer, *J. Am. Chem. Soc.*, 2009, **131**, 4143.
 - (a) F. Dahlmann, C. Lochmann, A. N. Marimuthu, M. Lara-Moreno, T. Stoecklin, P. Halvick, M. Raoult, O. Dulieu, R. Wild, S. Schlemmer and others, *J. Chem. Phys.*, 2021, 155; (b) F. Dahlmann, P. Jusko, M. Lara-Moreno, P. Halvick, A. N. Marimuthu, T. Michaelson, R. Wild, K. Geistlinger, S. Schlemmer, T. Stoecklin, R. Wester and S. Brünken, *Mol. Phys.*, 2022, 120; (c) M. Jia, J. Vanbuel, P. Ferrari, E. M. Fernández, S. Gewinner, W. Schöllkopf, M. T. Nguyen, A. Fielicke and E. Janssens, *J. Phys. Chem. C*, 2018, **122**, 18247; (d) I. Swart, F. M. F. de Groot, B. M. Weckhuysen, P. Gruene, G. Meijer and A. Fielicke, *J. Phys. Chem. A*, 2008, **112**, 1139.
 - J. Jin, T. Wulf, M. Jorewitz, T. Heine and K. R. Asmis, *Phys. Chem. Chem. Phys.*, 2023, **25**, 5262.
 - P. R. Kemper, P. Weis, M. T. Bowers and P. Maître, *J. Am. Chem. Soc.*, 1998, **120**, 13494.
 - (a) P. Maitre and C. W. Bauschlicher, *J. Phys. Chem.*, 1993, **97**, 11912; (b) R. H. Crabtree, *Chem. Rev.*, 2016, **116**, 8750; (c) G. J. Kubas, *Metal Dihydrogen and S-Bond Complexes: Structure, Theory, and Reactivity*, Springer Science & Business Media, 2001.
 - N. Heine and K. R. Asmis, *Int. Rev. Phys. Chem.*, 2015, **34**, 1.
 - K. R. Asmis, M. Brümmer, C. Kaposta, G. Santambrogio, G. von Helden, G. Meijer, K. Rademann and L. Wöste, *Phys. Chem. Chem. Phys.*, 2002, **4**, 1101.
 - P. D. Carnegie, A. B. McCoy and M. A. Duncan, *J. Phys. Chem. A*, 2009, **113**, 4849.
 - T. Ziegler and A. Rauk, *Theoret. Chim. Acta*, 1977, **46**, 1.
 - K. Kitaura and K. Morokuma, *Int. J. Quantum Chem.*, 1976, **10**, 325.
 - F. M. Bickelhaupt and E. J. Baerends, in *Reviews in Computational Chemistry*, ed. K. B. Lipkowitz and D. B. Boyd, Wiley, 2000, vol. 15, pp. 1–86.
 - L. Zhao, M. von Hopffgarten, D. M. Andrada and G. Frenking, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2018, **8**, e1345.
 - M. P. Mitoraj, A. Michalak and T. Ziegler, *J. Chem. Theory Comput.*, 2009, **5**, 962.
 - T. Ziegler and A. Rauk, *Theoret. Chim. Acta*, 1977, **46**, 1.
 - M. Mayer and K. R. Asmis, *J. Phys. Chem. A*, 2021, **125**, 2801.
 - W. R. Bosenberg and D. R. Guyer, *J. Opt. Soc. Am. B*, 1993, **10**, 1716.
 - N. Heine and K. R. Asmis, *Int. Rev. Phys. Chem.*, 2016, **35**, 507.
 - J. F. Stanton, J. Gauss, L. Cheng, M. E. Harding, D. A. Matthews and P. G. Szalay, *CFOUR, Coupled-Cluster techniques for Computational Chemistry, a quantum chemical program package*, 2024.
 - M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. W. Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16 (Revision C.01)*, Gaussian Inc., Wallingford CT, 2016.
 - G. te Velde, F. M. Bickelhaupt, E. J. Baerends, C. Fonseca Guerra, S. J. A. van Gisbergen, J. G. Snijders and T. Ziegler, *J. Comput. Chem.*, 2001, **22**, 931.



- 31 P. J. Stephens, F. J. Devlin, C. F. Chabalowski and M. J. Frisch, *J. Phys. Chem.*, 1994, **98**, 11623.
- 32 E. Caldeweyher, S. Ehlert, A. Hansen, H. Neugebauer, S. Spicher, C. Bannwarth and S. Grimme, *J. Chem. Phys.*, 2019, **150**, 154122.
- 33 E. van Lenthe and E. J. Baerends, *J. Comput. Chem.*, 2003, **24**, 1142.
- 34 E. van Lenthe, A. Ehlers and E.-J. Baerends, *J. Chem. Phys.*, 1999, **110**, 8943.

