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Assigning ^1H chemical shifts in paramagnetic mono- and bimetallic surface sites using DFT: a case study on the Union Carbide polymerization catalyst†

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The Union Carbide (UC) ethylene polymerization catalyst, based on silica-supported chromocene, is one of the first industrial catalysts prepared by surface organometallic chemistry, though the structure of the surface sites remains elusive. Recently, our group reported that monomeric and dimeric Cr(II) sites, as well as Cr(III) hydride sites, are present and that their proportion varies as a function of the Cr loading. While ^1H chemical shifts extracted from solid-state ^1H NMR spectra should be diagnostic of the structure of such surface sites, unpaired electrons centered on Cr atoms induce large paramagnetic ^1H shifts that complicate their NMR analysis. Here, we implement a cost-efficient DFT methodology to calculate ^1H chemical shifts for antiferromagnetically coupled metal dimeric sites using a Boltzmann-averaged Fermi contact term over the population of the different spin states. This method allowed us to assign the ^1H chemical shifts observed for the industrial-like UC catalyst. The presence of monomeric and dimeric Cr(II) sites, as well as a dimeric Cr(III)-hydride sites, was confirmed and their structure was clarified.

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1. Introduction

Polyethylene (PE) is one of the most commonly used plastics with an annual production exceeding 100 million tons.¹ Its manufacture relies mostly on Ziegler-Natta or Phillips catalysts, involving paramagnetic Ti(III) and Cr(III) active sites.^{1–3} Another notable example is the Union Carbide (UC) catalyst, based on silica-supported chromocene Cr^{II}Cp₂ (**1**), that has been used for specialized applications.^{4–6} The nature of its surface and active sites has been under investigation for more than 50 years due to the complexity of this system, though various types of Cr sites have been proposed based on IR and solid-state NMR spectroscopic analyses.^{5,7–9} For example, we have recently investigated the effect of the Cr loading on the IR response after CO adsorption in the UC catalyst. Complemented with detailed computational studies, it was possible to identify mono-grafted Cr(II) sites **I** as well as dimeric Cr(II) sites with weak (**III**) and strong (**II**) Cr–Cr interactions (Fig. 1A).¹⁰ At high Cr loading (1 and 2 wt% Cr), we reported that dimeric and monomeric Cr(III)-

hydrides (**IV** and **V**) are likely present, as evidenced by the appearance of low energy CO bands, indicative of Cr-formyl species formed upon CO insertion into a Cr-hydride bond.

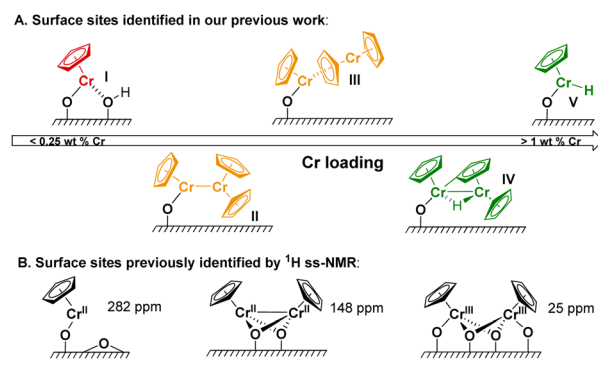


Fig. 1 (A) Identified surface sites in the UC catalyst as a function of the Cr loading based on CO-IR and EPR spectroscopy as well as DFT studies, as outlined in our previous work.¹⁰ The numbering of the surface species has been adapted accordingly. At low Cr loading, monomeric Cr(II) sites **I** are mainly present; these sites can possibly interact with residual surface OH groups (*vide infra*). When increasing the Cr loading, site **I** can interact with **1** to form dimeric sites with a bridging Cp ligand (**III**) or direct Cr–Cr interaction (**II**). Site **II** can yield a bridging Cr(III)-hydride complex (**IV**) via C–H activation at the Cp ring, which can further react with a surface hydroxyl group, leading to a monomeric Cr(III) hydride (**V**). (B) A mono-grafted Cr site and dimeric Cr(III) and Cr(II) surface sites were identified by Schnellbach *et al.* using temperature-dependent ^1H MAS NMR spectroscopy.¹⁹

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The formation of these sites is supported by DFT studies and their presence was identified by EPR spectroscopy and labeling studies. This study also indicated that the active sites are more likely to be monomeric Cr(III) sites, consistent with what was found in corresponding molecular systems.^{11,12}

With these results in hand, we decided to re-investigate the speciation of surface sites using solid-state NMR spectroscopy, used in pioneering studies on understanding the Cr-site structure.¹³ The advantage of this technique over IR spectroscopy is its high spectral resolution without the need for probe molecules that can influence or modify the surface site distributions and structures. However, the presence of unpaired electrons in Cr-based sites introduces additional complexity that impedes a straightforward interpretation of NMR spectra. Indeed, the interaction of unpaired electrons with nuclear spins yields so-called paramagnetic chemical shifts that can appear in a broad range and that are impossible to predict by way of basic qualitative reasoning.^{14–17} This makes the assignment of paramagnetically shifted signals exceptionally challenging, especially in the case of multimetric sites and antiferromagnetically coupled Cr atoms. While computational protocols have been optimized to calculate paramagnetic chemical shifts in the case of monomeric metal complexes,^{14,18} no work has been reported in the case of dimeric complexes to the best of our knowledge.

Previous ¹H solid-state magic-angle-spinning (MAS) NMR studies on the UC catalyst at low Cr loading highlighted the presence of two signals consistent with surface-attached chromium species around 282 ppm and 25 ppm.¹³ At higher Cr loadings, it was reported that the signal intensity of the peak at 282 ppm decreases with the appearance of a new resonance at 148 ppm. Furthermore, it was reported that the signal at 282 ppm shows a higher chemical shift upon lowering the temperature, while the signals at 148 ppm and 25 ppm are insensitive to temperature changes.¹³ Based on these findings, the signal at 282 ppm was assigned to a mono-grafted chromium site, while the resonances at 148 ppm and 25 ppm to dimeric Cr(III) and Cr(II) sites respectively (Fig. 1B).

In this work, we first investigated the effect of the Cr loading on the ¹H MAS NMR signatures of **1** supported on SiO₂₋₇₀₀ to parallel what we have recently carried out with IR spectroscopy.²⁰ Next, we optimized a computational approach to evaluate the chemical shift of weakly interacting spins in bimetallic complexes. More precisely, the chemical shift was calculated using a Boltzmann distribution of the Fermi contact terms averaged over the different spin states, calculated using a DFT methodology inspired by Rastrelli and Bagno.^{14,21} The spin–spin coupling constant *J*, which determines the splitting of the spin states, was estimated using the broken symmetry approach.²² This methodology was benchmarked on a series of paramagnetic metal monomers and dimers and later on used to decipher the structure of surface sites by assigning paramagnetic-shifted ¹H MAS NMR signals of the industrial-like UC catalyst at various Cr loadings.

2. Results and discussion

2.1. ¹H chemical shifts in the industrial-like UC catalyst

The industrial-like UC catalyst was prepared by reacting chromocene CrCp₂ (**1**) with silica, partially dehydroxylated at 700 °C, according to our previously reported procedure.¹⁰ The catalyst was prepared with four different Cr loadings: 0.25, 0.5, 1.0, and 2.0 wt% Cr, yielding respectively **1-SiO₂-0.25**, **1-SiO₂-0.5**, **1-SiO₂-1**, and **1-SiO₂-2**. The solid-state ¹H MAS NMR spectra were acquired at 16.4 T (700 MHz for ¹H), in flowing dry N₂ gas, and a temperature of 280 K. In addition, the temperature dependence of the NMR signals was investigated by recording spectra at 293, 273, and 253 K. ¹H MAS rates of 35 and 40 kHz were used to minimize the overlap of paramagnetic shifted ¹H signals and spinning sidebands.²³ For each sample, three ¹H MAS NMR spectra with different center frequencies of 300 ppm, 150 ppm, and 0 ppm were recorded to detect the different paramagnetic-shifted signals.

The ¹H MAS NMR spectrum of **1-SiO₂-1** in Fig. 2 shows a substantial sideband manifold, consistent with large ¹H chemical shift anisotropies associated with the paramagnetic nature of surface Cr sites.^{13,15,24} Besides the spinning sidebands, the ¹H MAS NMR spectrum of **1-SiO₂-1** shows a weak, highly deshielded signal at 318 ppm, together with a more intense signal around 155 ppm. The latter disappears after long acquisition times (~days) regardless of the temperature of the measurement, indicating the sensitivity of the associated surface species. Besides the two highly de-shielded signals, two additional peaks close to the diamagnetic region appear at 23 ppm and 12 ppm, as previously reported.¹³ In the diamagnetic region, the peaks at 2.1 ppm and 4.3 ppm are associated with the isolated and interacting OH groups of the silica surface²⁵ and the one at 6.7 ppm with a trace amount of cyclopentadiene adsorbed on silica, similar to what is observed for pentamethylcyclopentadiene.²⁶

Regarding temperature dependence, the most de-shielded peak at 318 ppm shifts downfield while cooling below room

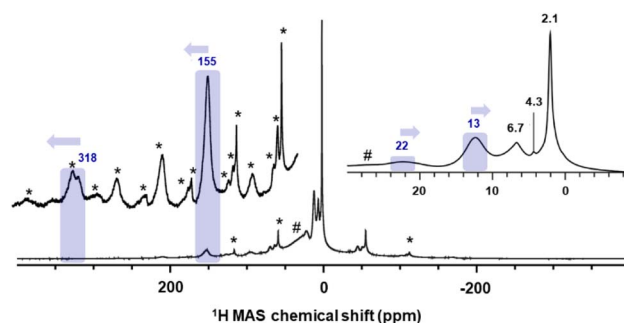


Fig. 2 1D relaxation-resolved ¹H MAS NMR echo spectrum of **1-SiO₂-1** acquired at 35 kHz MAS, 16.4 T, 0.005 s recycle delay, and a temperature of 280 K measured in the gas stream near the NMR rotor. The paramagnetic peaks are shown in blue, whereas the spinning sidebands and a background signal are indicated by "*" and "#" symbols, respectively. The blue arrows show the shift of the peaks upon cooling to 240 K. The peaks at 2.1, 4.3, and 6.7 ppm are indicative of isolated and interacting OH groups and trace amounts of cyclopentadiene adsorbed on silica respectively.



temperature ($\Delta T = -40$ K and $\Delta\delta = +46$ ppm). The peak at 155 ppm also shifts downfield while cooling below room temperature, but with a less pronounced temperature dependence ($\Delta T = -40$ K and $\Delta\delta = +16$ ppm) that is also indicative for monomeric sites, as reported by Schnellbach¹³ (see Fig. S6† for details). In contrast to the highly de-shielded resonances, the two peaks close to the diamagnetic region shift slightly up-field upon cooling below room temperature: $\Delta T = -40$ K and $\Delta\delta = -3$ (peak at 23 ppm) and -2 (peak at 12 ppm) as shown in Fig. S6.†

2.2. Effect of the Cr loading

To further investigate the attribution of the various resonances to specific surface sites, we next investigated the ¹H MAS NMR spectra of UC-like catalysts with different Cr loadings. In the ¹H MAS NMR spectrum of **1-SiO₂-0.25** (0.25 wt% Cr loading), only two highly de-shielded resonances at 318 ppm and 155 ppm (Fig. S4†) are observed. Based on our previous CO-IR investigation,¹⁰ this material mainly consists of monomeric Cr(II) sites **I** (Fig. 1). In addition, these peaks shift down-field upon cooling and are therefore consistent with monomeric sites or potentially dimeric Cr(II) sites with weak Cr–Cr interactions. This could result from the interaction between **I** and adsorbed chromocene (site **III**). Alternatively, we could expect an interaction between some of the Cr centers of **I** and nearby surface OH groups **I-OH**.

When increasing the Cr loading to 0.5 wt% (**1-SiO₂-0.5**, Fig. S5†), two additional resonances appear at 22 ppm and 13 ppm (see Fig. 3). For **1-SiO₂-1** and **1-SiO₂-2**, no additional resonances appear; however, the peak at 13 ppm gains in intensity compared to the peak at 22 ppm (see Fig. 3 and S6†). The appearance of these less deshielded signals parallels the formation of dimeric sites such as **II**, as evidenced earlier by CO-IR investigations.²⁰ The slightly up-field shift upon cooling is

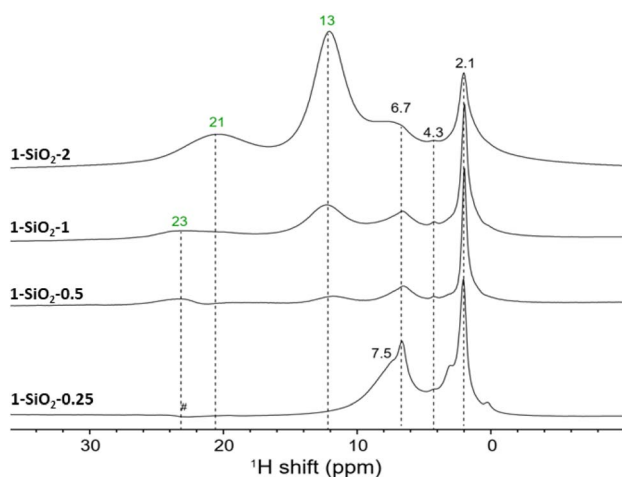


Fig. 3 Evolution of ¹H MAS NMR signal intensities in the diamagnetic region as a function of the Cr loading. A background signal in **1-SiO₂-0.25** is indicated by "#". While both peaks are absent in **1-SiO₂-0.25**, the signal at 13 ppm significantly increases in intensity by increasing the Cr loading (**1-SiO₂-2**).

also consistent with the presence of dimeric sites. The increase in intensity at 13 ppm at higher loading matches with the appearance of the low energy CO bands in IR, which we assigned to monomeric and dimeric Cr(III)-hydride sites (**V** and **IV** respectively).¹⁰

2.3. Paramagnetic proton shifts of monomeric and dimeric molecular complexes

After having qualitatively assigned the proton shifts based on our previous proposed structures and their temperature-dependent behavior, we next calculated the respective ¹H shifts to corroborate their assignment and determine the structural features of the specific surface sites. Towards this goal, we first evaluated the accuracy of the proposed calculation methodologies to predict the chemical shifts of Cp ligands by using **1** and [(CpCr^{II}OtBu)₂] (**2**) as molecular models for monomeric and dimeric sites.

Experimentally, chromocene (**1**) is characterized by a peak at 325 ppm at room temperature in d₈-toluene (see Fig. S7†). When lowering the temperature to 253 K, the signal shifts down-field to 383 ppm (Fig. S7†). The ¹H shift follows an inverse temperature dependence (Fig. S10†) in the temperature range from 323–238 K. Using the expression of the Fermi contact term (see the section Methodology and computational details), the hyperfine coupling constant was fitted to $A = 5.0$ MHz (assuming $g = g_e = 2$, Fig. S10†). A hyperfine coupling constant A of 3.8 MHz is predicted by DFT. From eqn (6), the ¹H resonance of **1** at 293 K is expected at 301 ppm with a down-field shift to 341 ppm at 253 K. This is in good agreement with the experimentally observed behavior, indicating that this approach is suitable for estimating the paramagnetic ¹H shifts of monomeric chromocene-based complexes.

With the dimeric [(CpCr^{II}OtBu)₂] complex **2**, we evaluate the method for ¹H shift calculations in the presence of electronic spin–spin interactions. The ¹H NMR spectrum of **2** shows a shift at 40 ppm corresponding to the H of the Cp ligand at room temperature in d₈-toluene. This signal moves up-field to 2 ppm ($\delta = -38$ ppm) when cooled at 253 K (Fig. 4A). This effect originates from the electronic structure of complex **2**, where each Cr^{II} center is in a local high spin state ($S = 2$, see Table S3†) and the weak spin–spin interaction between the two creates five global spin states ($S = 0, 1, 2, 3$, and 4).

These states split energetically as follows:

$$E_S = S(S + 1)J \quad (1)$$

with J (cm⁻¹) being the spin–spin coupling constant.²² These states are populated according to a Boltzmann distribution at a given temperature. The coupling constant for **2** was estimated to be $J = -208$ cm⁻¹ using the broken symmetry (BS) approach and the Yamaguchi formula (see the ESI†).^{22,27} This corresponds to an energy gap of 5.0 kJ mol⁻¹ between the singlet and triplet states for this complex. Thus, at 298 K, roughly 12% of **2** is in the global $S = 1$ state, while the rest (88%) adopts a $S = 0$ state. The global spin population varies significantly with temperature (see Fig. 4A). Assuming rapid exchange between spin states compared to the NMR time scale, the effective chemical shift for



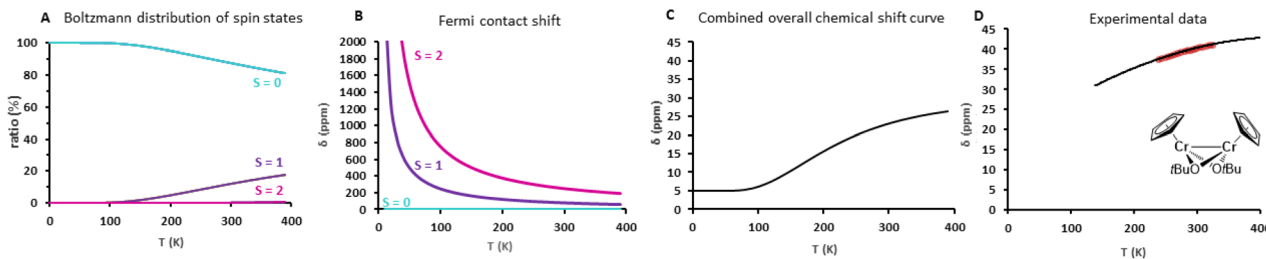


Fig. 4 Illustration of the different contributions to the chemical shift of complex 2. The calculated combined overall chemical shift curve in (C) results from a linear combination of the Boltzmann distribution of spin states (A) and the Fermi contact shifts δ_{FC} of the global spin states, as visualized in (B). For 2, a slight decrease in the chemical shift is expected while cooling below room temperature. The experimental data shown in (D) agree with the calculated ^1H shift curve of (C).

dimeric sites δ_{dim} should correspond to the Boltzmann-averaged linear combination of the Fermi contact shifts of the global spin states S_i ,²⁸ as shown in eqn (2):

$$\delta_{\text{dim}} = \delta_0 + \sum_{i=0}^n \delta_{FC_i} \left(\frac{\{N\}_i}{N} \right)$$

$$= \sigma_{\text{Fe}} - \sigma_{\text{orb}} + \sum_{i=0}^n \frac{2\pi}{\gamma_I} g \mu_B A \frac{S_i(S_i+1)}{3k_B T} \left\{ \frac{e^{-\frac{S_i(S_i+1)J}{k_B T}}}{\sum_i e^{-\frac{S_i(S_i+1)J}{k_B T}}} \right\} \quad (2)$$

The chemical shift calculated according to eqn (2) contains a double temperature dependence: the inverse temperature dependence of the Fermi contact shift for each global spin state, as well as the population of global spin states according to the Boltzmann distribution. For simplicity reasons, the hyperfine constant A between nuclear and electronic spins was calculated for the global high spin complex and is assumed to be constant for all other spin states (see Table S6†). Fig. 4C displays the ^1H shift curve of 2 as a combination of the Boltzmann distribution (Fig. 4A) and the individual Fermi contact shift curves (Fig. 4B) of the global spin states ($S = 0$ to 2 were considered). The predicted ^1H shift curve (18 ppm, 293 K, Fig. 4C) is in reasonable agreement with the experimental data obtained in a smaller temperature interval (40 ppm, 293 K, d_8 -toluene, see Fig. 4D). Although the calculated ^1H chemical shift is slightly underestimated, the temperature dependence is nicely reproduced. This difference could be due to an underestimation of the $S = 1$ population, because of a slightly elongated Cr–Cr bond distance in solution for instance. Indeed, the Cr–Cr bond distance strongly impacts the spin population and thus the chemical shift behavior. Fig. S14† illustrates the effect of changes in the global spin state populations on the shape of the ^1H shift curve of 2 when elongating the Cr–Cr distance. As seen from the J constant, the population of higher global spin states is favored at room temperature when the interatomic distance increases. An increase in the Cr–Cr distance by 0.25 Å (see ESI Section 4.3.3 and Fig. S14† for more details) can already cause an inversion in the temperature dependence of the chemical shift curve when cooling below room temperature. From these results, we can probably attribute the small deviations between

the measured and calculated chemical shifts to a small difference in the Cr–Cr distance between the crystal structure and the actual complex in solution (or on the surface).

The applicability of this method was then evaluated on a consistent series of dimeric complexes $[\text{CpCrX}_2 \text{ or } _3]_2$ (see Table 1). For the calculations of all molecular complexes in this work, the crystal structures of the complexes were used and only the hydrogen position was optimized. The experimental trend is nicely reproduced with an increase in the ^1H shift with increasing interatomic distance.

Overall, this methodology shows that the paramagnetic contribution to the ^1H chemical shielding primarily depends on the strength of the Cr–Cr interactions. Thus, the chemical shift is a direct descriptor of the spin state of the dimer and the Cr–Cr distance can be deduced *a posteriori* from the ^1H shift.

2.4. Method validation and applicability

We applied the aforescribed methodology for calculating the ^1H shifts of monomeric and dimeric sites on a set of paramagnetic monomeric and dimeric complexes based on 3d transition metals. For these calculations, we used the crystal structure of each complex and only optimized the proton positions (Fig. 5, see Table S3† for the references and experimental ^1H chemical shifts, Section S1† for the synthesis procedure of $[\text{Cp}_2\text{Cr}]$, $[(\text{Cp}_2\text{Cr}_2(\text{OSi}(\text{OtBu})_3)_2)]$ and $[(\text{Mes})_2\text{Co}]_2$ and S2.1† for the synthesis procedure and crystallographic details of $[(\text{Cp}_2\text{Cr}_2(\text{OSi}(\text{OtBu})_3)_3)^+(\text{Cp}_3\text{Cr}_2(\text{OSi}(\text{OtBu})_3)_4)^-]$). An overall good agreement ($R^2 = 0.969$ and slope = 1.014) is obtained between experimental and theoretical data. Given the complexity of the interpretation of paramagnetic shifts, this good agreement shows that the developed methodology can be

Table 1 Experimentally measured (δ_{exp} , benzene, 293 K) and theoretically calculated (δ_{calc}) ^1H shifts (ppm) for a series of dimeric Cr complexes with increasing interatomic distances ($d_{\text{Cr–Cr}}$, Å). See Tables S3 and S4 for details on the calculations

Complex	δ_{exp}	δ_{calc}	$d_{\text{Cr–Cr}}$
(CrCpOSi(OtBu) ₃) ₂	33 (ref. 10)	9	2.539
(CrCpOtBu) ₂	40 (ref. 29)	18	2.632
(CpCrOtBuCl) ₂	71 (ref. 30)	50	2.918
(CpCrCl ₂) ₂	159 (ref. 31)	234	3.459



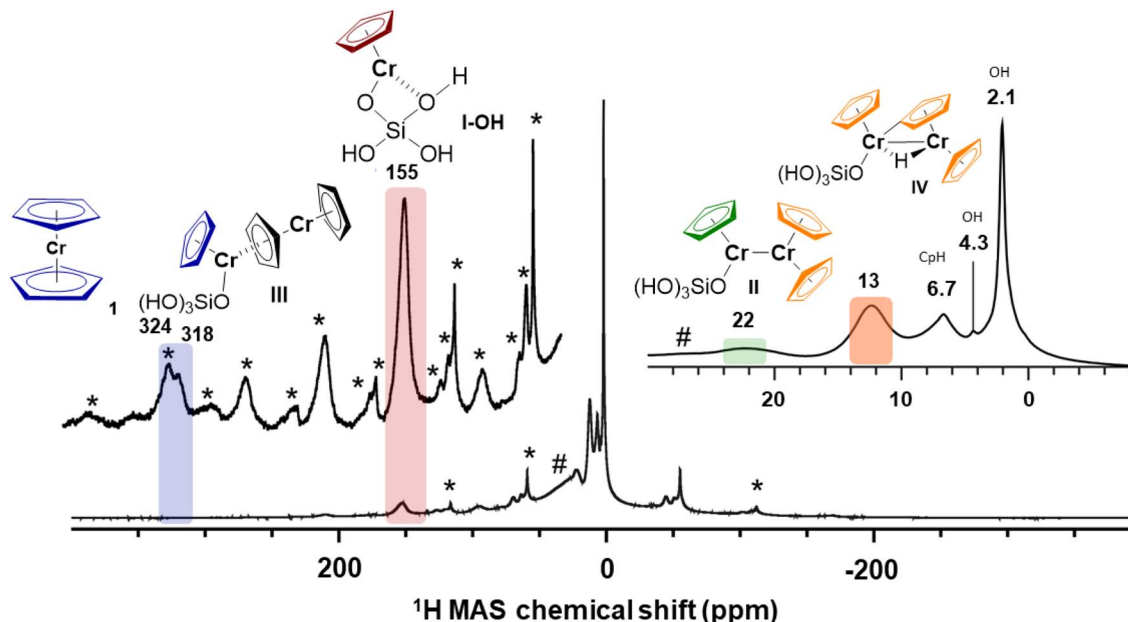


Fig. 6 ^1H MAS-NMR spectrum of 1-SiO₂-1 acquired at 40 kHz MAS, 16.4 T, and a temperature of 280 K measured in the gas stream near the NMR rotor. Spinning sidebands and a background signal are indicated by "*" and "#" symbols, respectively. The paramagnetic-shifted ^1H signals are assigned to monomeric sites interacting with nearby Si–O–Si bridges from the silica surface (155 ppm, red), as well as dimeric sites with strong (II, 22 ppm (green) and 13 ppm (orange)) and weak (III, 318 ppm, blue) Cr–Cr interactions. The presence of trace amounts of **1** strongly adsorbed on silica (318 ppm) cannot be excluded. C–H activated Cr(III)-hydrides (IV) appear at 13 ppm and are overlapping with the strongly interacting **1** unit of site II. The signals at 2.1–4.3 ppm and 6.7–7.5 ppm in the diamagnetic region are assigned respectively to isolated (2.1 ppm) and interacting (3.2 ppm and 4.3 ppm) surface hydroxyl groups, and trace amounts of physisorbed dicyclopentadiene (6.7–7.5 ppm).

3. Conclusions

In this work, we recorded the ^1H MAS NMR of Union Carbide catalysts at various Cr loadings and developed a computational methodology to calculate the ^1H NMR chemical shifts of weakly coupled dimeric complexes. This method takes into account the Boltzmann-averaged over the different spin states of the Fermi contact term. Combining the ^1H MAS NMR signatures of monomeric and dimeric model complexes and their temperature dependence augmented with theoretical calculations allows the assignment of monomeric and dimeric sites in the active UC-like catalyst as summarized in Fig. 6 above. This method was also applied to a range of molecular paramagnetic di-nuclear complexes, illustrating its potential in assignment ^1H spectra.

^1H MAS NMR investigations evidence the presence of mono-grafted sites interacting with nearby Si–O–Si bridges or surface OH groups at 155 ppm. In addition, the presence of dimeric surface sites with weak (III, 318 ppm) and strong Cr–Cr interactions (II, 22 ppm & 13 ppm), as well as C–H activated bridging Cr(III)-hydrides (IV, 13 ppm) is also proposed to be present at higher loadings. Our calculations also show that trace amounts of strongly adsorbed chromocene on silica (318 ppm) cannot be excluded. The proposed active site structure V with a predicted ^1H shift at 229 ppm is not detected experimentally, likely due to the low abundance (1–3%) and the potential overlap with other side bands.^{7,32} Overall, this proposal is consistent with previously published CO-IR studies, further supporting the nature of the surface species.¹⁰ This study also demonstrates that

computational approaches can now be of great help to decipher speciation in complex paramagnetic systems, by ascertaining the chemical shift assignments. This methodology would be applicable to elucidate the structure of relevant molecular and heterogeneous catalysts featuring paramagnetic centers.

4. Methodology and computational details

The isotropic chemical shift δ in NMR is given by

$$\delta = \sigma_{\text{ref}} - \sigma \quad (3)$$

where σ represents the isotropic part of the shielding tensor, and σ_{ref} represents the isotropic shielding constant of the observed nucleus in a reference compound. In contrast to diamagnetic species, the NMR spectra of paramagnetic systems are influenced by supplementary interactions between nuclear and electronic spins.^{14,33,34} Therefore, the shielding constant σ of a nucleus in a paramagnetic environment is described by the sum of the following 3 terms:

$$\sigma = \sigma_0 + \sigma_{\text{FC}} + \sigma_{\text{PC}} \quad (4)$$

where σ_0 is the shielding of the equivalent diamagnetic molecule, σ_{FC} the Fermi contact term and σ_{PC} the so-called pseudo-contact term. The Fermi contact shift (σ_{FC}) is reversely proportional to temperature (T) and proportional to the hyperfine coupling constant (A) according to the following equation:



$$\sigma_{\text{FC}} = -\frac{2\pi}{\gamma_1} g \mu_{\text{B}} A \frac{S(S+1)}{3k_{\text{B}}T} \quad (5)$$

where γ_1 is the gyromagnetic ratio of the nucleus, g is the g factor of the complex, μ_{B} is the Bohr magneton and $k_{\text{B}}T$ is the thermal energy. This term is the dominant contribution to the paramagnetic chemical shift of 3d-transition metal complexes and organic radicals.†

All calculations were performed using Gaussian 09. The diamagnetic shielding constant σ_0 , the g factor, and the hyperfine coupling constant A of Cr complexes were estimated at the B3LYP^{35–38}/cc-pVTZ³⁹ level, which was proved appropriate by Bagno and co-workers,¹⁴ and using the structures previously optimized at the B3LYP/BS1 level of theory (see ESI Section 4.1†) with BS1 being 6-31+G(d) for O and Cl, 6-31G(d) for C, H, and Si, and LANL2TZ/LANL2 for Cr.^{40,41} When crystal structures were available, only the positions of protons were optimized. The same approach was used for the prediction of supported monomeric and dimeric sites, where the silica surface was modeled by using RO = –Si(OH)₃.

Isotropic magnetic shielding tensors σ_{orb} were calculated using the gauge-independent atomic orbital (GIAO) approach.^{42–44} Isotropic hyperfine coupling constants A ^{45,46} between nuclear and electronic spins and σ_{orb} were calculated as weighted average values for different sets of equivalent protons. Paramagnetic chemical shifts for monomeric sites δ_{mon} are calculated by employing σ_0 , A and the molecular g factor in the equation below while using ferrocene as a reference complex (σ_{Fe}) and neglecting the pseudo-contact term δ_{PC} :

$$\delta_{\text{mon}} = \delta_0 + \delta_{\text{FC}} = \sigma_{\text{Fe}} - \sigma_0 + \frac{2\pi}{\gamma_1} g \mu_{\text{B}} A \frac{S(S+1)}{3k_{\text{B}}T} \quad (6)$$

For dimeric sites presenting electronic spin–spin interactions, we account for the Boltzmann distribution of global spin states besides the Fermi contact term (see Section 2.3 for a more detailed explanation). At first, the Fermi contact shifts are calculated for each global spin state using the A hyperfine coupling constant obtained on the high spin state. To get access to the energy splitting between global spin states ($E = S(S+1)J$), spin–spin coupling constants J were calculated using the high-spin and broken symmetry (BS)^{22,27,47} energies according to the Yamaguchi formula,¹⁴ which was calculated at the same level of theory used for the structure optimization (see ESI Section 1† for details).

Data availability

All data are provided in the ESI† and additional data can be available upon request.

Author contributions

A. N. and D. T. contributed equally and performed the synthesis of paramagnetic materials, Z. J. B. performed the NMR experiments, M. W. carried out the crystallographic analysis,

and A. N. carried out most of the calculations under the guidance of P.-A. P. P.-A. P. and C. C. designed and lead the project.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

† In the case of 3d-transition metal complexes, the contribution from the pseudo-contact term is usually negligible compared to δ_{FC} and often within the 1–10 ppm range for ¹H NMR. Only a few Å around the location of the spin density are affected. In the case of chromocene, we found that the contribution is not larger than 0.2 ppm and can therefore be neglected.

- M. P. McDaniel, in *Advances in Catalysis*, ed. B. C. Gates and H. Knözinger, Academic Press, 2010, vol. 53, pp. 123–606.
- M. F. Delley, F. Núñez-Zarur, M. P. Conley, A. Comas-Vives, G. Siddiqi, S. Norsic, V. Monteil, O. V. Safonova and C. Copéret, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 11624–11629.
- A. Ashuiev, F. Allouche, N. Wili, K. Searles, D. Klose, C. Copéret and G. Jeschke, *Chem. Sci.*, 2021, **12**, 780–792.
- H. R. Sailors and J. P. Hogan, *J. Macromol. Sci., Part A*, 1981, **15**, 1377–1402.
- F. J. Karol, G. L. Karapinka, C. Wu, A. W. Dow, R. N. Johnson and W. L. Carrick, *J. Polym. Sci., Part A: Polym. Chem.*, 1972, **10**, 2621–2637.
- F. J. Karol and C. Wu, *J. Polym. Sci., Part A: Polym. Chem.*, 1974, **12**, 1549–1558.
- S. L. Fu and J. H. Lunsford, *Langmuir*, 1990, **6**, 1774–1783.
- S. L. Fu and J. H. Lunsford, *Langmuir*, 1990, **6**, 1784–1792.
- A. Zecchina, G. Spoto and S. Bordiga, *Faraday Discuss. Chem. Soc.*, 1989, **87**, 149–160.
- D. Trummer, A. G. Nobile, P.-A. Payard, A. Ashuiev, Y. Kakiuchi, D. Klose, G. Jeschke and C. Copéret, *Chem. Sci.*, 2022, **13**(31), 11091–11098.
- K. H. Theopold, *Eur. J. Inorg. Chem.*, 1998, 15–24.
- K. H. Theopold, R. A. Heintz, S. K. Noh and B. J. Thomas, in *Homogeneous Transition Metal Catalyzed Reactions*, American Chemical Society, 1992, ch. 41, vol. 230, pp. 591–602.



- 13 M. Schnellbach, F. H. Köhler and J. Blümel, *J. Organomet. Chem.*, 1996, **520**, 227–230.
- 14 F. Rastrelli and A. Bagno, *Chem.–Eur. J.*, 2009, **15**, 7990–8004.
- 15 L. Peng, R. J. Clément, M. Lin and Y. Yang, in *NMR and MRI of Electrochemical Energy Storage Materials and Devices*, The Royal Society of Chemistry, 2021, pp. 1–70, DOI: [10.1039/9781839160097-00001](https://doi.org/10.1039/9781839160097-00001).
- 16 A. J. Pell and G. Pintacuda, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2015, **84–85**, 33–72.
- 17 A. J. Pell, G. Pintacuda and C. P. Grey, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2019, **111**, 1–271.
- 18 P. Fernández, H. Pritzkow, J. J. Carbó, P. Hofmann and M. Enders, *Organometallics*, 2007, **26**, 4402–4412.
- 19 D. E. DeRoshia, N. A. Arnet, B. Q. Mercado and P. L. Holland, *Inorg. Chem.*, 2019, **58**, 8829–8834.
- 20 D. Trummer, A. G. Nobile, P.-A. Payard, A. Ashuiev, Y. Kakiuchi, D. Klose, G. Jeschke and C. Copéret, *Chem. Sci.*, 2022, **13**, 11091–11098.
- 21 A. Borgogno, F. Rastrelli and A. Bagno, *Dalton Trans.*, 2014, **43**, 9486–9496.
- 22 *Metalloproteins: Methods and Protocols*, ed. J. C. Fontecilla-Camps, N. Yvain and Humana, Springer, 2014.
- 23 T. Polenova, R. Gupta and A. Goldbourt, *Anal. Chem.*, 2015, **87**, 5458–5469.
- 24 J. Novotný, L. Jeremias, P. Nimax, S. Komorovsky, I. Heinmaa and R. Marek, *Inorg. Chem.*, 2021, **60**, 9368–9377.
- 25 D. R. Kinney, I. S. Chuang and G. E. Maciel, *J. Am. Chem. Soc.*, 1993, **115**, 6786–6794.
- 26 M. D. Korzyński, Z. J. Berkson, B. Le Guennic, O. Cador and C. Copéret, *J. Am. Chem. Soc.*, 2021, **143**, 5438–5444.
- 27 N. Ferré, N. Guihéry and J.-P. Malrieu, *Phys. Chem. Chem. Phys.*, 2015, **17**, 14375–14382.
- 28 J. Blahut, A. L. Lejeune, S. Ehrling, I. Senkovska, S. Kaskel, F. M. Wisser and G. Pintacuda, *Angew. Chem., Int. Ed.*, 2021, **60**, 21778–21783.
- 29 M. H. Chisholm, F. A. Cotton, M. W. Extine and D. C. Rideout, *Inorg. Chem.*, 1979, **18**, 120–125.
- 30 S. E. Nefedov, A. A. Pasynskii, I. L. Eremenko, B. Orazsakhov, O. G. Ellert, V. M. Novotortsev, S. B. Katser, A. S. Antsyshkina and M. A. Porai-Koshits, *J. Organomet. Chem.*, 1988, **345**, 97–104.
- 31 K. Angermund, A. Doehring, P. W. Jolly, C. Krueger and C. C. Romao, *Organometallics*, 1986, **5**, 1268–1269.
- 32 S. L. Fu, M. P. Rosynek and J. H. Lunsford, *Langmuir*, 1991, **7**, 1179–1187.
- 33 F. H. Köhler, *Paramagnetic Complexes in Solution: The NMR Approach*, *eMagRes*, 2011, DOI: [10.1002/9780470034590.emrstm1229](https://doi.org/10.1002/9780470034590.emrstm1229).
- 34 P. Hrobárik, R. Reviakine, A. V. Arbuznikov, O. L. Malkina, V. G. Malkin, F. H. Köhler and M. Kaupp, *J. Chem. Phys.*, 2007, **126**, 024107.
- 35 S. H. Vosko, L. Wilk and M. Nusair, *Can. J. Phys.*, 1980, **58**, 1200–1211.
- 36 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
- 37 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 38 P. J. Stephens, F. J. Devlin, C. F. Chabalowski and M. J. Frisch, *J. Phys. Chem.*, 1994, **98**, 11623–11627.
- 39 N. B. Balabanov and K. A. Peterson, *J. Chem. Phys.*, 2005, **123**, 064107.
- 40 P. J. Hay and W. R. Wadt, *J. Chem. Phys.*, 1985, **82**, 299–310.
- 41 L. E. Roy, P. J. Hay and R. L. Martin, *J. Chem. Theory Comput.*, 2008, **4**, 1029–1031.
- 42 J. Gauss, *Chem. Phys. Lett.*, 1992, **191**, 614–620.
- 43 J. Gauss, *Berichte der Bunsengesellschaft für physikalische Chemie*, 1995, **99**, 1001–1008.
- 44 K. Wolinski, J. F. Hinton and P. Pulay, *J. Am. Chem. Soc.*, 1990, **112**, 8251–8260.
- 45 V. Barone, *Chem. Phys. Lett.*, 1996, **262**, 201–206.
- 46 N. Rega, M. Cossi and V. Barone, *J. Chem. Phys.*, 1996, **105**, 11060–11067.
- 47 P. W. Atkins, *Quanta: A Handbook of Concepts*, W. H. Freeman and Company, New York, 2nd edn, 2010.

