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## Two pathways of proton transfer reaction to (triphos)Cu( $\eta^1$ -BH<sub>4</sub>) *via* a dihydrogen bond [triphos = 1,1,1-tris(diphenylphosphinomethyl)ethane]<sup>†</sup>

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The interaction of the  $\eta^1$ -tetrahydroborate copper(i) complex (triphos)Cu( $\eta^1$ -BH<sub>4</sub>) (**1**) with proton donors [CF<sub>3</sub>CH<sub>2</sub>OH (TFE), (CF<sub>3</sub>)<sub>2</sub>CHOH (HFIP), (CF<sub>3</sub>)<sub>3</sub>COH (PFTB), PhOH, *p*-NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>OH (PNP), *p*-NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>N=NC<sub>6</sub>H<sub>4</sub>OH (PNAP), CF<sub>3</sub>OH] was a subject of a combined IR spectroscopic and theoretical investigation. Spectral ( $\Delta\nu$ ) and thermodynamic ( $\Delta H$ ) parameters of dihydrogen bond (DHB) formation were determined experimentally. The terminal hydride ligand (characterized by the basicity factor  $E_b(\text{BH}) = 0.87 \pm 0.01$ ) is found to be a site of proton transfer which begins with nucleophilic substitution of BH<sub>4</sub><sup>−</sup> by the alcohol oxygen atom on the copper center (*BH pathway*). The activation barrier computed for (CF<sub>3</sub>)<sub>2</sub>CHOH in CH<sub>2</sub>Cl<sub>2</sub> –  $\Delta G_{273\text{ K}}^\ddagger = 20.6 \text{ kcal mol}^{-1}$  – is in good agreement with the experimental value ( $\Delta G_{270\text{ K}}^\ddagger = 20.0 \text{ kcal mol}^{-1}$ ). An abnormal dependence of the reaction rate on the proton donor strength found experimentally in dichloromethane is explained computationally on the basis of the variation of the structural and energetic details of this process with the proton donor strength. In the second reaction mechanism found (*CuH pathway*), DHB complexes with the initial ROH coordination to the bridging hydride lead to B–H<sub>br</sub> bond cleavage with BH<sub>3</sub> elimination. “Copper assistance” *via* the Cu...O interaction is not involved. This mechanism can be evoked to explain the occurrence of proton transfer in coordinating solvents.

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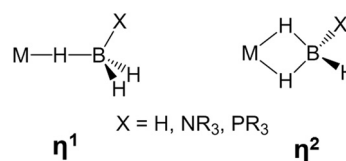
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## Introduction

The activation of E–H (E = H, C, Si, B, N) bonds by transition metal complexes is of great interest to many research fields and constitutes a rapidly developing area of main group element chemistry, organometallic chemistry and catalysis. B–H bond activation occurs in a large variety of reactions involving boron hydrides, such as metal catalyzed hydroboration,<sup>1,2</sup> and dehydrocoupling of amine- and phosphine-boranes.<sup>3–10</sup> When B–H bonds coordinate to transition metal atoms,  $\sigma$ -complexes may be characterized in solution and even isolated in the solid state (Scheme 1).<sup>5,11–15</sup> Such complexes are generally considered as intermediates of the B–H bond activation process.<sup>10,12,13,15,16</sup>



Scheme 1 Types of metal borohydride  $\sigma$ -complexes.

In previous studies carried out on (Ph<sub>3</sub>P)<sub>2</sub>Cu( $\eta^2$ -BH<sub>4</sub>)<sup>17</sup> and (PP<sub>3</sub>)Ru( $\eta^1$ -BH<sub>4</sub>),<sup>18</sup> PP<sub>3</sub> =  $\kappa^4$ -P(CH<sub>2</sub>CH<sub>2</sub>PPh<sub>2</sub>)<sub>3</sub> (Scheme 2), we have shown that coordination of the BH<sub>4</sub><sup>−</sup> ligand to the metal decreases the proton accepting ability of the tetrahydroborate ligand (if compared with the non-coordinated BH<sub>4</sub><sup>−</sup> anion).

In the presence of protic reagents, the simultaneous existence of multiple hydride centres (bridged and terminal BH and MH hydride ligands) may lead to the formation of a large variety of dihydrogen-bonded complexes (DHB), but only a few of them are real intermediates of the proton transfer reaction.<sup>17,18</sup>

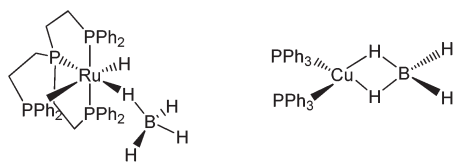
As part of an ongoing systematic investigation addressing the reactivity of transition metal tetrahydroborates, we report here a combined theoretical (DFT) and spectroscopic study (IR, NMR) of the interaction of the known copper(i)

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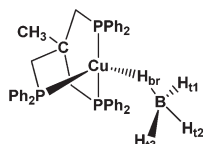
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Scheme 2



Scheme 3

tetrahydroborate complex (triphos)Cu( $\eta^1$ -BH<sub>4</sub>) (**1**) (Scheme 3), [triphos =  $\kappa^3$ -H<sub>3</sub>CC(CH<sub>2</sub>PPh<sub>2</sub>)<sub>3</sub>] with a number of proton donors (OH acids). This study is aimed at unraveling the proton transfer mechanism to monodentate tetrahydroborate ( $\eta^1$ -BH<sub>4</sub>) complexes, accompanied by the release of molecular hydrogen and at assessing the influence of the DHB complex type and the metal atom nature on the H<sub>2</sub> evolution reaction.

## Results and discussion

### Theoretical investigation of DHB complexes

To study and model the electronic structure of the DHB complexes, *in silico* analysis was carried out using both the “real” complex **1** and the related copper complex bearing the methylated analogue of triphos, *i.e.* (triphos<sup>Me</sup>)Cu( $\eta^1$ -BH<sub>4</sub>) (**2**) (Fig. 1), where triphos<sup>Me</sup> is  $\kappa^3$ -H<sub>3</sub>CC(CH<sub>2</sub>PMe<sub>2</sub>)<sub>3</sub>.

Replacement of triphos with triphos<sup>Me</sup> does not have any meaningful effect on the initial hydride geometry, as the optimized structures of complexes **1** and **2** do not differ by more than 0.08 Å for Cu–P bond distances and 0.03 Å for the B–H bond distances (Table S1†). The angles  $\angle$ P–Cu–P and  $\angle$ H–B–H do not differ more than  $\pm 5^\circ$ . The complexes **1** and **2** form the same set of DHB complexes with all alcohols, featuring similar geometries (Fig. S3–S6†). This validates the use of complex **2** as a model.

Coordination of the tetrahydroborate anion to the metal atom leads to a variety of possible DHB complexes involving BH ligands.<sup>17,18</sup> The geometry optimization (DFT/M06) for 2-HOR gives five different types of DHB complexes (Scheme 4): bifurcated DHB complexes with preferential coordination to BH<sub>br</sub> (**IIbc**) (this type was found for all alcohols except for MeOH); complex with coordination to the Cu atom (**IIcb**); bifurcated DHB complexes on two BH<sub>term</sub> (**IIa**) and BH<sub>term</sub> and BH<sub>br</sub> (**IIab**) and trifurcated DHB complexes on BH<sub>br</sub> and two BH<sub>term</sub> (**IIIab**).

Trifurcated DHB complexes such as **IIIab** were previously described for (PP<sub>3</sub>)RuH( $\eta^1$ -BH<sub>4</sub>),<sup>18</sup> so we can assume that for-

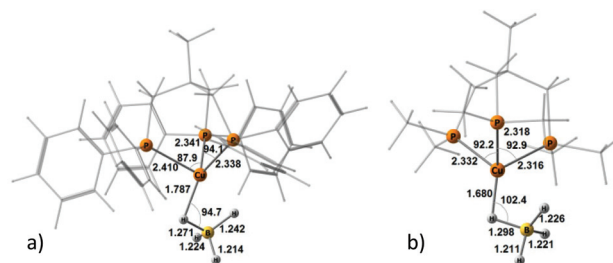
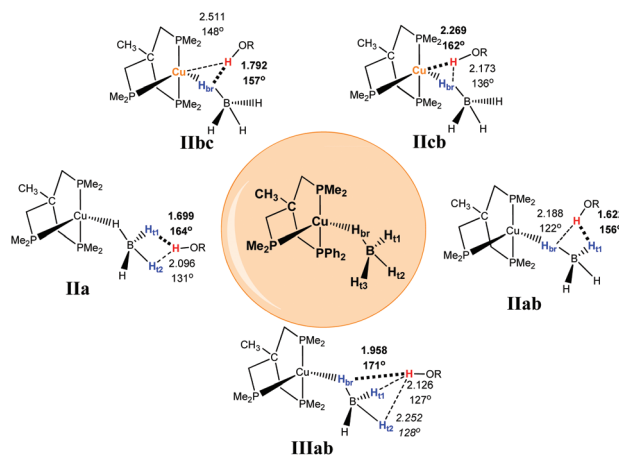


Fig. 1 M06-optimized geometries of “real” molecule **1** (a) and Me-model **2** (b).



Scheme 4 Possible types of DHB complexes between **2** and ROH. The structural parameters reported on the scheme as a representative example are coming from the adducts between **2** and (CF<sub>3</sub>)<sub>2</sub>CHOH (HFIP).

mation of trifurcated DHB complexes is possible only in the case of a monodentate tetrahydroborate ligand ( $\eta^1$ -BH<sub>4</sub>).

### Structural analysis

The intermolecular distances O–H...H–B (Table S2†) are in the range 1.622–1.985 Å for primary contacts (shorter) and 2.062–2.382 Å for secondary (longer) interactions, but, irrespective of the DHB complex type, the H...H contacts are shorter than the sum of van der Waals radii for the two hydrogen atoms [ $\Sigma$ vdW(H–H) = 2.4 Å]. For primary interactions the O–H...H(B) angles range from 171 to 152°. A Cu...H separation smaller than  $\Sigma$ vdW(Cu–H) = 2.6 Å and the O–H...Cu angle falling between 140 and 170° were chosen as a criterion for classifying the Cu...HO contact as a hydrogen-bonded (HB) interaction. Regardless of the DHB complex type, the elongation of both the OH bond [0.008–0.020 Å] and the CuH bond [0.007–0.032 Å] was observed (Table S3†). The maximum O–H bond elongation was observed in complexes with coordination of ROH to BH<sub>term</sub> (**IIa**, **IIab** and **IIIab**), while the biggest Cu–H bond elongation was found for bifurcated DHB complexes **IIa** and **IIab**. Thus, coordination to BH<sub>term</sub> induces a general weakening of the Cu<sup>+</sup>...BH<sub>4</sub><sup>−</sup> interaction with concomitant lengthening of the BH<sub>term</sub> bond. Elongation of the BH<sub>br</sub> bond



was observed only in complexes **IIbc** and **IIcb** (with the exception of complex **IIcb\_TFE**).

### Electron-density analysis

Electron density redistribution that occurs upon DHB formation was characterized in terms of the number of electrons shared between the atoms involved in the interaction. Different approaches were used: natural population analysis (NPA),<sup>19</sup> Wiberg bond indices (WBIs)<sup>20</sup> and Bader's Quantum Theory of "Atoms in Molecules" (QTAIM).<sup>21–23</sup>

Analysis of the molecular electrostatic potential (MEP) of **2** reveals an enhanced electron density on the  $\text{BH}_4^-$  ligand with two minima ( $V_{\text{min}}$ ) at  $-52.1$  and  $-54.3$  kcal mol<sup>-1</sup>, located at  $\text{BH}_{\text{term}}$  ligands (Fig. 2). This fact suggests the preferred proton donor coordination on  $\text{BH}_{\text{term}}$  ligands.

As expected, upon DHB formation the charge on the proton of ROH becomes more positive, whereas the charge on the interacting hydridic hydrogen(s) becomes more negative (Table S8†).

Within the framework of the QTAIM theory, a hydrogen bond is characterized by the presence of a (3, -1) critical point; this allows for its easy identification and differentiation from other types of interactions.<sup>24</sup> Despite the existence of several short intermolecular contacts in all DHB complexes **2-HOR**, a critical point (3, -1) was found only for the shortest contact with the most linear O–H...H(B) and O–H...Cu arrangement (Table 1, Fig. S8–S13, Table S11†). The presence of additional interactions causes deviation of the OH...X moiety from linearity and mirrors the values of the H...H and H...Cu bond ellipticity. The maximum ellipticity value was

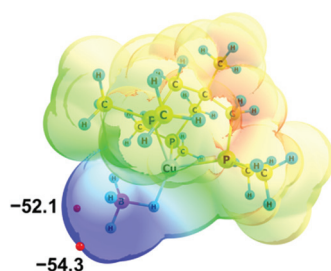


Fig. 2 Molecular electrostatic potential map of **2** (at a distance 1.5 vdW) in the RGB scale 0.06 ÷ -0.09 a.u. Electron density minima ( $V_{\text{min}}$  in kcal mol<sup>-1</sup>) marked as red spheres.

Table 1 Energy of the H...X bond critical point ( $E_{\text{H...X}}$ ), Laplacian of the electron density at the critical point ( $\nabla^2\rho_{\text{c}}$ ), electron densities at the H...H bond critical point ( $\rho_{\text{c}}$ ) and bond ellipticity ( $\epsilon$ ) for **2-HFIP** DHB complexes

	$E_{\text{H...X}}$ , kcal mol <sup>-1</sup>	$\nabla^2\rho_{\text{c}}$ , au	$\rho_{\text{c}}$ , au	$\epsilon$	Contact
<b>IIbc</b>	-3.2	0.047	0.019	0.25	$\text{BH}_{\text{br}}\cdots\text{HO}$
<b>IIcb</b>	-3.0	0.046	0.019	0.17	$\text{Cu}\cdots\text{HO}$
<b>IIa</b>	-5.0	0.064	0.026	0.35	$\text{BH}_{\text{t1}}\cdots\text{HO}$
<b>IIab</b>	-5.6	0.067	0.028	0.20	$\text{BH}_{\text{t1}}\cdots\text{HO}$
<b>IIIab</b>	-3.2	0.051	0.018	1.39	$\text{BH}_{\text{br}}\cdots\text{HO}$

found in complexes **HFIP-IIIab** ( $\epsilon = 1.39$ ), **TFE-IIcb** ( $\epsilon = 1.03$ ) and **MeOH-IIab** ( $\epsilon = 0.97$ ).

WBI (bond population) is a parameter that characterizes the order of the bond between two atoms.<sup>20</sup> The values of WBI for the primary H...H contact in the DHB complexes **2-HOR** range from 0.008 to 0.058, while for secondary interactions the WBI values are less than 0.007 (Table S9†). In **IIbc** complexes, for TFE and HFIP, the WBI value of the OH...Cu contact is about 0.031, whereas in the **IIcb** complex WBI values range from 0.039 to 0.061. The WBI values for interacting OH and BH bonds decrease according to its elongation.

The electron density shift that takes place during the DHB formation was analyzed using the "base-to-acid" donation energy estimated from 2<sup>nd</sup>-order perturbative analysis ( $E^2$ ) of donor-acceptor interactions as implemented in NBO.<sup>25,26</sup> Typically, the hydrogen bond entails the transfer of the electron density from the HOMO orbital of the base to an empty  $\sigma_{\text{XH}}^*$  orbital (LUMO) of the acid. In all the complexes analyzed herein, the main  $\sigma_{\text{base}} \rightarrow \sigma_{\text{OH}}^*$ -donation corresponds to the shortest H...H contact. Secondary contacts possess weaker but still valuable donation and notably only the energy of primary donation is sensitive to the proton donor strength (Table S10†). For DHB complexes **IIa**, the identified donations are from two bonding molecular orbitals (MO) of BH ligands to the anti-bonding MO of the OH-group:  $\sigma_{\text{BH}_{\text{t1}}} \rightarrow \sigma_{\text{OH}}^*$  and  $\sigma_{\text{BH}_{\text{t2}}} \rightarrow \sigma_{\text{OH}}^*$ . For complexes **IIab**, the main donations are from the bonding MO of the BH ligand to the three-centered MO of the Cu–H–B bond:  $\sigma_{\text{BH}_{\text{t1}}} \rightarrow \sigma_{\text{OH}}^*$  and  $\tau_{\text{Cu-H-B}} \rightarrow \sigma_{\text{OH}}^*$ . Complexes **IIbc** and **IIcb** are both characterized by  $\tau_{\text{Cu-H-B}} \rightarrow \sigma_{\text{OH}}^*$  and donation from electron pairs on the Cu d-orbital to the alcohol ( $n_{\text{Cu}}^{\text{d}} \rightarrow \sigma_{\text{OH}}^*$ ). For **IIbc** the  $\tau_{\text{Cu-H-B}} \rightarrow \sigma_{\text{OH}}^*$  energy is larger, whereas for **IIcb** the  $n_{\text{Cu}}^{\text{d}} \rightarrow \sigma_{\text{OH}}^*$  has a dominant impact on the total "base-to-acid" donation energy. Trifurcated complex **IIIab** with TFE and HFIP is characterized by a strong stabilization energy  $E^2$  of the primary  $\tau_{\text{Cu-H-B}} \rightarrow \sigma_{\text{OH}}^*$  interaction and two weak  $\sigma_{\text{BH}_{\text{t1}}} \rightarrow \sigma_{\text{OH}}^*$  and  $\sigma_{\text{BH}_{\text{t2}}} \rightarrow \sigma_{\text{OH}}^*$  donations. For all the complexes studied, the secondary interactions provide over 15% of the total amount of stabilization energy ( $E^2$ ). Thus, NBO analysis confirms the multifurcated nature of DHB complexes through the existence of several donor-acceptor interactions.

### Interaction energies

Table 2 shows the formation energies of the DHB complexes, determined as the energy difference between the adduct and the isolated reactants (Table S4†) as well as the DHB energies derived from the AIM data. When the basis set superposition error (BSSE) was taken into account, a significant lowering of the complexation energy was obtained (up to -20%), especially for TFE and HFIP. The inclusion of nonspecific solvent effects ( $\text{CH}_2\text{Cl}_2$  (DCM), THF) significantly lowers the formation energy, with bifurcated DHB complexes **IIa** and **IIab** remaining the most stable. The formation enthalpies of the DHB complexes in the gas phase, calculated from the  $\Delta\nu_{\text{OH}}$  (difference between stretching vibration frequencies of the OH-group of free and bonded alcohol),  $\Delta H_{\text{theor}}^{\circ}$  ( $\Delta\nu_{\text{OH}} = 3.7\text{--}6.0$  kcal mol<sup>-1</sup> (Table 2) are comparable to the experimental enthalpy values



**Table 2** Formation energies of DHB complexes 2-HFIP (in kcal mol<sup>-1</sup>)

	$\Delta E$	$\Delta E_{\text{ZPVE}}$	$\Delta E_{\text{BSSE}}$	$\Delta H_{\text{DCM}}$	$\Delta H_{\text{THF}}$	$\Delta H_{\text{theor}}^{\circ} (\Delta \nu_{\text{OH}})$	$E_{\text{H}\cdots\text{X}}^a$
<b>IIbc</b>	-23.4	-19.8	-17.2	-5.2	-5.9	-3.7	-3.2
<b>IIcb</b>	-23.7	-21.6	-17.3	-5.9	-6.5	-3.9	-3.0
<b>IIa</b>	-25.9	-23.4	-20.4	-8.8	-9.3	-6.0	-5.0
<b>IIab</b>	-25.3	-22.8	-19.0	-8.8	-9.4	-5.7	-5.6
<b>IIIab</b>	-23.8	-20.7	-18.4	-6.8	-7.3	-4.4	-3.2

$$^a E_{\text{H}\cdots\text{X}} = 0.5 \times V(r).$$

determined in solution,  $\Delta H_{\text{exp}}^{\circ} (\Delta \nu_{\text{OH}}) = 3.2\text{--}5.4$  kcal mol<sup>-1</sup>. The energies  $E_{\text{H}\cdots\text{X}}$ , derived from the AIM data, are also larger for complexes **IIa** and **IIab**.

### Frequency analysis

All four stretching vibrations of the BH<sub>4</sub><sup>-</sup> group in (triphos)Cu(η<sup>1</sup>-BH<sub>4</sub>) are IR-active. The frequency analysis for the “real” molecule **1** gives three stretching vibrations for BH<sub>term</sub>:  $\nu_{\text{BHterm}}^{\text{as1}}$  at 2492 cm<sup>-1</sup>,  $\nu_{\text{BHterm}}^{\text{as2}}$  at 2419 cm<sup>-1</sup>,  $\nu_{\text{BHterm}}^{\text{s}}$  at 2313 cm<sup>-1</sup> as well as  $\nu_{\text{BHbr}}$  at 2114 cm<sup>-1</sup>. Frequency analysis for Me-model **2** gives lower energy for the  $\nu_{\text{BHterm}}^{\text{as}}$  stretching vibrations (2468, 2385 cm<sup>-1</sup>) as well as for  $\nu_{\text{BHterm}}^{\text{s}}$  (2362 cm<sup>-1</sup>) and  $\nu_{\text{BHbr}}$  (1919 cm<sup>-1</sup>) modes. Despite this frequency difference, the relative band shifts ( $\Delta \nu$ ) entailed by DHB formation are of the same order.

Formation of intermolecular complexes leads to the low-frequency shift of the stretching vibrations involved in DHB (Tables 3 and S5<sup>†</sup>). The low-frequency shift of  $\nu_{\text{OH}}$  falls in the range from 144 to 357 cm<sup>-1</sup>, while the low-frequency shift of  $\nu_{\text{BHterm}}^{\text{as1}}$  is between -4 and -17 cm<sup>-1</sup>. The largest  $\Delta \nu_{\text{BH}}$  values (from -11 to -42 cm<sup>-1</sup>) were found for the complexes **IIa** and **IIab**. The low-frequency shifts  $\nu_{\text{BHbr}}$  ranging from -13 to -39 cm<sup>-1</sup> were computed for the complexes of **IIbc** and **IIcb** types only.

### Spectral investigation of DHB

In order to assess the proton accepting ability of the BH ligand in (triphos)Cu(η<sup>1</sup>-BH<sub>4</sub>) (**1**), its interaction with fluorinated alcohols [CF<sub>3</sub>CH<sub>2</sub>OH (TFE), (CF<sub>3</sub>)<sub>2</sub>CHOH (HFIP) and (CF<sub>3</sub>)<sub>3</sub>COH (PFTB)] and phenols [PhOH, *p*-NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>OH (PNP) and *p*-NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>N=NC<sub>6</sub>H<sub>4</sub>OH (PNAP)] was investigated in low polar media (DCM, THF) by means of IR spectroscopy in a wide temperature range (190–310 K). Upon DHB formation in DCM, the band assigned to OH stretching vibration ( $\nu_{\text{OH}}$ ) decreases

and in the same time a new broad low-frequency band ( $\nu_{\text{OH}}^{\text{bond}}$ ) appears, corresponding to stretching vibrations of the OH group linked by DHB (OH<sup>δ+</sup>...<sup>δ-</sup>HB) (Table 4).

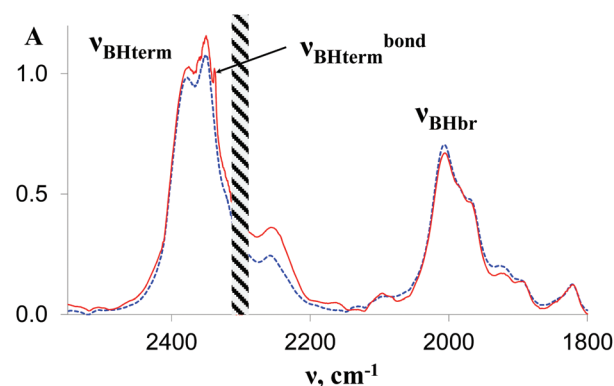
These data allow for the calculation of the basicity factor of the BH ligand  $E_j(\text{BH}) = 0.87 \pm 0.01$ . The  $E_j(\text{BH})$  values obtained from the computed frequencies are in good agreement with the experimental outcomes ( $E_j(\text{BH}) = 0.87 \pm 0.01$  for **IIa** complexes and  $0.84 \pm 0.01$  for **IIab**) (Tables S6 and S7, Fig. S7<sup>†</sup>). This estimation shows that the proton accepting ability of the BH ligand in (triphos)Cu(η<sup>1</sup>-BH<sub>4</sub>) is lower than that of BH in (Ph<sub>3</sub>P)<sub>2</sub>Cu(η<sup>2</sup>-BH<sub>4</sub>) ( $E_j = 0.91$ )<sup>17</sup> and (PP<sub>3</sub>)RuH(η<sup>1</sup>-BH<sub>4</sub>) ( $E_j = 0.98$ ).<sup>18</sup>

The IR spectra of complex **1** alone and in the presence of an excess of TFE (5–10 equiv.) in THF at 190 K (Fig. 3) show a change of the symmetry of BH stretching vibration bands of **1** ( $\nu_{\text{BHterm}}^{\text{as}} = 2387$  cm<sup>-1</sup> and new bands  $\nu_{\text{BHterm}}^{\text{s}} = 2351$  cm<sup>-1</sup>) and

**Table 4** Spectral (in cm<sup>-1</sup>) and energetic (in kcal mol<sup>-1</sup>) characteristics of DHBs in DCM

Proton donor	$P_i^a$	$\nu_{\text{OH}}^{\text{free}}$	$\nu_{\text{OH}}^{\text{bond}}$	$\Delta \nu_{\text{OH}}$	$\Delta H_{\text{exp}}^{\circ} (\Delta \nu_{\text{OH}})$
TFE	0.90	3600	3419	-181	-3.8
PhOH	1.00	3569	3363	-206	-4.0
HFIP	1.07	3576	3372	-204	-4.0
PNAP	1.23	3546	3267	-279	-5.0
PNP	1.27	3546	3245	-301	-5.3
PFTB	1.33	3525	3245	-290	-5.2

<sup>a</sup>  $P_i$  – acidity factors as characteristics of the acid proton-donating ability were taken from ref. 17 and 27.

**Fig. 3** IR spectra in the  $\nu_{\text{BH}}$  region of **1** (0.027 M, blue dashed line) and in the presence of TFE (10 equiv.) in THF;  $l = 0.4$  mm,  $T = 190$  K.**Table 3** Spectral parameters (OH band shifts,  $\Delta \nu$ , in cm<sup>-1</sup>, and changes of its integral intensity,  $\Delta A_{\text{OH}}$ , in km mol<sup>-1</sup>) upon DHB formation for 2-HFIP

	$\Delta \nu_{\text{OH}}$	$\Delta A_{\text{OH}}$	$\Delta \nu_{\text{BHterm}}^{\text{as1}}$	$\Delta \nu_{\text{BHterm}}^{\text{as2}}$	$\Delta \nu_{\text{BHterm}}^{\text{s}}$	$\Delta \nu_{\text{BHbr}}$
<b>IIbc</b>	-194	361	-11	55	50	-26
<b>IIcb</b>	-211	453	-16	46	33	-26
<b>IIa</b>	-367	544	8	30	-34	106
<b>IIab</b>	-342	496	-7	44	-42	128
<b>IIIab</b>	-240	453	9	50	46	46



the appearance of bonded BH-groups vibration,  $\nu_{\text{BHterm}}^{\text{bond}} = 2338 \text{ cm}^{-1}$  ( $\Delta\nu = -49 \text{ cm}^{-1}$ ).

### Spectral investigation of the proton transfer reaction

Complex **1** reacts with excess proton donors (TFE, HFIP) in DCM even at low temperatures (200–250 K). Above 250 K,  $\text{H}_2$  evolution is observed even when a stoichiometric amount of proton donors is used. The initial bands  $\nu_{\text{BHterm}}$  and  $\nu_{\text{BHbr}}$  of both **1** and its DHB complexes decrease in intensity and the spectral picture characteristic of  $[(\text{RO})\text{BH}_3]^-$  species formation is observed. At the same time, new low-frequency bands at 2109 and 2069  $\text{cm}^{-1}$  are observed, ascribed to products of further alcoholysis,  $[(\text{RO})_n\text{BH}_{(4-n)}]^-$  ( $n = 2-3$ ) (Fig. 4). The identity of the final metal containing reaction product  $[(\text{triphos})\text{Cu}]^+[(\text{RO})_4\text{B}]^-$  is confirmed by NMR spectroscopy ( $^{31}\text{P}\{^1\text{H}\}$ : 31.29 ppm,  $^{11}\text{B}\{^1\text{H}\}$ : 2.49 ppm). A similar position of  $^{11}\text{B}$  resonance was reported for  $[(\text{RO})_4\text{B}]^-$  species characterized by NMR spectroscopy and X-ray diffraction.<sup>28,29</sup> The low-field position of phosphorus resonance supports the formation of the  $[(\text{triphos})\text{Cu}]^+$  cation characterized in ref. 30 in the presence of weakly coordinating counter ions. This behavior is different from that observed for  $(\text{Ph}_3\text{P})_2\text{Cu}(\eta^2\text{-BH}_4)$ , which gives a dimeric product  $\{[(\text{Ph}_3\text{P})_2\text{Cu}]_2(\mu, \eta^2: \eta^2\text{-BH}_4)\}^+[(\text{RO})_4\text{B}]^-$  upon reaction with alcohols.<sup>17</sup> It should be noted that alcoholysis of **1** does proceed in THF in contrast to  $(\text{Ph}_3\text{P})_2\text{Cu}(\eta^2\text{-BH}_4)$ ,<sup>17</sup> which reacts with weak acids only in DCM. Protonation of **1** in THF takes more time and requires higher temperatures (>250 K) than in DCM, suggesting a different reaction mechanism (*vide infra*).

The reaction scheme and its kinetic equation (Scheme 5 and eqn (1) and (2)) are similar to that obtained previously for hydride protonation in  $(\text{Ph}_3\text{P})_2\text{Cu}(\eta^2\text{-BH}_4)$ .<sup>17</sup> The observed rate constants (270–315 K) range from  $(3.5 \pm 0.1) \times 10^{-4}$  to

**Table 5** Kinetic data for protonation of **1** by different alcohols in DCM

Alcohol	Ratio	$k_{\text{obs}}$ , ( $\text{M s}^{-1}$ )	Temperature
TFE	1 : 1	$(4.9 \pm 0.1) \times 10^{-4}$	270 K
	1 : 2	$(6.4 \pm 0.1) \times 10^{-4}$	
HFIP	1 : 1	$(3.5 \pm 0.1) \times 10^{-4}$	270 K
	1 : 2	$(5.4 \pm 0.1) \times 10^{-4}$	
HFIP	1 : 1	$(4.0 \pm 0.1) \times 10^{-4}$	290 K
HFIP	1 : 1	$(8.5 \pm 0.1) \times 10^{-4}$	300 K
HFIP	1 : 1	$(4.7 \pm 0.1) \times 10^{-3}$	315 K

$(4.7 \pm 0.1) \times 10^{-3} \text{ s}^{-1}$  (Table 5). From the Eyring analysis of the temperature dependence the activation parameters were obtained for the reaction with HFIP:  $\Delta H^\ddagger = 9.0 \pm 0.5 \text{ kcal mol}^{-1}$  and  $\Delta S^\ddagger = -44 \pm 2 \text{ cal (mol K)}^{-1}$ . A high negative entropy value indicates a highly organized transition state as was found previously for  $(\text{Ph}_3\text{P})_2\text{Cu}(\eta^2\text{-BH}_4)$ .<sup>17</sup>

$$-\frac{d[a]}{dt} = k_{\text{obs}}[a] = \frac{k_2 \cdot k_3}{k_{-2} + k_3} \frac{K_1 \cdot [a][b]}{1 + K_1[b]} \quad (1)$$

where  $[a] = (\text{triphos})\text{Cu}(\eta^1\text{-BH}_4)$ ,  $[b] = [\text{HOR}]$

$$k_3 \gg k_{-2}; K_1[b] \ll 1 \Rightarrow k_{\text{obs}} \sim K_1 k_2 [b] \quad (2)$$

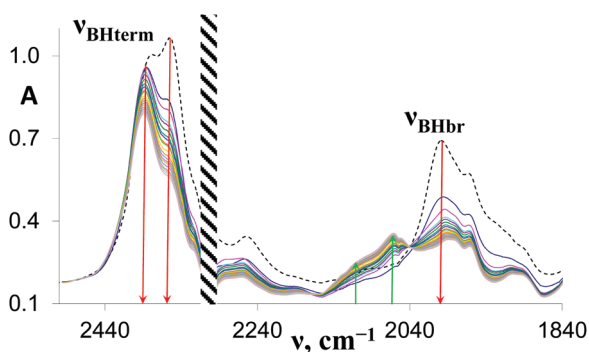
### Quantum-chemical investigation of the proton transfer reaction mechanism

The quantum-chemical analysis of the proton transfer mechanism was performed at the DFT/M06 level of theory for the model compound  $(\text{triphos}^{\text{Me}})\text{Cu}(\eta^1\text{-BH}_4)$  (**2**). The proton transfer reaction at the  $\text{BH}_{\text{term}}$  site mediated by the DHB complex of **IIa** type (BH pathway) can be divided into three elementary steps: (A) nucleophilic substitution of the  $\text{BH}_4^-$  ligand by the alcohol oxygen atom on the copper center with concomitant Cu–H bond dissociation and O–H bond activation due to the oxygen coordination to the Lewis acidic site (Cu); (B) facilitated by the O–H bond activation, the proton transfer *via* the DHB intermediate  $(\text{triphos}^{\text{Me}})\text{Cu}(\eta^1\text{-O}^+(\text{R}))\text{H}\cdots\text{H}_4\text{B}$  that leads to the formation of the  $\eta^2\text{-H}_2$  complex  $(\text{triphos}^{\text{Me}})\text{Cu}(\eta^1\text{-O}(\text{R}))\cdots(\eta^2\text{-H}_2)\text{BH}_3$ ; (C) elimination of the  $\text{H}_2$  molecule and formation of the reaction product  $(\text{triphos}^{\text{Me}})\text{Cu}(\eta^1\text{-O}(\text{R})\text{BH}_3)$ .

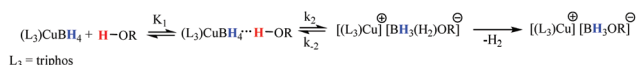
Remarkably, the course of proton transfer *via* the BH pathway depends on the proton donor strength. Thus, for a strong proton donor like  $\text{CF}_3\text{OH}$  the first transition state ( $\text{TS1}_{\text{BH}}^{\text{A+B}}$ ) is concerted, combining the first two elementary steps: nucleophilic substitution (A) and proton transfer (B) (Fig. 5 and Fig. S14 and S15†) followed by low-barrier  $\text{H}_2$  evolution ( $\text{TS1}_{\text{BH}}^{\text{C}}$ ). The transition state  $\text{TS1}_{\text{BH}}^{\text{A+B}}$  is a six-membered cycle  $[(\text{B})\text{-H-Cu}\cdots\text{O}\cdots\text{H}\cdots\text{H}(\text{B})]$  (Fig. S16†), similar to what was found for  $(\text{Ph}_3\text{P})_2\text{Cu}(\eta^2\text{-BH}_4)$ .<sup>17</sup>

For weaker proton donors (HFIP, TFE, MeOH) the first transition state ( $\text{TS1}_{\text{BH}}^{\text{A}}$ ) is simple nucleophilic substitution, while the second transition state is concerted ( $\text{TS2}_{\text{BH}}^{\text{B+C}}$ ) and includes simultaneous proton transfer (B) and  $\text{H}_2$  dissociation (C) stages (Fig. 6).

The concerted stage (B+C) for HFIP, TFE and MeOH is the rate determining step, and it resembles simple protonation of



**Fig. 4** Time-evolution of IR spectra in the  $\nu_{\text{BH}}$  region during protonation of **1** (0.05 M, dotted line) by HFIP (0.10 M) in DCM,  $l = 1.0 \text{ mm}$ ,  $T = 270 \text{ K}$ . Shaded area denotes the region masked by solvent absorption. Total reaction time: 4 h. Spectra were recorded every 5–10 min.



**Scheme 5**



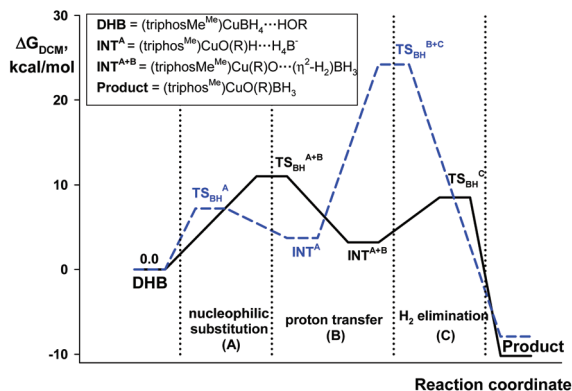


Fig. 5 Energy profiles for the protonation of 2 by CF<sub>3</sub>OH (black solid line) and HFIP (blue dashed line) via the BH pathway.

BH<sub>4</sub><sup>−</sup> by a medium strength acid like TFE.<sup>31</sup> Comparing the activation energies of BH<sub>4</sub><sup>−</sup> protonation by ROH<sup>31</sup> (Table 6) and by (triphos<sup>Me</sup>)Cu(η<sup>1</sup>-O(R)H) (TS<sub>BH</sub><sup>B+C</sup>, Fig. 5), a 15–25 kcal mol<sup>−1</sup> barrier lowering is recorded when the alcohol is coordinated to copper. The calculated value of the activation barrier for TFE is lower than that for HFIP (Table 6), in full agreement with the experimental observations (Table 5). This is obviously connected with the different “acidity enhancement” upon alcohol coordination to copper and formation of the active (triphos<sup>Me</sup>)Cu(η<sup>1</sup>-O(R)H) species. We propose that TFE has the best balance between the initial OH acidity and the basicity of the oxygen atom interacting with Cu. The combination of these two factors leads to the maximal lowering of the activation barrier of the rate determining step.

A second possible proton transfer mechanism is the protonation of the CuH site (CuH pathway) going through the formation of DHB complexes at the BH<sub>br</sub> site (IIbc and IIab). The first stage of the process (TS1<sub>CuH</sub>) implies the concerted B–H<sub>br</sub> bond dissociation and transfer of the H<sub>O</sub> proton to the Cu–H<sub>br</sub> yielding the [(triphos<sup>Me</sup>)Cu(η<sup>2</sup>-H<sub>2</sub>)]<sup>+</sup> complex weakly bound to the [(RO)BH<sub>3</sub>]<sup>−</sup> counterion (Fig. 7). The subsequent H<sub>2</sub> elimination gives the (triphos<sup>Me</sup>)Cu(η<sup>1</sup>-O(R)BH<sub>3</sub>) product *via* a negli-

Table 6 A comparison of the activation barriers for the BH pathway and the CuH pathway (ΔG<sub>DCM</sub><sup>†</sup>, kcal mol<sup>−1</sup>)

	BH pathway			CuH pathway	
	TS1 <sub>BH</sub> <sup>a</sup>	TS2 <sub>BH</sub> <sup>b</sup>	TS <sub>BH4</sub> <sup>−c</sup>	TS1 <sub>CuH</sub>	TS2 <sub>CuH</sub>
CF <sub>3</sub> OH	11.0	5.4	—	12.5	2.1
HFIP	7.2	20.6	35.1	33.2	1.3
TFE	10.4	18.9	44.5	33.3	0.9
MeOH	11.3	35.7	53.3	36.4	0.6

<sup>a</sup> TS1<sub>BH</sub> is TS1<sub>BH</sub><sup>A+B</sup> for CF<sub>3</sub>OH or TS1<sub>BH</sub><sup>A</sup> for other alcohols. <sup>b</sup> TS2<sub>BH</sub> is TS2<sub>BH</sub><sup>C</sup> for CF<sub>3</sub>OH or TS2<sub>BH</sub><sup>B+C</sup> for other alcohols. <sup>c</sup> Activation barriers of concerted proton transfer and hydrogen evolution to BH<sub>4</sub><sup>−</sup> as suggested in ref. 31, calculated on the M06/6-311++G(d,p) level.

gible barrier TS2<sub>CuH</sub> (less than 2.1 kcal mol<sup>−1</sup> for all proton donors, Table 6). This CuH pathway is similar for all the proton donors studied, with an activation barrier (TS1<sub>CuH</sub>) in the 33.2–36.4 kcal mol<sup>−1</sup> range for HFIP, TFE and MeOH. In the case of CF<sub>3</sub>OH this activation energy is much smaller: 12.5 kcal mol<sup>−1</sup>. The large energetic difference between strong and weak proton donors is somewhat similar to that found for (PP<sub>3</sub>)Ru(η<sup>1</sup>-BH<sub>4</sub>).<sup>18</sup>

The reaction with CF<sub>3</sub>OH features a higher impact of O–H bond dissociation whereas HFIP, TFE and MeOH have a predominant contribution of B–H<sub>br</sub> bond dissociation to the TS geometry (Fig. S16–S25†). For this reason, the activation barriers of proton transfer from CF<sub>3</sub>OH are comparable for both pathways (TS1<sub>CuH</sub>, ΔG<sub>theor</sub><sup>†</sup> = 12.5 kcal mol<sup>−1</sup>; TS1<sub>BH</sub><sup>A+B</sup>, ΔG<sub>theor</sub><sup>†</sup> = 11.0 kcal mol<sup>−1</sup>) (Fig. S14†).

One more peculiarity of the CuH pathway is the absence of direct involvement of the copper atom in the reaction, at odds with the BH pathway. This results in the ability of the title complex to undergo proton transfer also in coordinating solvents like THF (*via* the CuH pathway), unlike (Ph<sub>3</sub>P)<sub>2</sub>Cu(η<sup>2</sup>-BH<sub>4</sub>).<sup>17</sup> The latter does not react with proton donors in THF due to the Cu...O interaction with the solvent which prevents the “copper assisted” lower barrier proton transfer. Thus, the copper assistance along the CuH pathway

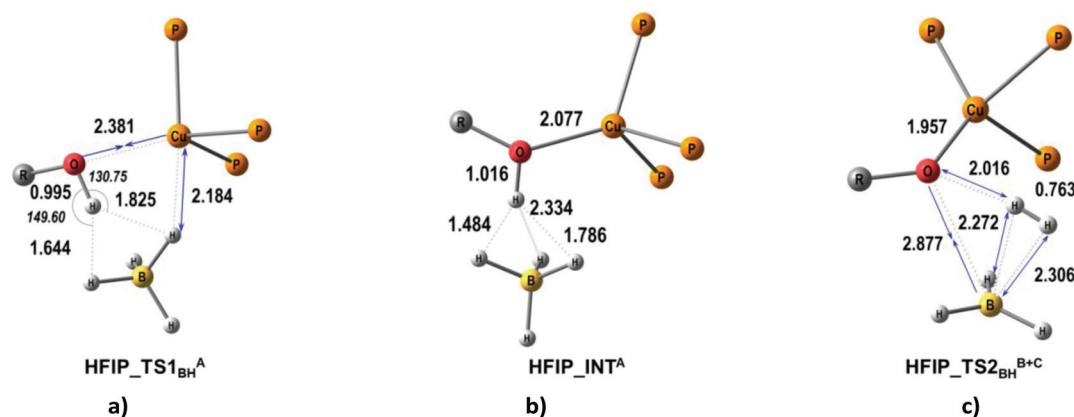


Fig. 6 M06-optimized structures of TSs (a, c) and intermediates (b) for the BH pathway of proton transfer from HFIP to (triphos<sup>Me</sup>)Cu(η<sup>1</sup>-BH<sub>4</sub>).



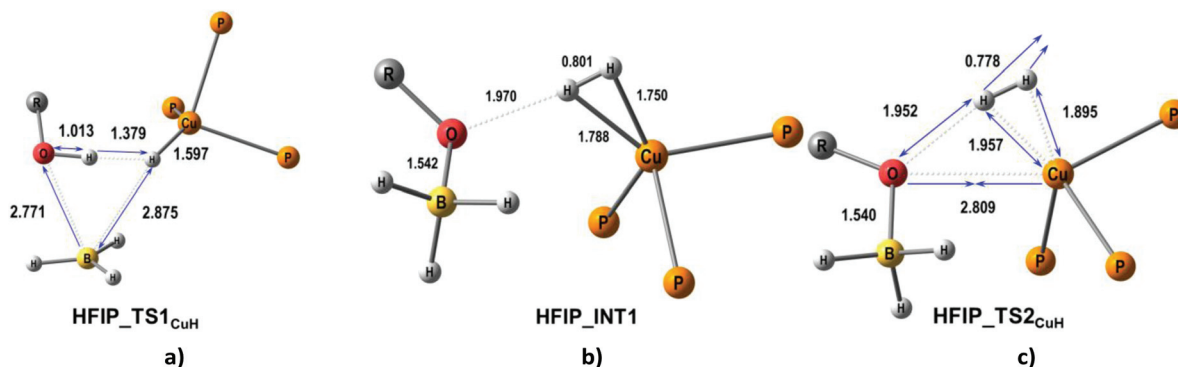


Fig. 7 M06-optimized structures of TSs (a, c) and intermediates (b) for the CuH pathway of proton transfer from HFIP to (triphos)<sup>Me</sup>Cu(η<sup>1</sup>-BH<sub>4</sub>).

for (triphos)Cu(η<sup>1</sup>-BH<sub>4</sub>) is not needed, and this reaction route becomes feasible also in the presence of coordinating solvents despite its higher activation barriers (TS1<sub>CuH</sub>,  $\Delta G_{\text{theor}}^{\ddagger} = 33.2 \text{ kcal mol}^{-1}$  for HFIP) when compared with the BH pathway (TS2<sub>BH</sub><sup>B+C</sup>,  $\Delta G_{\text{theor}}^{\ddagger} = 20.6 \text{ kcal mol}^{-1}$  for HFIP). In DCM, the operating mechanism for all proton donors should be the BH pathway. The calculated activation barrier for HFIP ( $\Delta G_{273 \text{ K}}^{\ddagger} = 20.6 \text{ kcal mol}^{-1}$ ) is in good agreement with the experimental data ( $\Delta G_{270 \text{ K}}^{\ddagger} = 20.0 \text{ kcal mol}^{-1}$  and  $\Delta G_{315 \text{ K}}^{\ddagger} = 21.8 \text{ kcal mol}^{-1}$ ).

## Experimental

### General considerations

All manipulations were performed under a dry argon atmosphere using a standard Schlenk technique. Commercially available argon (99.9%) was additionally purified from traces of oxygen and moisture by sequential passage through the Ni/Cr catalyst column and 4 Å molecular sieves.

HPLC grade solvents (Acros Organics) were used for sample preparation after additional purification by standard procedures. Dichloromethane (DCM) and tetrahydrofuran (THF) were dehydrated over CaH<sub>2</sub> and Na/benzophenone, respectively. All solvents were freshly distilled under argon prior to use. Deuterated solvent (CD<sub>2</sub>Cl<sub>2</sub>) was dried on CaH<sub>2</sub> and degassed by three freeze-pump-thaw cycles prior to use. Fluorinated alcohols were obtained from P&M (Moscow, Russia) and Fluka Analytical, were dried over anhydrous K<sub>2</sub>CO<sub>3</sub> and distilled under argon prior to use. Other reagents were from Sigma-Aldrich and were used as received.

### Variable-temperature infrared (IR) measurements

IR spectra were recorded on FTIR Nicolet 6700 and FTIR Shimadzu IR Prestige-21 spectrometers using 0.04–0.22 cm CaF<sub>2</sub> cells. Low temperature IR studies were carried out in the 190–300 K temperature range using a home-modified cryostat (Carl Zeiss Jena). The accuracy of the experimental temperature was  $\pm 0.5 \text{ K}$ . The cryostat modification allows for transfer of the reagents (premixed at either low or room temperature) under an inert atmosphere directly into the cells.

### NMR experiments

NMR spectra were recorded on a Bruker Avance II 400 MHz spectrometer. <sup>1</sup>H chemical shifts are reported in parts per million (ppm) relative to tetramethylsilane (TMS) and were calibrated against the residual solvent resonance, while <sup>31</sup>P {<sup>1</sup>H} NMR was referenced to 85% H<sub>3</sub>PO<sub>4</sub> with the downfield shift taken as positive and <sup>11</sup>B{<sup>1</sup>H} NMR was referenced to BF<sub>3</sub>·Et<sub>2</sub>O.

### Preparation of 1,1,1-tris(diphenylphosphinomethyl)ethane copper(i) tetrahydroborate (1)

Complex 1 was synthesized according to a previously described procedure.<sup>32</sup> Here, full spectroscopic (IR, NMR) characterization is reported, for the sake of literature data completeness. **Anal. calcd** for C<sub>41</sub>H<sub>43</sub>BCuP<sub>3</sub>: C, 70.04; H, 6.16; B, 1.54; P, 13.22. **Found**: C, 69.47; H, 6.20; B, 1.50; P, 13.06. **IR** (Nujol, cm<sup>-1</sup>) 2321 (s, BH<sub>4</sub><sup>as</sup><sub>term</sub>), 2354 (sh, BH<sub>4</sub><sup>as</sup><sub>term</sub>); 2233 (sh, BH<sub>4</sub><sup>s</sup><sub>term</sub>); 1988, 1967 (m, BH<sub>br</sub>). **<sup>1</sup>H NMR** (400 MHz, CD<sub>2</sub>Cl<sub>2</sub>, 295 K): 7.57–7.30 ppm (m, 6C<sub>6</sub>H<sub>5</sub>, 30 H); 2.50 ppm (s, 3CH<sub>2</sub>, 6 H); 0.87 ppm (s, 1CH<sub>3</sub>, 3 H); 0.98 ppm (q, BH<sub>4</sub><sup>-</sup>, 4 H); **<sup>31</sup>P{<sup>1</sup>H} NMR** (400 MHz, CD<sub>2</sub>Cl<sub>2</sub>, 295 K): –25.8 ppm (s, br, 3 P); **<sup>11</sup>B{<sup>1</sup>H} NMR** (400 MHz, CD<sub>2</sub>Cl<sub>2</sub>, 295 K): –32.80 ppm (s).

### Computational details

Full geometry optimizations were carried out with the GAUSSIAN09 (Revision C.01)<sup>33</sup> package at the density functional theory (DFT) level using the M06 functional.<sup>34</sup> The basis sets used were spin-state-corrected s6-31G(d) for the metal center (Cu),<sup>35</sup> 6-311++G(d,p) for atoms of the BH<sub>4</sub><sup>-</sup> and alcohol OH-groups,<sup>36,37</sup> 6-31G(d) for the phosphorus atoms,<sup>38</sup> and 6-31G for all the other atoms.

Frequency calculations were performed for all the optimized complexes in the gas phase and are reported without the use of scaling factors. The nature of all the stationary points on the potential energy surfaces was confirmed by a vibrational analysis.<sup>39</sup> Transition state (TS) structures showed only one negative eigenvalue in their diagonalized force constant matrices, and their associated eigenvectors were confirmed to correspond to the motion along the reaction



coordinate under consideration using the Intrinsic Reaction Coordinate (IRC) method.<sup>40</sup>

The complex formation energy was calculated in the gas phase taking into account the basis sets superposition error (by the Bernardi and Boys method, BSSE),<sup>41</sup> and ZPVE correction was determined from the unscaled harmonic frequencies.<sup>42,43</sup>

Inclusion of nonspecific solvent effects in the calculations was performed by using the SMD method.<sup>44</sup> The interaction energy was calculated in THF ( $\epsilon = 7.4$ ) and  $\text{CH}_2\text{Cl}_2$  ( $\epsilon = 8.9$ ) for the gas phase optimized geometries. Changes in Gibbs energies and enthalpies in the solvent were determined using the corresponding corrections obtained for the gas phase:<sup>45</sup>

$$\Delta H_{\text{Solv.}} = \Delta E_{\text{Solv.}} + \Delta H_{\text{gas}}^{\text{corr}} \quad (3)$$

$$\Delta G_{\text{Solv.}} = \Delta E_{\text{Solv.}} + \Delta G_{\text{gas}}^{\text{corr}} \quad (4)$$

Natural atomic charges and Wiberg bond indices<sup>20</sup> were calculated using the natural-bond orbital (NBO) analysis<sup>19</sup> implemented in Gaussian09. Topological analysis of the electron-density distribution function  $\rho(r)$  was performed using the AIMALL<sup>46</sup> program package based on the wave function obtained by the M06 calculations. The energies of  $\text{H}\cdots\text{H}$  interactions were calculated using the correlation between the binding energy ( $E_{\text{H}\cdots\text{H}}$ ) and the value of the density-functional potential energy  $V(r)$  in the corresponding critical point (3, -1):  $E_{\text{H}\cdots\text{H}} = 0.5 \cdot V(r)$ .<sup>47,48</sup> Hydrogen bond ellipticity,  $\epsilon_{\text{H}\cdots\text{H}}$ , was defined as  $\epsilon = (\lambda_1/\lambda_2 - 1)$ , where  $\lambda_1$  and  $\lambda_2$  are the negative eigenvalues of the Hessian of the electron density at the bond critical point ordered such that  $\lambda_1 < \lambda_2 < 0$ .<sup>21–23</sup>

## Conclusions

The interaction of alcohols of variable strength with the copper(i) borohydride complex (triphos) $\text{Cu}(\eta^1\text{-BH}_4)$  results in a great variety of DHB complexes which encompass different mechanisms involving M–H and E–H bond (E = B, O) activation steps. The proton transfer at the terminal B–H bond begins with nucleophilic substitution of the  $\text{BH}_4^-$  ligand by the alcohol oxygen atom on the copper center that further activates the O–H bond for the subsequent proton transfer *via* the DHB intermediate  $(\text{triphos}^{\text{Me}})\text{Cu}(\eta^1\text{-O}^+(\text{R}))\text{H}\cdots\text{H}_4\text{B}^-$ . A variation of the structural and energetic details of this process with the proton donor strength is found through computation, and the theoretical results are fully consistent with the experimental observations in dichloromethane showing an abnormal dependence of the reaction rate on the proton donor strength.

In contrast, DHB complexes with the initial ROH coordination to the bridging B–H bond lead to B– $\text{H}_{\text{br}}$  bond cleavage with subsequent  $\text{BH}_3$  elimination and formation of the related copper hydride (triphos) $\text{CuH}$ . This pathway does not involve any kind of “copper assistance” *via* the  $\text{Cu}\cdots\text{O}$  interaction, so this mechanism can be evoked to explain the occurrence of proton transfer also in coordinating solvents. As copper badly

stabilizes molecular hydrogen complexes (non-classical hydrides), the protonation of copper hydride is disfavored with respect to protonation of boron hydride even for strong proton donors (exemplified by  $\text{CF}_3\text{OH}$ ). This behavior differs from that shown by  $(\text{PP}_3)_2\text{RuH}(\eta^1\text{-BH}_4)$ ,<sup>6</sup> where the preferred pathway involves the  $\text{BH}_{\text{br}}$  bond dissociation followed by the ruthenium hydride protonation. The mechanism of proton transfer to monodentate tetrahydroborate ( $\eta^1\text{-BH}_4$ ) complexes depends on the metal ability to stabilize  $\eta^2\text{-H}_2$  complexes.

## Acknowledgements

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## Notes and references

- 1 G. C. Fu, M. Beller and C. Bolm, in *Transition Metals for Organic Synthesis*, Wiley-VCH Verlag GmbH, 2008, pp. 141–146.
- 2 G. Povie, M. Marzorati, P. Bigler and P. Renaud, *J. Org. Chem.*, 2013, **78**, 1553–1558.
- 3 A. Staubitz, A. P. M. Robertson, M. E. Sloan and I. Manners, *Chem. Rev.*, 2010, **110**, 4023–4078.
- 4 A. Staubitz, A. P. M. Robertson and I. Manners, *Chem. Rev.*, 2010, **110**, 4079–4124.
- 5 G. Alcaraz and S. Sabo-Etienne, *Angew. Chem., Int. Ed.*, 2010, **49**, 7170–7179.
- 6 N. E. Stubbs, A. P. M. Robertson, E. M. Leitao and I. Manners, *J. Organomet. Chem.*, 2013, **730**, 84–89.
- 7 A. John, K. I. Goldberg and D. M. Heinekey, in *Organometallic Pincer Chemistry*, ed. G. van Koten and D. Milstein, Springer Berlin Heidelberg, Berlin, Heidelberg, 2013, pp. 271–287.
- 8 H. C. Johnson, T. N. Hooper and A. S. Weller, in *Synthesis and Application of Organoboron Compounds*, ed. E. Fernández and A. Whiting, Springer International Publishing, 2015, vol. 49, ch. 6, pp. 153–220.
- 9 R. L. Melen, *Chem. Soc. Rev.*, 2015, **45**, 775–788.
- 10 A. Rossin and M. Peruzzini, *Chem. Rev.*, 2016, DOI: 10.1021/acs.chemrev.6b00043.
- 11 Y. Kawano, K. Yamaguchi, S.-Y. Miyake, T. Kakizawa and M. Shimoi, *Chem. – Eur. J.*, 2007, **13**, 6920–6931.
- 12 R. N. Perutz and S. Sabo-Etienne, *Angew. Chem., Int. Ed.*, 2007, **46**, 2578–2592.
- 13 C. J. Stevens, R. Dallanegra, A. B. Chaplin, A. S. Weller, S. A. Macgregor, B. Ward, D. McKay, G. Alcaraz and S. Sabo-Etienne, *Chem. – Eur. J.*, 2011, **17**, 3011–3020.
- 14 A. E. Nako, A. J. P. White and M. R. Crimmin, *Dalton Trans.*, 2015, **44**, 12530–12534.
- 15 A. Kumar, I. K. Priest, T. N. Hooper and A. Weller, *Dalton Trans.*, 2016, **45**, 6183–6195.



- 16 Y. Kawano, M. Uruichi, M. Shimoi, S. Taki, T. Kawaguchi, T. Kakizawa and H. Ogino, *J. Am. Chem. Soc.*, 2009, **131**, 14946–14957.
- 17 I. E. Golub, O. A. Filippov, E. I. Gutsul, N. V. Belkova, L. M. Epstein, A. Rossin, M. Peruzzini and E. S. Shubina, *Inorg. Chem.*, 2012, **51**, 6486–6497.
- 18 N. V. Belkova, E. V. Bakhmutova-Albert, E. I. Gutsul, V. I. Bakhmutov, I. E. Golub, O. A. Filippov, L. M. Epstein, M. Peruzzini, A. Rossin, F. Zanobini and E. S. Shubina, *Inorg. Chem.*, 2014, **53**, 1080–1090.
- 19 E. D. Glendening, J. K. Badenhop, A. E. Reed, J. E. Carpenter, J. A. Bohman, C. Morales and F. Weindhold, *NBO 5.0*, Theoretical Chemistry Institute, University of Wisconsin, Madison, WI, 2001.
- 20 K. B. Wiberg, *Tetrahedron*, 1968, **24**, 1083–1096.
- 21 R. F. W. Bader, *Atoms in Molecules: A Quantum Theory (International Series of Monographs on Chemistry)*, Oxford University Press, USA, 1994.
- 22 P. L. Popelier, *Atoms in Molecules: An Introduction*, Prentice Hall, London, 2000.
- 23 C. Matta and R. J. Boyd, *Quantum Theory of Atoms in Molecules: Recent Progress in Theory and Application*, Wiley-VCH, New York, 2007.
- 24 S. J. Grabowski, W. A. Sokalski and J. Leszczynski, *J. Phys. Chem. A*, 2005, **109**, 4331–4341.
- 25 L. M. Epstein and E. S. Shubina, *Coord. Chem. Rev.*, 2002, **231**, 165–181.
- 26 N. V. Belkova, E. S. Shubina and L. M. Epstein, *Acc. Chem. Res.*, 2005, **38**, 624–631.
- 27 A. V. Iogansen, *Theor. Exp. Chem.*, 1973, **7**, 249–256.
- 28 I. E. Golub, E. S. Gulyaeva, O. A. Filippov, V. P. Dyadchenko, N. V. Belkova, L. M. Epstein, D. E. Arkhipov and E. S. Shubina, *J. Phys. Chem. A*, 2015, **119**, 3853–3868.
- 29 W. G. Henderson and E. F. Mooney, in *Annu. Rep. NMR Spectrosc.*, ed. E. F. Mooney, Academic Press, 1969, vol. 2, pp. 219–291.
- 30 C. Bianchini, C. A. Ghilardi, A. Meli, S. Midollini and A. Orlandini, *Inorg. Chem.*, 1985, **24**, 932–939.
- 31 O. A. Filippov, A. M. Filin, V. N. Tsupreva, N. V. Belkova, A. Lledós, G. Ujaque, L. M. Epstein and E. S. Shubina, *Inorg. Chem.*, 2006, **45**, 3086–3096.
- 32 P. Dapporto, S. Midollini, A. Orlandini and L. Sacconi, *Inorg. Chem.*, 1976, **15**, 2768–2774.
- 33 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision C.01*, Gaussian, Inc., Wallingford CT, 2009.
- 34 Y. Zhao and D. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215–241.
- 35 M. Swart, M. Güell, J. M. Luis and M. Solà, *J. Phys. Chem. A*, 2010, **114**, 7191–7197.
- 36 W. J. Hehre, R. Ditchfield and J. A. Pople, *J. Chem. Phys.*, 1972, **56**, 2257–2262.
- 37 J. D. Dill and J. A. Pople, *J. Chem. Phys.*, 1975, **62**, 2921–2923.
- 38 R. Krishnan, J. S. Binkley, R. Seeger and J. A. Pople, *J. Chem. Phys.*, 1980, **72**, 650–654.
- 39 J. Fritsch and G. Zundel, *J. Phys. Chem.*, 1981, **85**, 556–561.
- 40 R. Kramer and G. Zundel, *J. Chem. Soc., Faraday Trans.*, 1990, **86**, 301–305.
- 41 S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553–566.
- 42 C. Tuma, D. A. Boese and N. C. Handy, *Phys. Chem. Chem. Phys.*, 1999, **1**, 3939–3947.
- 43 G. Buemi, in *Hydrogen Bonding—New Insights*, ed. S. Grabowski, Springer, Netherlands, 2006, pp. 51–107.
- 44 A. V. Marenich, C. J. Cramer and D. G. Truhlar, *J. Phys. Chem. B*, 2009, **113**, 6378–6396.
- 45 M. Sumimoto, N. Iwane, T. Takahama and S. Sakaki, *J. Am. Chem. Soc.*, 2004, **126**, 10457–10471.
- 46 T. A. Keith, *AIMAll (Version 15.05.18)*, TK Gristmill Software, Overland Park KS, USA, 2015.
- 47 E. Espinosa, I. Alkorta, I. Rozas, J. Elguero and E. Molins, *Chem. Phys. Lett.*, 2001, **336**, 457–461.
- 48 E. Espinosa, E. Molins and C. Lecomte, *Chem. Phys. Lett.*, 1998, **285**, 170–173.

