

Cite this: *Chem. Sci.*, 2023, 14, 889

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

# Radical-like reactivity for dihydrogen activation by coinage metal–aluminyl complexes: computational evidence inspired by experimental main group chemistry†

Diego Sorbelli,<sup>ID</sup> \*<sup>ab</sup> Leonardo Belpassi<sup>ID</sup> \*<sup>b</sup> and Paola Belanzoni<sup>ID</sup> \*<sup>ab</sup>

The computational study of an unprecedented reactivity of coinage metal–aluminyl complexes with dihydrogen is reported. In close resemblance to group 14 dimetallenes and dimetallynes, the complexes are predicted to activate H<sub>2</sub> under mild conditions. Two different reaction pathways are found disclosing a common driving force, *i.e.*, the nucleophilic behavior of the electron-sharing M–Al (M = Cu, Ag, Au) bond, which enables a cooperative and diradical-like mechanism. This mode of chemical reactivity emerges as a new paradigm for dihydrogen activation and calls for experimental feedback.

Received 20th October 2022  
Accepted 15th December 2022

DOI: 10.1039/d2sc05815d

rsc.li/chemical-science

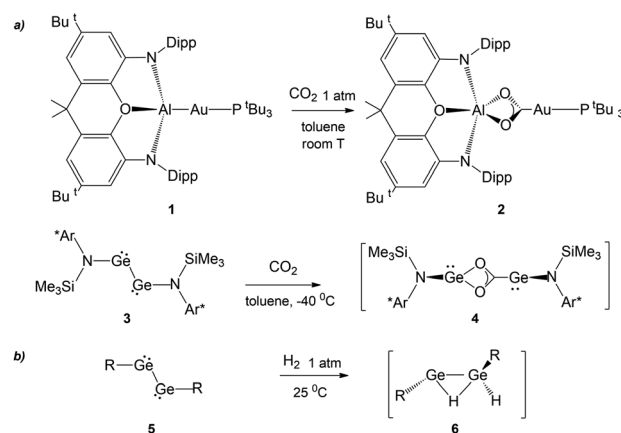
## Introduction

The recent synthesis of the gold–aluminyl complex **1** (Scheme 1) and its outstanding reactivity in capturing CO<sub>2</sub> under mild experimental conditions represent a unique case in the framework of gold chemistry.<sup>1</sup> We have thoroughly discussed that the driving force of this reactivity is the presence of an electron-sharing, weakly polarized Au–Al bond acting as a nucleophilic site inducing a cooperative diradical-like reactivity.<sup>2–5</sup> Shortly after this, analogous reactivity was reported to occur for copper– and silver–aluminyl complexes, for which, however, the reaction proceeds towards the formation of carbonate complexes with CO extrusion.<sup>6</sup>

A surprising feature concerning the reaction of **1** with carbon dioxide is that it bears strict analogies with amido-digermyne compound **3** (Scheme 1), which, in reducing CO<sub>2</sub> to CO, proceeds through an intermediate **4** that is structurally very similar to insertion product **2**.<sup>7</sup> Particularly intriguing, aryl- and amido-digermynes RGeGer (R = [C<sub>6</sub>H<sub>3</sub>-2,6(C<sub>6</sub>H<sub>3</sub>-2,6<sup>i</sup>Pr<sub>2</sub>)<sub>2</sub>],<sup>8</sup> [N(SiMe<sub>3</sub>)(C<sub>6</sub>H<sub>2</sub>Me(C(H)Ph<sub>2</sub>)<sub>2</sub>)]<sup>9</sup> and [N(Si<sup>i</sup>Pr<sub>3</sub>)(2,6-[C(H)Ph<sub>2</sub>]<sub>2</sub>-4-<sup>i</sup>PrC<sub>6</sub>H<sub>2</sub>)]<sup>10</sup>) **5** (Scheme 1) are reported to easily activate H<sub>2</sub> under mild experimental conditions both in solution<sup>8,10</sup> and the solid state at temperatures as low as –10 °C.<sup>9</sup> In all cases, experiments and theoretical investigations outline a reaction mechanism that proceeds *via* a singly bridged intermediate [RGe(μ-H)GeHR] species **6**, which subsequently, upon isomerization, yields different hydrogenation products, which are experimentally

revealed depending on the steric hindrance of the substituents.<sup>11,12</sup> Notably, the facile reactivity with H<sub>2</sub> has also been reported for other group 14 dimetallenes and dimetallynes.<sup>13</sup> The digermene is suggested to possess substantial diradical character and therefore to react through H atom abstraction from H<sub>2</sub>, followed by recombination of the resultant radical pair.<sup>8</sup>

As bare ligands, aluminyls resemble singlet carbenes in possessing an electron lone pair and an accessible vacant p orbital, thus potentially showing a similar reactivity. Interestingly, it has been reported that acyclic and cyclic (alkyl)(amino) carbenes **7** and **8a, b** can activate H<sub>2</sub> under mild



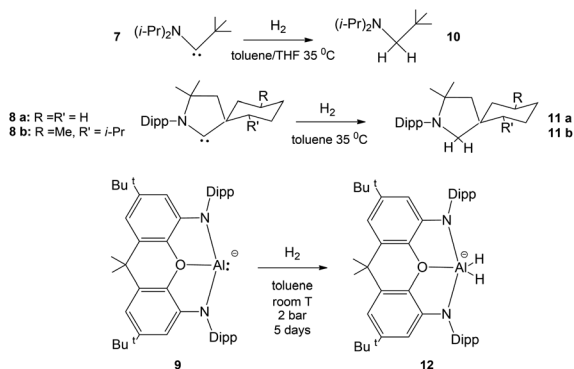
Scheme 1 (a) Similar reactivity of [tBu<sub>3</sub>PAuAl(NON)] (NON = 4,5-bis(2,6-diisopropylanilido)-2,7-di-*tert*-butyl-9,9-dimethylxantheno) complex **1** and (Ar\*)(SiMe<sub>3</sub>)GeGe(Ar\*)(SiMe<sub>3</sub>) (Ar\* = C<sub>6</sub>H<sub>2</sub>(C(H)Ph<sub>2</sub>)<sub>2</sub>Me-2,6,4) complex **3** with CO<sub>2</sub>.<sup>7</sup> (b) First step in the mechanism of the reaction of several aryl- and amido-digermyne compounds **5** with H<sub>2</sub>. (R = [C<sub>6</sub>H<sub>3</sub>-2,6(C<sub>6</sub>H<sub>3</sub>-2,6<sup>i</sup>Pr<sub>2</sub>)<sub>2</sub>],<sup>8</sup> [N(SiMe<sub>3</sub>)(C<sub>6</sub>H<sub>2</sub>Me(C(H)Ph<sub>2</sub>)<sub>2</sub>)]<sup>9</sup> and [N(Si<sup>i</sup>Pr<sub>3</sub>)(2,6-[C(H)Ph<sub>2</sub>]<sub>2</sub>-4-<sup>i</sup>PrC<sub>6</sub>H<sub>2</sub>)]<sup>10</sup>).

<sup>a</sup>Department of Chemistry, Biology and Biotechnology, University of Perugia, Via Elce di Sotto, 8 – 06123, Perugia, Italy. E-mail: diegosorbelli00@gmail.com; paola.belanzoni@unipg.it

<sup>b</sup>CNR Institute of Chemical Science and Technologies “Giulio Natta” (CNR-SCITEC), Via Elce di Sotto, 8 – 06123, Perugia, Italy. E-mail: leonardo.belpassi@cnr.it

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2sc05815d>





Scheme 2 Similar reactivity of (alkyl)(amino)carbenes **7** and **8a, b** and aluminyl **9** with H<sub>2</sub>.

conditions, by behaving as nucleophiles (Scheme 2).<sup>14</sup> In analogy, the ability of [M<sub>2</sub>Al(NON)]<sub>2</sub> (M = Li, Na, K) species to react with H<sub>2</sub> has been also reported (Scheme 2, complex **9**)<sup>15–17</sup> together with detailed mechanistic studies.<sup>18</sup> Despite these similarities, when used as coordination ligands, aluminyls and carbenes have a strikingly different chemical behaviour. In a recent work, we have shown that gold–aluminyl complexes, unlike gold carbene analogues featuring a dative Au–C bond, are able to react with CO<sub>2</sub>.<sup>4</sup>

In this scenario the interest in exploring the reactivity of gold–aluminyl complexes towards H<sub>2</sub> naturally arises. The electron-sharing nature of the Au–Al bond in complex **1** allowed the rationalization of its reactivity with CO<sub>2</sub>,<sup>2</sup> in close resemblance with that of digermene complex **3**, and thus, we reasoned that the same bonding model may also favor the reaction of complex **1** with H<sub>2</sub> analogous to that of complexes **5**. The possibility for H<sub>2</sub> activation to occur with aluminyl complexes is very attractive, especially in light of potential applications in hydrogen storage and catalysis, beside H<sub>2</sub> being vital in several industrial processes, organic synthesis and also in biological functions.<sup>19,20</sup>

In addition, given the similar reactivity of gold–copper- and silver–aluminyl complexes with CO<sub>2</sub>, a possible H<sub>2</sub> activation by copper- and silver–aluminyl complexes is certainly worth exploring.

Herein we report that [tBu<sub>3</sub>PMAl(NON)] (M = Cu, Ag, Au) complexes should indeed react with dihydrogen, with the Cu–Al complex featuring the most kinetically favored and exergonic reaction with H<sub>2</sub>. The calculations predict the experimentally accessible formation of a singly bridged [tBu<sub>3</sub>PM(μ-H)Al(H)(NON)] species for all the metal–aluminyl complexes and an additional doubly bridged [tBu<sub>3</sub>PM(μ-H)<sub>2</sub>Al(NON)] product for copper and silver. Detailed electronic structure calculations highlight the central role of the electron-sharing M–Al bond in inducing a diradical-like reactivity towards H<sub>2</sub>, in close resemblance to digermynes, as surmised.

## Results and discussion

The free energy profiles for the reaction of [tBu<sub>3</sub>PMAl(NON)] complexes with H<sub>2</sub> have been calculated using the same

computational protocol employed in ref. 2 for the reaction of [tBu<sub>3</sub>PAuAl(NON)] with CO<sub>2</sub> (see Computational details). For [tBu<sub>3</sub>PAuAl(NON)], the free energy profile is depicted in Fig. 1 together with the schematic structure of all stationary points. The optimized structures of all the stationary points in Fig. 1 are reported in the ESI (Fig. S1).†

Starting from the reactant complex (RC<sub>Au</sub>), H<sub>2</sub> activation can occur both at the Au (blue line profile, ΔG = 28.4 kcal mol<sup>−1</sup>) and the Al (red line profile, ΔG = 32.2 kcal mol<sup>−1</sup>) sites, with activation at Au being favoured. From the Al site, the reaction leads directly to the formation of the *cis*-[tBu<sub>3</sub>PAu(μ-H)Al(H)(NON)] singly bridged species PC<sub>Au</sub> via the concerted transition state TS<sub>Au</sub><sup>IV</sup>. From the Au site, a first step (formally a H<sub>2</sub> “oxidative addition” to Au via TS<sub>Au</sub><sup>I</sup>) yields an unstable intermediate species INT<sub>Au</sub> which rearranges (via TS<sub>Au</sub><sup>II</sup>, ΔG = 29.0 kcal mol<sup>−1</sup>) by a hydrogen transfer to Al, leading to the singly bridged *trans*-[tBu<sub>3</sub>PAu(μ-H)Al(H)(NON)] species PC<sub>Au</sub> which is in equilibrium with the more stable PC<sub>Au</sub> (via TS<sub>Au</sub><sup>III</sup>, ΔG = 8.3 kcal mol<sup>−1</sup>). Interestingly, the overall RC<sub>Au</sub>-to-PC<sub>Au</sub> conversion is practically thermoneutral, suggesting a possibly reversible reaction and a virtually ideal condition for the use of complex **1** as a hydrogen-transfer catalyst. The free energy activation barriers for the two paths roughly fit in the 25–30 kcal mol<sup>−1</sup> range,

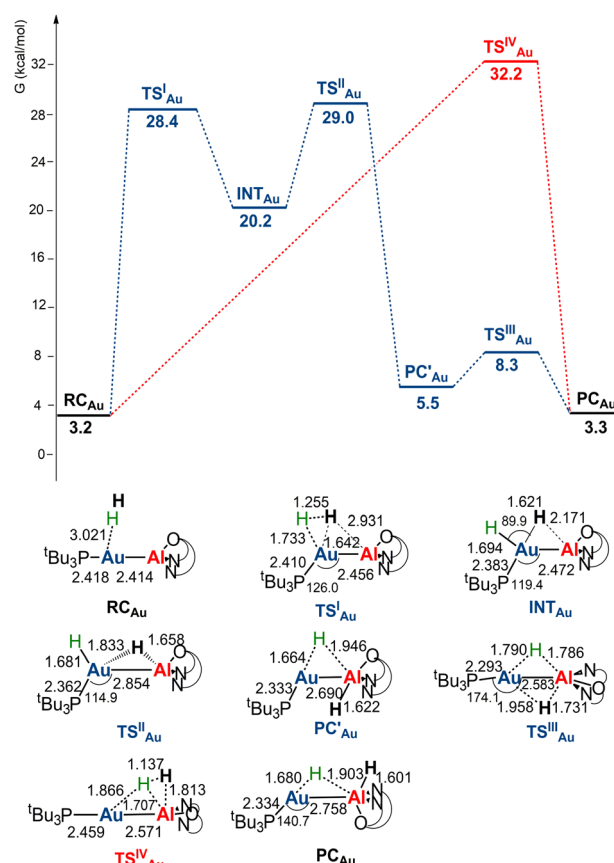


Fig. 1 Free energy profile for the reaction of [tBu<sub>3</sub>PAuAl(NON)] with H<sub>2</sub> at the Au (blue line) and the Al (red line) sites. ΔG values refer to the energy of the separated reactants taken as zero. Selected interatomic distances (Å) and bond angles (degrees) are shown with all the stationary point structures.



typically observed for experimental H<sub>2</sub> activation processes under mild conditions (see ref. 18 and 21–24 for some recent examples).

The structural analogy between the gold–aluminyl PC<sub>Au</sub> and digermine **6** (Scheme 1) singly bridged species is striking.

Notably, at variance with digermine **6**, PC<sub>Au</sub> is not expected to easily undergo isomerization (see Scheme S1 and Discussion in the ESI†) and, in particular, dissociation into two separate hydride complexes (*i.e.* [t-Bu<sub>3</sub>PAuH] and [HAL(NON)]) (PC<sub>Au</sub>-HH, see Fig. S1†).

Given the thermoneutrality of this reaction, experimental evidence for H<sub>2</sub> cleavage could not be straightforward. In light of the reported more efficient reaction of copper- and silver-aluminyl complexes with carbon dioxide, it is interesting at this point to investigate the reactivity of [t-Bu<sub>3</sub>PMAl(NON)] (M = Cu, Ag) complexes with H<sub>2</sub>. The reaction profiles are depicted in Fig. 2. The structures of all stationary points for both copper- and silver-aluminyl complexes are shown in Fig. S2–S4 in the ESI.†

For the silver- and copper-aluminyls, H<sub>2</sub> activation can occur symmetrically at the Ag (blue line profile, ΔG = 33.2 kcal mol<sup>-1</sup>) and the Al (red line profile, ΔG = 33.6 kcal mol<sup>-1</sup>) sites, and remarkably unsymmetrically at the Cu (blue line profile, ΔG

= 18.7 kcal mol<sup>-1</sup>) and the Al (red line profile, ΔG = 32.0 kcal mol<sup>-1</sup>) sites. From the Al site, the reaction leads directly to the formation of the singly bridged aluminyl species PC<sub>Cu</sub>/PC<sub>Ag</sub> for both copper and silver complexes *via* the concerted transition state TS<sub>Cu</sub><sup>IV</sup>/TS<sub>Ag</sub><sup>IV</sup>. Although both are found to have similar energy to that of TS<sub>Au</sub><sup>IV</sup>, the respective products are found to be more stable. In particular, the formation of PC<sub>Ag</sub> is slightly exergonic (−1.1 kcal mol<sup>-1</sup>) and the formation of PC<sub>Cu</sub> is even more exergonic (−5.0 kcal mol<sup>-1</sup>).

The alternative path, where H<sub>2</sub> approaches the complexes closer to the metal site (blue paths in Fig. 2), highlights appreciable differences between lighter coinage metals and gold, instead. Qualitatively, the first step is analogous in all cases: H<sub>2</sub> approaches the metal site, yielding intermediates INT<sub>Cu</sub>/INT<sub>Ag</sub>. In the second step, however, copper and silver complexes remarkably differ from gold, since the intermediate is converted unequivocally to a stable doubly bridged product [t-Bu<sub>3</sub>PM(μ-H)<sub>2</sub>Al(NON)] (M = Cu, Ag) in both cases (PC<sub>Cu</sub><sup>′</sup>/PC<sub>Ag</sub><sup>′</sup>).

From a quantitative perspective, differences between the three metals are even more pronounced. Indeed, the silver-aluminyl complex displays a higher activation barrier for TS<sub>Ag</sub><sup>IV</sup> (33.2 kcal mol<sup>-1</sup>) and a slightly exergonic formation of PC<sub>Ag</sub><sup>′</sup> (ΔG = −1.9 kcal mol<sup>-1</sup>) *via* isomerization from the unstable INT<sub>Ag</sub>. On the other hand, the path for the copper-aluminyl complex shows much more favorable energetics overall with respect to silver and gold complexes. In the first step, a more stable intermediate (INT<sub>Cu</sub>) is formed *via* a much lower activation barrier at TS<sub>Cu</sub><sup>I</sup> (ΔG = 18.7 kcal mol<sup>-1</sup>). Subsequently, the INT<sub>Cu</sub> isomerization leads to the highly stable doubly bridged product PC<sub>Cu</sub><sup>′</sup> (ΔG = −12.3 kcal mol<sup>-1</sup>). Remarkably, PC<sub>Cu</sub><sup>′</sup>, PC<sub>Ag</sub><sup>′</sup> and TS<sub>Au</sub><sup>III</sup> structures are similar, even though for gold the doubly bridged species is not a minimum energy point. Thus, preferential formation of singly bridged aluminyl species with the second hydrogen bound to Al is found for gold.

The reaction profiles described above clearly indicate that coinage metal–aluminyl complexes can activate H<sub>2</sub>, and a particularly facile H<sub>2</sub> cleavage is predicted with the copper-aluminyl complex. Based on the RC<sub>Cu</sub> – TS<sub>Cu</sub><sup>I</sup> – INT<sub>Cu</sub> – TS<sub>Cu</sub><sup>II</sup> – PC<sub>Cu</sub><sup>′</sup> path, due to the relatively low activation barrier and high exergonicity, dihydrogen activation could be experimentally demonstrated more easily. Under suitable conditions, isolation and characterization of the doubly bridged copper species should be achievable. Furthermore, as reported extensively in the literature,<sup>25–30</sup> this type of reaction may be affected by the presence of tunnel effects that contribute to the lowering of the activation barriers. With the aim of assessing if such effects may be important in this context, we used transition state theory (TST)<sup>31</sup> and the Eckart tunneling correction<sup>32</sup> (see Computational details) to estimate the tunnel effect on the Eyring fitted activation barrier associated with TS<sub>M</sub><sup>I</sup>. The results show that, indeed, a moderate tunnel-related lowering of the activation barriers associated with TS<sub>M</sub><sup>I</sup> may be expected (in the range of 1.3–3.7 kcal mol<sup>-1</sup>, see Table S1†). A proper assessment of this effect is beyond the scope of this work, but this estimate suggests that the tunnel effects may impact, to some extent, the reactivity reported here.

A common feature of the reaction profiles in Fig. 1 and 2 is the possible H<sub>2</sub> activation at two different sites (M and Al),

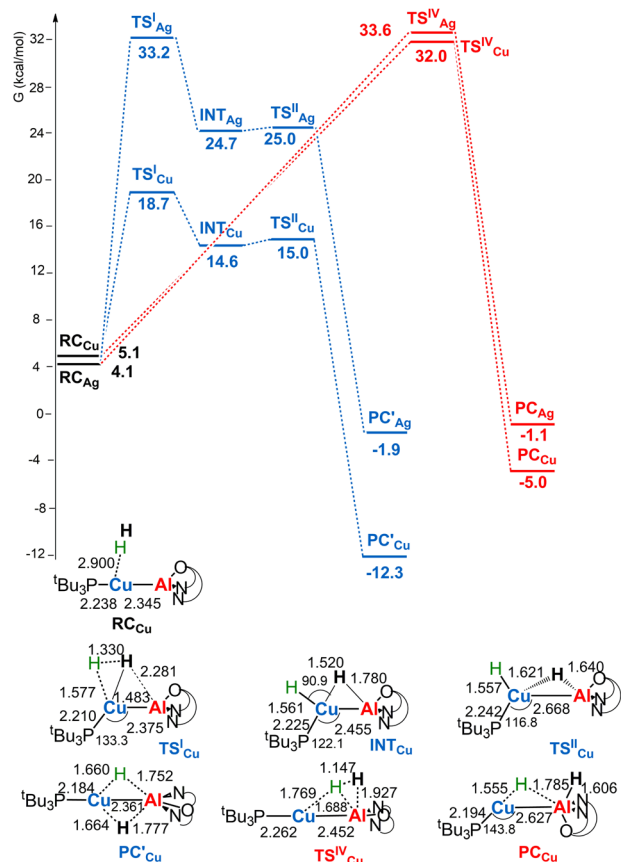


Fig. 2 Free energy profile for the reaction of [t-Bu<sub>3</sub>PMAl(NON)] (M = Cu, Ag) with H<sub>2</sub> at the M (blue line) and the Al (red line) sites. ΔG values refer to the energy of the separated reactants taken as zero. Selected interatomic distances (Å) and bond angles (degrees) are shown with all the stationary point structures for the [t-Bu<sub>3</sub>PCuAl(NON)] complex (for the [t-Bu<sub>3</sub>PAGAl(NON)] complex see Fig. S3 in the ESI.†).



apparently suggesting that the two distinct paths are initiated by interactions of a different nature when H<sub>2</sub> approaches the complexes (transition states TS<sub>M</sub><sup>I</sup> and TS<sub>M</sub><sup>IV</sup>). To investigate the nature of the interaction between H<sub>2</sub> and [t-Bu<sub>3</sub>PMAl(NON)] (M = Cu, Ag, Au) at the corresponding TS<sub>M</sub><sup>I</sup> and TS<sub>M</sub><sup>IV</sup>, natural orbitals for chemical valence (NOCV) analysis,<sup>33,34</sup> energy decomposition analysis (EDA)<sup>35–37</sup> and charge displacement (CD)<sup>38–40</sup> approaches have been applied (see Table S2 and Fig. S5–S16 in the ESI†).

As shown in Fig. 3, the NOCV analysis at TS<sub>Au</sub><sup>IV</sup> for the gold-aluminyll complex reveals a main component ( $\Delta\rho'_1$ ) of the H<sub>2</sub>–[t-Bu<sub>3</sub>PAuAl(NON)] interaction, which can be depicted, upon decomposition of the corresponding donor and acceptor NOCVs (see Fig. S5 and S6 in the ESI†), as mainly dominated by a charge transfer from the HOMO of the complex (a  $\sigma$  MO representing the electron-sharing Au–Al bond) towards the LUMO of H<sub>2</sub> (Fig. 2a and S5†). A second non-negligible component can be envisaged ( $\Delta\rho'_2$ , Fig. 2b and S6†), representing a charge transfer from the H<sub>2</sub> HOMO to the [t-Bu<sub>3</sub>PAuAl(NON)] LUMO, mainly composed of the empty 3p<sub>z</sub> orbital of Al. Notably, this NOCV description of the H<sub>2</sub>–[t-Bu<sub>3</sub>PAuAl(NON)] interaction is strongly reminiscent of both the CO<sub>2</sub>–[t-Bu<sub>3</sub>PAuAl(NON)] interaction (see ref. 2–5) and the H<sub>2</sub>–[RGeGeR] interaction reported by Frenking and coworkers, where the digermene, by employing the  $\sigma$  Ge–Ge HOMO and the 4p<sub>z</sub>-centred LUMO, interacts with the H<sub>2</sub> LUMO and HOMO, respectively.<sup>12</sup>

Surprisingly, despite remarkable structural and energetic differences, the H<sub>2</sub>–[t-Bu<sub>3</sub>PAuAl(NON)] interactions at TS<sub>Au</sub><sup>I</sup> are qualitatively similar to those at TS<sub>Au</sub><sup>IV</sup>. As shown in Fig. 2c and d (Fig. S7 and S8 in the ESI†), the NOCV analysis analogously reveals two dominant components:  $\Delta\rho'_1$  consisting of a charge transfer from the  $\sigma$  Au–Al HOMO of the complex towards the

LUMO of H<sub>2</sub>, and  $\Delta\rho'_2$  featuring a charge transfer from the H<sub>2</sub> HOMO towards the LUMO of the complex.

Notably, analogous results arise upon NOCV analysis of the H<sub>2</sub>-complex interaction at TS<sub>M</sub><sup>IV</sup> and TS<sub>M</sub><sup>I</sup> for copper- and silver-aluminyll complexes. As can be inferred from the results reported in Fig. S9–S16 in the ESI,† at both TS<sub>Cu</sub><sup>I</sup>/TS<sub>Ag</sub><sup>I</sup> and TS<sub>Cu</sub><sup>IV</sup>/TS<sub>Ag</sub><sup>IV</sup> the main component of the H<sub>2</sub>-complex interaction is represented by a charge transfer from the  $\sigma$  bonding M–Al molecular orbital towards the H<sub>2</sub> antibonding LUMO. This finding is fully consistent with the nature of the M–Al bond. As previously reported for the Cu–Al bond<sup>2</sup> and based on Fig. S17, S18 and Table S3 in the ESI and Discussion therein,† despite small changes in the bond polarization, in these complexes the M–Al (M = Cu, Al, Au) is an electron-sharing bond, acting as a nucleophilic site for the activation of H<sub>2</sub>.

Furthermore, similar to the gold complex, both copper- and silver-aluminyll complexes feature a second driving force in the interaction with H<sub>2</sub> at the two TSs, that is a charge transfer from the H<sub>2</sub> HOMO towards the LUMO of the complex. So apparently, for all the metals and at both TSs, the nature of the H<sub>2</sub>-complex interaction is qualitatively analogous.

Differences between TS<sub>M</sub><sup>IV</sup> and TS<sub>M</sub><sup>I</sup> and between copper and its heavier homologues silver and gold become evident on a quantitative ground. In particular, the ETS-NOCV and CD analyses reveal that  $\Delta\rho'_1$  and  $\Delta\rho'_2$  have different relative weights in the H<sub>2</sub>–[t-Bu<sub>3</sub>PMAl(NON)] interaction. Indeed, for the gold complex, the associated  $\Delta E_{oi}^1/\Delta E_{oi}^2$  and CT<sub>1</sub>/CT<sub>2</sub> values are respectively –108.3/–19.2 kcal mol<sup>–1</sup> and 0.18/0.07 e at TS<sub>Au</sub><sup>IV</sup>, while these values are –59.7/–27.9 kcal mol<sup>–1</sup> and 0.18/0.12 for TS<sub>Au</sub><sup>I</sup> (Table S2 in the ESI†). An analogous trend is also observed for the  $\Delta E_{oi}^1/\Delta E_{oi}^2$  (CT<sub>1</sub>/CT<sub>2</sub>) values associated with TS<sub>M</sub><sup>IV</sup> and TS<sub>M</sub><sup>I</sup> (M = Cu, Ag), where the weight of the H<sub>2</sub>-to-complex charge transfer increases at TS<sub>M</sub><sup>I</sup> (Table S2 in the ESI†).

This result clearly shows that (i) the H<sub>2</sub>-complex main interaction is quantitatively similar upon changing the coinage metal and (ii) the relative weight of  $\Delta\rho'_2$  at TS<sub>M</sub><sup>I</sup> is larger for all the complexes, which can be explained based on the frontier molecular orbitals (FMO) analyses. From the M and Al atomic orbital (AO) contributions to the LUMO of the [t-Bu<sub>3</sub>PMAl(NON)] complex at the corresponding TS<sub>M</sub><sup>IV</sup> and TS<sub>M</sub><sup>I</sup> structures (Table S4 in the ESI†), differences become evident. The LUMO at TS<sub>M</sub><sup>IV</sup> displays a smaller coinage metal contribution compared to Al (see for instance 17.7% Cu vs. 23.5% Al at TS<sub>Cu</sub><sup>IV</sup>). On the other hand, at TS<sub>M</sub><sup>I</sup> the metal contribution is significantly increased (see for instance 43.9% Cu at TS<sub>Cu</sub><sup>I</sup>), with contributions from Al remaining almost unchanged. These increased metal contributions to the LUMO complex at TS<sub>M</sub><sup>I</sup> are responsible for a larger  $\Delta\rho'_2$  (*i.e.* an enhanced electrophilicity of the complexes), which should arise from the highly bent P–M–Al angle at all the TS<sub>M</sub><sup>I</sup>, favouring the LUMO delocalization over the metal and also explaining the preferred H<sub>2</sub> interaction at the metal site (and the corresponding lower activation barrier).

The extent of the P–M–Al bending also rationalizes the more energetically favourable pathway for the copper complex with respect to its silver and gold analogues. While the P–M–Al angle is below 130° for both TS<sub>Ag</sub><sup>I</sup> and TS<sub>Au</sub><sup>I</sup> (129.1° and 126.0°, respectively),

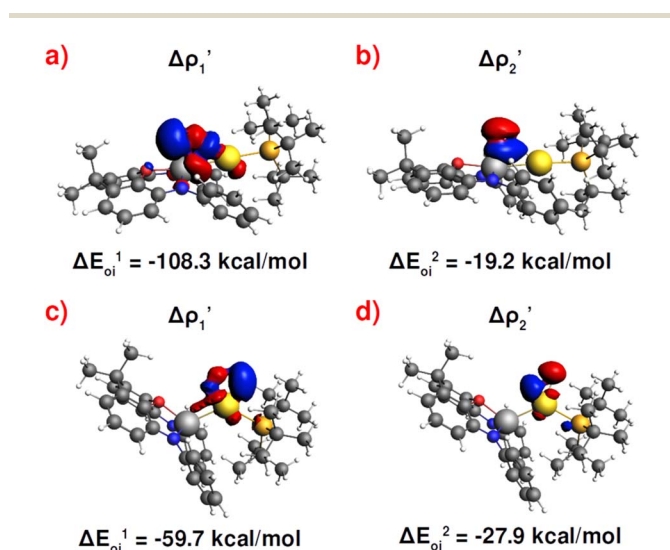


Fig. 3 Results of the NOCV analysis of the [H<sub>2</sub>]-[t-Bu<sub>3</sub>PAuAl(NON)] interaction at TS<sup>IV</sup> (top) and TS<sup>I</sup> (bottom). The isosurfaces for the  $\Delta\rho'_1$  and  $\Delta\rho'_2$  components for TS<sup>IV</sup> ((a) and (b), respectively) and TS<sup>I</sup> ((c) and (d), respectively) are reported together with the associated orbital interaction energy. The charge flux is red → blue. The isovalue is  $3me/a_0^3$  for all isosurfaces. See Fig. S5–S8 in the ESI† for the complete NOCV analysis.



respectively), the Cu–Al complex features a larger P–Cu–Al angle at  $TS_{Cu}^I$  ( $133.0^\circ$ ). This apparently marginal difference has a significant effect on the activation barrier, as shown by the activation strain model analysis<sup>41–43</sup> (ASM, see Tables S5–S7 in the ESI and Discussion therein<sup>†</sup>) of the reaction pathways. The less bent copper structure actually reduces the distortion penalty associated with the complex rearrangement from  $18.7 \text{ kcal mol}^{-1}$  at  $TS_{Au}^I$  to  $5.2 \text{ kcal mol}^{-1}$  at  $TS_{Cu}^I$ , thus lowering the activation barrier. The analysis of the variation of the metal AO contributions to the LUMO of the complex at different P–M–Al angles (see Fig. S19 and Table S8 in the ESI<sup>†</sup>) shows that the bending required to efficiently enhance the metal contribution to the LUMO is significantly lower for copper with respect to that of silver and gold, thus rationalizing the reduced bending deviation and therefore the lower distortion penalty for the copper–aluminum system.

The coinage metal–aluminum complex reactivity described here is very different from that of general transition metal–Lewis acid (TM–LA) compounds. For TM–LA, the presence of dative polarized  $TM(\delta^-)–LA(\delta^+)$  bonds usually favours the polarization of  $H_2$  and, thus, its dissociation and oxidative addition to the metal center.<sup>44</sup> A quantitative assessment of the different reactivities of M–Al (M = Cu, Ag, Au) vs. TM–LA is given in the ESI (see Fig. S20, Table S9 and Discussion therein),<sup>†</sup> where a platinum–aluminum complex featuring a dative  $Pt \rightarrow Al$  bond, reported by Bourissou and coworkers<sup>21</sup> to experimentally activate  $H_2$ , has been selected by us as a test case.

On the other hand, further analogies between gold–aluminum (and copper– and silver–aluminum) and digermene compounds can be inferred by investigating the radical pair recombination mechanism for the formation of product  $PC_{Au}$  in comparison with that of **6**. As mentioned earlier, this mechanism (radical  $H^\cdot$  abstraction followed by radical pair recombination) has been proposed for digermenes.<sup>8</sup> As shown in Fig. S21 in the ESI<sup>†</sup> and in Fig. 4, we modelled homolytic and heterolytic breaking of the two substrate–H bonds in  $PC_{Au}$  for probing which of these two mechanisms is the most favourable in this case.

The most suitable fragmentation (*i.e.* the one featuring the lowest associated energy for the dissociation of the substrate–H bonds) is the radical one (see Fig. 4 and S21 in the ESI<sup>†</sup>), meaning that the most favorable mechanism would be  $H^\cdot$  abstraction followed by a radical pair recombination (the associated  $\Delta E$  values are  $83.6$  and  $84.6 \text{ kcal mol}^{-1}$ , while  $\Delta E$  values of  $362.4/179.9 \text{ kcal mol}^{-1}$  and  $195.2/388.8 \text{ kcal mol}^{-1}$  are calculated for hydride/proton abstraction and recombination). Interestingly, the dissociation of the remaining hydrogen from the radicals to yield the gold–aluminum complex still features positive dissociation energies (Fig. 4a), suggesting that the complex is able to stabilize the radical pair. Such an ability is preserved even when accounting for the geometrical relaxation of the gold–aluminum complex, which is, in absolute value, smaller than the dissociation energies.

Even more interestingly, this model indirectly reveals the diradical-like nature of this mechanism. By inspection of the spin densities on the radical  $[^tBu_3PAu(\mu-H)Al(NON)]^\cdot$  and  $[^tBu_3PAuAlH(NON)]^\cdot$  fragments (Fig. 4b), no spin polarization is observed on the bound hydrogen, with the spin density being

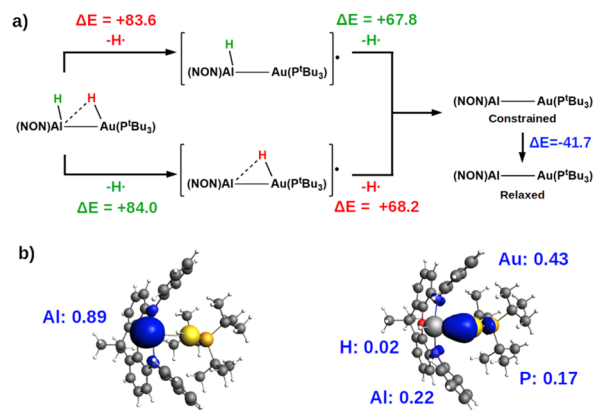


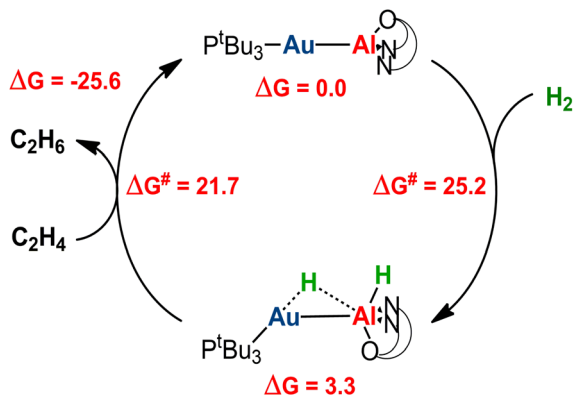
Fig. 4 (a) Pathways for the homolytic dissociation of the two H-substrate bonds in the singly bridged product  $[^tBu_3PAu(\mu-H)AlH(NON)]$  ( $PC_{Au}$ ) with relative associated energies ( $\text{kcal mol}^{-1}$ ). (b) Spin density (in blue) associated with the  $[^tBu_3PAu(\mu-H)Al(NON)]^\cdot$  (left) and  $[^tBu_3PAuAlH(NON)]^\cdot$  (right) radicals. The isovalue for the surface is  $5me/a_0^3$ . The most relevant atomic spin polarization values are reported.

mostly localized either on Al (spin polarization of  $0.89 \text{ e}$ ) or in the region between Au and Al atoms, (spin polarization values of  $0.43$  and  $0.22 \text{ e}$  for Au and Al, respectively). This finding demonstrates that the radical-like reactivity of these fragments has a fundamental role in favoring the activation of  $H_2$  *via* homolytic H–H dissociation and the consequent formation of the thermodynamically accessible  $PC_{Au}$ . This picture is strongly supported by the analogous dissociation paths and spin densities (Fig. S22 in the ESI<sup>†</sup>) observed for the model digermene reported in ref. 12.

Analogously, the same model applied for studying the formation of  $PC_{Cu}/PC_{Ag}$  and  $PC'_{Cu}/PC'_{Ag}$  supports a similar radical-pair stabilization mechanism for both copper- and silver-aluminum complexes. As displayed in Fig. S23–S26 in the ESI,<sup>†</sup>  $H^\cdot$  abstraction followed by radical pair recombination is favored for both  $PC_{Cu}$  and  $PC_{Ag}$  (the associated  $\Delta E$  values are  $88.7/85.9$  and  $83.3/85.0 \text{ kcal mol}^{-1}$  respectively, see Fig. S24<sup>†</sup>), while the  $\Delta E$  values associated with the hydride/proton abstraction and recombination mechanism are much higher (see Fig. S23<sup>†</sup>). Analogous values have also been calculated for the same mechanisms in the case of  $PC'_{Cu}$  and  $PC'_{Ag}$  (see Fig. S25 and S26<sup>†</sup>).

Notably, monitoring the charge migration along the reaction paths with the evolution of Voronoi deformation density (VDD)<sup>45</sup> atomic charges on the metal ( $q_M$ ) and aluminium ( $q_{Al}$ ) atoms, as well as on the two hydrogen atoms of  $H_2$  ( $q_{H_1}$  and  $q_{H_2}$ ), indirectly confirms this mechanistic picture. As shown in Table S10 in the ESI,<sup>†</sup> both  $q_M$  and  $q_{Al}$  become increasingly positive along the reaction paths (indicating charge depletion), while  $q_{H_1}$  and  $q_{H_2}$  become increasingly negative (indicating charge accumulation on  $H_2$ ). Remarkably, both increases of  $q_M/q_{Al}$  and  $q_{H_1}/q_{H_2}$  are almost symmetrical for all the stationary points, indicating the absence of  $H_2$  polarization and further confirming the bimetallic cooperative radical pair stabilization mechanism. Furthermore, the VDD atomic charge evolution along the reaction profiles suggests that, starting from what we





Scheme 3 Schematic computational catalytic cycle for the hydrogenation of ethylene catalyzed by the  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$  complex **1**. Relative free energies and free energy activation barriers are given in  $\text{kcal mol}^{-1}$ .

expect to be a formal  $\text{M}(0)$  species in the initial complexes based on previous results on  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$ ,<sup>46</sup> the oxidation state at  $\text{M}$  (and  $\text{Al}$ ) remains formally unaltered.

Catalytic application of the hydrogenated  $\text{PC}'_{\text{M}}/\text{PC}_{\text{M}}$  species is an exciting prospect. In particular, the almost thermoneutral formation of  $\text{PC}_{\text{Au}}$  is virtually an ideal condition for the  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$  complex to be used as a catalyst in hydrogenation reactions.<sup>47</sup> Selecting ethylene as a substrate, we modeled the alkene hydrogenation process catalyzed by  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$ , as shown in Scheme 3 (the full path and structures are reported in Fig. S27 and S28 in the ESI†).

Starting from  $\text{PC}_{\text{Au}}$ , hydrogenation of ethylene occurs in a single, exergonic step ( $-25.6 \text{ kcal mol}^{-1}$ ), *via* a concerted transition state ( $\text{TS}_{\text{Au}}^{\text{cat}}$ , see Fig. S28 in the ESI†), where both hydrogens are simultaneously transferred to the substrate, forming ethane and regenerating the gold–aluminum catalyst, with a kinetically accessible barrier ( $21.7 \text{ kcal mol}^{-1}$ ). Given the enhanced stability of  $\text{PC}_{\text{Cu}}$  and  $\text{PC}_{\text{Ag}}$  (thus representing larger thermodynamic sinks), this catalytic mechanism is predicted to be less efficient for copper- and silver–aluminum complexes. Indeed, as shown in Fig. S27,† while the ethane formation is found to be exergonic for both species ( $\Delta G = -15.7$  and  $-20.3 \text{ kcal mol}^{-1}$  for  $\text{Cu}$  and  $\text{Ag}$ , respectively), the activation barrier is higher for both metals ( $29.4$  and  $26.8 \text{ kcal mol}^{-1}$  for  $\text{Cu}$  and  $\text{Ag}$ , respectively), thus reducing the possible catalytic performance of the copper- and silver–aluminum complexes.

## Conclusions

In conclusion, computational evidence for the activation of  $\text{H}_2$  by coinage metal–aluminum complexes *via* a radical pair stabilization mechanism is provided, in close resemblance with experimentally observed dihydrogen activation with group 14 dimetallenes and dimetallynes, and at a variance with the reactivity of general  $\text{TM-LA}$  complexes. This offers an alternative paradigm for small molecule activation by electron-rich and highly covalent metal–metal bonds which are able to induce a radical-like reactivity. The kinetically accessible and almost thermoneutral formation of product species

appears certainly as an ideal condition for the use of the  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$  complex as a catalyst for hydrogenation of, for instance, unsaturated C–C bonds.<sup>47</sup>

This work falls within the sustainable “catalysis by design” as a green key technology in continuing the search for innovative strategies for small molecule activation, and strongly calls for systematic experimental feedback.<sup>48</sup>

## Computational details

Complexes  $[\text{Bu}_3\text{PMAl}(\text{NON})]$  ( $\text{M} = \text{Cu}, \text{Ag}, \text{Au}$ ) have been slightly simplified at the  $\text{NON}$  site by replacing the two *tert*-butyl groups at the peripheral positions of the dimethylxanthene moiety with hydrogen atoms and the two Dipp substituents on the nitrogen atoms with phenyl groups. The effect of modelling on this class of complexes has been extensively evaluated in ref. 2 and 4 where the same computational set up as that used in the present work was applied. Good agreement with experimental data was found for geometries, and in general, both the reaction mechanism and the electronic structure calculations show negligible deviations due to the structural simplifications used.

All geometry optimizations and frequency calculations on optimized structures (minima with zero imaginary frequencies and transition states with one imaginary frequency) for the  $\text{H}_2$  reaction with the  $[\text{Bu}_3\text{PMAl}(\text{NON})]$  ( $\text{M} = \text{Cu}, \text{Ag}, \text{Au}$ ) complexes have been carried out using the Amsterdam density functional (ADF) code<sup>49,50</sup> in combination with the related Quantum-regions Interconnected by Local Description (QUILD) program.<sup>51</sup> The PBE<sup>52</sup> GGA exchange–correlation (XC) functional, the TZ2P basis set with a small frozen core approximation for all atoms, the ZORA Hamiltonian<sup>53–55</sup> for treating scalar relativistic effects and the Grimme’s D3-BJ dispersion correction were used.<sup>56,57</sup> Solvent effects were modelled employing the conductor-like screening model (COSMO) with the default parameters for toluene as implemented in the QUILD code.<sup>58</sup> Effects of the exchange–correlation functional and solvation in this framework have been recently evaluated and are found, overall, to negligibly affect the results.<sup>4</sup> The effect of different exchange–correlation functionals on the reaction energetics has been also investigated in this work, highlighting an overall marginal impact on both the kinetics and thermodynamics of the reported reactions (see Table S11 and Fig. S29 in the ESI†).

The same computational setup has also been used for the AMS, EDA, CD-NOCV and ETS-NOCV analyses, as well as for the calculation of Voronoi deformation density (VDD)<sup>45</sup> charges along the reaction paths. This computational protocol has been used in ref. 1 and 2 to study the  $[\text{Bu}_3\text{PAuAl}(\text{NON})]$  and  $[\text{Bu}_3\text{PAuCO}_2\text{Al}(\text{NON})]$  complexes and to investigate the mechanisms of the  $\text{CO}_2$  insertion reaction in similar compounds featuring gold and group 13 elements.<sup>3–5</sup>

The analysis of the transition state  $\text{TS}_2$  for the reaction of the aluminum–platinum complex with  $\text{H}_2$  has been carried out with the same computational protocol on the geometry optimized in the original work.<sup>21</sup> Similarly, the radical recombination pathways for the model digermene compounds



have been investigated at the same computational level using the geometries provided in ref. 12.

The transition state theory (TST)<sup>31</sup> calculations for the evaluation of tunnel effects have been carried out with the kinetic and statistical thermodynamic package (KiSThelp)<sup>59</sup> computer code. Activation energies have been computed from the rate constant with the three-parameter fitted Arrhenius equation. The non-corrected activation energy ( $E_a^{\text{TST}}$ ) has been obtained from fitting of the conventional TST-calculated rate constant, while the tunneling-corrected activation energy ( $E_a^{\text{TST-Eckart}}$ ) has been obtained *via* fitting of the TST rate constant corrected with the Eckart tunneling correction.<sup>32</sup>

## Data availability

All the relevant data (including methodology description, and mechanistic, CD-NOCV, ASM, EDA, VDD and TST data discussed in the manuscript) are provided within the ESI.†

## Author contributions

D. S., L. B. and P. B. conceived the project, D. S., L. B. and P. B. performed the research, and D. S., L. B. and P. B. wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by the Ministero dell'Università e della Ricerca (MUR, project AMIS, through the program "Dipartimenti di Eccellenza – 2018–2022").

## Notes and references

- J. Hicks, A. Mansikkamäki, P. Vasko, J. M. Goicoechea and S. Aldridge, *Nat. Chem.*, 2019, **11**, 237–241.
- D. Sorbelli, L. Belpassi and P. Belanzoni, *J. Am. Chem. Soc.*, 2021, **143**, 14433–14437.
- D. Sorbelli, L. Belpassi and P. Belanzoni, *Inorg. Chem.*, 2022, **61**, 1704–1716.
- D. Sorbelli, L. Belpassi and P. Belanzoni, *Chem. Sci.*, 2022, **13**, 4623–4634.
- D. Sorbelli, E. Rossi, R. W. A. Havenith, J. E. M. N. Klein, L. Belpassi and P. Belanzoni, *Inorg. Chem.*, 2022, **61**, 7327–7337.
- C. McManus, J. Hicks, X. Cui, L. Zhao, G. Frenking, J. M. Goicoechea and S. Aldridge, *Chem. Sci.*, 2021, **12**, 13458–13468.
- J. Li, M. Hermann, G. Frenking and C. Jones, *Angew. Chem., Int. Ed.*, 2012, **51**, 8611–8614.
- G. H. Spikes, J. C. Fettinger and P. P. Power, *J. Am. Chem. Soc.*, 2005, **127**, 12232–12233.
- J. Li, C. Schenk, C. Goedecke, G. Frenking and C. Jones, *J. Am. Chem. Soc.*, 2011, **133**, 18622–18625.
- T. J. Hadlington, M. Hermann, J. Li, G. Frenking and C. Jones, *Angew. Chem., Int. Ed.*, 2013, **52**, 10199–10203.
- L. Zhao, F. Huang, G. Lu, Z.-X. Wang and P. v. R. Schleyer, *J. Am. Chem. Soc.*, 2012, **134**, 8856–8868.
- M. Hermann, C. Goedecke, C. Jones and G. Frenking, *Organometallics*, 2013, **32**, 6666–6673.
- F. Hanusch, L. Groll and S. Inoue, *Chem. Sci.*, 2021, **12**, 2001–2015.
- G. D. Frey, V. Lavallo, B. Donnadieu, W. W. Schoeller and G. Bertrand, *Science*, 2007, **316**, 439–441.
- J. Hicks, P. Vasko, J. M. Goicoechea and S. Aldridge, *Nature*, 2018, **557**, 92–95.
- A. Hinz and F. Breher, *Angew. Chem., Int. Ed.*, 2018, **57**, 8818–8820.
- M. J. Evans, M. D. Anker, C. L. McMullin, S. E. Neale and M. P. Coles, *Angew. Chem., Int. Ed.*, 2021, **60**, 22289–22292.
- N. Villegas-Escobar, A. Toro-Labbé and H. F. Schaefer, *Chem.–Eur. J.*, 2021, **27**, 17369–17378.
- G. J. Kubas, *Chem. Rev.*, 2007, **107**, 4152–4205.
- P. E. M. Siegbahn, *Adv. Inorg. Chem.*, 2004, **56**, 101–125.
- M. Devillard, R. Declercq, E. Nicolas, A. W. Ehlers, J. Backs, N. Saffon-Merceron, G. Bouhadir, J. C. Slootweg, W. Uhl and D. Bourissou, *J. Am. Chem. Soc.*, 2016, **138**, 4917–4926.
- M. Xu, A. R. Jupp, Z. W. Qu and D. W. Stephan, *Angew. Chem., Int. Ed.*, 2018, **57**, 11050–11054.
- K. L. Bamford, Z. W. Qu and D. W. Stephan, *J. Am. Chem. Soc.*, 2019, **141**, 6180–6184.
- Z.-W. Qu, H. Zhu, S. Grimme, Z. Qu, H. Zhu and S. Grimme, *ChemCatChem*, 2021, **13**, 3401–3404.
- R. J. McMahon, *Science*, 2003, **299**, 833–834.
- R. A. More O'Ferrall, *J. Phys. Org. Chem.*, 2010, **23**, 572–579.
- D. Ley, D. Gerbig and P. R. Schreiner, *Org. Biomol. Chem.*, 2012, **10**, 3781–3790.
- J. Meisner and J. Kästner, *Angew. Chem., Int. Ed.*, 2016, **55**, 5400–5413.
- J. E. M. N. Klein, D. Mandal, W. M. Ching, D. Mallick, L. Que and S. Shaik, *J. Am. Chem. Soc.*, 2017, **139**, 18705–18713.
- P. R. Schreiner, *Trends Chem.*, 2020, **2**, 980–989.
- H. Eyring, *Chem. Rev.*, 1935, **17**, 65–77.
- C. Eckart, *Phys. Rev.*, 1930, **35**, 1303.
- M. Mitoraj and A. Michalak, *J. Mol. Model.*, 2007, **13**, 347–355.
- A. Michalak, M. Mitoraj and T. Ziegler, *J. Phys. Chem. A*, 2008, **112**, 1933–1939.
- K. Morokuma, *J. Chem. Phys.*, 1971, **55**, 1236–1244.
- T. Ziegler and A. Rauk, *Theor. Chim. Acta*, 1977, **46**, 1–10.
- L. Zhao, M. von Hopffgarten, D. M. Andrada and G. Frenking, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2018, **8**, 1345.
- L. Belpassi, I. Infante, F. Tarantelli and L. Visscher, *J. Am. Chem. Soc.*, 2008, **130**, 1048–1060.
- G. Bistoni, S. Rampino, F. Tarantelli and L. Belpassi, *J. Chem. Phys.*, 2015, **142**, 084112.
- GitHub, BERTHA-4c-DKS/pycubesd, <https://github.com/BERTHA-4c-DKS/pycubesd>, accessed, 10 October 2022.
- I. Fernández and F. M. Bickelhaupt, *Chem. Soc. Rev.*, 2014, **43**, 4953–4967.



- 42 F. M. Bickelhaupt and K. N. Houk, *Angew. Chem., Int. Ed.*, 2017, **56**, 10070–10086.
- 43 P. Vermeeren, S. C. C. van der Lubbe, C. F. Guerra, F. M. Bickelhaupt and T. A. Hamlin, *Nat. Protoc.*, 2020, **15**, 649–667.
- 44 R. M. Charles and T. P. Brewster, *Coord. Chem. Rev.*, 2021, **433**, 213765.
- 45 C. F. Guerra, J.-W. Handgraaf, E. J. Baerends and F. M. Bickelhaupt, *J. Comput. Chem.*, 2004, **25**, 189–210.
- 46 I. F. Leach, D. Sorbelli, L. Belpassi, P. Belanzoni, R. W. A. Havenith and J. E. M. N. Klein, *Dalton Trans.*, 2023, **52**, 11–15.
- 47 P. Su, Y. Li and Z. Ke, *Chem.-Asian J.*, 2021, **16**, 3427–3436.
- 48 We would like to make the readership aware that these authors have been informed by email from Prof. Simon Aldridge that some experiments, carried out in his group in Oxford, seem to show no evidence of basic reactivity of the Au/Al (and Cu/Al) system with H<sub>2</sub>. Unfortunately, to date, these experimental results have not been published (or made available in any public repository), to the best of our knowledge. These authors themselves are not aware of any experimental information, such as, for instance, experimental conditions (solvent, temperature, pressure, etc.) employed.
- 49 *ADF Manual ADF Program System Release 2014*, 1993.
- 50 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**, 931–967.
- 51 M. Swart and F. M. Bickelhaupt, *J. Comput. Chem.*, 2008, **29**, 724–734.
- 52 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 53 E. Van Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1993, **99**, 4597–4610.
- 54 E. Van Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1994, **101**, 9783–9792.
- 55 E. Van Lenthe, *J. Chem. Phys.*, 1999, **110**, 8943–8953.
- 56 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.
- 57 S. Grimme, S. Ehrlich and L. Goerigk, *J. Comput. Chem.*, 2011, **32**, 1456–1465.
- 58 C. C. Pye and T. Ziegler, *Theor. Chem. Acc.*, 1999, **101**, 396–408.
- 59 S. Canneaux, F. Bohr and E. Henon, *J. Comput. Chem.*, 2014, **35**, 82–93.

