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# Robust dicopper(I) $\mu$ -boryl complexes supported by a dinucleating naphthyridine-based ligand†

Pablo Rios,<sup>a</sup> Matthew S. See,<sup>a</sup> Rex C. Handford,<sup>a</sup> Simon J. Teat<sup>b</sup> and T. Don Tilley<sup>b,\*a</sup>

Copper boryl species have been widely invoked as reactive intermediates in Cu-catalysed C–H borylation reactions, but their isolation and study have been challenging. Use of the robust dinucleating ligand DPFN (2,7-bis(fluoro-di(2-pyridyl)methyl)-1,8-naphthyridine) allowed for the isolation of two very thermally stable dicopper(I) boryl complexes, [(DPFN)Cu<sub>2</sub>( $\mu$ -Bpin)][NTf<sub>2</sub>] (**2**) and [(DPFN)Cu<sub>2</sub>( $\mu$ -Bcat)][NTf<sub>2</sub>] (**4**) (pin = 2,3-dimethylbutane-2,3-diol; cat = benzene-1,2-diol). These complexes were prepared by cleavage of the corresponding diborane *via* reaction with the alkoxide [(DPFN)Cu<sub>2</sub>( $\mu$ -O<sup>t</sup>Bu)][NTf<sub>2</sub>] (**3**). Reactivity studies illustrated the exceptional stability of these boryl complexes (thermal stability in solution up to 100 °C) and their role in the activation of C(sp)–H bonds. X-ray diffraction and computational studies provide a detailed description of the bonding and electronic structures in these complexes, and suggest that the dinucleating character of the naphthyridine-based ligand is largely responsible for their remarkable stability.

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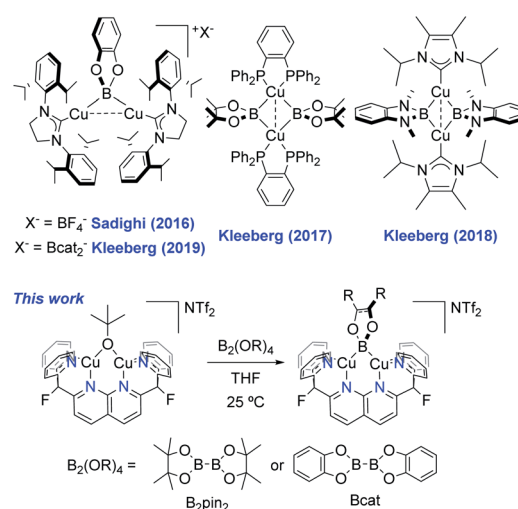
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## Introduction

Copper-catalysed C–H borylation is a powerful tool in organic chemistry due to the stability and versatility of the resulting organoboron species as synthetic building blocks.<sup>1</sup> This catalysis is thought to involve Cu(I) boryl (Cu–BR<sub>2</sub>) complexes as key reactive intermediates. However, the chemistry of Cu(I) boryls has only recently been observed; while the first proposal of a Cu–BR<sub>2</sub> intermediate stems from a 2000 report by Miyaura<sup>2</sup> and Hosomi,<sup>3</sup> the first well-defined Cu(I) boryl species was described by Sadighi and coworkers in 2005.<sup>4</sup> Since then, only a few have been isolated.<sup>5</sup> The paucity of examples is likely due to their inherent instability, given that most of those reported decompose in solution at or below room-temperature to elemental Cu.<sup>4,5</sup> Notably, these cases involve monodentate *N*-heterocyclic carbene (NHC) or phosphine supporting ligands. The reported examples stable at room-temperature often make use of apolar solvents (*e.g.* benzene), such as the ring-expanded NHC-supported (6-Dipp)CuBpin complex reported by Liptrout *et al.*<sup>5h</sup> In this example, limited decomposition is observed at 25 °C after several weeks. In other cases, the empty p orbital on boron is stabilised by means of adjacent nitrogen atoms, as in the systems described by Yamashita, Nozaki *et al.* Their synthetic procedure involves mixing an anionic boryl ligand

with copper halides, giving the corresponding boryl complexes, generally in *ca.* 25–30% yield.<sup>5e,f</sup>

In 2016, Sadighi reported the first dinuclear boryl, {(SIPr)Cu<sub>2</sub>( $\mu$ -Bcat)}{BF<sub>4</sub>} (Scheme 1, top left, SIPr = 1,3-bis(2,6-diisopropylphenyl)imidazolin-2-ylidene), from reaction of {(SIPr)Cu<sub>2</sub>( $\mu$ -OSiMe<sub>3</sub>)}{BF<sub>4</sub>} with bis(catecholato)diboron (B<sub>2</sub>cat<sub>2</sub>) at –35 °C.<sup>6</sup> The same dicopper cation was obtained by Kleeberg in 2019 upon treatment of mononuclear (SIPr)CuO<sup>t</sup>Bu with B<sub>2</sub>cat<sub>2</sub> in THF. Interestingly, mononuclear (SIPr)CuBcat was observed when the reaction was carried out in toluene.<sup>7</sup> By modifying the boryl and supporting ligands, the Kleeberg group obtained additional examples of neutral dicopper boryls



Scheme 1 Examples of dicopper(I)  $\mu$ -boryl complexes.

<sup>a</sup>Department of Chemistry, University of California, Berkeley, USA

<sup>b</sup>Advanced Light Source, Lawrence Berkeley National Laboratory, Berkeley, CA 94720-1460, USA. E-mail: tdttilley@berkeley.edu

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(Scheme 1).<sup>5c,8</sup> However, the latter species readily decompose in solution at or below room-temperature and require storage in the solid-state under an inert atmosphere, complicating their characterisation and study as catalytic intermediates.

Here we report the synthesis and characterisation of two dicopper(i) boryl complexes supported by a 1,8-naphthyridine-based dinucleating ligand. These species are persistent in solution and tolerate heating to at least 100 °C. In addition, their role in the activation of C(sp)–H bonds and the origins of their stability are described.

## Results and discussion

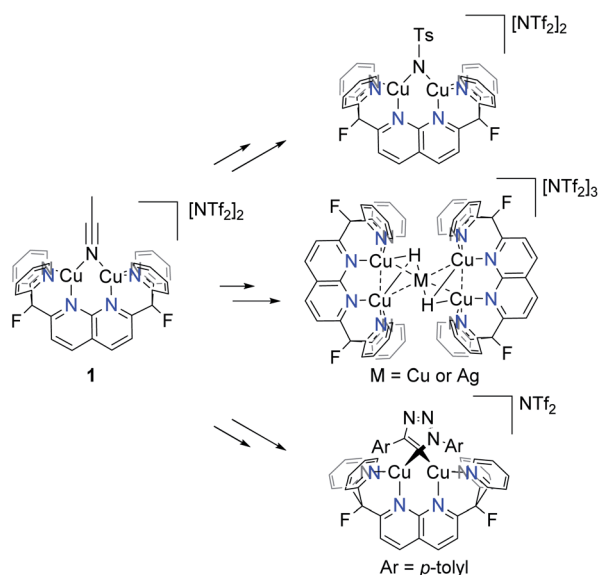
Previous results from this laboratory established 2,7-bis(fluoro-di(2-pyridyl)methyl)-1,8-naphthyridine (DPFN) as an effective platform for stabilisation of bimetallic units. Dicopper(i) [(DPFN)Cu<sub>2</sub>]<sup>+</sup> complexes, accessed *via* the convenient starting material [(DPFN)Cu<sub>2</sub>(μ-NCMe)][NTf<sub>2</sub>]<sub>2</sub> (**1**),<sup>9</sup> exhibit remarkable bimetallic electrophilicity and have allowed for the study of several elusive reactive intermediates such as a dicopper(i,ii) nitrenoid,<sup>10</sup> pentanuclear metal hydrides,<sup>11</sup> and an unprecedented dicopper(i) bridging triazolidine (Scheme 2).<sup>12–14</sup> Thus, [(DPFN)Cu<sub>2</sub>]<sup>n+</sup> seemed a promising framework for the synthesis and stabilisation of dicopper boryl units.

Due to the electrophilic character of **1**, introduction of a boryl fragment requires an anionic borylation reagent. Whereas few synthons for boryl anions are available, a Lewis base adduct of a tetraalkoxy diboron (with *e.g.*, alkoxide, fluoride or NHC) gives an isolable sp<sup>2</sup>–sp<sup>3</sup> anionic diboron compound capable of transferring a boryl group to an electrophile under metal-free conditions.<sup>15</sup> To pursue this possibility, one equivalent of K[B<sub>2</sub>pin<sub>2</sub>O<sup>−</sup>Bu] was added to complex **1** in *ortho*-difluorobenzene (*o*-DFB) at 25 °C to give a color change from orange to dark green after 3 h. A <sup>1</sup>H NMR analysis revealed

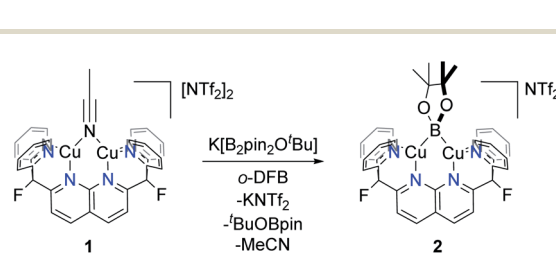
formation of a single new symmetrical species (**2**) with a set of resonances corresponding to the DPFN ligand. This was corroborated by <sup>19</sup>F{<sup>1</sup>H} NMR spectroscopy, which showed that the singlet at −175.1 ppm for **1** had been replaced by a singlet at −177.8 ppm. The <sup>1</sup>H NMR spectrum of the reaction mixture revealed the presence of free MeCN and <sup>t</sup>BuOBpin, as well as a singlet at 1.44 ppm associated with **2**. This assignment is supported by the <sup>11</sup>B NMR spectrum containing a single broad resonance for <sup>t</sup>BuOBpin at 21.7 ppm, in agreement with reported values (Fig. S13†).<sup>16</sup> Altogether, these data suggest the formation of boryl complex [(DPFN)Cu<sub>2</sub>(μ-Bpin)][NTf<sub>2</sub>]<sub>2</sub> (**2**, Scheme 3).

Scale-up of the synthesis of **2**, or use of THF as solvent, led to formation of several side-products as ascertained by <sup>1</sup>H NMR analysis (Fig. S14†). One possible side product could result from the transfer of −O<sup>t</sup>Bu from K[B<sub>2</sub>pin<sub>2</sub>O<sup>−</sup>Bu]. To test this hypothesis, one equivalent of KO<sup>t</sup>Bu was added to a solution of **1** in *o*-DFB at 25 °C, after which the reaction mixture became dark orange (Scheme 4). The <sup>1</sup>H NMR spectrum after 1 h exhibited a new set of resonances resulting from the DPFN ligand, along with a new singlet at 1.76 ppm indicating the clean formation of [(DPFN)Cu<sub>2</sub>(μ-O<sup>t</sup>Bu)][NTf<sub>2</sub>]<sub>2</sub> (**3**). The resonances for **3** correspond to those of one of the aforementioned side-products formed in the synthesis of **2**, and demonstrates −O<sup>t</sup>Bu transfer to some extent.

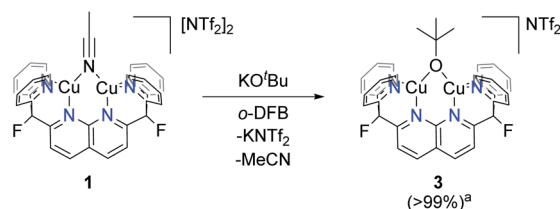
Complex **3** was isolated by precipitation from *o*-DFB or THF with pentane, but removal of volatile components from the resulting powder *in vacuo* resulted in conversion to an insoluble, intractable material. Nonetheless, dark brown X-ray quality crystals were grown by layer diffusion of O(SiMe<sub>3</sub>)<sub>2</sub> into an *o*-DFB solution of **3** (generated in solution) after 3 days at 25 °C.<sup>17</sup> The solid-state structure, as determined by single crystal diffraction analysis, confirms the proposed assignment (Fig. 1). While the Cu–O bond distances are similar to those observed in previously reported dicopper complexes containing a 1,8-naphthyridine diphosphine ligand (*ca.* 1.9 Å),<sup>18</sup> the Cu...



Scheme 2 Examples of detected and/or isolated reactive intermediates containing the [(DPFN)Cu<sub>2</sub>]<sup>n+</sup> scaffold.



Scheme 3 Initial synthesis of complex **2**.



Scheme 4 Synthesis of complex **3**. <sup>a</sup>Conversion determined by <sup>1</sup>H NMR spectroscopy.



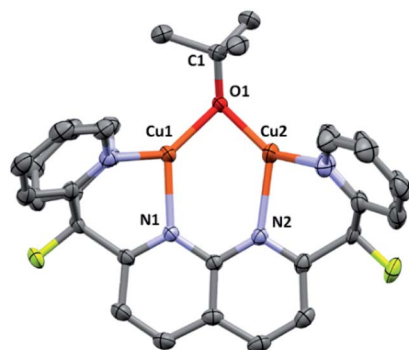


Fig. 1 Solid-state molecular structure of **3** (50% probability ellipsoids); H atoms are omitted for clarity. Only the major disorder component of the *tert*-butoxide fragment is shown.

Cu distance in **3** is much shorter (2.687(1) Å vs. 2.9–3.0 Å) and the Cu–O–Cu angle is considerably more acute (87.6(2)° vs. 100–108°), indicating a closer metal–metal contact enforced by the dipyridyl (vs. phosphinyl) side-arms. However, the metrical parameters for **3** are in good agreement with previous results from this laboratory on a dicopper(I) DPFN complex with a bridging aryloxide ligand.<sup>19</sup>

With a reliable route to complex **3**, reactions with diboranes were explored. Addition of B<sub>2</sub>pin<sub>2</sub> to complex **3** in THF (prepared *in situ* from **1**) resulted in a dark green solution after stirring at 25 °C for 15 h. Clean formation of boryl species **2** and <sup>t</sup>BuOBpin (Scheme 5) was observed by <sup>1</sup>H, <sup>19</sup>F{<sup>1</sup>H} and <sup>11</sup>B {<sup>1</sup>H} NMR spectroscopy, and no side-products were observed upon increasing the concentration of the reaction solution. Complex **2** was isolated as green crystals by layering pentane over a *o*-DFB/THF solution which gave analytically pure **2** in 75% yield. The identity of the complex (Fig. 2, top) was confirmed by single-crystal X-ray diffraction. Similarly, B<sub>2</sub>cat<sub>2</sub> reacted with complex **3** generated *in situ* to give the corresponding dicopper(I) boryl complex [(DPFN)Cu<sub>2</sub>(μ-Bcat)][NTf<sub>2</sub>] (**4**) after 2 h at 25 °C (Scheme 5), as judged by multinuclear

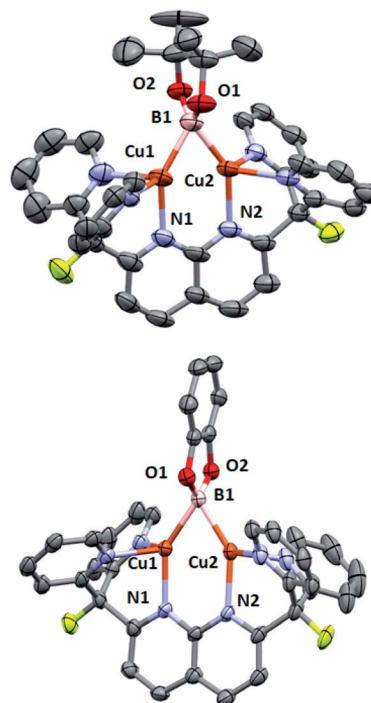
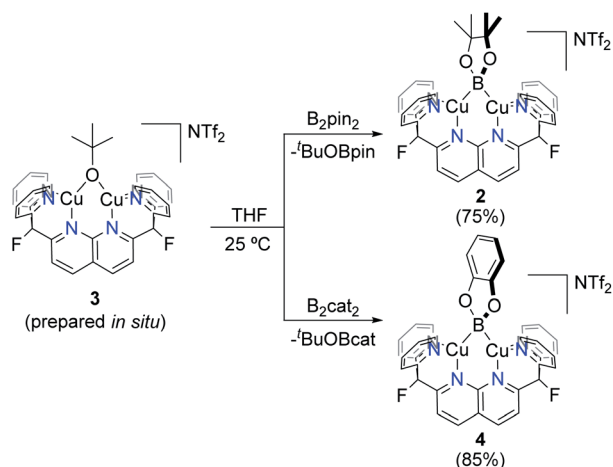


Fig. 2 Solid-state molecular structures (50% probability ellipsoids) of **2** (top) and **4** (bottom); H atoms are omitted for clarity.

NMR analysis on the resulting purple THF solution. The much shorter reaction time compared to that for the formation of **2** might be due to the higher Lewis acidity of B<sub>2</sub>cat<sub>2</sub> in comparison to B<sub>2</sub>pin<sub>2</sub>.<sup>20</sup> Similar work-up conditions afforded pure **4** as green X-ray quality crystals in 85% yield (Fig. 2, bottom). Unlike most previously described mono- or dicopper boryl complexes, **2** and **4** were synthesised and isolated at 25 °C, indicating a considerably higher stability. This is also reflected in mass spectra (Fig. S28 and S29†), which indicate persistence of the dinuclear structures under the ionisation conditions.

The increased stability of these boryl complexes in solution raises interest in their bond metrics compared to those of reported, thermally unstable analogues. However, **2** and **4** possess geometrical parameters that are within the range of those reported for dicopper μ-boryl complexes. The Cu–B bond distances (2.07–2.09 Å) are considerably similar to those in cationic {[(SIPr)Cu]<sub>2</sub>(μ-Bcat)}{BF<sub>4</sub>} (2.04–2.05 Å),<sup>6</sup> but are substantially shorter than those in the dinuclear neutral complexes synthesised by Kleeberg and co-workers (>2.17 Å).<sup>5c,8</sup> Similarly, **2** and **4** feature Cu–B–Cu bond angles (67–68°) that are more acute than that in Sadighi's dinuclear complex (72.1(2)°) but more obtuse than in Kleeberg's complexes (*ca.* 60°). Finally, the Cu⋯Cu distances (*ca.* 2.32 Å) are between those of the cationic (2.4083(9) Å)<sup>6</sup> and neutral (*ca.* 2.22–2.27 Å)<sup>5c,8</sup> μ-boryl examples previously reported. Nevertheless, all such distances are significantly shorter than the sum of covalent radii for Cu (2.64 Å),<sup>21</sup> implying that the short metal–metal distances are due to the bridging boryl ligands and/or cuprophilic interactions.<sup>22</sup> Interestingly, the increased stability of **2**



Scheme 5 Synthesis of complexes **2** and **4** from complex **3** (generated *in situ*). Isolated yields in parentheses.

and **4** in solution does not seem to be attributable to peculiar binding metrics.

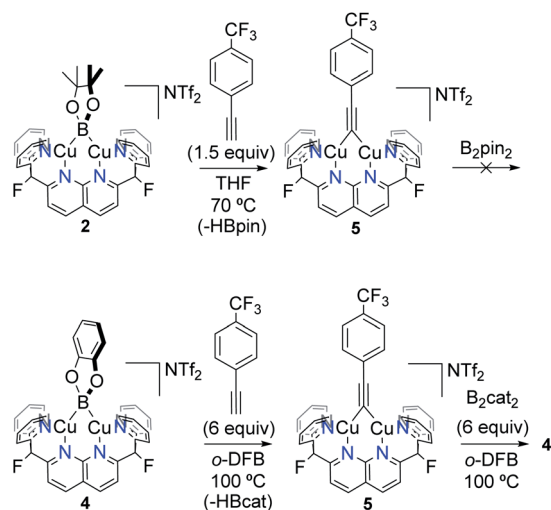
The robust character of these boryl species is also manifested in their reactivity. Both **2** and **4** exhibited no reaction with a number of small molecules including internal alkynes, olefins, silanes, azides or organic molecules possessing weak C–H bonds (*e.g.* fluorene), over a range of reaction conditions (THF or *o*-DFB as solvents and temperatures up to 70 or 100 °C, respectively; Table S1/ Fig. S15 and S16†). In all cases, these bimetallic complexes coexist in solution with the added reagent even at reaction times of 24 h. In contrast to results reported by Kleeberg *et al.* for dicopper boryls, **2** and **4** are inert towards 4-iodotoluene.<sup>8</sup> Aldehydes (1,3,5-trioxane or mesitylaldehyde) did not react with **2** or **4**, in stark contrast to results observed for (IPr)CuBpin (IPr = 1,3-bis(2,6-diisopropylphenyl)imidazol-2-ylidene),<sup>23</sup> suggesting that nuclearity might be one of the reasons behind this difference in reactivity (*vide infra*). Warming of **2** or **4** to 75 °C in the presence of either CO<sub>2</sub> or O<sub>2</sub> gave rise to a complex mixture of products, as shown in the NMR spectra of Fig. S17 and S18,† whereas H<sub>2</sub> was unreactive. In contrast, HC≡C(*p*-CF<sub>3</sub>-C<sub>6</sub>H<sub>4</sub>) cleanly reacted with both bridging boryl complexes to yield [(DPFN)Cu<sub>2</sub>(μ-C≡C(C<sub>6</sub>H<sub>4</sub>)CF<sub>3</sub>)] [NTf<sub>2</sub>] (**5**),<sup>19</sup> as summarised in Scheme 6 (Fig. S19–S21†). Concomitant formation of HBpin is observed in the case of complex **2**. In the case of **4**, HBcat seems to decompose over time to catBOBcat under the reaction conditions. This has been confirmed by warming HBcat in *o*-DFB at 100 °C (see Fig. S22 and S27† for more information). The increased Lewis basicity of boryl **2** compared to **4** is evident from the reaction conditions necessary to achieve high (>80%) conversions: whereas **2** reacted with 1.5 equiv of alkyne at 70 °C in 21 hours, boryl **4** required more forcing conditions (6 equiv of alkyne and warming for 90 hours at 100 °C). This difference was further confirmed by monitoring reactions of **2** and **4** with a different alkyne, PhC≡CH, in parallel (by <sup>1</sup>H NMR spectroscopy). For this alkyne, both

reactions were slower than those observed for *p*-CF<sub>3</sub>C<sub>6</sub>H<sub>4</sub>C≡CH, as expected, and whereas the reaction of **2** with 1.5 equiv of alkyne reached 45% completion at 70 °C after 22 h, the reaction of **4** required a higher concentration of the alkyne (6 equiv) and higher temperature (100 °C) to reach 56% conversion after 24 h.

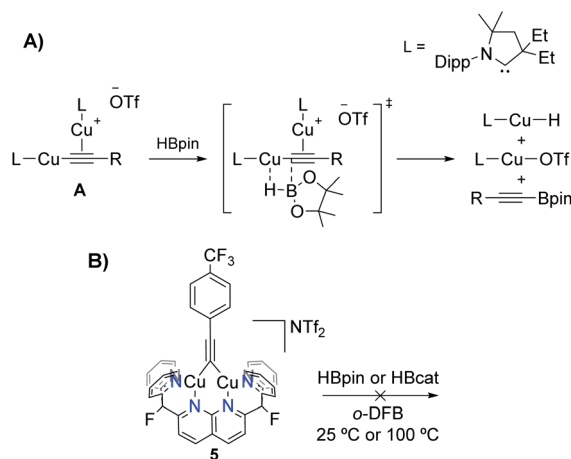
The proposed higher basicity of **2** finds additional support from DFT calculations (see Fig. S40†). Interestingly, bridging alkyne **5** further reacted with B<sub>2</sub>cat<sub>2</sub> (but not with B<sub>2</sub>pin<sub>2</sub>) to regenerate Bcat complex **4** (Fig. S21†), and this reactivity trend is consistent with the observed reactions of **3** with diboranes (*vide supra*). The boron-containing product is not readily identified in the reaction mixture involving **5** and B<sub>2</sub>cat<sub>2</sub>, as its NMR resonances are obscured by the solvent peak. Attempts to cleanly isolate and fully characterize this product have not been successful (see ESI†).

The clean deprotonation of HC≡C(*p*-CF<sub>3</sub>-C<sub>6</sub>H<sub>4</sub>) to give **5** at high temperatures is surprising, given the potential side-reactions that are possible in the presence of free hydroborane, such as alkyne hydroboration<sup>24</sup> or dehydrogenative borylation. Bertrand and coworkers recently reported catalytic C(sp)–H dehydrogenative borylation using carbene-stabilised Cu complexes, presumably by way of a catalytically active σ,π-dicopper acetylide intermediate **A** (Scheme 7A).<sup>25</sup> Unlike species **A**, the triple bond of bridging alkynyl **5** is not engaged in π-bonding to copper, making it susceptible to hydroboration processes. Control experiments involved the independent synthesis of **5** by treating the previously reported bridging phenyl complex [(DPFN)Cu<sub>2</sub>(μ-Ph)] [NTf<sub>2</sub>]<sup>26</sup> with HC≡C(*p*-CF<sub>3</sub>-C<sub>6</sub>H<sub>4</sub>). Addition of HBpin or HBcat to **5** did not result in reaction at 25 or 100 °C over 48 h (Fig. S25 and S26†). This divergence in reactivity from that observed by Bertrand *et al.* could be due to the different binding mode of the alkyne fragment to the dicopper core (σ,σ *vs.* σ,π), or to the rigidity imposed by the dinucleating DPFN ligand.

Computational methods were used to gain insights into the unusual robustness of bridging boryls **2** and **4**. First, steric



**Scheme 6** Reactivity of complexes **2** and **4** with HC≡C(*p*-CF<sub>3</sub>-C<sub>6</sub>H<sub>4</sub>), yielding complex **5**, which can further react with B<sub>2</sub>cat<sub>2</sub> to regenerate boryl species **4**.



**Scheme 7** (A) Terminal alkyne dehydrogenative borylation mechanism proposed by Bertrand and coworkers.<sup>25</sup> (B) Control experiments using **5** and hydroboranes that rule out alkyne hydroboration processes.





**Table 1** Natural charge and bond order values for selected computed dicopper(i)  $\mu$ -boryl complexes

Parameter	$[(\text{DPFN})\text{Cu}_2(\mu\text{-Bpin})]^+$	$[(\text{DPFN})\text{Cu}_2(\mu\text{-Bcat})]^+$	$\{[(\text{SIPr})\text{Cu}]_2(\mu\text{-Bcat})\}^{+a}$
Nat. charge on Cu	0.32	0.31	0.37/0.40
Nat. charge on B	0.58	0.59	0.50
Cu–B bond order	0.65	0.67	0.58
Cu–Cu bond order	0.73	0.72	0.32

<sup>a</sup> Values obtained from ref. 6.

congestion around the boron atom was quantified by determining its percent buried volume ( $\%V_{\text{bur}}$ ), described as the volume of a sphere centered on boron that is occupied by the  $[(\text{DPFN})\text{Cu}_2]^+$  scaffold and the substituents on B (Fig. S32–S35†).<sup>27,28</sup> As expected,  $\%V_{\text{bur}}$  values for **2** and **4** are remarkably large (*ca.* 80%), yet intermediate between values observed for unstable species in solution, namely mononuclear  $(\text{IPr})\text{CuBpin}$  ( $\%V_{\text{bur}} = 65.3\%$ ) and  $\{[(\text{SIPr})\text{Cu}]_2(\mu\text{-Bcat})\}\{\text{BF}_4\}$  ( $\%V_{\text{bur}} = 83.4\%$ ). In light of this, steric hindrance around boron is not likely to be a strong contributor to the stability.

DFT analyses were performed on **2** and **4** to understand their electronic structures, using the PBE0-D3/6-31g(d,p)/SDD level of theory on the cationic fragments.<sup>29</sup> The optimised structures were found to be local minima, with geometries in excellent agreement with metrics observed for the solid state structures (Table S2†). Natural charges<sup>30</sup> on Cu and B reflect minimal differences with those of related cationic systems, as displayed in Table 1. Likewise, Wiberg Bond Order analysis exhibited similar values for all Cu–B bonds, albeit the Cu $\cdots$ Cu interactions seem to be stronger in the cationic fragments of **2** and **4**, as expected based on the observed, short Cu $\cdots$ Cu distances. In addition, no bond critical point between the Cu atoms was observed by QTAIM calculations (Fig. S38 and S39†).<sup>31,32</sup>

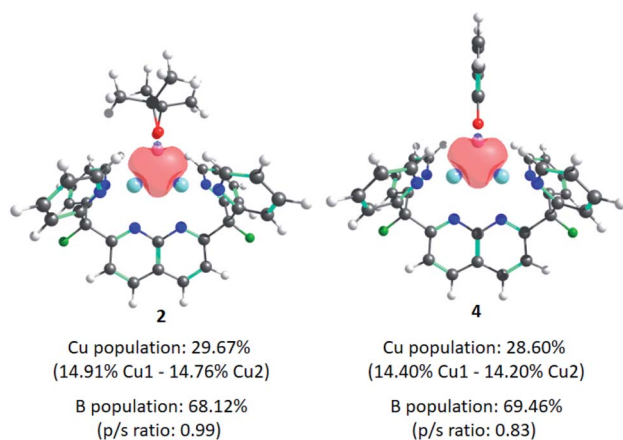
Finally, the electron distribution in the Cu–B bonds was investigated, since studies attribute the high nucleophilicity and reactivity exhibited by these complexes to the Cu-boryl  $\sigma$ -bonding electrons. Computational work by Carbó and Fernández in 2012,<sup>33</sup> and Sheong and Lin in 2021,<sup>34</sup> revealed that the

contribution of B-based orbitals to the M–BR<sub>2</sub> bond is remarkably high (*ca.* 70%, with a p/s ratio  $\approx$  0.9)<sup>35</sup> in the case of mononuclear Cu-boryl species, whereas in other examples like Au- or Pd-boryl compounds, the contribution of B is 56% and 53%, respectively. Therefore, the Cu–B bond tends to be substantially more polarised towards the B atom. Natural Localised Molecular Orbital (NLMO)<sup>30</sup> analysis of **2** and **4** reveals  $\sigma$ -donation of the boryl ligand to the empty 4s orbitals on the Cu atoms (as previously observed for  $[(\text{DPFN})\text{Cu}_2]^{n+}$  complexes)<sup>9</sup> with negligible back-donation from the metals to boron. This 3c–2e bonding scenario is illustrated in Fig. 3. Surprisingly, the contributions of Cu- and B-based orbitals to the bonding are almost identical to those previously reported for reactive, mononuclear Cu-boryl complexes (*ca.* 30% Cu and 70% B), with a considerably smaller p/s ratio on B in the case of **4**.

These data indicate that the extreme stability and chemical inertness of boryls **2** and **4** in polar solvents does not stem from peculiar geometrical or electronic factors, since these properties are similar to those of much less stable analogues. Rather, the rigid framework imposed by the dinucleating 1,8-naphthyridine-based ligand seems to be responsible for the observed stability, by precluding dynamic behaviour in solution that might produce an equilibrium concentration of more reactive, monomeric species.<sup>4,5,7</sup> This hypothesis finds additional support from a recent study by Ito and coworkers, which concludes that a borylcopper(i) dimer is the dormant species in a catalytic asymmetric borylation process, whereas the mononuclear Cu-boryl complex is the active form of the catalyst.<sup>36</sup>

## Conclusions

The  $\mu\text{-O}^t\text{Bu}$  complex **3** provides a convenient synthetic pathway to a new type of dicopper boryl derivative, by a method that may provide more general access to complexes featuring reactive fragments stabilised by a dicopper core. In reactions of **3** with diboranes  $\text{B}_2\text{pin}_2$  and  $\text{B}_2\text{cat}_2$ , mild conditions lead to the dinuclear boryl compounds **2** and **4** by cleavage of a B–B bond. Striking properties of the new boryl complexes reflect a high stability for the  $\text{Cu}_2\text{B}$  cores. This stability appears to relate to strong, delocalised bonding across the bridging interaction supported by an effective, rigid dinucleating ligand that suppresses access to reactive monocopper boryl units. These results are therefore consistent with earlier observations concerning the ability of the  $[(\text{DPFN})\text{Cu}_2]^{n+}$  platform to stabilise reactive intermediates. Notably, the stability/reactivity balance in  $\mu$ -boryl dicopper complexes should be tunable by way of modifications to the dinucleating ligand. For example, side



**Fig. 3** Natural Localised Molecular Orbitals (NLMO) of complexes **2** and **4**, depicting  $\sigma$ -donation of the boryl fragment to the dicopper core.



arms that enforce a lower coordination number at the metal could lead to higher reactivity, and such possibilities are currently being explored in this laboratory.

## Data availability

Crystallographic data for complexes 2, 3 and 4 have been deposited at the CCDC under 2130621–2130623

## Author contributions

P. R. and T. D. T. conceived of the idea. P. R. and M. S. performed the experiments and analysed the data for synthesis, reactivity and characterization. The crystallographic data were analysed by R. C. H. (complexes 2, 3 and 4) and S. J. T. (complex 2). P. R. performed the DFT studies. P. R., M. S., R. C. H. and T. D. T. wrote and edited the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

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