

Cite this: *Chem. Sci.*, 2025, 16, 14518

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Driving diverse bond functionalisation with *N*-heterocyclic silylene-coinage metal–aryl complexes†

Moushaki Ghosh,^a Kumar Gaurav,^a Prakash Panwaria,^{†a} Rishukumar Panday,^a Srinu Tothadi^b and Shabana Khan^{†a}

Anchoring homolytic and heterolytic bond functionalisation at low-coordinate coinage metal centres is important due to their potential use as active catalysts in organic transformations. In the realm of carbene-stabilised coinage metal chemistry, heteroatom functionalised coinage metal precursors synthesised from such bond activations have long been explored. Interestingly, *N*-heterocyclic silylene, being an equally potent neutral donor ligand, has not been used for the same. Of note, carbene-stabilised heteroatom functionalised coinage metal precursors are vastly developed with copper centres only, while silver has been underexplored. This work reports the isolation of a variety of [PhC{N(^tBu)}₂SiN(SiMe₃)₂] (1) coordinated aryl-copper(i) and aryl-silver(i) complexes (2–8). We have also examined the reactivity pattern of organo-copper with differently substituted silylenes (9–11). These complexes were then utilised to cleave various homolytic and heterolytic bonds to access silylene-coordinated heteroatom functionalised coinage metal complexes (12–24). We have shown the reaction of reactive aryl-coinage metal precursors towards homolytic bonds, having B–B and Se–Se bonds, which led to the formation of an NHSi-supported dimeric μ -boryl bound Cu(i) complex (12) and a new class of unprecedented NHSi-supported coinage metal-selenogenolates (14–16). These aryl-coinage metal precursors also smoothly afforded several elusive NHSi-copper and silver amides (17–22) via N–H bond cleavage. A heterolytic cleavage of the P–Si bond resulted in the formation of NHSi stabilised copper and silver phosphide complexes (23 and 24), among which the latter is the first precedent of the dimeric Ag-phosphide complex. Lastly, we have utilised NHSi $\rightarrow$ copper–aryl complexes as aryl transfer reagents in C–C coupling reactions, which led to the formation of products in excellent yields with a high TON. The analogous silver complex was employed in the three-component α -aminonitrile synthesis efficiently. Our report establishes NHSi coordinated aryl copper and silver complexes as a perfect and robust platform for accessing a diverse array of reactive coinage metal precursors that were hitherto unknown.

Received 3rd February 2025
Accepted 28th June 2025

DOI: 10.1039/d5sc00879d

rsc.li/chemical-science

1 Introduction

The *N*-heterocyclic silylenes (NHSis), known as heavier congeners of *N*-heterocyclic carbenes (NHCs), are now considered one of the formidable classes of neutral donor ligand systems.¹ Over the years, several NHSis have been isolated and utilised in stabilising reactive main group species,² coordinating transition metals for homogeneous catalysis,³ and activating small molecules.^{1c,4} Groundbreaking studies by various pioneering scientists have empowered NHSis as one of the promising ligand motifs for stabilising numerous elusive and reactive species.^{1e,5} Despite their superior catalytic activities in several organic transformations, the NHSi stabilised coinage metal complexes are comparatively less studied than the NHC ones.⁶ However, recent years have witnessed a growth in isolating differently functionalised NHSi-coordinated coinage metal complexes and exploring their catalytic activities (Chart 1a).⁷ Given our

^aDepartment of Chemistry, Indian Institute of Science Education and Research Pune, Dr Homi Bhabha Road, Pashan, Pune-411008, India. E-mail: shabana@iiserpune.ac.in

^bAnalytical and Environmental Sciences Division and Centralized Instrumentation Facility, CSIR-Central Salt and Marine Chemicals Research Institute, Gijubhai Badheka Marg, Bhavnagar-364002, India

† Electronic supplementary information (ESI) available. CCDC 2304275 (2), 2304278 (3), 2419144 (4), 2450951 (5), 2304292 (6), 2419145 (7), 2419147 (8), 2304280 (9), 2450954 (10), 2450956 [PhC{N(^tBu)}₂Si{N(Dipp)SiMe₃}], 2450959 (11), 2450960 (12), 2304289 (13), 2304286 (14), 2325641 (15), 2419148 (16), 2450961 (17), 2304281 (18), 2304284 (19), 2304285 (20), 2304283 (21), 2419162 (22), 2419149 (23) and 2419163 (24). For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d5sc00879d>

‡ Both authors contributed equally.

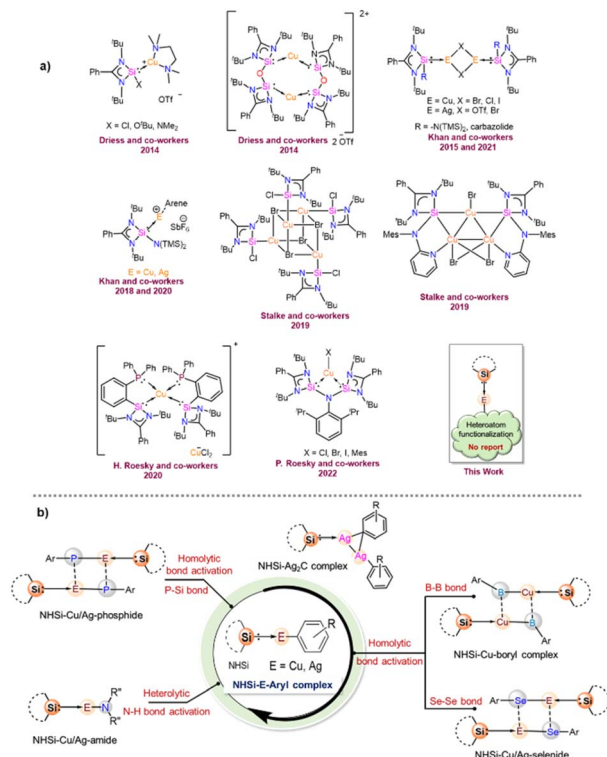


Chart 1 (a) Reported benzamidinato silylene coordinated Cu(I) complexes to date. No NHSE-coordinated Cu(I)/Ag(I)-aryl complex is reported;^{7e,g,i,j,11} (b) overview of the present work.

particular interest in this field,^{7a-g} we continue to be intrigued by the future possibilities of NHSE-based coinage metal complexes as useful synthons beyond the coordination of coinage metal halides to encompass a broader range of synthetic possibilities. The coordination of NHSEs with coinage metals having heteroatom functionalisation remains uncharted territory until now. Many NHC-based copper [Cu(I)] synthons have been isolated and studied for bond activation and homogeneous catalysis.⁸ Interestingly, the analogous situation with silver [Ag(I)] is far more underexplored. This is primarily because of the higher thermal decomposition of Ag(I) complexes under ambient conditions. Also, carbene-Ag(I) complexes are chosen precursors for transmetallation reactions with other transition metals, such as Cu, Au, Fe, Ni, and Co.⁹ The bond strength between carbenes and transition metals is greater than that of carbene-Ag(I) complexes, facilitating a smoother transmetallation process. In this context, aryl-coinage metal precursors(I) are a very reactive class of complexes,^{8c,10} which can be explored in bond activations due to the lability of E-aryl bonds (E = Cu and Ag). Also, NHC-based Cu(I) complexes with anionic C-donor atoms have been explored for their catalytic application and functionalisation reactions, such as the formation of Cu-X bonds (X = -SR, -PR₂, -OR, and -NR₂), intramolecular hydroalkoxylation of alkynes, the CuAAC reaction, and defluoroborylation of fluoroalkenes.^{8p} However, such utilisation of carbene-Ag(I)-aryl complexes is missing in the literature. Very recently, the NHC-coordinated Ag(I)-mesityl complex has been used as a precursor for crafting graphitic carbon nitride (g-

C₃N₄),¹² paving the way for their further exploration in organometallic and material chemistry. All these findings inspired us to isolate the NHSE-based Cu(I) and Ag(I)-aryl complexes (2–8), creating a platform for the further functionalisation of the labile Cu/Ag-aryl bonds (Chart 1b). In this work, we have demonstrated the coordination behaviour of aryl-coinage metal precursors exclusively with benzamidinato silylene, [PhC{N(^tBu)}₂SiN(SiMe₃)₂] (1), and then showcased the reactivities of these precursors towards various homolytic and heterolytic bond cleavage reactions (12–24) (Chart 1b). We have also briefly checked the reactivity pattern of different NHSEs (9–11) with organocopper complexes to establish the breadth of the methodology. While the reactions of NHSE with Cu(I)-aryl precursors afforded di-coordinated NHSE → Cu-aryl complexes (2–5) smoothly, the reaction of Ag(I)-mesityl afforded a unique Ag₂C ring with argentophilic interaction (6). The stability of the complex is interesting because of the high level of angle strains and torsional strains of the thermodynamically unstable three-membered Ag₂C ring. Interestingly, there is no precedent of such a complex with NHSEs to date. However, the reactions with other Ag(I)-aryl precursors yielded nearly linear coordinated complexes 7 and 8 due to the enhanced steric effects of aryl groups. The isolation of these complexes created a perfect platform for bond functionalisation. Inspired by Sadighi's isolation of NHC-supported Cu(I)-boryl complexes [(IPr)Cu(Bpin), (ICy)Cu(Bpin)], catalytically active for CO₂ reduction and a few isolated carbene-stabilised copper-boryl complexes,¹³ we isolated NHSE-Cu-boryl complexes *via* diboron bond cleavage, which yielded complex 12 with a dimeric μ -boryl framework. Additionally, given the scarcity of NHC/CAAC-supported chalcogenides¹⁴ and the absence of NHSE-coordinated coinage metal chalcogenolates, we also reacted NHSE → Cu(I)/Ag(I)-aryl complexes with different diselenides, leading to the isolation of NHSE-Cu/Ag selenogenolate (14–16). Very recently, we have shown that the NHSE → Cu-amides can demonstrate fascinating luminescence properties.¹⁵ Along with this, N-H bond cleavage is also an important part of cross-coupling reactions; hence, we attempted syntheses of Cu/Ag-amides (17–22).¹⁶ Lastly, employing the σ -bond metathesis reaction of Cu/Ag-C and P-Si bonds, we were able to isolate rarely occurring Cu(I) and Ag(I) phosphide complexes (23–24). Meanwhile, NHC-Cu-phosphides have been reported only in limited instances.⁸ⁱ To the best of our knowledge, the precedence of low-coordinate Ag-phosphide is still elusive. Overall, this vast piece of work intends to highlight the synthetic versatility and facile bond activation potential of NHSE-Cu/Ag-aryl complexes towards various energy demanding bonds such as B-B (~462.86 kJ mol⁻¹), Se-Se (~239.48 kJ mol⁻¹), N-H (~387.41 kJ mol⁻¹), and P-Si (~263.58 kJ mol⁻¹) (Table S4, see the ESI for the details[†]). To understand the bonding nature of all the newly synthesised complexes, density functional theory (DFT) calculations were performed using the Gaussian 09 program package (see the ESI for details[†]), and the description of theoretical input is provided at the relevant places.

To establish the significance of such organo-coinage metal precursors in catalytic reactions, we performed a few brief proof-of-concept catalytic model reactions, such as C-C cross-



coupling reactions and a three-component α -aminonitrile synthesis, which afforded the desired compounds in excellent yields (*vide infra*).

2 Results and discussion

2.1 Synthesis of NHSi-E-aryl (E = Cu and Ag) complexes

2.1.1 Synthesis of NHSi-based Cu(I)-aryl complexes (2–5).

We started our investigation by reacting bis-(trimethylsilyl) amide substituted benzamidinato silylene, $[\text{PhC}\{\text{N}(\text{tBu})\}_2\text{SiN}(\text{SiMe}_3)_2]$ (**1**), with different aryl copper precursors¹⁷ in toluene at room temperature, which furnished complexes 2–5 (Scheme 1) in good yield.

The $^{29}\text{Si}\{^1\text{H}\}$ spectra display a broad peak at δ 6.5 and δ 6.6 ppm for the Si(II) $\rightarrow$ Cu centre for complexes 2 and 3–5, respectively. The molecular structures of 2–5 demonstrate that the Si(II) centres adopt a distorted tetrahedral geometry (Fig. 1). The Si–Cu bond lengths are 2.2590(8) Å (complex 2), 2.2407(8) Å (complex 3), 2.246(2) Å (complex 4), and 2.242(1) Å (complex 5), which are shorter than the previously reported Si–Cu bond length in the carbazole-substituted benzamidinato silylene stabilised dimeric Cu_2Br_2 complex (2.203(2) and 2.212(2) Å),^{7c} and longer than that in $[\text{PhC}\{\text{N}(\text{tBu})\}_2\text{SiN}(\text{SiMe}_3)_2]$ stabilised Cu_2Br_2 (2.222(2) Å). But they match closely with that in the dimeric Cu_2I_2 complex of $[\text{PhC}\{\text{N}(\text{tBu})\}_2\text{SiN}(\text{SiMe}_3)_2]$ (2.243(3) and 2.250(3) Å).^{6a} They are significantly longer than the predicted bond length for the NHSi $\rightarrow$ Cu(I) complex (2.061 Å) by Frenking and co-workers.¹⁸ It was observed that the Si–Cu bond lengths decrease with increasing steric bulk around the aryl group, following the trend $2 > 4 > 5 > 3$. They feature almost linear geometry around the Cu(I) centre featuring a bond angle Si1–Cu1–C1 of 173.28(6)° (complex 2), 170.72(7)° (complex 3), 172.7(1)° (complex 4) and 172.5(1)° (complex 5) which is in accordance with the previously reported IMes (1,3-bis-(2,4,6-trimethylphenyl)imidazole-2-ylidene) stabilised Cu-mesityl complex [173.53(9)°]¹⁹ but the Si–Cu–C angles of complexes 2–5,



Fig. 1 Molecular structures of 2 (a), 3 (b), 4 (c), and 5 (d). The bond distance (Å) and bond angle (°) are discussed in the ESI.†

are shorter than that of the carbene coordinated copper-mesityl complexes (in the range of [174.53(6)–178.06(12)°]).^{19,20} To understand the implications of these bonding patterns on the electronic structure, the frontier molecular orbital (FMO) analysis was carried out (Fig. S111†). This reveals that the highest occupied molecular orbital (HOMO) is primarily localised over the Cu d orbital and the π orbital of the mesityl group. Conversely, the electron density in HOMO–1 predominantly resides over the Cu(I)– C_{ipso} bond, with a marginal distribution extending over the Si(II) $\rightarrow$ Cu bond and the mesityl group. Notably, HOMO–3 is majorly localised over the Si(II) $\rightarrow$ Cu bond, exhibiting a minor dispersion over the Cu(I)– C_{ipso} bond and the amidinate fragment. The HOMOs of complexes 3 and 4 (see the ESI for computational details†) show a predominant contribution from the Cu(I)– C_{ipso} bond and a minor involvement from the Si(II) $\rightarrow$ Cu bond and the aryl group. It is important to note that the electron density on C_{ipso} is more in complex 3 (35%) and 4 (35%) in the HOMO than in complex 2 (27%), indicating the stronger electron-donating effect of Dipp (2,6-diisopropylphenyl) and Trip(2,4,6-triisopropylphenyl) over the Mes (2,4,6-trimethylphenyl) group. However, the lowest unoccupied molecular orbital (LUMO) for all complexes is predominantly localised over the benzamidinato fragment.

2.1.2 Synthesis of NHSi-based Ag(I)-aryl complexes (6–8).

Unlike organocopper(I) complexes, the chemistry of organo-silver(I) complexes has not been explored much, as we already discussed. There are a few examples of ligated silver(I)-mesityl moieties in the literature.²¹ Upon reaction of compound **1** with mesityl silver^{17c} in toluene at room temperature, an interesting μ -mesityl bound dimeric silver(I) complex (**6**) was obtained (Scheme 1). The $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum of **6** indicated the formation of the desired complex by showing two doublets at δ 14.7 and 11.5 ppm due to the presence of two NMR active nuclei of Ag (109 and 107). Complex **6** features a 3c–2e bond involving the C_{ipso} atom of the mesityl group with two silver



Scheme 1 The reaction of compound **1** with various aryl Cu(I) and Ag(I) compounds.



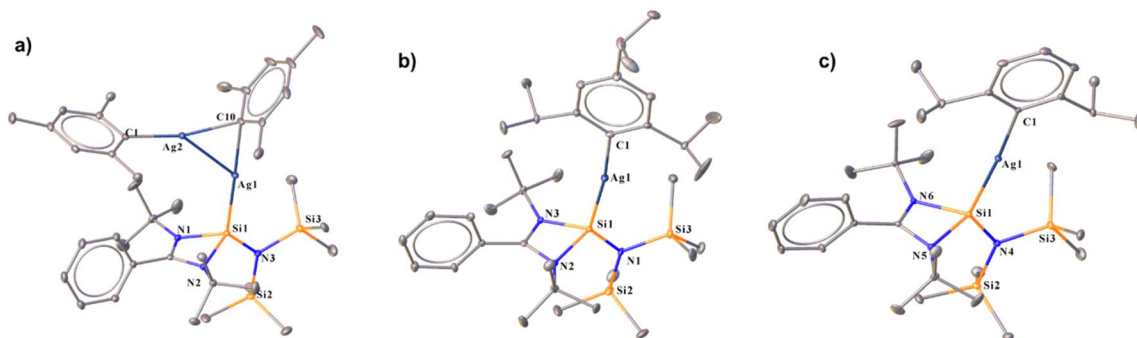


Fig. 2 Molecular structures of **6** (a), **7** (b) and **8** (c). The bond distance (Å) and bond angle (°) are discussed in the ESI.†

atoms (Fig. 2), resulting in close proximity of Ag1–Ag2 (2.7453(5) Å). The Ag1–Ag2 bond length of **6** is much shorter than the sum of the van der Waals radii (3.440 Å).²² The aryl group bound to coinage metal complexes often shows μ -bonding in the aggregated form to form a polynuclear complex (Fig. 2).²³ The Si1–Ag1 bond length in **6** is 2.3900(9) Å, which is similar to that of the carbazole-substituted benzamidinato silylene coordinated tetrameric Ag₄I₄ complex.^{7c} Kays and co-workers reported an *m*-terphenyl group-bound dimeric Ag₂ core with a Ag $\cdots$ Ag distance of 2.6706(3) Å, which is shorter than that of complex **6**.²⁴ While the Ag1–C1 bond in **6** is shorter (2.106 Å), that of the μ -bridged mesityl-bound Ag atoms are longer (Ag1–C10 2.215(3) and Ag2–C10 2.211(3) Å) and form almost symmetrical bonds with the C10 atom. This is the first NHC-supported three-membered Ag₂C ring system. Sadighi and co-workers isolated a triangular [Ag₂H]⁺ core stabilised by 1,3-bis(2,6-diisopropylphenyl)imidazolin-2-ylidene (SIDipp),²⁵ but such a molecular framework with NHCs is not known. Furthermore, we also tried to validate whether such a heteroleptic complex is present in the solution; thus, we performed diffusion ordered NMR spectroscopy (DOSY). In this method, molecules are differentiated according to their diffusion coefficient (*D*), which correlates with their hydrodynamic radius.²⁶ We found the hydrodynamic radius of complex **6** to be 6.24 Å in toluene-*d*₈, which suggests that it stays as a monomeric complex in solution. Our attempt to crystallise them in the monomeric form in other solvents like THF and DCM was unsuccessful. The FMO analysis shows that the HOMO of complex **6** shows the contribution over the delocalised mesityl–Ag–C_{ipso} and d orbital of the Ag atom bonded to NHC, whereas HOMO–1 shows the contribution over the localised mesityl group and d orbital of the Ag atom (Fig. S112†). HOMO–2 is localised over the mesityl group, whereas HOMO–3 is delocalised over the Si(II) $\rightarrow$ Ag₂C ring connected to the μ -bridged mesityl group (see the ESI for computational details†). Furthermore, natural bond orbital (NBO) analysis was performed to understand the presence of metallophilic interaction in **6**, which confirms the presence of argentophilic interaction (Fig. S120†). The second-order perturbative energy [*E*⁽²⁾] was hence calculated to quantify the bonding–antibonding interactions. The *E*⁽²⁾ value for the delocalisation of electron density from the Ag(i) centre to the σ^* orbital of the Ag(i)–Si bond was 2.10 kcal mol^{–1}. However, the *E*⁽²⁾ value for the Ag(i) centre attached to the NHC to the σ^* orbital of the Ag(i)–C_{ipso}

bond was found to be 1.71 kcal mol^{–1}. The Wiberg bond index (WBI) of the Ag $\cdots$ Ag bond was found to be 0.072, which is in the range (0.022–0.094) of reported literature on argentophilic interaction.²⁷ Interestingly, treatment of AgTrip and AgDipp with **1** yielded NHC-bound linear Ag(i)–aryl complexes (**7** and **8**) (Scheme 1). The ²⁹Si{¹H} NMR spectra for complexes **7** and **8** show two doublets appearing at δ 14.3 and 10.9; 14.1 and 10.8 ppm, respectively, due to the coupling of ¹⁰⁹Ag and ¹⁰⁷Ag with the ²⁹Si centre. The Si–Ag bond lengths for complexes **7** and **8** are 2.400(1) and 2.387(6) Å, respectively, whereas the Si–Ag–C bond angles for complex **7** [172.21(6)°] deviated more from linearity than that of complex **8** [173.44(5)°] due to more steric bulk for the former than the latter, as also observed in complexes **3–4** (Fig. 1). The HOMO of both these complexes resides on the NHC $\rightarrow$ Ag–C_{aryl} fragment (Fig. S113†). The rest of the MOs for complexes **7** and **8** are similar to those of complexes **3** and **4**.

2.1.3 Reaction of organocopper towards differently functionalised benzamidinato silylene. To monitor the reactivity pattern of organo-copper reagents with differently substituted NHCs, we further probed the reactivity pattern of [PhC{N(^{*t*}Bu)}₂SiCl], [PhC{N(^{*t*}Bu)}₂SiO^{*t*}Bu], and newly synthesised [PhC{N(^{*t*}Bu)}₂Si{N(Dipp)SiMe₃}] with CuMes (Scheme 2).²⁸ For that, we have treated [PhC{N(^{*t*}Bu)}₂SiCl], [PhC{N(^{*t*}Bu)}₂SiO^{*t*}Bu], and [PhC{N(^{*t*}Bu)}₂Si{N(Dipp)SiMe₃}] with mesityl copper in a 1 : 1 molar ratio, which furnished complexes **9**, **10**, and **11**,



Scheme 2 Syntheses of complexes **9–11**.



respectively.²⁹ We selected these variations based on their differences in electronic and steric parameters, anticipating diverse reactivities. The formation of **9** proceeded through σ -bond metathesis at the labile Si–Cl moiety, and the mesityl group migrated to the Si(II) centre to form a dimeric Cu₂Cl₂ core (Fig. S136†). The peak at δ 32.7 ppm in the ²⁹Si{¹H} NMR spectrum is more deshielded than the previously reported dimeric copper(I) chloride of compound **1** (δ 4.79 ppm),^{14f} which might be due to the presence of the σ -donating mesityl group at the silicon(II) centre. **9** crystallises in the monoclinic *P*2₁/*n* space group (see Fig. S136†) and is isostructural to the dimeric Cu₂Cl₂ complex of **1**.^{14f}

The silicon(II) centre of **9** attains a distorted tetrahedral geometry with a Si–Cu bond length of 2.213(1) Å.^{14f} On the other hand, [PhC{N(^tBu)}₂SiO^tBu] furnishes μ -dimeric mesityl-bridged Cu(I) complex, **10** (Scheme 2). The ²⁹Si{¹H} NMR spectrum shows a broad peak at δ 3.8 ppm, deshielded from [PhC{N(^tBu)}₂SiO^tBu] (δ –5.0 ppm),^{28b} but comparable to the dimeric Cu₂Cl₂ complex of **1** as mentioned above. Complex **10** crystallises in the triclinic *P* $\bar{1}$ space group with a Si–Cu bond length of 2.249(1) Å, and the Cu–C bond lengths are 2.170(4) and 2.132(4) Å, suggesting the μ -bridging fashion (Fig. 3). The tri-coordinated Cu(I) adopts a Si1–Cu1–C1 bond angle of 119.8(1)°. The Cu...Cu distance in **10** is 2.335(6) Å, which is shorter than the Cu...Cu distance (2.46 Å) in mesityl copper.¹⁰ To broaden the scope of the reactivity pattern, we performed a reaction of mesityl copper with newly synthesised [PhC{N(^tBu)}₂Si{N(Dipp)SiMe₃}] and obtained complex **11**, similar to complex **2**. Formation of complex **11** was confirmed by ²⁹Si{¹H} NMR, which shows a broad peak at δ 11.8 ppm, which is further deshielded than [PhC{N(^tBu)}₂Si{N(Dipp)SiMe₃}] (δ 2.98 ppm). The Si–Cu distance in complex **11** is observed to be 2.227(2) Å, shorter than that in complex **2** {2.2590(8) Å}, and the Cu–C distance is 1.934(3) Å, with a bond angle of 179.68(9)° around Si–Cu–C centres (Fig. 3). The abovementioned reactivity of different silylenes clearly outlines the influence of steric and electronic properties on the structure of the final product.

2.2 Reactivities of NHCs-based E-aryl complexes (**2** and **7**) towards various bond cleavage reactions

Based on various structural motifs obtained (**2**–**11**) through the coordination ability of differently functionalised silylene with organo-coinage metal precursors, complexes **2** and **7** were



Fig. 3 Molecular structures of **10** (a) and **11** (b). The bond distance (Å) and bond angle (°) are discussed in the ESI.†



Scheme 3 Reaction of complex **2** with a diboron compound (B₂cat₂).

chosen for further exploration in a range of bond activation reactions.

2.2.1 Synthesis of the NHCs-based Cu(I)–boryl complex (12**) via B–B bond cleavage.** Various NHC and CAAC-coordinated low-valent boron species have found application in optoelectronics.³⁰ The coinage metal complexes of diborene and boron–boron triple-bonded systems also showcased interesting photophysical behaviour.³¹ Thus, we were interested in probing the reaction of **2** with the diboron (B–B bond) reagents³² in the quest of homolytic bond cleavage.³³

Thus, we performed a reaction of less sterically bulky bis(catecholato)diboron (B₂cat₂) with complex **2** (Scheme 3) in toluene. In this case, we could obtain dimeric [PhC{N(^tBu)}₂SiN(SiMe₃)₂Cu₂B₂Cat₂] (**12**) and Mes–Bcat (**13**) as a mixture of the products. The isolation of complex **12** also indicated the higher reactivity of sterically less hindered B₂cat₂, which was previously pointed out.³⁴ The molecular structure of complex **12** (Fig. 4) established the μ -coordination mode of the boryl linkage to Cu(I) coordinated with NHCs unambiguously. However, the poor quality of the crystals prevented us from discussing the metrical parameters. Analogous NHC stabilised μ -boryl Cu(I) complexes were reported by Sadhigi, Kleeberg, and co-workers, which show a μ -boryl Cu(I) core (Chart 2).³⁵ We could not get a clean ¹H NMR spectrum of **12** as it was isolated as a mixture with **13** (see the ESI†). However, ²⁹Si{¹H} NMR spectra revealed two sharp singlets at δ 6.3 and 7.1 ppm, corresponding to the –SiMe₃ moiety, and a singlet at δ 3.7 ppm, presumably indicating the Si(II)–Cu(I) bond. Although there are a few Cu– μ -boryl



Fig. 4 Molecular structure of **12**.



Chart 2 Previously reported examples of NHC-supported μ -boryl Cu(I) complexes.^{34,35}

complexes reported in the literature, the understanding of the associated FMOs is largely missing, apart from the work by Tilley and co-workers.^{35e} Drawing inspiration from this, we analysed its electronic structure, which reveals that the HOMO of the dimeric Cu(I)- μ^2 -Boryl complex (**12**) (Fig. S114†) is localised over the Cu₂B₂ ring. The orbital composition of the HOMO shows that the localisation of electron densities at Cu centres is 19 and 13%, while at B centres, it is 30 and 22%, respectively. This 3c–2e bonding situation was further confirmed by the natural localised molecular orbital (NLMO) analysis, showing that the contribution of each Cu centre and B orbital in the Cu–B–Cu bond is 13, 6, and 73%, respectively. HOMO–1 also has a contribution over the Cu₂B₂ ring along with the Si(II) $\rightarrow$ Cu bond, whereas the electron density in the HOMO–2 is predominantly localised over the Si(II) $\rightarrow$ Cu bond, along with the contribution from the catechol-boryl moiety.

The reaction pathway shows that complex **2** forms an intermediate (**Int**) involving non-covalent interaction between boron atoms of B₂pin₂ and the mesityl ring (Fig. 5), proceeding *via* an

intermediate to give complex **12** (detailed analyses provided in the ESI†). Interestingly, the preparation of aryl boronate esters either involves Suzuki–Miyaura cross-coupling for C–B bond formation³⁶ or base-mediated ArB(OH)₂ formation.³⁷ Thus, the formation of **13** as a byproduct of this reaction in the crystalline form gives easy access to mesityl boronic ester (see the ESI†). It is important to note that a similar reaction with complex **7** failed to yield the NHC $\rightarrow$ Ag–boryl complex, likely due to poor thermal stability and rapid dissociation of the Ag–B bond. Although we attempted to monitor the fate of the product using ²⁹Si{¹H} NMR spectroscopy, we were able to detect a Si(II)–bound Ag(I) species. However, the exact nature of the species formed alongside Trip–Beat remains unclear at this stage.

We also tried the reaction of complex **2** with other diboron agents discussed in the ESI.†

2.2.2 Synthesis of NHC-based Cu(I) and Ag(I) selenogenolate complexes (14–16) *via* Se–Se bond cleavage. To access NHC-supported coinage metal chalcogenolate complexes, we treated complex **2** with diphenyl diselenide (Ph₂Se₂) and sterically demanding bis(2,4,6-trimethylphenyl) diselenide (Mes₂Se₂) in



Scheme 4 The reaction of complexes **2** and **7** with diaryl diselenide (Ar'₂Se₂).



Fig. 5 Relative energy (kcal mol^{–1}) (not drawn to scale) reaction profile for the formation of complex **12** from complex **2** with B₂cat₂.

toluene at room temperature (Scheme 4). This led to the formation of immediate white precipitates, which were characterised by ^1H and $^{77}\text{Se}\{^1\text{H}\}$ NMR spectra as MesSePh^{38} and Mes_2Se ,³⁹ respectively (see the ESI†). The filtrate part of the reaction mixture afforded suitable single crystals of **14** and **15**. The reaction of complex **7** with Mes_2Se_2 also afforded NHSi stabilised $\text{Ag}(\text{I})$ mesitylselenenolate (**16**) (Scheme 4). The $^{29}\text{Si}\{^1\text{H}\}$ NMR spectra display two sharp singlets for the $-\text{SiMe}_3$ groups at δ 4.4 and 5.9 ppm for **14**, and δ 5.3 and 6.7 ppm for **15**, respectively. A broad peak for $\text{Si}(\text{II})-\text{Cu}$ was observed in $^{29}\text{Si}\{^1\text{H}\}$ NMR spectra at δ 5.9 and 3.2 ppm for **14** and **15**, respectively. Complex **16** shows a doublet at δ 4.12–4.47 and 9.13–9.49 ppm, similar to complex **7**, discussed earlier. Complexes **14**–**16** display a dimeric E_2Se_2 ($\text{E} = \text{Cu}$ and Ag) core with the $\text{Si}-\text{E}$ bond lengths of 2.248(3), 2.2631(6), and 2.401(1) Å, for **14**, **15**, and **16**, respectively. The $\text{E}-\text{Se}$ distances in **14**–**16** are 2.456(1), 2.5076(6), and 2.6245(7) Å, respectively (Fig. 6). They are marginally longer than those in the $\text{Ph}_3\text{PCu}(\mu-\text{SePh})_2\text{Cu}(\text{PPh}_3)_2 \cdot \text{CH}_3\text{CN}$ complex {2.406(1) Å} by Oliver and co-workers and $[\text{Ag}_4(\mu-\text{SePh})_4(\text{Pr}_2\text{-bimy})_4]$ {2.649(1), 2.748(1) Å} by Corrigan and co-workers.^{14d,40} The $\text{Cu}\cdots\text{Cu}$ distances in **14** and **15** are 2.938(2) and 3.4398(6) Å, respectively, which are longer than the $\text{Cu}\cdots\text{Cu}$ distance in the $\text{Ph}_3\text{PCu}(\mu-\text{SePh})_2\text{Cu}(\text{PPh}_3)_2 \cdot \text{CH}_3\text{CN}$ complex {2.738(1) Å},⁴¹ but shorter than the $\text{Cu}\cdots\text{Cu}$ distance [3.1378(9) and 3.255(1) Å] in $[\text{PhC}(\text{N}^t\text{Bu})_2\text{N}(\text{TMS})_2\text{SiCu}_2\text{X}_2]$ ($\text{X} = \text{Cl}$ and Br).^{14f} The $\text{Ag}\cdots\text{Ag}$ distance is 3.8110(7) Å in complex **16**, significantly longer than the argentophilic interaction range and complex **6**.²² Noteworthy to mention that there is no report of such dimeric carbene-coordinated $\text{Ag}(\text{I})$ selenenolates in the literature.⁴⁰ In a similar line, we also tried similar reactions with diphenyl sulfide (Ph_2S_2) and diphenyl telluride (Ph_2Te_2) for consecutive bond activation, but could not obtain the desired compounds. FMO analysis shows that the HOMO of complexes **14** and **15** (Fig. S115†) is prominently concentrated over the dimeric Cu_2Se_2 ring, demonstrating substantial contributions from the $\text{Si}(\text{II})$ centres and the phenyl rings, whereas HOMO–3 is localised over the $\text{Si}(\text{II}) \rightarrow \text{Cu}$ bond for complex **15**. As we observed the presence of cuprophilic interaction in the molecular structure of **14**, we went on to examine its $E^{(2)}$ value, which shows 0.62 kcal mol^{−1} strength for $\text{Cu}\cdots\text{Cu}$ interaction (Fig. S121†), less than the reported strength.⁴² It thus denotes the weaker nature of cuprophilic interaction. We have also calculated the

change in the Gibbs free energy (ΔG) of **14** in the temperature range of 0–450 K to investigate the thermal stability.

It was found that **14** is stable in the dimeric form up to the temperatures of 380 and 420 K in the solution and gas phase, respectively (see the ESI for computational details†). The **Int** observed in the reaction pathway for the formation of complex **14** (Fig. 7) is stabilized by the interaction between Cu and Se centers, $\text{C}-\text{H}\cdots\text{Se}$, $\text{C}-\text{H}\cdots\pi$, and tetrel bonding. These non-covalent interactions favour the formation of the $\text{Cu}-\text{Se}$ bond and the breaking of the $\text{Cu}-\text{C}_{\text{ipso}}$ bond, which is visualised in the **TS**, and confirmed by the relevant geometrical parameters and NBO analysis (see the ESI for details†).

2.2.3 Synthesis of NHSi-based $\text{Cu}(\text{I})/\text{Ag}(\text{I})$ amide complexes (17–22) via N–H bond cleavage. There are prodigious examples of NHC-coordinated copper alkyl complexes that work as intermediates to different $\text{X}-\text{H}$ ($\text{X} = \text{N}$, O , and C) bond functionalised products, which even led to the isolation of the first NHC-based copper amido and anilido complexes.^{8p,43} Recently, carbene metal amide (CMA) complexes have attracted special attention due to their promising photophysical properties.^{16,44} Motivated by the fact that NHSi-metal amide complexes had not been isolated to date until very recently by us,¹⁵ we delved into the reactivity of the NHSi-ligated $\text{Cu}(\text{I})/\text{Ag}(\text{I})$ aryl complex towards N–H bond cleavage, which facilitated a straightforward route for NHSi-coordinated metal amide complexes (Scheme 5a). The stoichiometric reaction of complexes **2** and **7**, with N–H containing compounds, afforded NHSi-metal amide complexes, which are stable for days under inert conditions at room temperature and months at 0 °C.

Complex **17**, formed by reacting **2** with pyrrole, shows a broad peak at δ 7.8 ppm in the $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum, and its structure was confirmed by single-crystal X-ray diffraction. The $\text{Si1}-\text{Cu1}$ bond length for complex **17** was found to be 2.231(3) Å, and the $\text{Cu}-\text{N}$ bond length was 1.878(7) Å. Complex **17** features an almost linear $\text{Si}-\text{Cu}-\text{N}$ bond angle of 172.2(5)° (Fig. 8). Similarly, reacting **2** with 2-mesityl-1H-pyrrole yielded complex **18**, displaying a $^{29}\text{Si}\{^1\text{H}\}$ peak at δ 7.4 ppm, deshielded relative to **2** due to the π -electron-donating mesityl group. Complex **18** has a $\text{Si1}-\text{Cu1}$ bond length of 2.203(1) Å and a $\text{Cu1}-\text{N1}$ bond length of 1.874(3) Å (Fig. 8), with a nearly linear $\text{Si1}-\text{Cu1}-\text{N1}$ angle of 177.5(1)°. The $\text{Si}-\text{Cu}$ bond distance is comparable to that of the heteroleptic $[\text{Cu}(\text{tmeda})(\text{PhC}(\text{N}^t\text{Bu})_2\text{Si}\{\text{O}^t\text{Bu}\})][\text{OTf}]$ ($\text{tmeda} =$

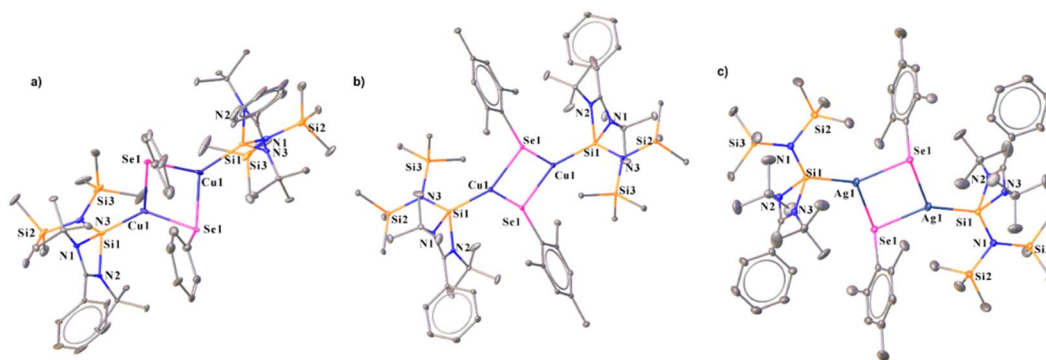


Fig. 6 Molecular structures of **14** (a), **15** (b), and **16** (c). The bond distance (Å) and bond angle (°) are discussed in the ESI.†



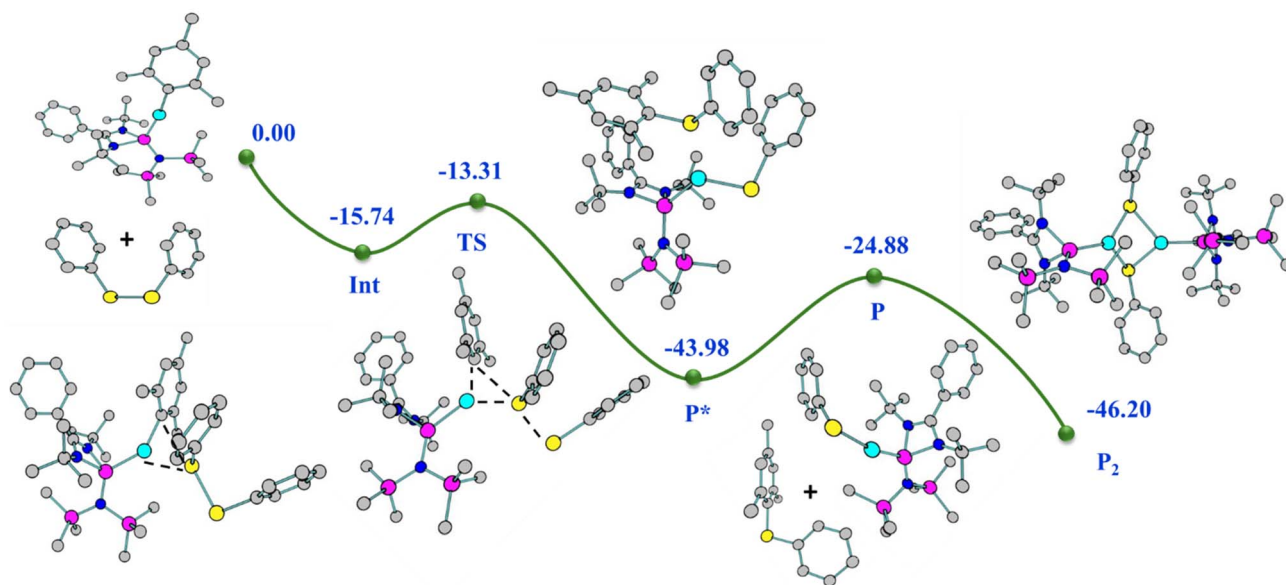


Fig. 7 Relative energy (kcal mol^{-1}) (not drawn to scale) reaction profile for the formation of complex **14** from complex **2** with Ph_2Se_2 .



Scheme 5 Syntheses of various NHSi-coordinated metal amide complexes [monodentate amides (a), bidentate amides (b), and perfluorinated amides (c)] from **2** and **7**.

N,N,N,N-tetramethylethylenediamine) complex [$2.2003(6)$ Å]. Furthermore, we have performed FMO visualisation along with NBO analysis for all the isolated NHSi $\rightarrow$ E-amide complexes (**17**–**22**) to elucidate the nature of the bonding. The HOMO of complex **17** is located over the pyrrolato ring whereas HOMO–1

and HOMO–2 have a major contribution from the Si(II) $\rightarrow$ Cu(I) bond (Fig. S116[†]). For **18**, the HOMO is localised on the 2-mesitylpyrrolide moiety, HOMO–1 on the Cu(I)–N_{amide} fragment, and HOMO–2 across the Si(II) $\rightarrow$ Cu bond (Fig. S116[†]). After isolating complexes **17** and **18**, we were curious to investigate the coordinative variability of N-donor functionalised secondary amines. To explore the coordination ability of N–H systems containing auxiliary N-donor atoms, we obtained a few coordinated Cu(I) complexes (**19** and **20**) (Scheme 5b). The three-coordinate NHSi-copper(I) complex (**19**) was isolated by treating **2** with *N*-mesitylpyridin-2-amine, which gives a broad peak at δ 6.9 ppm for the Si(II) centre in the $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum. Complex **19** features a Si1–Cu1 bond length of $2.1964(8)$ Å (Fig. 8), which is shorter than the Si(II) $\rightarrow$ Cu(I) bond in complex **18**. The Cu1–N5 bond length of **19** is $2.654(2)$ Å, which is significantly longer than the reported IPr coordinated three copper(I) complexes attached to the 2-(2,3,4,5-tetrafluorophenyl) pyridine group ($2.185(4)$ Å).⁴⁵ Thompson and co-workers isolated a few (NHC)Cu(py₂BMe₂) {py₂BMe₂ = di(2-pyridyl)dimethylborate} complexes, which feature N⁺N donation to the Cu(I) centre. These complexes possess Cu–N bond lengths ranging from $1.9929(16)$ to $2.0288(15)$ Å.⁴⁶ The Si1–Cu1–N1 bond angle in **19** is $172.44(7)^\circ$, which deviates from linearity due to the additional N-donation. Stalke and co-workers isolated *N*-mesitylpyridin-2-amine substituted benzamidinato silylene coordinated trimeric Cu(I) halide complexes,⁴⁷ which also feature pyridyl N-donation to the Cu(I) centre with N $\rightarrow$ Cu bond lengths from $1.995(4)$ to $2.009(2)$ Å. These bonds are significantly shorter than the observed N $\rightarrow$ Cu bond in **19**, denoting a greater extent of electron donation in the former one. The amidinate ligands are infinitely versatile in terms of potential structure and substitution patterns.⁴⁸ This prompted us to isolate complex **20** containing the Dipp group substituted formamidine moiety (Scheme 5b) as an example of auxiliary N-



Fig. 8 Molecular structures of complexes 17 (a), 18 (b), 19 (c), 20 (d), 21 (e), and 22 (f). The bond distance (Å) and bond angle (°) are discussed in the ESI.†

donor coordinated secondary amines. This features a broad peak at δ 6.6 ppm for Si(II) attached to three-coordinate Cu(I) centres in the $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum. Due to an additional N-donor atom, the Si(II) centre is more shielded than the complexes 17 and 18. Complex 20 is structurally reminiscent of the CAAC stabilised three-coordinate formamidinate copper(I) complex (Fig. 8).⁴⁹ The molecular structure of 20 reveals a Si1–Cu1–N1 angle of $171.24(9)^\circ$, whereas, for the CAAC one, the C–Cu–N angle is $175.09(7)^\circ$.^{49,50} The Si1–Cu1 and Cu1–N1 bond lengths are 2.194(1) and 1.903(3) Å, respectively. The Cu1–N2 bond length is 2.712(3) Å, shorter than the sum of their van der Waals radii (2.94 Å), suggesting the coordinative nature. This is also shorter than the formamidinate Cu(I) complex of CAAC (Cu–N 2.912(14) Å).⁴⁹ The C–N bond lengths in the formamidinate backbone are different, with long N1–C13 and short N2–C13 distances of 1.337(4) and 1.294(4) Å, respectively. FMO visualisation of 19 and 20 shows that the HOMO of complex 19 shows that there is a major contribution from the 2-mesitylpyridyl moiety and d orbital of the Cu atom (Fig. S117†). The electron density at the HOMO–1 is predominantly

localised over the Cu–N_{amide} bond with a minor contribution over the pyridine ring, whereas the electron density is delocalised over the Si(II) $\rightarrow$ Cu–N–mesityl moiety in HOMO–2 with a minor contribution from the pyridine ring (Fig. S117†). However, for complex 20 the HOMO is majorly located over the formamidinate moiety whereas HOMO–2 and HOMO–4 are spread over the Si(II) $\rightarrow$ Cu bond (Fig. S117†). NBO calculation further shows that the N $\rightarrow$ Cu bond is stabilised by $E^{(2)}$ values of 3.80 and 4.54 kcal mol^{–1} for 19 and 20, respectively (Fig. S122a and b†). While there could be possibilities of numerous secondary amide groups in the literature, N–H bond activation of bis-perfluoroamine is of particular interest because of its weak donation ability to the metal centre.⁵¹ Treatment of complex 2 with bis-pentafluorophenylamine in toluene yielded complex 21 (Scheme 5c), with a Si1–Cu1 bond length of 2.220(1) Å and a Si1–Cu1–N1 angle of $169.3(1)^\circ$ (Fig. 8), deviating from linearity due to *ortho*-fluorine Cu $\cdots$ F contacts (Cu1–F10: 2.791(3) Å and Cu1–F5: 2.727(2) Å). These interactions elongate the corresponding C–F bonds [C–F5: 1.350(6) Å and C–F2: 1.358(3) Å]. Complex 21 shows the peak at δ 7.1 ppm for the Si(II)

Table 1 Bond lengths of NHSi-coordinated Cu–mesityl and Cu–amide complexes

Complex	$^{29}\text{Si}\{^1\text{H}\}$ chemical shift (δ) ppm	Si(II) $\rightarrow$ Cu(I) bond length (Å)	Cu–C/N bond length (Å)
2	6.5	2.2509(8)	1.946(2)
17	7.8	2.231(3)	1.878(7)
18	7.4	2.203(1)	1.874(3)
19	6.9	2.1964(8)	1.906(2)
20	6.6	2.194(1)	1.903(3)
21	7.1	2.220(1)	1.929(4)



centre in the $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum. As a proof of concept, the reaction of complex 7 with bis-perfluoroarylamine produces a structurally identical complex of 22 (Scheme 5c). The Si–Ag bond length in complex 22 is 2.3543(8) Å, longer than the Si–Cu bond length in complex 21. The intermolecular π -stacking between the phenyl group of the amidinate moiety and the perfluoro-substituted benzene ring further stabilises complexes 21 and 22. Also, complex 21 features intermolecular F $\cdots$ F interaction (see the ESI†). The MOs of these complexes are reminiscent of complex 18 (Fig. S118†). However, the Cu $\cdots$ F interactions in complex 21/22 are supported by the $E^{(2)}$ values of 0.70/2.26 and 3.06/1.04 kcal mol $^{-1}$ for F5 and F10, respectively (Fig. S122c and d†).

An apparent trend emerges after observing the chemical shifts of $^{29}\text{Si}\{^1\text{H}\}$ NMR and bonding parameters of the Cu–amide complexes (17–21) with 2 (Table 1). We observe a deshielding effect on the Si(II) centres for complexes 17–21 that can be attributed to the electrophilic nature of the amide group. Furthermore, the Si(II) $\rightarrow$ Cu and Cu–N_{amide} bond lengths of complexes 17–21 are shorter than the corresponding Si(II) $\rightarrow$ Cu and Cu–C_{ipso} bond lengths in complex 2. This discrepancy suggests an enhanced electropositive character of the Cu(I) centre in complexes 17–21, as also realised from the Mulliken charge analysis (Table 2) on the Si, Cu, C_{ipso}, and N_{amide}. It was found that the electropositive character of the Cu(I) centre indeed increases in 17–21 with a gradual increase in

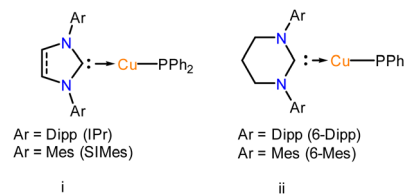


Chart 3 Structurally defined monomeric NHC $\rightarrow$ Cu–PPh₂ complexes reported in the literature.

the electrophilic nature of the amide ligand as compared to the mesityl group of 2. As a proof of concept, we investigated the reaction pathway of complex 17. Interestingly, the reaction profile (Fig. 9) for the formation of complex 17 shows the hydrogen transfer of the pyrrole group to the mesityl ipso carbon in the transition state (TS), which was confirmed by the imaginary frequency at 1373 cm $^{-1}$ along these reaction coordinates. Furthermore, the covalent nature of the C_{ipso}–H and N–H bonds is confirmed by the enormously strong C–H $\cdots$ N hydrogen bond in the TS with an $E^{(2)}$ energy of 134.62 kcal mol $^{-1}$ (details in the ESI†).

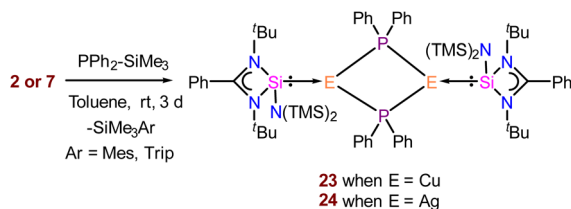
2.2.4 Syntheses of NHSi-based Cu(I) and Ag(I) phosphide complexes (23–24) via P–Si bond cleavage. Low-coordinate coinage metal phosphide complexes stabilised by electron-donating neutral ligands are rare due to their propensity to form oligomers.⁵² This is also reflected by the availability of the limited examples of structurally characterised NHC-coordinated Cu(I)–PR₂ species in the literature (Chart 3).^{8f} The metal centre must have at least one vacant frontier orbital with suitable symmetry to accept π -donation from the p-character lone pair of the phosphide ligand to form an E–PR₂ type bond. However, late transition metals like Cu and Ag, with their higher d-electron counts, are more susceptible to interactions

Table 2 Mulliken charges on Si(II), Cu(I), C_{ipso}, and N_{amide} centres

	2	17	18	19	20	21	22
q_{Si}	0.617	0.830	0.793	0.857	0.837	0.820	0.599
q_{M}	−0.474	−0.342	−0.193	−0.214	−0.203	−0.227	−0.060
$q_{\text{C/N}}$	−0.183	−0.115	−0.209	−0.248	−0.240	−0.239	−0.354



Fig. 9 Relative energy (kcal mol $^{-1}$) (not drawn to scale) reaction profile for the formation of complex 17 from complex 2 with pyrrole.



Scheme 6 Syntheses of NHSi-coordinated metal phosphide complexes from 2 and 7.

between their filled metal orbitals and the phosphorus lone pair. Such interactions often induce a pyramidal geometry at the phosphorus atom. Thus, understanding their bonding situation is an important aspect. To our knowledge, low-coordinate Ag(I)-PR₂ complexes are still structurally unknown. However, polynuclear [Ag₁₂(PSiMe₃)₆(ⁱPr₂-bimy)₆] and [Ag₂₆P₂(-PSiMe₃)₁₀(ⁱPr₂-bimy)₈] were isolated.⁵³ Thus, stabilisation of such coinage metal phosphides in low-nuclearity is challenging. This inspired us to proceed with NHSi → Cu/Ag-PR₂ isolation with complexes 2 and 7 as suitable precursors for the safer handling of Ph₂P-SiMe₃. Adding one equivalent of PPh₂-SiMe₃ to the pale-yellow solution of complexes 2 and 7 resulted in an instantaneous cloudy solution (Scheme 6). Stirring the resulting solution for 2 days, we could isolate complexes 23 and 24 with ~30% yield. The two sets of doublets at δ 2.80–3.23 and 9.30–8.87 ppm in ²⁹Si{¹H} NMR indicate the Si(II)-Ag centre in complex 24. ³¹P{¹H} NMR shows the peak at δ –13.3 ppm for complex 24.

The molecular structures of both complexes reveal a dimeric μ-bridged Cu(I)/Ag(I)-PPh₂ core coordinated with two NHSi moieties from each end (Fig. 10). The formation of 23 and 24 resulted from eliminating Mes-SiMe₃ (for 23) and Trip-SiMe₃

(for 24). However, these eliminations are not very facile under ambient conditions, hence dropping the yield of the desired product. The dimeric core represents a near rhomboid geometry with bond angles for 23 [P1–Cu1–P1 86.16° and Si1–Cu1–P1 136.98°] and 24 [P1–Ag1–P1 84.48° and Si1–Ag1–P1 137.49°]. The Si–Cu(I)/Ag(I) bond distances are 2.2758(8) and 2.401(1) Å for 23 and 24, respectively, which are longer than that in the NHC → Cu–PR₂ complexes [1.9272(18) Å for the monomeric (6-Dipp)CuPPh₂ complex and 1.9234(15) Å for the dimeric IDippCuPPh₂ complex] reported in the literature.^{8f} The Cu–P distance [2.3799(8) Å] in complex 23, is longer than that in the monomeric (6-Dipp)CuPPh₂ complex [2.2113(5) Å] and is in line with that of the dimeric IDippCuPPh₂ complex [2.3298(5) Å].^{8h,i,54} The Ag–P bond distance [2.537(1) Å] in complex 20 is in line with the Ag–P bond distances reported in [Ag₁₂(PSiMe₃)₆(ⁱPr₂-bimy)₆] and [Ag₂₆P₂(PSiMe₃)₁₀(ⁱPr₂-bimy)₈] [2.579(3) and 2.605(5) Å].⁵³ The intermetallic distances (E⋯E) are significantly longer [3.458 Å (complex 23) and 3.791 Å (complex 24)], suggesting the absence of metallophilic interactions in these complexes, unlike those in complex 14. Since such low-coordinate Cu(I)-phosphide complexes are limited only to their structure elucidation, it became customary for us to understand the electronic structures of complexes 23 and 24 using FMO visualisation. The HOMO and HOMO–1 of both complexes were composed of the Cu/Ag–P bonding interaction with d–p orbital overlap, whereas HOMO–4 resides over Si(II)–Cu/Ag(I) bonds (Fig. S119†). Isolation and characterisation of NHSi → coinage metal–phosphides are the first of their kind, thus indicating future possibilities of fine-tuning and utilisation in homogeneous catalysis. The reaction pathway shows that C–H⋯π and C–H⋯P hydrogen bonding stabilises the **Int** observed in the reaction profile (Fig. 11) for the formation of complex 23. The strengthening of the interaction between P and Cu centers in the **TS** suggests the bond formation between P⋯Cu centers. Apart from their brief mention here, a complete discussion of the computed pathways, including energetic profiles and structural details, is provided in the revised ESI (pages S102–S110†). We performed detailed DFT calculations on four representative complexes of each bond type among 12–24, as already mentioned. These complexes were selected to reflect the four distinct types of bond activations investigated: homolytic B–B (12) and Se–Se (14), and heterolytic N–H (17) and P–Si (23) bond activations. We believe the selected cases will provide insight into the general reactivity patterns. All the mechanistic pathways show a similar fashion of nucleophilic attack of the reactant to the Cu(I) centre in the transition states, followed by the mesityl group migration to the electrophilic substituent, affording the desired complexes.

2.3 The catalytic utility of NHSi-organo-coinage metal complexes towards different organic transformations

The examples of the NHSi ligated transition metal mediated catalysis are limited to a handful of catalytic reactions.⁵⁵ Recently, our group has demonstrated the scope of NHSi ligated coinage metal complexes in a few different organic reactions, like azide–alkyne cycloaddition,^{6a,7e,47} glycosidation,^{7f} and three-



Fig. 10 Molecular structures of complexes 23 (a) and 24 (b). The bond distance (Å) and bond angle (°) are discussed in the ESI.†





Fig. 11 Relative energy (kcal mol^{-1}) (not drawn to scale) reaction profile for the formation of complex **23** from complex **2** with $\text{PPh}_2\text{SiMe}_3$.

component coupling reactions.^{7d} Motivated by our previous results on catalysis, we aimed to probe the catalytic performances of the isolated NHSi-organocoinage metal complexes. Our brief attempt at these complexes towards catalysing organic transformations is equally promising.

2.3.1 Use of NHSi-organocopper complexes as an aryl transfer agent in the C–C coupling reaction. Over the years, the prodigious use of aryl boronic esters and aryl magnesium precursors has dominated the area of C–C cross-coupling reactions.⁵⁶ A recent report by Uchiyama and co-workers has shown the applicability of aryl copper in Pd-catalysed C–C coupling with sterically demanding substrates.⁵⁷ Moreover, there are only two reports where NHSis is used as a ligand (catalysts **A** and **B**, Chart 4) for $\text{sp}^3\text{--sp}^2$ and $\text{sp}^2\text{--sp}^2$ C–C coupling reactions.⁵⁸

Since we already had well-characterised NHSi–Cu–aryl systems (**2** and **5**), we utilised them as an efficient aryl group transfer agent for Pd-catalysed C–C coupling reactions with aryl iodides (Scheme 7).

The C–C coupling reactions with a few aryl iodides were carried out under the optimised conditions with complex **2**. We observed good to excellent product yields for the hetero-aromatic (**IV** and **VIII**) and fused aromatic (**V** and **IX**) systems (Scheme 7). Using the NHSi coordinated organo-copper complex as an alternative to conventional aryl boronic esters provides a lower Pd catalyst loading (0.5 mol%) compared to the conventional C–C coupling reactions.⁵⁹ It is important to note that the Pd catalyst loading (5 mol%) for the C–C coupling reaction by Uchiyama and co-workers remains much higher with the use of additional ligand loading (up to 15 mol%) (Table 3).

Previous Work

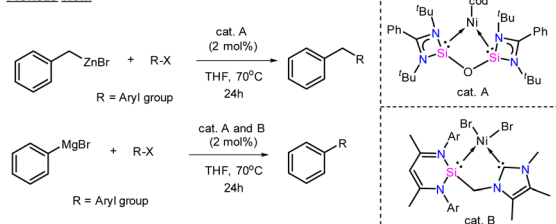


Chart 4 Previous reports on C–C coupling reactions with NHSi as a ligand on the metal centre. (cod = cyclooctadiene).



Scheme 7 Pd-catalysed $\text{sp}^2\text{--sp}^2$ C–C coupling reaction using **2** and **5** as an aryl transfer reagent. *Isolated yields.

Table 3 Pd-catalysed C–C coupling reaction using mesityl copper



Serial no.	Pd-based catalyst (mol%)	Temperature (°C)	Yield (%)	TON
1	5	80–140	91 (ref. 57)	18.2
This work	0.5	110	93	186

The TON for the formation of compound **1** is observed to be 186 with our catalytic method, whereas it is 18.2 by utilising the method of Uchiyama and co-workers (Table 3). Hence, our method utilising **2** and **5** as the aryl transfer reagent works much more efficiently, leading to scope for future exploration of more NHSi-coordinated organocopper compounds in various other organic transformations. This enhanced catalytic performance is presumably due to the strong σ -donor properties of NHSi, which are also observed in other previous reports.^{4c}

2.3.2 Use of NHSi–organosilver complexes as catalysts for three-component reactions for various α -aminonitrile synthesis. Following our success in C–C coupling reactions with the NHSi–organocopper complexes mentioned above, we wanted to explore the catalytic performance of the organosilver analogues as well. We chose the one-pot synthesis of α -aminonitriles as these are an important class of bioactive molecules, and the Strecker reaction is one of the convenient methods to synthesise them.⁶⁰ Until today, these reactions are catalysed by a large excess of Lewis acids as catalysts under vigorous reaction conditions. Thus, we opted for microwave-assisted α -aminonitrile preparation for our purpose. We optimised the best-performing reaction conditions with 5 mol% loading of catalyst **6** at 80 °C for 3 h under solvent-free conditions (Scheme 8). We obtained good to excellent yields as documented in Scheme 8. The future scope can be extended to the number of substrates

with specific utility. This short study highlights the potential of these organo–silver complexes for such important catalytic transformations.

3 Conclusions

In summary, we have established a versatile route to utilise the NHSi–organocoinage metal complexes as an efficient precursor for heteroatom functionalisation. The derivatisation of NHC or CAAC-based coinage metal complexes has become important lately because of the efficient optoelectronic applications of carbene–metal–amide complexes and several organic transformations. However, the functionalisation of NHSi–Cu(I) and Ag(I) complexes was unknown to date. Our adopted methodology allows facile formation of NHSi-supported Cu–B (**12**), Cu/Ag–Se (**14–16**), Cu/Ag–N (**17–22**), and Cu/Ag–P (**23** and **24**) bonds. Moreover, utilising N-donor attached secondary amines gives three-coordinate copper(I) complexes (**21** and **22**). This synthetic protocol helped us to isolate rarely occurring coinage metal phosphides, **23** and **24**. We have also demonstrated the potential of these NHSi–organocoinage metal complexes in two important catalytic organic transformations, which led to the isolation of desired products in excellent yields, paving the way for more exploration in this area.

Data availability

The data supporting this article have been included as part of the ESI.† The ESI contains experimental data, X-ray data, NMR spectra, computational details, and related references.

Author contributions

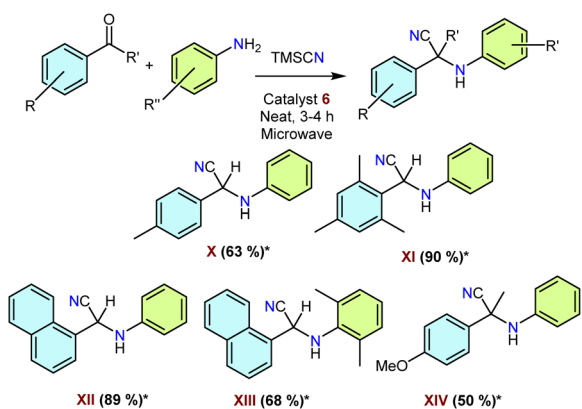
S. K. and M. G. conceived the idea of the project. M. G. contributed to the design, development, execution, and compilation of the project. M. G. performed all the syntheses and characterisation studies and wrote the original draft with the help of P. P. P. performed all the theoretical calculations, prepared the corresponding figures, and provided insightful feedback. K. G. carried out catalytic reactions and curated spectroscopic details in the ESI,† and prepared the figures of characterisation methods along with M. G. S. T. refined the crystallographic data provided by M. G. R. K. P. put significant efforts into refining additional crystal data during the revision. All authors contributed to the discussion. S. K. contributed to fund acquisition, project administration, offered critical insights, coordinated the research, and finalised the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

S. K. thanks SERB-CRG for the financial support. S. K. also thanks DST-FIST for a single-crystal X-ray diffractometer



Scheme 8 General schematic for NHSi–organosilver catalysed α -aminonitrile synthesis. *Isolated yields are reported after taking an average of three runs. # Catalyst loading 5 mol%.



facility. M. G. and P. P. thank IISER Pune for providing a fellowship. K. G. acknowledges the Prime Minister Research Fellowship (PMRF) for providing a fellowship. S. T. is grateful to AESD&CIF and CSIR-CSMCRI. M.G. thanks Dr Sandeep Mishra for his insightful help and discussion regarding NMR. We express our gratitude for the support and resources extended by the “PARAM Brahma Facility” under the National Supercomputing Mission, Government of India, located at IISER Pune.

Notes and references

- (a) B. Blom, D. Gallego and M. Driess, *Inorg. Chem. Front.*, 2014, **1**, 134–148; (b) B. Blom, M. Stoelzel and M. Driess, *Chem.–Eur. J.*, 2013, **19**, 40–62; (c) Y. P. Zhou and M. Driess, *Angew. Chem., Int. Ed.*, 2019, **58**, 3715–3728; (d) S. Yao, A. Saddington, Y. Xiong and M. Driess, *Acc. Chem. Res.*, 2023, **56**, 475–488; (e) L. Wang, Y. Li, Z. Li and M. Kira, *Coord. Chem. Rev.*, 2022, **457**, 214413–214441; (f) S. S. Sen, S. Khan, P. P. Samuel and H. W. Roesky, *Chem. Sci.*, 2012, **3**, 659–682; (g) M. Haaf, T. A. Schmedake and R. West, *Acc. Chem. Res.*, 2000, **33**, 704–714; (h) S. Takahashi, E. Bellan, A. Baceiredo, N. Saffon-Merceron, S. Massou, N. Nakata, D. Hashizume, V. Branchadell and T. Kato, *Angew. Chem., Int. Ed.*, 2019, **58**, 10310–10314; (i) I. Alvarado-Beltran, A. Baceiredo, N. Saffon-Merceron, V. Branchadell and T. Kato, *Angew. Chem., Int. Ed.*, 2016, **55**, 16141–16144.
- C. K. Shan, S. L. Yao and M. Driess, *Chem. Soc. Rev.*, 2020, **49**, 6733–6754.
- (a) D. Gallego, A. Brück, E. Irran, F. Meier, M. Kaupp, M. Driess and J. F. Hartwig, *J. Am. Chem. Soc.*, 2013, **135**, 15617–15626; (b) Y.-P. Zhou, S. Raoufmoghaddam, T. Szilvási and M. Driess, *Angew. Chem., Int. Ed.*, 2016, **55**, 12868–12872.
- (a) Y. Wang, M. Karni, S. Yao, A. Kaushansky, Y. Apeloig and M. Driess, *J. Am. Chem. Soc.*, 2019, **141**, 12916–12927; (b) Y. Wang, T. Szilvási, S. Yao and M. Driess, *Nat. Chem.*, 2020, **12**, 801–807; (c) M. P. Luecke, L. Giarrana, A. Kostenko, T. Gensch, S. Yao and M. Driess, *Angew. Chem., Int. Ed.*, 2022, **61**, e202110398; (d) Y. Xiong, S. Dong, S. Yao, J. Zhu and M. Driess, *Angew. Chem., Int. Ed.*, 2022, **61**, e202205358.
- (a) P. Jutzi, U. Holtmann, D. Kanne, C. Krüger, R. Blom, R. Gleiter and I. Hyla-Kryspin, *Chem. Ber.*, 1989, **122**, 1629–1639; (b) M. Denk, R. Lennon, R. Hayashi, R. West, A. V. Belyakov, H. P. Verne, A. Haaland, M. Wagner and N. Metzler, *J. Am. Chem. Soc.*, 1994, **116**, 2691–2692; (c) C. W. So, H. W. Roesky, J. Magull and R. B. Oswald, *Angew. Chem., Int. Ed.*, 2006, **45**, 3948–3950; (d) M. Kira, S. Ishida, T. Iwamoto and C. Kabuto, *J. Am. Chem. Soc.*, 1999, **121**, 9722–9723; (e) M. Asay, S. Inoue and M. Driess, *Angew. Chem., Int. Ed.*, 2011, **50**, 9589–9592; (f) B. D. Rekken, T. M. Brown, J. C. Fettingner, H. M. Tuononen and P. P. Power, *J. Am. Chem. Soc.*, 2012, **134**, 6504–6507; (g) A. V. Protchenko, K. H. Birj Kumar, D. Dange, A. D. Schwarz, D. Vidovic, C. Jones, N. Kaltsoyannis, P. Mountford and S. Aldridge, *J. Am. Chem. Soc.*, 2012, **134**, 6500–6503; (h) A. Rosas-Sanchez, I. Alvarado-Beltran, A. Baceiredo, N. Saffon-Merceron, S. Massou, V. Branchadell and T. Kato, *Angew. Chem., Int. Ed.*, 2017, **56**, 10549–10554; (i) S. Fujimori and S. Inoue, *Eur. J. Inorg. Chem.*, 2020, **2020**, 3131–3142; (j) R. Guthardt and C. Jones, *Chem. Commun.*, 2024, **60**, 1583–1586; (k) M. M. Roy, M. J. Ferguson, R. McDonald, Y. Zhou and E. Rivard, *Chem. Sci.*, 2019, **10**, 6476–6481; (l) P. Garg, A. Carpentier, I. Douair, D. Dange, Y. Jiang, K. Yuvaraj, L. Maron and C. Jones, *Angew. Chem., Int. Ed.*, 2022, **61**, e202201705.
- (a) M. Ghosh and S. Khan, *Dalton Trans.*, 2021, **50**, 10674–10688; (b) C. Weetman and S. Inoue, *ChemCatChem*, 2018, **10**, 4213–4228; (c) H. Zhu, A. Kostenko, D. Franz, F. Hanusch and S. Inoue, *J. Am. Chem. Soc.*, 2023, **145**, 1011–1021; (d) D. Wendel, D. Reiter, A. Porzelt, P. J. Altmann, S. Inoue and B. Rieger, *J. Am. Chem. Soc.*, 2017, **139**, 17193–17198.
- (a) J. Hossain, J. S. Gopinath, S. Tothadi, P. Parameswaran and S. Khan, *Organometallics*, 2022, **41**, 3706–3717; (b) M. Ghosh and S. Khan, in *Advances in Inorganic Chemistry*, Elsevier, 2023, vol. 81, pp. 279–303; (c) M. Ghosh, S. Tothadi and S. Khan, *Organometallics*, 2021, **40**, 3201–3210; (d) N. Parvin, N. Sen, S. Tothadi, S. Muhammed, P. Parameswaran and S. Khan, *Organometallics*, 2021, **40**, 1626–1632; (e) N. Parvin, J. Hossain, A. George, P. Parameswaran and S. Khan, *Chem. Commun.*, 2020, **56**, 273–276; (f) N. Parvin, B. Mishra, A. George, M. Neralkar, J. Hossain, P. Parameswaran, S. Hotha and S. Khan, *Chem. Commun.*, 2020, **56**, 7625–7628; (g) N. Parvin, S. Pal, J. Echeverria, S. Alvarez and S. Khan, *Chem. Sci.*, 2018, **9**, 4333–4337; (h) M. Nazish, H. Bai, C. M. Legendre, R. Herbst-Irmer, L. Zhao, D. Stalke and H. W. Roesky, *Chem. Commun.*, 2022, **58**, 12704–12707; (i) S. Khan, S. Pal, N. Kathewad, I. Purushothaman, S. De and P. Parameswaran, *Chem. Commun.*, 2016, **52**, 3880–3882; (j) R. Yadav, X. Sun, R. Köppe, M. T. Gamer, F. Weigend and P. W. Roesky, *Angew. Chem., Int. Ed.*, 2022, **61**, e202211115.
- (a) W. Xie and S. Chang, *Angew. Chem., Int. Ed.*, 2016, **55**, 1876–1880; (b) A. Bonet, V. Lillo, J. Ramírez, M. M. Díaz-Requejo and E. Fernández, *Org. Biomol. Chem.*, 2009, **7**, 1533–1535; (c) T. Ohishi, M. Nishiura and Z. Hou, *Angew. Chem., Int. Ed.*, 2008, **47**, 5792–5795; (d) J. R. Herron and Z. T. Ball, *J. Am. Chem. Soc.*, 2008, **130**, 16486–16487; (e) L. Zhang, Z. Li, M. Takimoto and Z. Hou, *Chem. Rec.*, 2020, **20**, 494–512; (f) S. Bagherzadeh and N. P. Mankad, *J. Am. Chem. Soc.*, 2015, **137**, 10898–10901; (g) L. E. English, T. M. H. Downie, C. L. Lyall, M. F. Mahon, C. L. McMullin, S. E. Neale, C. M. Saunders and D. J. Liptrot, *Chem. Commun.*, 2023, **59**, 1074–1077; (h) T. M. Horsley Downie, M. F. Mahon, J. P. Lowe, R. M. Bailey and D. J. Liptrot, *ACS Catal.*, 2022, **12**, 8214–8219; (i) T. M. H. Downie, J. W. Hall, T. P. C. Finn, D. J. Liptrot, J. P. Lowe, M. F. Mahon, C. L. McMullin and M. K. Whittlesey, *Chem. Commun.*, 2020, **56**, 13359–13362; (j) S. Ostrowska, P. Arnaut, D. J. Liptrot, C. S. Cazin and S. P. Nolan, *Chem. Commun.*, 2023, **59**, 9126–9129; (k) L. Zhang and Z. Hou, *Chem. Sci.*,



- 2013, **4**, 3395–3403; (l) F. Lazreg, F. Nahra and C. S. J. Cazin, *Coord. Chem. Rev.*, 2015, **293**, 48–79; (m) J. Hossain, R. Akhtar and S. Khan, *Polyhedron*, 2021, **201**, 115151–115187; (n) S. Díez-González, E. D. Stevens, N. M. Scott, J. L. Petersen and S. P. Nolan, *Chem.–Eur. J.*, 2008, **14**, 158–168; (o) L. Yang and H. Wang, *ChemSusChem*, 2014, **7**, 962–998; (p) A. A. Danopoulos, T. Simler and P. Braunstein, *Chem. Rev.*, 2019, **119**, 3730–3961.
- 9 J. C. Garrison and W. J. Youngs, *Chem. Rev.*, 2005, **105**, 3978–4008.
- 10 M. Stollenz and F. Meyer, *Organometallics*, 2012, **31**, 7708–7727.
- 11 (a) A. G. Avent, B. Gehrhus, P. B. Hitchcock, M. F. Lappert and H. Maciejewski, *J. Organomet. Chem.*, 2003, **686**, 321–331; (b) Y. Inagawa, S. Ishida and T. Iwamoto, *Chem. Lett.*, 2014, **43**, 1665–1667; (c) G. Tan, B. Blom, D. Gallego, E. Irran and M. Driess, *Chem.–Eur. J.*, 2014, **20**, 9400–9408; (d) G. Tan, B. Blom, D. Gallego and M. Driess, *Organometallics*, 2014, **33**, 363–369; (e) T. Troadec, A. Prades, R. Rodriguez, R. Mirgalet, A. Baceiredo, N. Saffon-Merceron, V. Branchadell and T. Kato, *Inorg. Chem.*, 2016, **55**, 8234–8240; (f) N. Parvin, R. Dasgupta, S. Pal, S. S. Sen and S. Khan, *Dalton Trans.*, 2017, **46**, 6528–6532; (g) N. Parvin, S. Pal, S. Khan, S. Das, S. K. Pati and H. W. Roesky, *Inorg. Chem.*, 2017, **56**, 1706–1712; (h) S. Abe, Y. Inagawa, R. Kobayashi, S. Ishida and T. Iwamoto, *Organometallics*, 2022, **41**, 874–882; (i) M. Nazish, M. M. Siddiqui, S. K. Sarkar, A. Munch, C. M. Legendre, R. Herbst-Irmer, D. Stalke and H. W. Roesky, *Chem.–Eur. J.*, 2021, **27**, 1744–1752.
- 12 S. Amanullah, W. Cao, E. Brack, M. Plodinec and C. Copéret, *Angew. Chem., Int. Ed.*, 2025, **64**, e202417428.
- 13 (a) D. S. Laitar, P. Müller and J. P. Sadighi, *J. Am. Chem. Soc.*, 2005, **127**, 17196–17197; (b) Y. Segawa, M. Yamashita and K. Nozaki, *Angew. Chem., Int. Ed.*, 2007, **46**, 6710–6713; (c) R. S. Charman, J. A. Hobson, R. A. Jackson, M. F. Mahon, S. E. Neale and D. J. Liptrot, *Chem.–Eur. J.*, 2024, **30**, e202302704; (d) D. J. Babula, R. S. Charman, T. H. Jerome, T. M. Horsley Downie, M. F. Mahon and D. J. Liptrot, *Eur. J. Inorg. Chem.*, 2023, **26**, e202300043; (e) W. Drescher, C. Borner and C. Kleeberg, *New J. Chem.*, 2021, **45**, 14957–14964.
- 14 (a) A. S. Romanov, C. R. Becker, C. E. James, D. Di, D. Credgington, M. Linnolahti and M. Bochmann, *Chem.–Eur. J.*, 2017, **23**, 4625–4637; (b) W. Zou, Q. Zhu, J. C. Fettinger and P. P. Power, *Inorg. Chem.*, 2021, **60**, 17641–17648; (c) M. Azizpoor Fard, A. M. Polgar and J. F. Corrigan, *Organometallics*, 2020, **39**, 2900–2906; (d) J. Kampf, R. Kumar and J. P. Oliver, *Inorg. Chem.*, 1992, **31**, 3626–3629.
- 15 M. Ghosh, J. Chatterjee, P. Panwaria, A. Kudlu, S. Tothadi and S. Khan, *Angew. Chem., Int. Ed.*, 2024, **63**, e202410792.
- 16 (a) R. Jazzar, M. Soleilhavoup and G. Bertrand, *Chem. Rev.*, 2020, **120**, 4141–4168; (b) R. Hamze, J. L. Peltier, D. Sylvinson, M. Jung, J. Cardenas, R. Haiges, M. Soleilhavoup, R. Jazzar, P. I. Djurovich, G. Bertrand and M. E. Thompson, *Science*, 2019, **363**, 601–606; (c) P. J. Conaghan, C. S. B. Matthews, F. Chotard, S. T. E. Jones, N. C. Greenham, M. Bochmann, D. Credgington and A. S. Romanov, *Nat. Commun.*, 2020, **11**, 1758; (d) M. Gernert, L. Balles-Wolf, F. Kerner, U. Müller, A. Schmiedel, M. Holzapfel, C. M. Marian, J. Pflaum, C. Lambert and A. Steffen, *J. Am. Chem. Soc.*, 2020, **142**, 8897–8909; (e) A. S. Romanov, S. T. Jones, Q. Gu, P. J. Conaghan, B. H. Drummond, J. Feng, F. Chotard, L. Buizza, M. Foley, M. Linnolahti, D. Credgington and M. Bochmann, *Chem. Sci.*, 2020, **11**, 435–446; (f) R. Tang, S. Xu, T.-L. Lam, G. Cheng, L. Du, Q. Wan, J. Yang, F.-F. Hung, K.-H. Low, D. L. Phillips and C.-M. Che, *Angew. Chem., Int. Ed.*, 2022, **61**, e202203982.
- 17 (a) T. Tsuda, T. Yazawa, K. Watanabe, T. Fujii and T. Saegusa, *J. Org. Chem.*, 1981, **46**, 192–194; (b) S. Gambarotta, C. Floriani, A. Chiesi-Villa and C. Guastini, *J. Chem. Soc., Chem. Commun.*, 1983, 1156–1158; (c) E. M. Meyer, S. Gambarotta, C. Floriani, A. Chiesi-Villa and C. Guastini, *Organometallics*, 1989, **8**, 1067–1079; (d) S. S. Sen, J. Hey, R. Herbst-Irmer, H. W. Roesky and D. Stalke, *J. Am. Chem. Soc.*, 2011, **133**, 12311–12316.
- 18 C. Boehme and G. Frenking, *Organometallics*, 1998, **17**, 5801–5809.
- 19 L. Kuehn, A. F. Eichhorn, D. Schmidt, T. B. Marder and U. Radius, *J. Organomet. Chem.*, 2020, **919**, 121249.
- 20 M. Riethmann, S. A. Föhrenbacher, H. Keiling, N. V. Ignat'ev, M. Finze and U. Radius, *Inorg. Chem.*, 2024, **63**, 8351–8365.
- 21 M. Dahlen, T. P. Seifert, S. Lebedkin, M. T. Gamer, M. M. Kappes and P. W. Roesky, *Chem. Commun.*, 2021, **57**, 13146–13149.
- 22 H. Schmidbaur and A. Schier, *Angew. Chem., Int. Ed.*, 2015, **54**, 746–784.
- 23 B. K. Tate, A. J. Jordan, J. Bacsá and J. P. Sadighi, *Organometallics*, 2017, **36**, 964–974.
- 24 Y. Liu, L. J. Taylor, S. P. Argent, J. McMaster and D. L. Kays, *Inorg. Chem.*, 2021, **60**, 10114–10123.
- 25 B. K. Tate, C. M. Wyss, J. Bacsá, K. Kluge, L. Gelbaum and J. P. Sadighi, *Chem. Sci.*, 2013, **4**, 3068–3074.
- 26 D. Li, I. Keresztes, R. Hopson and P. G. Williard, *Acc. Chem. Res.*, 2009, **42**, 270–280.
- 27 (a) C. M. Lemon, D. C. Powers, M. Huynh, A. G. Maher, A. A. Phillips, B. P. Triplet and D. G. Nocera, *Inorg. Chem.*, 2022, **62**, 3–17; (b) M. P. Carranza, B. R. Manzano, F. A. Jalón, A. M. Rodriguez, L. Santos and M. Moreno, *New J. Chem.*, 2013, **37**, 3183–3194; (c) M. P. Carranza, B. R. Manzano, F. A. Jalon, A. M. Rodriguez, L. Santos and M. Moreno, *Inorg. Chem.*, 2010, **49**, 3828–3835; (d) L. Ray, M. M. Shaikh and P. Ghosh, *Inorg. Chem.*, 2008, **47**, 230–240; (e) J. E. Aguado, M. J. Calhorda, P. J. Costa, O. Crespo, V. Félix, M. C. Gimeno, P. G. Jones and A. Laguna, *Eur. J. Inorg. Chem.*, 2004, **2004**, 3038–3047.
- 28 (a) S. S. Sen, H. W. Roesky, D. Stern, J. Henn and D. Stalke, *J. Am. Chem. Soc.*, 2010, **132**, 1123–1126; (b) R. Azhakar, R. S. Ghadwal, H. W. Roesky, H. Wolf and D. Stalke, *Organometallics*, 2012, **31**, 4588–4592.
- 29 R. Waterman, *Organometallics*, 2013, **32**, 7249–7263.



- 30 (a) K. K. Hollister, A. Molino, G. Breiner, J. E. Walley, K. E. Wentz, A. M. Conley, D. A. Dickie, D. J. Wilson and R. J. Gilliard Jr, *J. Am. Chem. Soc.*, 2022, **144**, 590–598; (b) K. E. Wentz, A. Molino, L. A. Freeman, D. A. Dickie, D. J. Wilson and R. J. Gilliard Jr, *J. Am. Chem. Soc.*, 2022, **144**, 16276–16281; (c) S. K. Sarkar, K. K. Hollister, A. Molino, A. D. Obi, C.-L. Deng, B. Y. E. Tra, B. M. Stewart, D. A. Dickie, D. J. D. Wilson and R. J. Gilliard Jr, *J. Am. Chem. Soc.*, 2023, **145**, 21475–21482.
- 31 (a) P. Bissinger, A. Steffen, A. Vargas, R. D. Dewhurst, A. Damme and H. Braunschweig, *Angew. Chem., Int. Ed.*, 2015, **54**, 4362–4366; (b) H. Braunschweig, T. Dellermann, R. D. Dewhurst, B. Hupp, T. Kramer, J. D. Mattock, J. Mies, A. K. Phukan, A. Steffen and A. Vargas, *J. Am. Chem. Soc.*, 2017, **139**, 4887–4893.
- 32 (a) E. C. Neeve, S. J. Geier, I. A. Mkhalid, S. A. Westcott and T. B. Marder, *Chem. Rev.*, 2016, **116**, 9091–9161; (b) R. D. Dewhurst, E. C. Neeve, H. Braunschweig and T. B. Marder, *Chem. Commun.*, 2015, **51**, 9594–9607.
- 33 A. F. Eichhorn, L. Kuehn, T. B. Marder and U. Radius, *Chem. Commun.*, 2017, **53**, 11694–11696.
- 34 L. Dang, H. Zhao, Z. Lin and T. B. Marder, *Organometallics*, 2008, **27**, 1178–1186.
- 35 (a) C. M. Wyss, J. Bitting, J. Bacsá, T. G. Gray and J. P. Sadighi, *Organometallics*, 2016, **35**, 71–74; (b) C. Kleeberg and C. Borner, *Organometallics*, 2018, **37**, 4136–4146; (c) C. Borner, L. Anders, K. Brandhorst and C. Kleeberg, *Organometallics*, 2017, **36**, 4687–4690; (d) P. M. Rutz, J. r. Grunenberg and C. Kleeberg, *Organometallics*, 2022, **41**, 3044–3054; (e) P. Rios, M. S. See, R. C. Handford, S. J. Teat and T. D. Tilley, *Chem. Sci.*, 2022, **13**, 6619–6625.
- 36 G. A. Molander, S. L. Trice and S. D. Dreher, *J. Am. Chem. Soc.*, 2010, **132**, 17701–17703.
- 37 N. Oka, T. Yamada, H. Sajiki, S. Akai and T. Ikawa, *Org. Lett.*, 2022, **24**, 3510–3514.
- 38 R. An, L. Liao, X. Liu, S. Song and X. Zhao, *Org. Chem. Front.*, 2018, **5**, 3557–3561.
- 39 S. Kolay, A. Wadawale, D. Das, H. K. Kisan, R. B. Sunoj and V. K. Jain, *Dalton Trans.*, 2013, **42**, 10828–10837.
- 40 W. J. Humenny, S. Mitzinger, C. B. Khadka, B. K. Najafabadi, I. Vieira and J. F. Corrigan, *Dalton Trans.*, 2012, **41**, 4413–4422.
- 41 N. V. S. Harisomayajula, S. Makovetskyi and Y. C. Tsai, *Chem.–Eur. J.*, 2019, **25**, 8936–8954.
- 42 R. Bühler, R. M. Wolf, C. Gemel, J. Stephan, S. N. Deger, S. Kahlal, R. A. Fischer and J.-Y. Saillard, *Inorg. Chem.*, 2024, **63**, 17617–17625.
- 43 (a) M. Stollenz, M. John, H. Gehring, S. Dechert, C. Grosse and F. Meyer, *Inorg. Chem.*, 2009, **48**, 10049–10059; (b) J. I. van der Vlugt, *Chem. Soc. Rev.*, 2010, **39**, 2302–2322; (c) P. A. Chase and D. W. Stephan, *Angew. Chem., Int. Ed.*, 2008, **47**, 7433–7437.
- 44 (a) T. Scattolin, A. A. Logvinov, N. V. Tzouras, C. S. J. Cazin and S. P. Nolan, *Organometallics*, 2023, **42**, 2692–2730; (b) L. A. Goj, E. D. Blue, C. Munro-Leighton, T. B. Gunnoe and J. L. Petersen, *Inorg. Chem.*, 2005, **44**, 8647–8649; (c) L. A. Goj, E. D. Blue, S. A. Delp, T. B. Gunnoe, T. R. Cundari, A. W. Pierpont, J. L. Petersen and P. D. Boyle, *Inorg. Chem.*, 2006, **45**, 9032–9045.
- 45 R. Molteni, K. Edkins, M. Haehnel and A. Steffen, *Organometallics*, 2016, **35**, 629–640.
- 46 V. A. Krylova, P. I. Djurovich, B. L. Conley, R. Haiges, M. T. Whited, T. J. Williams and M. E. Thompson, *Chem. Commun.*, 2014, **50**, 7176–7179.
- 47 A. N. Paesch, A. K. Kreyenschmidt, R. Herbst-Irmer and D. Stalke, *Inorg. Chem.*, 2019, **58**, 7000–7009.
- 48 S. T. Barry, *Coord. Chem. Rev.*, 2013, **257**, 3192–3201.
- 49 A. S. Romanov, F. Chotard, J. Rashid and M. Bochmann, *Dalton Trans.*, 2019, **48**, 15445–15454.
- 50 F. Chotard, V. Sivchik, M. Linnolahti, M. Bochmann and A. S. Romanov, *Chem. Mater.*, 2020, **32**, 6114–6122.
- 51 J. Kögel, D. Sorokin, A. Khvorost, M. Scott, K. Harms, D. Himmel, I. Krossing and J. Sundermeyer, *Chem. Sci.*, 2018, **9**, 245–253.
- 52 B. K. Najafabadi and J. F. Corrigan, *Dalton Trans.*, 2015, **44**, 14235–14241.
- 53 B. K. Najafabadi and J. F. Corrigan, *Chem. Commun.*, 2015, **51**, 665–667.
- 54 (a) G. C. Fortman, A. M. Slawin and S. P. Nolan, *Organometallics*, 2010, **29**, 3966–3972; (b) J. Yuan, L. Zhu, J. Zhang, J. Li and C. Cui, *Organometallics*, 2017, **36**, 455–459.
- 55 (a) H. Ahuja, H. Kaur and R. Arevalo, *Inorg. Chem. Front.*, 2023, **10**, 6067–6076; (b) S. Kalra, D. Pividori, D. Fehn, C. Dai, S. Dong, S. Yao, J. Zhu, K. Meyer and M. Driess, *Chem. Sci.*, 2022, **13**, 8634–8641.
- 56 (a) L. Yin and J. Liebscher, *Chem. Rev.*, 2007, **107**, 133–173; (b) M. Busch, M. D. Wodrich and C. Corminboeuf, *ACS Catal.*, 2017, **7**, 5643–5653.
- 57 M. Oi, R. Takita, J. Kanazawa, A. Muranaka, C. Wang and M. Uchiyama, *Chem. Sci.*, 2019, **10**, 6107–6112.
- 58 (a) W. Wang, S. Inoue, S. Yao and M. Driess, *J. Am. Chem. Soc.*, 2010, **132**, 15890–15892; (b) C. I. Someya, M. Haberberger, W. Wang, S. Enthaler and S. Inoue, *Chem. Lett.*, 2013, **42**, 286–288.
- 59 R. Martin and S. L. Buchwald, *Acc. Chem. Res.*, 2008, **41**, 1461–1473.
- 60 J. Wang, X. Liu and X. Feng, *Chem. Rev.*, 2011, **111**, 6947–6983.

