



Cite this: *Dalton Trans.*, 2018, **47**, 769

## Mono- and dinuclear Ni(I) products formed upon bromide abstraction from the Ni(I) ring-expanded NHC complex [Ni(6-Mes)(PPh<sub>3</sub>)Br]<sup>†</sup>

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Bromide abstraction from the three-coordinate Ni(I) ring-expanded N-heterocyclic carbene complex [Ni(6-Mes)(PPh<sub>3</sub>)Br] (**1**; 6-Mes = 1,3-bis(2,4,6-trimethylphenyl)-3,4,5,6-tetrahydropyrimidin-2-ylidene) with TlPF<sub>6</sub> in THF yields the T-shaped cationic solvent complex, [Ni(6-Mes)(PPh<sub>3</sub>)(THF)][PF<sub>6</sub>] (**2**), whereas treatment with NaBAr<sub>4</sub> in Et<sub>2</sub>O affords the dimeric Ni(I) product, [(Ni(6-Mes)(PPh<sub>3</sub>)<sub>2</sub>(μ-Br)][BAr<sub>4</sub>] (**3**). Both **2** and **3** act as latent sources of the cation [Ni(6-Mes)(PPh<sub>3</sub>)]<sup>+</sup>, which can be trapped by CO to give [Ni(6-Mes)(PPh<sub>3</sub>)(CO)]<sup>+</sup> (**5**). Addition of [(Et<sub>3</sub>Si)<sub>2</sub>(μ-H)][B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>] to **1** followed by work up in toluene results in the elimination of phosphine as well as halide to afford a co-crystallised mixture of [Ni(6-Mes)(η<sup>2</sup>-C<sub>6</sub>H<sub>5</sub>Me)][B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>] (**4**), and [6MesH...C<sub>6</sub>H<sub>5</sub>Me][B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>]. Treatment of **1** with sodium salts of more strongly coordinating anions leads to substitution products. Thus, NaBH<sub>4</sub> yields the neutral, diamagnetic dimer [(Ni(6-Mes))<sub>2</sub>(BH<sub>4</sub>)<sub>2</sub>] (**6**), whereas NaBH<sub>3</sub>(CN) gives the paramagnetic monomeric cyanotrihydroborate complex [Ni(6-Mes)(PPh<sub>3</sub>)(NCBH<sub>3</sub>)] (**7**). Treatment of **1** with NaO<sup>t</sup>Bu/NHPh<sub>2</sub> affords the three-coordinate Ni(I) amido species, [Ni(6-Mes)(PPh<sub>3</sub>)(NPh<sub>2</sub>)] (**8**). The electronic structures of **2**, **5**, **7** and **8** have been analysed in comparison to that of previously reported **1** using a combination of EPR spectroscopy and density functional theory.

Received 6th November 2017,  
Accepted 1st December 2017

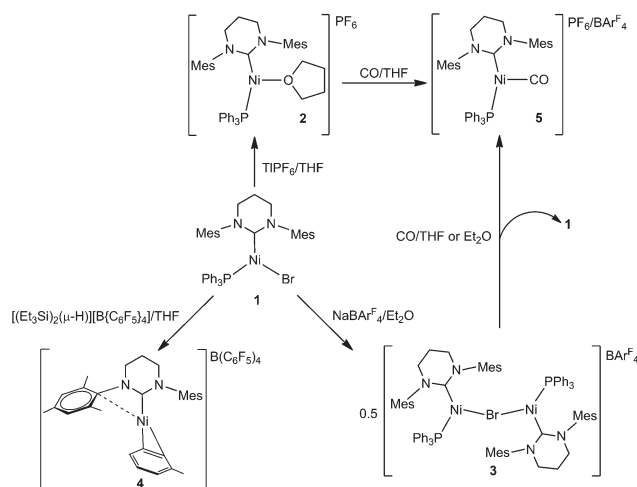
DOI: 10.1039/c7dt04187j

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## Introduction

In a very recent review, Lin and Power referred to Ni(I) as a ‘... “rare” oxidation state of growing importance’.<sup>1,2</sup> In terms of monodentate ligands, the early dependence on tertiary phosphines to stabilise Ni(I)<sup>3,4</sup> has largely been superseded by the use of N-heterocyclic carbenes (NHCs) and these have facilitated the isolation of a wide range of fully characterised four-, three- and even two-coordinate Ni(I) species.<sup>5,6</sup>

Over the last few years, we have used so-called ring-expanded NHCs (RE-NHCs; carbenes with ring sizes >5) for the preparation of three- and two-coordinate Ni(I) complexes with interesting stoichiometric<sup>7</sup> and catalytic chemistry,<sup>8</sup> as well as novel magnetic properties.<sup>9</sup> In all cases, the starting point for our chemistry has been the three-coordinate species



Scheme 1

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<sup>†</sup>Electronic supplementary information (ESI) available. CCDC 1578636–1578644, 1582301 and 1584193. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7dt04187j

[Ni(RE-NHC)(PPh<sub>3</sub>)Br].<sup>10</sup> The first of these to be prepared, [Ni(6-Mes)(PPh<sub>3</sub>)Br] (**1**, Scheme 1),<sup>8</sup> has continued to be the focus of much of our attention as it tends to yield readily isolable products.



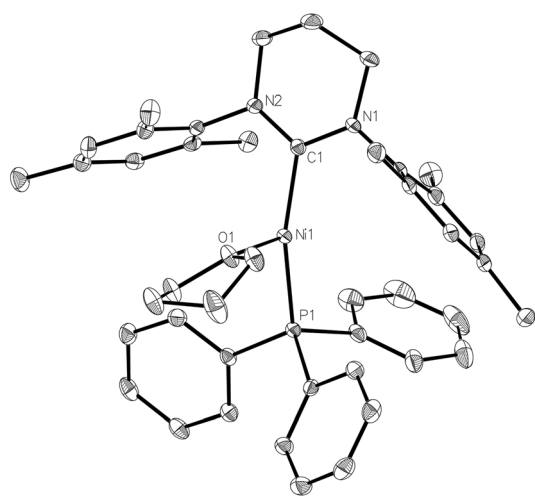
Herein, we describe the stoichiometric reactivity of **1** with a range of bromide abstracting agents to afford seven new Ni(I) complexes. Five of these are monomeric (cationic as well as neutral) and their adoption of T- or Y-shaped structures has been probed using DFT calculations.

## Results and discussion

### Bromide abstraction from **1** by $[\text{Ti}]^+$ , $\text{NaBAR}^{\text{F}}_4$ and $[(\text{Et}_3\text{Si})_2(\mu\text{-H})][\text{B}(\text{C}_6\text{F}_5)_4]$

We have previously shown that the addition of free 6-Mes to **1** results in transfer of the bromide ligand to the outer-sphere to give the two-coordinate, cationic product  $[\text{Ni}(\text{6-Mes})_2]\text{Br}^{\ominus}$ .<sup>9</sup> Initial efforts to abstract bromide from **1** with more typical halide abstractors such as  $\text{AgX}$  reagents ( $\text{X} = \text{BF}_4, \text{NO}_3, \text{OTf}$ ) yielded only mixtures of products containing the pyrimidinium salt  $[\text{6-MesH}]\text{X}$  and the plating out of what appeared to be metallic nickel. However, when **1** was treated with  $\text{TiPF}_6$  in THF, the three-coordinate cationic THF complex  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{THF})][\text{PF}_6]$  (**2**) was isolated as a pale yellow solid in 85% yield (Scheme 1).

The X-ray crystal structure of **2** (Fig. 1) revealed a distorted T-shaped geometry at the Ni(I) centre, with C–Ni–P and C–Ni–O angles of  $156.58(4)^\circ$  and  $103.74(4)^\circ$  respectively. In contrast to the precursor complex **1**, the Ni–C<sub>6-Mes</sub> bond length was slightly elongated ( $1.9601(12)$  Å *cf.*  $1.942(2)$  Å), although the Ni–P bond was unchanged. The Ni–O distance of  $2.0603(9)$  Å was intermediate between those reported for the neutral  $\beta$ -diketiminato species  $[\text{L}^{\text{R}}\text{Ni}(\text{THF})]$  ( $\text{L}^{\text{R}} = [\text{HC}(\text{C}^{\text{t}}\text{Bu})\text{NC}_6\text{H}_3(\text{t}^{\text{Pr}})_2]^-$ ,  $2.000(1)$  Å)<sup>11–13</sup> and the cationic, bis-THF complex  $[(\text{THF})_2\text{Ni}(\text{CNAr}^{\text{Mes}2})_3][\text{OTf}]$  ( $\text{Ar}^{\text{Mes}2} = 2,6\text{-}(2,4,6\text{-Me}_3\text{C}_6\text{H}_2)_2\text{C}_6\text{H}_3$ ;  $2.174(2)$  Å,  $2.1935(19)$  Å).<sup>14</sup>

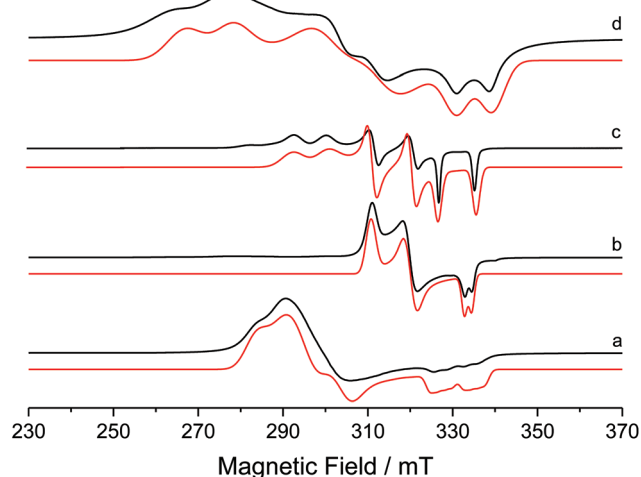


**Fig. 1** Molecular structure of the cation in **2**. Ellipsoids are shown at 30% probability. Hydrogen atoms have been omitted for clarity. Selected bond lengths (Å) and angles ( $^\circ$ ): Ni(1)–C(1)  $1.9601(12)$ , Ni(1)–P(1)  $2.2117(3)$ , Ni(1)–O(1)  $2.0603(9)$ , C(1)–Ni(1)–P(1)  $156.58(4)$ , C(1)–Ni(1)–O(2)  $103.73(4)$ , P(1)–Ni(1)–O(2)  $99.69(3)$ .

The  $^1\text{H}$  NMR spectrum of **2** displayed a series of broad resonances between *ca.*  $\delta$  17–0 which could not be integrated. As the signals for the bound THF could not be assigned, we were unable to establish spectroscopically the lability of the THF ligand. However, X-ray crystallography repeatedly revealed the presence of THF following recrystallization of **2** from a number of solvents ( $\text{CH}_2\text{Cl}_2, \text{C}_6\text{H}_5\text{F}, \text{C}_6\text{H}_6$ ) suggesting that the THF cannot be easily dissociated from the nickel.

The formation of  $\text{TiBr}$  as a side-product in the synthesis of **2** proved problematic, as even following multiple recrystallisations, complete removal was not always achievable. This manifested itself in EPR spectra of **2** ( $\text{ESI}^+$ ), but more obviously in reactions with CO (*vide infra*). Fig. 2 shows the EPR spectrum of a ‘clean’ sample of complex **2** (Fig. 2d). The spin Hamiltonian parameters of the EPR spectra of all of the species shown in Fig. 2 are listed in Table 1, and are discussed further below.

In an attempt to circumvent the problem of  $\text{TiBr}$  contamination, **1** was reacted instead with  $\text{NaBAR}^{\text{F}}_4$  in THF. No bromide abstraction resulted. However, a reaction between **1** and  $\text{NaBAR}^{\text{F}}_4$  took place upon changing the solvent to  $\text{Et}_2\text{O}$ , affording large orange crystals. These proved to be of the unusual cationic, mono-bromide bridged dimer,  $[\{\text{Ni}(\text{6-Mes})(\text{PPh}_3)_2(\mu\text{-Br})\}][\text{BAR}^{\text{F}}_4]$  (**3**, Scheme 1) rather than **2**. The structure of **3** (Fig. 3) comprised of two  $\{\text{Ni}(\text{6-Mes})(\text{PPh}_3)\}$  fragments and a close to symmetrically Ni bound  $\mu$ -bromide ligand (Ni1–Br1  $2.3803(6)$  Å, Ni2–Br1  $2.3688(6)$  Å) in an arrangement distorted from linearity ( $\angle\text{Ni1–Br1–Ni2}$   $167.64(3)^\circ$ ). This is adopted presumably to minimise the steric demands of the ligand substituents in the solid-state. Unsurprisingly, these steric demands also preclude the ligands from eclipsing each other relative to the Ni...Ni axis. Thus, there is  $46.8(1)^\circ$  angle between the mean planes containing atoms Br1, Ni1, P1, C1 and Br1, Ni2, P2, C41, respectively. The Ni...Ni separation exceeds  $4.7$  Å.

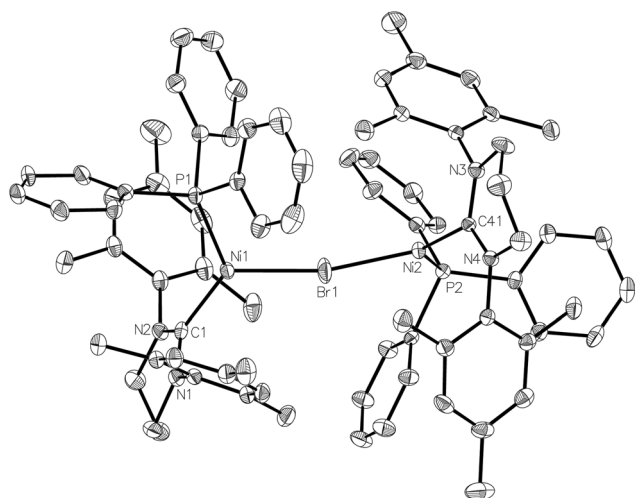


**Fig. 2** Experimental (black) and simulated (red) X-band CW EPR spectra of (a) **1**, (b) **5**, (c) **8** and (d) **2** in frozen THF solution at 140 K.



**Table 1** Spin Hamiltonian parameters for  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)\text{X}]^{0/+}$  dissolved in THF: experimental and computed  $g$ -tensors, and phosphorus and bromide (indicated by #) superhyperfine coupling constants (MHz), with Euler angles derived from DFT calculations

	$g$ values				Euler angles/rad			$A$ values/MHz				Euler angles/rad		
	$g_1$	$g_2$	$g_3$	$g_{\text{iso}}$	$\alpha$	$\beta$	$\gamma$	$A_1$	$A_2$	$A_3$	$a_{\text{iso}}$	$\alpha$	$\beta$	$\gamma$
<b><math>[\text{Ni}(\text{6-Mes})(\text{PPh}_3)\text{Br}]</math> 1</b>														
Expt	2.050	2.265	2.365	2.227	-2.270	2.619	1.643	184	194	250	209	0.035	1.580	1.936
								-6 <sup>#</sup>	-27 <sup>#</sup>	70 <sup>#</sup>	12 <sup>#</sup>	1.580 <sup>#</sup>	1.566 <sup>#</sup>	-1.328 <sup>#</sup>
DFT	2.055	2.252	2.285	2.197	-2.276	2.246	1.628	173	173	204	183	0.176	1.590	1.968
								-19 <sup>#</sup>	-21 <sup>#</sup>	51 <sup>#</sup>	4 <sup>#</sup>	0.093 <sup>#</sup>	1.578 <sup>#</sup>	1.859 <sup>#</sup>
<b><math>[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{THF})][\text{PF}_6]</math> 2</b>														
Expt	2.025	2.210	2.490	2.242	1.497	2.540	1.458	292	210	419	307	0.646	2.969	-0.474
DFT	2.013	2.315	2.389	2.239	-1.580	1.203	1.622	-7	-15	-19	-14	1.700	1.635	-0.436
<b><math>[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{CO})][\text{PF}_6]</math> 5</b>														
Expt	2.035	2.121	2.185	2.114	-3.114	3.064	-1.605	21	29	48	33	1.738	1.560	1.287
DFT	2.044	2.117	2.155	2.105	3.100	3.051	-1.654	14	24	42	26	1.728	1.558	1.235
<b><math>[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{NCBH}_3)]</math> 7</b>														
Expt	2.028	2.225	2.373	2.209	1.176	2.554	-1.575	260	300	260	273	-0.815	1.427	1.780
DFT	2.020	2.286	2.303	2.203	-1.950	1.051	1.568	185	187	220	197	-0.865	1.423	1.778
<b><math>[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{NPh}_2)]</math> 8</b>														
Expt	2.050	2.150	2.290	2.163	-1.459	0.209	0.877	230	265	300	265	1.621	1.602	-1.216
DFT	2.059	2.162	2.238	2.152	-1.513	0.227	0.930	248	249	280	259	1.554	1.610	-1.190

**Fig. 3** Molecular structure of the cation in **3**. Ellipsoids are shown at the 30% probability level. Hydrogen atoms have been omitted for clarity. Selected bond lengths (Å) and angles (°): Ni(1)–C(1) 1.937(4), Ni(1)–P(1) 2.2172(12), Ni(1)–Br(1) 2.3803(6), Ni(2)–C(41) 1.935(4), Ni(2)–P(2) 2.2186(11), Ni(2)–Br(1) 2.3688(6), C(1)–Ni(1)–P(1) 124.15(11), C(1)–Ni(1)–Br(1) 122.16(11), P(1)–Ni(1)–Br(1) 113.68(4), Ni(1)–Br(1)–Ni(2) 167.64(3).

**3** exhibited a paramagnetic NMR spectrum in  $\text{Et}_2\text{O}$  solution, as well as a room temperature magnetic moment ( $\mu_{\text{eff}}$ , Evans method) of  $2.51 \mu_{\text{B}}$ , consistent with the presence two Ni(I) centres.<sup>15</sup> Density functional theory (DFT) calculations based on the crystal structure coordinates with optimised hydrogen atom positions afforded Mulliken spin populations of 0.78 at each Ni centre, thus correctly representing the formal Ni(I) oxidation states. Broken-symmetry DFT predicted antiferromagnetic coupling between the two nickel centres,

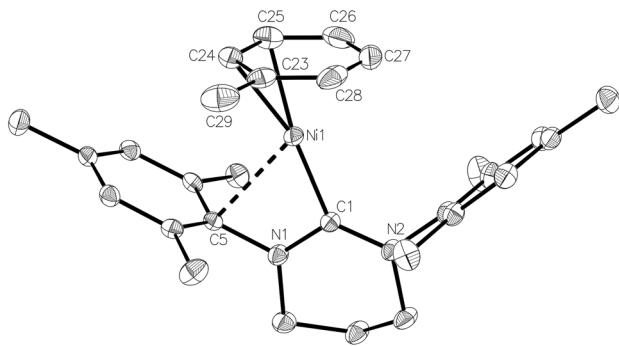
with a medium strength negative exchange coupling constant (density functional dependent: TPSSH:  $-97.6 \text{ cm}^{-1}$ , B3LYP:  $-76.1 \text{ cm}^{-1}$ , PBE0:  $-64.3 \text{ cm}^{-1}$ , M06:  $-69.3 \text{ cm}^{-1}$ ). This coupling appears weak enough to allow significant population of the high-spin state at room temperature: indeed, a Boltzmann population distribution analysis showed that *ca.* 40% of the triplet state would be populated at 300 K (ESI<sup>†</sup>).

EPR spectroscopy revealed that, at least in diethyl ether solution,<sup>16</sup> the two Ni(I) centres were  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)\text{Br}]$  (**1**) and (solvated)  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)]^+$ . Thus, the CW EPR spectrum (140 K, frozen  $\text{Et}_2\text{O}$  glass; ESI<sup>†</sup>) clearly contained signal intensity from **1**, as well as a second Ni(I) centre. The similar profile of this second species to that of **2** suggests it is the diethyl ether complex,  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{OEt}_2)]^+$ . Dissociation of **3** was also supported by the appearance of  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)(\text{CO})][\text{BAR}^{\text{F}}_4]$  (**5-BAR<sup>F</sup>**, *vide infra*) by IR spectroscopy following treatment of the dimer with CO in  $\text{Et}_2\text{O}$  solution.

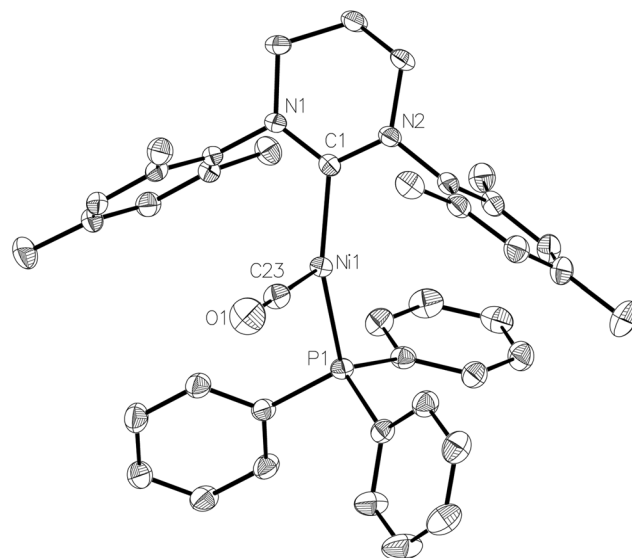
Treatment of **1** with an equimolar amount of  $[(\text{Et}_3\text{Si})_2(\mu\text{-H})][\text{B}(\text{C}_6\text{F}_5)_4]^{17}$  in fluorobenzene resulted in an instantaneous colour change from yellow to red. Upon layering with toluene, light green crystals formed, which consisted of a co-crystallised mixture of the Ni(I) toluene salt,  $[\text{Ni}(\text{6-Mes})(\eta^2\text{-C}_6\text{H}_5\text{Me})][\text{B}(\text{C}_6\text{F}_5)_4]$  (**4**), and  $[(\text{6-MesH})\cdots\text{C}_6\text{H}_5\text{Me}][\text{B}(\text{C}_6\text{F}_5)_4]$ .

The X-ray structure of the metal-containing cation is shown in Fig. 4. The metrics of the coordinated toluene ligand revealed short Ni–C24/C25 distances (2.054(3) Å and 2.092(3) Å), intermediate Ni–C23/C26 distances (2.152(3) Å, 2.202(3) Å) and two substantially longer interactions (Ni–C27 2.271(3) Å, Ni–C28 2.241(3) Å), consistent with an  $\eta^2$  rather than  $\eta^6$  bound arene ligand.<sup>5m,18</sup> To overcome electron deficiency, this then formally 13-electron nickel centre exhibits a close interaction with the *ipso*-C of one of the mesityl rings (Ni–C5 2.525(2) Å; *cf.* Ni–C14 3.367(2) Å). Comparable stabilising close contacts have been





**Fig. 4** Structure of the metal-containing, cationic component in compound **4**. Ellipsoids are shown at the 30% probability level. Hydrogen atoms have been omitted for clarity. Selected bond lengths (Å) and angles (°): Ni(1)–C(1) 1.914(2), Ni(1)–C(23) 2.152(3), Ni(1)–C(24) 2.054(3), Ni(1)–C(25) 2.092(3), Ni(1)–C(26) 2.202(3), Ni(1)–C(27) 2.271(3), Ni(1)–C(28) 2.241(3), N(1)–C(1)–Ni(1) 104.72(15), N(2)–C(1)–Ni(1) 134.13(17).



**Fig. 5** Molecular structure of the cation in **5**. Ellipsoids are shown at the 30% probability level. Hydrogen atoms have been omitted for clarity. Selected bond lengths (Å) and angles (°): Ni(1)–C(1) 1.943(3), Ni(1)–C(23) 1.787(3), Ni(1)–P(1) 2.2374(8), C(23)–O(1) 1.132(4), C(1)–Ni(1)–P(1) 151.93(9), C(1)–Ni(1)–C(23) 106.24(13), P(1)–Ni(1)–C(23) 101.76(11), O(1)–C(23)–Ni(1) 178.0(3).

seen in other coordinatively unsaturated metal complexes bearing bulky NHCs.<sup>19</sup>

Further analysis revealed that the toluene ligand lies almost parallel to one of the fluoroaryl ligands of the  $[B(C_6F_5)_4]^-$  anion. A value of  $9.9^\circ$  for the angle between the relevant least-squares aromatic ring planes,  $3.78 \text{ \AA}$  for the centroid–centroid distance between these rings and  $3.27 \text{ \AA}$  for the shortest distance from the centroid of one ring to the mean plane of the other support the presence of offset  $\pi$ – $\pi$  stacking and additional  $\pi$ -stabilisation of the complex.

The ratio of **4**:toluene stabilised pyrimidinium cation varied from one synthesis to another. The ‘best’ ratio, determined crystallographically, in terms of optimising the percentage of nickel complex yielded, was 65 : 35. This was achieved by (i) performing the complete reaction in a glovebox and (ii) washing the  $[(Et_3Si)_2(\mu-H)][B(C_6F_5)_4]$  five times with hexane and drying overnight. Although the irreproducibility in yield of **4** frustrated efforts to further characterise the complex, the synthetic approach was validated by isolation of the corresponding mesitylene analogue  $[Ni(6-Mes)(\eta^2-C_6H_3Me_3)][B(C_6F_5)_4]$  ( $ESI^+$ ) through reaction of **1** with  $[(Et_3Si)_2(\mu-H)][B(C_6F_5)_4]$  in  $C_6H_5F$ , followed by crystallisation from  $C_6H_5F$ /mesitylene. Again co-crystallisation with pyrimidinium salt  $\{[(6-MesH)\cdots C_6H_3Me_3][B(C_6F_5)_4]\}$  was found.

#### Synthesis and characterisation of $[Ni(6-Mes)(PPh_3)(CO)]^+$

Exposure of a THF solution of **2** to 1 atm CO led to an almost instantaneous colour change from yellow to pale green. Removal of the CO atmosphere after *ca.* 1 min, followed by recrystallization of the residue from THF/hexane gave light green/yellow crystals of the Ni(I) carbonyl complex,  $[Ni(6-Mes)(PPh_3)(CO)][PF_6]$  (**5**- $[PF_6]$ ). The EPR spectrum of **5**- $[PF_6]$  (Fig. 2b) and the corresponding DFT calculations of the spin Hamiltonian parameters is discussed in further detail below.

The X-ray crystal structure (Fig. 5) revealed a similarly distorted T-shaped geometry to that of **2** ( $\angle C-Ni-P = 151.93(9)^\circ$ ). Both the Ni–CO bond length (Ni1–C23 =  $1.787(3) \text{ \AA}$ ) as well as

the  $\nu(CO)$  in the IR spectrum ( $2032 \text{ cm}^{-1}$ ) showed good agreement with the few other (predominantly neutral) reported Ni(I)–CO complexes.<sup>20–23</sup> As expected, **5**- $[PF_6]$  displayed broad, paramagnetic  $^1H$  and  $^{13}C$  NMR spectra, although upon subjecting a THF solution to 1 atm  $^{13}CO$ , we observed the rapid appearance of an isotopically enhanced carbonyl resonance in the  $^{13}C$  NMR spectrum at  $\delta$  198.1, consistent with reversible coordination of the carbonyl ligand.<sup>22</sup> Since the SOMO is an orbital with antibonding character between the Ni centre and the CO ligand, one would expect a weakened Ni–CO bond, and it may be this that facilitates the facile exchange with  $^{13}CO$ .<sup>22,23</sup>

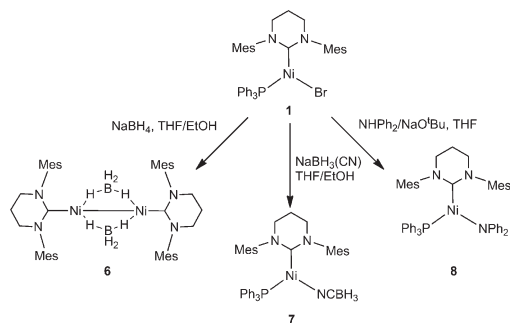
As aforementioned, the presence of residual TlBr in samples of **2** was apparent from reactions with CO, particularly after prolonged periods. At times  $>1$  min, the initial pale green solution of **5**- $[PF_6]$  became orange, and then over *ca.* 48 h, yellow. A small number of crystals were isolated from this yellow solution and structurally characterised as the Ni(II) salt,  $[Ni(6-Mes)(PPh_3)(CO)Br][PF_6]$  ( $ESI^+$ ).

#### Reactions of **1** with $NaBH_3X$ ( $X = H, CN$ ) and $NaNPh_2$

Efforts to abstract the bromide ligand from **1** using sodium salts of more coordinating anions generated the Ni(I) products **6**–**8** shown in Scheme 2.

$NaBH_4$  addition to a yellow THF suspension of **1** in the presence of EtOH rapidly generated a green solution, from which dark green crystals of the dimeric borohydride complex,  $\{[Ni(6-Mes)]_2(\mu-BH_4)_2\}$  (**6**) were isolated in 90% yield. The X-ray crystal structure of **6** is shown in Fig. 6 and revealed asymmetry across the  $\{Ni_2B_2\}$  moiety. Thus, B1 is closer to Ni2 than to





Scheme 2

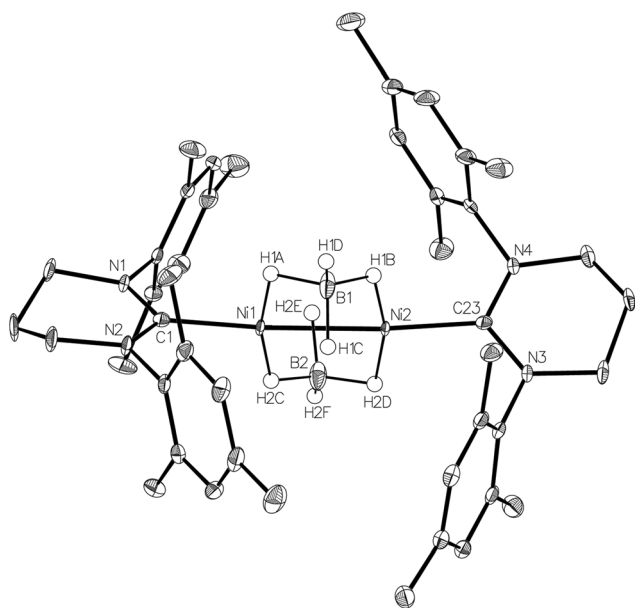


Fig. 6 Molecular structure of compound 6. Ellipsoids are displayed at 50% probability. The minor disordered component of C25 and hydrogen atoms (with the exception of those bonded to boron centres) have been omitted for clarity.

Ni1, and the reverse situation prevails for B2 (Ni1...B1 2.180(2), Ni1...B2 2.143(3), Ni2...B1 2.144(2), Ni2...B2 2.181(3) Å). Overall, the data suggest a rare  $\mu^2, \eta^1: \eta^1$  coordination mode<sup>24,25</sup> for the borohydride based on B1, with H1C being equidistant from both metal centres (Ni1–H1C 2.16(4) Å; Ni2–H1C 2.11(4) Å). The B2 based borohydride has a similar coordination mode once experimental errors are taken into consideration. However, H2E may be closer to Ni1 (1.92(4) Å) than to Ni2 (2.12(4) Å), which would indicate a tendency towards an even more unusual  $\mu^2, \eta^2: \eta^1$  coordination mode.<sup>25</sup> In an effort to further probe the bonding of the borohydrides, a neutron dataset was collected, but a phase transition hampered acquisition of any additional insights (see Experimental). IR spectroscopy provided little in the way of diagnostic characterisation of any particular coordination mode, as only a single,

broad  $\nu(\text{B-H})$  absorption band was measured at  $2378 \text{ cm}^{-1}$  in KBr.

6 adds to the surprisingly few examples of structurally characterised nickel borohydride complexes,<sup>26–28</sup> in amongst which there is just a single example of a bridging borohydride species<sup>29</sup> and a Ni(i)–BH<sub>4</sub> complex.<sup>27,30</sup>

The 2.4218(4) Å separation of the two Ni centres is suggestive of a Ni–Ni bond,<sup>1</sup> which explains the diamagnetism of the complex in solution.<sup>31</sup> The borohydride groups appear fluxional, with just a single broad resonance apparent in both the <sup>1</sup>H (*ca.*  $\delta$  –5.1) and <sup>11</sup>B{<sup>1</sup>H} (*ca.*  $\delta$  –30) NMR spectra. The proton signal sharpened slightly with <sup>11</sup>B decoupling, but remained broad even down to 198 K.

NaBH<sub>4</sub> was replaced by NaBH<sub>3</sub>(CN) in an effort to prepare a Ni–Ni dimer analogous to 6 but with an asymmetric and potentially simpler coordination mode. Instead, the paramagnetic, monomeric Ni(i) complex, [Ni(6-Mes)(PPh<sub>3</sub>)(NCBH<sub>3</sub>)] (7, Fig. 7) was formed. Cyanotrihydroborate complexes remain (like their [BH<sub>4</sub>]<sup>–</sup> counterparts) extremely rare for nickel,<sup>32</sup> and unknown for Ni(i). Trigonal planar 7 exhibited a Ni–N bond length of 1.924(2) Å which, although shorter than that reported in [(tren)Ni( $\mu$ -NCBH<sub>3</sub>)<sub>2</sub>]<sup>2+</sup> (tren = 2,2',2''-tri-aminoethylamine),<sup>32a</sup> is consistent with values reported for a number of monomeric cobalt derivatives.<sup>33</sup> The EPR spectrum of 7 (see ESI†) confirms the paramagnetism of this complex, but is again poorly resolved due to overlapping intensity originating from the precursor complex 1.

We have previously reported that 1 reacts with NaO<sup>t</sup>Bu to provide a low yielding route to the Ni(0) product, [Ni(6-Mes)(PPh<sub>3</sub>)<sub>2</sub>].<sup>34</sup> A repeat of this reaction in the presence of diphenylamine<sup>35</sup> afforded the three-coordinate Ni(i) amido

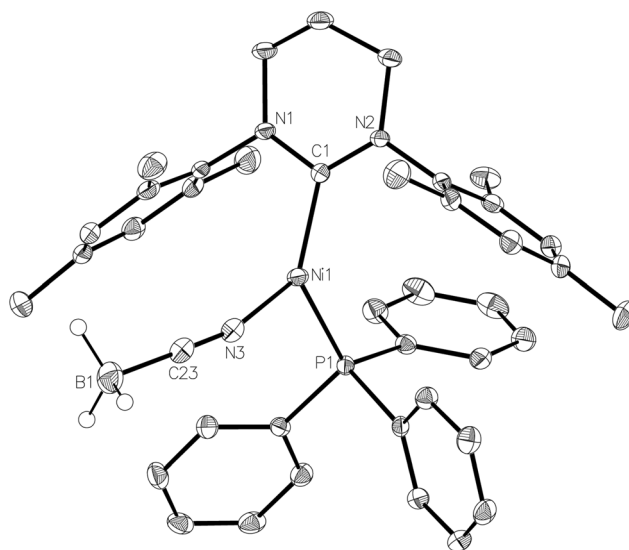
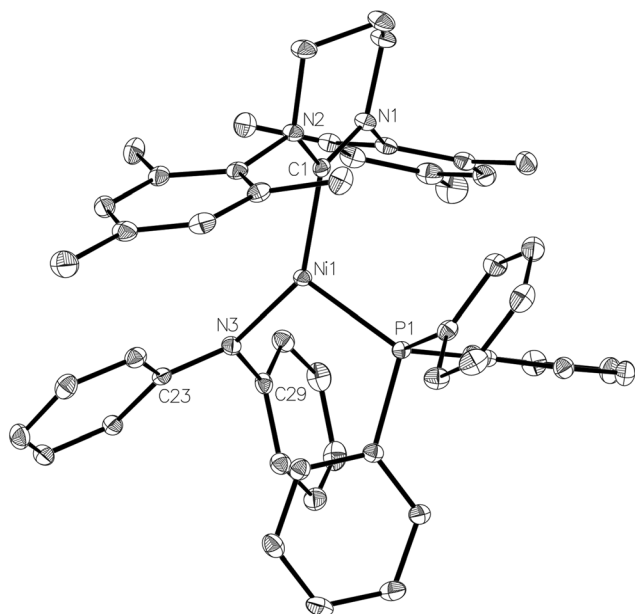


Fig. 7 Molecular structure of compound 7. Ellipsoids are displayed at 30% probability. The minor disordered component of C3 and hydrogen atoms (with the exception of those bonded to boron) have been omitted for clarity. Selected bond lengths (Å) and angles (°): Ni(1)–C(1) 1.957(2), Ni(1)–P(1) 2.2201(6), Ni(1)–N(3) 1.924(2), C(1)–Ni(1)–P(1) 125.14(7), C(1)–Ni(1)–N(3) 132.38(9), P(1)–Ni(1)–N(3) 102.47(7).





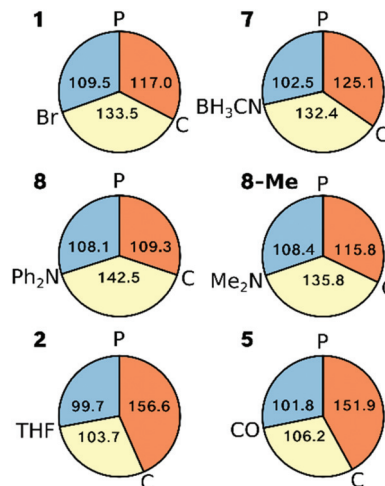
**Fig. 8** Molecular structure of **8**. Ellipsoids are shown at 30% probability. Hydrogen atoms and the minor disordered component of C3 have been omitted for clarity. Selected bond lengths (Å) and angles (°): Ni(1)–C(1) 1.9612(14), Ni(1)–P(1) 2.2337(4), Ni(1)–N(3) 1.9350(12), C(1)–Ni(1)–P(1) 109.25(4), C(1)–Ni(1)–N(3) 142.47(5), P(1)–Ni(1)–N(3) 108.13(4).

complex, [Ni(6-Mes)(PPh<sub>3</sub>)(NPh<sub>2</sub>)] (**8**), as a deep-red solid that could be isolated in high (72%) yield.

The X-ray crystal structure of the complex (Fig. 8) revealed a more acute C–Ni–P angle (109.25(4)°) than that found in the starting bromide complex (117.01(6)°), which most likely results from the need to alleviate steric clashes between the P- and N-bound phenyl groups. Indeed, replacing the phenyl groups with methyl groups *in silico* and fully relaxing the geometry showed that steric effects play a role in shaping the geometry of **8**: in the case of the (hypothetical) [Ni(6-Mes)(PPh<sub>3</sub>)(NMe<sub>2</sub>)] complex **8-Me** with a less bulky NMe<sub>2</sub> ligand, the C–Ni–N angle decreased by 6.7° while the P–Ni–C angle increased by 6.5°. Concomitantly, the Ni–N bond decreased from 1.94 Å in the crystal structure to 1.84 Å in **8-Me** (fully relaxing the crystal structure of **8** results in a bond length of 1.90 Å). Sterics may also account for the non-planarity of the amido group (dihedral angle between Ni1, N3, C23 and C29 of *ca.* 160°), as well as the elongation of the Ni–N distance (1.9350(12) Å) compared to those in either [Ni(d<sup>t</sup>bpe){N(2,6-<sup>i</sup>Pr<sub>2</sub>C<sub>6</sub>H<sub>3</sub>H)}] (1.881(2) Å; d<sup>t</sup>bpe = <sup>t</sup>Bu<sub>2</sub>P(CH<sub>2</sub>)<sub>2</sub>P<sup>t</sup>Bu<sub>2</sub>)<sup>36</sup> or [Ni(PPh<sub>3</sub>)<sub>2</sub>{N(SiMe<sub>3</sub>)<sub>2</sub>}] (1.88(1) Å).<sup>4,37</sup>

### Electronic structure analysis of complexes **1**, **2**, **5**, **7** and **8**

Inspection of the angles around Ni in complexes **1**, **2**, **5**, **7** and **8** (as well as **8-Me**) shows that the L–Ni–P angle varies least across all complexes (<10°, Fig. 9). Complexes **1** and **7** can be classed as Y-shaped (∠C–Ni–L, ∠P–Ni–C > ∠L–Ni–P, Fig. 9), whereas complexes **2** and **5** are T-shaped (∠P–Ni–C > ∠C–Ni–L, ∠L–Ni–P angles, see Fig. 9). Complex **8** represents a more sym-



**Fig. 9** Pie diagrams representing the angles around the Ni ion in the crystal structures for **1**, **2**, **5**, **7**, **8** and the geometry optimised structure of **8-Me**; orange: ∠P–Ni–C, yellow: ∠C–Ni–L, blue: ∠L–Ni–P, where L stands for the respective ligand.

metric Y-shaped case with a larger C–Ni–L angle than in **1** and **7** (142.5° vs. 133.5°, 132.4°) and a smaller P–Ni–C angle (109.2° vs. 117.0°, 125.1°). Although one may also view complex **8** as a T-shaped complex with ∠C–Ni–L as the largest angle, this appears to be purely due to steric and not electronic effects, as is seen by **8-Me** adopting a more Y-shaped geometry.

The overall geometric changes in fully relaxing the crystal structures are small (see ESI†). Most importantly, the striking consistency of the Ni–P and Ni–C bond lengths in the crystal structures across the series (variation <0.03 Å and <0.02 Å, respectively) is preserved upon geometry optimisation (variation <0.03 Å and <0.04 Å, respectively; ESI†).

The tendency of three-coordinate transition metal d<sup>9</sup> complexes to form either T- or Y-shaped geometries is due to the Jahn–Teller effect, thus lifting orbital degeneracy (d<sub>xy</sub>, d<sub>x<sup>2</sup>–y<sup>2</sup></sub>) at the ideal D<sub>3h</sub> symmetry (Fig. 10a). MO theory predicts that the SOMO in a T-shaped d<sup>9</sup> complex will be of d<sub>x<sup>2</sup>–y<sup>2</sup></sub> character, whereas in a Y-shaped d<sup>9</sup> complex, it will be of d<sub>xy</sub> character (Fig. 10a), in agreement with the dominant character of the DFT-calculated orbitals (Fig. 10b). For Ni(I) complexes, this was most recently discussed by the groups of Holland and Lee,<sup>13,21</sup> and prior to that, by Pietrzyk.<sup>38</sup> Holland and co-workers<sup>21</sup> rationalised the formation of T- vs. Y-shaped complexes with a charge donation analysis (natural bond orbital analysis).<sup>39</sup> Their findings indicated that a T-shape is inherently favoured by d<sup>9</sup> complexes, but a Y-shape can result when there is increased donation of charge from the ligands to the metal centre, thus effectively partially reducing the metal centre. In the present case, the analysis of Mulliken<sup>40</sup> and Chelpg<sup>41</sup> charges did not reveal a clear connection between charges and geometry. Likewise, the Mulliken spin populations on the Ni ion do not show a significant variation across the series (ESI†).

The CW X-band EPR spectra of complexes **1**, **2**, **5** and **8** were shown in Fig. 2. The resulting spin Hamiltonian parameters,





$A_3(^{31}\text{P})$  component is found on  $g_1$  because of almost exact and complete alignment between the  $A_z$  axis and the  $g_x$  axis, as deducible by Fig. 11. The extent of anisotropy and rhombicity for **5** is much lower than for the other complexes, and is similar to that reported previously for  $[\text{L}^{\text{Me}}\text{Ni}(\text{CO})]$  ( $\text{L}^{\text{Me}} = [\text{HC}(\text{C}(\text{Me})\text{NC}_6\text{H}_3(\text{iPr})_2)_2]^-$ ),<sup>21</sup> with  $g$  values of 2.01, 2.17 and 2.19 and T-shape geometry.

Notably, the calculated  $^{31}\text{P}$  superhyperfine coupling for the two cationic complexes (**2** and **5**) are an order of magnitude smaller than for the series of neutral complexes reported. Whilst the experimental and calculated values for **5** are in reasonable agreement, the experimentally observed hyperfine for **2** does not match the DFT-derived values and bears closer resemblance to the neutral complexes; currently, we do not have an explanation for this observation.

Fig. 12 shows the computed SOMO d-character of the different complexes as a function of the difference between the largest and the two smaller angles (e.g.  $(\angle\text{C-Ni-Br}) - (\angle\text{P-Ni-C}) - (\angle\text{Br-Ni-P})$  for **1**, denoted as  $\Delta\Delta(\text{bond angle})$ ), taken as an index for the deviation from ideal  $D_{3h}$  symmetry. Noticeable in Fig. 12a is the smaller difference in  $d_{xy}$  and  $d_{x^2-y^2}$  contributions to the SOMO for complexes **1** and **7** (closest examples to  $D_{3h}$  symmetry where  $d_{xy}$  and  $d_{x^2-y^2}$  are degenerate), and the increasingly higher  $d_{x^2-y^2}$  character (simultaneously to  $d_{xy}$  contributions approaching zero) when moving away from ideal  $D_{3h}$  symmetry towards T-shape symmetry. Both these observations seem to be in good agreement with what was described

previously and represented in Fig. 9. Very interesting is the case of complex **8**, which as we noted above may be regarded as a T-shape complex with  $\angle\text{C-Ni-L}$  as the largest angle. However, a fully geometry optimised version of the same complex where the amido phenyl substituents were replaced by methyl substituents (**8-Me**, *vide supra*) showed angles that are similar to the Y-shape complexes **1** and **7**. Our interpretation is that this compound is *electronically* inclined to be a Y-shape (similar to the other neutral compounds of the present series), however large steric strain pushes the amido group towards the carbene ligand, thus *geometrically* distorting it towards a T-shape. Orbital distribution and coordination geometry should reflect the shape and magnitude of the  $\mathbf{g}$  tensor associated with the paramagnetic centre. In Fig. 12b, experimental  $\Delta g_{\text{rel}}$ , a parameter used to evaluate the shape of the diagonalised  $\mathbf{g}$  tensor and calculated according to eqn (1), is also reported as a function of the  $\Delta\Delta(\text{bond angle})$ .

$$\Delta g_{\text{rel}} = \frac{|g_3 - g_2|}{|g_3 - g_1|} \times 100 \quad (1)$$

It can be seen that an increase in the  $d_{x^2-y^2}$  contribution to the SOMO corresponds to a shift of the  $g_2$  value away from  $g_3$  towards  $g_1$ , and indeed for T-shape complexes  $g_2$  is closer to  $g_1$  than to  $g_3$ , highlighting a geometry induced shape shifting of the  $\mathbf{g}$  tensor.

## Conclusions

Treatment of the three-coordinate Ni(I) complex  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)\text{Br}]$  (**1**) with a variety of bromide abstracting reagents has yielded a series of new mono- and dinuclear nickel products. Of most interest are the three-coordinate  $d^9$  complexes, **2**, **5**, **7** and **8**, of general formula  $[\text{Ni}(\text{6-Mes})(\text{PPh}_3)\text{X}]^{0/+}$  that distort from ideal  $D_{3h}$  symmetry by forming either T-shaped or Y-shaped geometries. These structural differences manifest in different electronic structure characteristics, namely that the SOMO for a T-shape complex is expected to be of  $d_{x^2-y^2}$  character, whereas for a Y-shape complex, it will be of  $d_{xy}$  character. Electron paramagnetic resonance spectroscopy was used to derive spin Hamiltonian parameters for this series of three-coordinate Ni(I) complexes, which showed that all complexes have a rhombic  $\mathbf{g}$ -tensor profile and that the  $^{31}\text{P}$  superhyperfine couplings are predominantly isotropic. The much lower magnitude of  $^{31}\text{P}$  superhyperfine coupling constants observed for the CO-containing complex **5** was explained with a smaller spin density found at the phosphorus ligand as predicted by density functional theory calculations. The overall computed spin densities in this series are polarised differently for the Y- and T-shaped complexes, namely with a larger lobe *trans* to the phosphine ligand in the former case as opposed to a larger lobe *trans* to the ligand in the latter case. This directly affects the shape and magnitude of the  $\mathbf{g}$ -tensor: while all complexes have a rhombic  $\mathbf{g}$ -tensor with  $g_1 \ll g_2 < g_3$ , a larger  $d_{x^2-y^2}$  contribution to the SOMO shifts  $g_2$  closer to  $g_1$ .

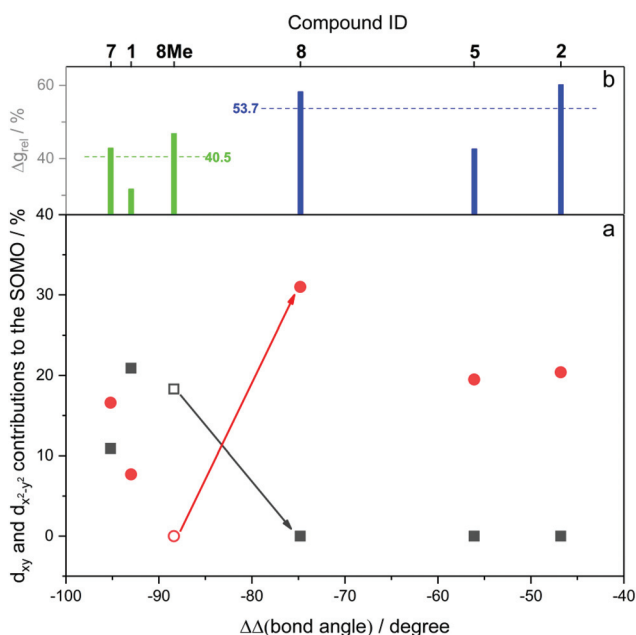


Fig. 12 (a)  $d_{xy}$  and  $d_{x^2-y^2}$  contributions to the spin up SOMO orbital as a function of the difference between the largest and the two smaller angles ( $\Delta\Delta(\text{bond angle})$ ). Black squares refer to  $d_{xy}$  and red circles to  $d_{x^2-y^2}$ . Empty symbols refers to d orbital contributions from fully optimised structures rather than X-ray crystal structures. (b)  $\Delta g_{\text{rel}}$  values as a function of the double bond angle variation. Average  $\Delta g_{\text{rel}}$  values for Y- and T-shape geometries are also reported.







## X-ray crystallography

Using Mo(K $\alpha$ ) radiation, single crystals of compounds **2**, **3** and **7** were analysed using an Agilent Xcalibur diffractometer, while datasets for **4** and **5**, as well as **2a** (*vide infra*; ESI $\dagger$  only) were collected on a Nonius kappaCCD machine. An Agilent Supernova diffractometer was used to study **6**, **8** and [Ni(6-Mes)( $\eta^2$ -C<sub>6</sub>H<sub>3</sub>Me<sub>3</sub>)]B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub> (ESI $\dagger$  only) using Cu(K $\alpha$ ) radiation. All experiments were conducted at 150 K, with the exception of that for **6**, which was achieved at 100 K. Details of the data collections and refinements are given in Table 2. The structures were uniformly solved using SHELXS,<sup>42</sup> and refined using full-matrix least squares in SHELXL<sup>43</sup> *via* the Olex-2<sup>44</sup> software suite. Only noteworthy refinement details follow.

A small amount of racemic twinning was accounted for in the refinement of **2a** (ESI $\dagger$  only). This structure represents a *P*<sub>2</sub><sub>1</sub> polymorph of compound **2**, the latter solving in space group *P*<sub>2</sub><sub>1</sub>/*c*. In **3**, the asymmetric was seen to contain one anion, one cation and one molecule of diethyl ether. While the cation and solvent were both ordered, disorder prevailed for four of the [Bar<sub>4</sub><sup>F</sup>]<sup>−</sup> trifluoromethyl substituents. In particular, the fluorine atoms attached to C88, C111 and C103 were each modelled over two sites in 65 : 35, 55 : 45 and 65 : 35 ratios, respectively, while the entire CF<sub>3</sub> group containing C87 exhibited 65 : 35 disorder. In **3** (and all subsequent structures containing disordered [Bar<sub>4</sub><sup>F</sup>]<sup>−</sup> trifluoromethyl groups) C–F and F...F distances within each disordered region were restrained to being similar in the final least squares. In addition, the ADPs for fractional occupancy atoms were also restrained, to assist convergence.

The cation in the asymmetric unit of **4** also fell prey to disorder. In particular, there is a 50 : 50 ratio of the tolyl-Ni-carbene moiety present *versus* the tolyl...pyrimidinium pair, the latter being stabilised by a C–H... $\pi$  interaction. In **5**, the asymmetric unit was seen to comprise one cationic nickel containing species, one [PF<sub>6</sub>]<sup>−</sup> anion and one THF molecule. The crystal was small, which contributed to weak diffraction at higher Bragg angles. Hence, data were truncated to a  $\theta$  value of 24.7°.

The borohydride hydrogen atoms in the structure of compound **6** were readily located and refined with a common *U*<sub>iso</sub> in each [BH<sub>4</sub>]<sup>−</sup> moiety. No distance restraints employed. C25 was modelled for 87 : 13 disorder, and the minor component of this atom was refined isotropically. A data collection was also performed on this compound, at room temperature (designated **6a**, ESI $\dagger$ ), in which the asymmetric unit was seen to consist of one half of a dimer molecule, wherein the metal centres and carbene carbon atoms were noted to coincide with a crystallographic 2-fold rotation axis. This necessarily means that the apical NHC carbons (C3 and C15) are each disordered in a 50 : 50 ratio. This disorder precluded addition of the hydrogen atoms bound to C2 and C14 using the riding model; hence, they were omitted from the refinement. The borohydride hydrogens were located, and refined without restraints, but their credibility is somewhat questionable given their associated *U*<sub>iso</sub> values and the overall atomic displace-

ment parameters. The reason for implementing a room temperature data collection for **6a** was to resolve a phase transition that arose in the course of a neutron experiment conducted on **6**, using VIVALDI, at the ILL. The rationale for doing a neutron experiment arose because, at 100 K, the borohydride moieties appeared to coordinate unsymmetrically to the nickel centres. Unfortunately, during cooling at the neutron source, the large crystals cracked. This ultimately resulted in collection of a neutron data set at room temperature, which suggested a different space group (*C*<sub>2</sub>/*c*) to that for the structure determined at 100 K using X-rays (*P*<sub>2</sub><sub>1</sub>/*c*).

This phase transition, from a diffraction perspective, results in averaging the electron density that arises from the borohydrides across the sample and, overall, the ambient temperature neutron data did not afford any additional insight into the bonding subtleties which the experiment aimed to probe.

The asymmetric unit in **7** was seen host to one molecule of THF in addition to one molecule of the nickel complex. C3 in the latter was equally disordered over two sites, and the four chemically equivalent C–C distances involving C3/C3A were restrained to being similar in the final least squares. Three of the five atoms in the solvent were also refined to take account of 75 : 25 disorder. Once again, the chemically equivalent distances involving fractional occupancy atoms in this moiety were restrained to being similar, and ADP restraints were also incorporated to assist convergence.

In addition to one molecule of the complex, the asymmetric unit in **8** was noted to contain one molecule of guest diethyl ether.

Analysing the crystal structure of [Ni(6-Mes)( $\eta^2$ -C<sub>6</sub>H<sub>3</sub>Me<sub>3</sub>)]B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub> (ESI $\dagger$  only) was nothing short of excruciating, and it involved three data collections, some 350 refinements and approximately 10 data integrations. The issue appears to be that the compound is undergoing a phase transition. A more detailed treatment is presented in the ESI $\dagger$ .

Crystallographic data for all compounds have been deposited with the Cambridge Crystallographic Data Centre as supplementary publications CCDC 1578636–1578644, 1582301 and 1584193 for **2**, **3**, **4**, **5**, **6**, **7**, **8**, **2a** (ESI $\dagger$ ), [Ni(6-Mes)( $\eta^2$ -C<sub>6</sub>H<sub>3</sub>Me<sub>3</sub>)]B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub> (ESI $\dagger$ ), **6a** (ESI $\dagger$ ) and [Ni(6-Mes)(PPh<sub>3</sub>)(CO)Br][PF<sub>6</sub>] (ESI $\dagger$ ) respectively.

## EPR spectroscopy

Samples for EPR measurements were prepared under an N<sub>2</sub> atmosphere in a glovebox. A solution of each complex was prepared by dissolving *ca.* 4 mg of **1–3**, **5**, **7** and **8** in 200  $\mu$ L of dry THF (in all cases, a small quantity of dry toluene was also added to improve the quality of the polycrystalline glass formed in frozen solution, and thereby enhance the quality of the EPR spectra). The solutions were transferred to an EPR tube, sealed in the glove box and then cooled to 77 K before rapid transfer to the pre-cooled EPR cavity. The X-band CW EPR measurements were performed on a Bruker EMX spectrometer utilizing an ER4119HS resonator, 100 kHz field modulation at 140 K.





**Table 2** Crystal data and structural refinement details for compounds 2–8

	2	3	4	5	6	7	8
Empirical formula	C <sub>44</sub> H <sub>51</sub> F <sub>6</sub> N <sub>2</sub> NiOP <sub>2</sub>	C <sub>116</sub> H <sub>108</sub> BBrF <sub>24</sub> N <sub>4</sub> Ni <sub>2</sub> OP <sub>2</sub>	C <sub>53</sub> H <sub>36.5</sub> BF <sub>20</sub> N <sub>2</sub> Ni <sub>0.5</sub>	C <sub>45</sub> H <sub>51</sub> F <sub>6</sub> N <sub>2</sub> NiO <sub>2</sub> P <sub>2</sub>	C <sub>44</sub> H <sub>64</sub> B <sub>2</sub> N <sub>4</sub> Ni <sub>2</sub>	C <sub>45</sub> H <sub>54</sub> BN <sub>3</sub> NiOP	C <sub>56</sub> H <sub>63</sub> N <sub>3</sub> NiOP
Formula weight	858.52	2300.14	1121.50	886.53	788.03	753.40	883.77
Crystal system	Monoclinic	Triclinic	Monoclinic	Monoclinic	Monoclinic	Monoclinic	Monoclinic
Space group	<i>P</i> 2 <sub>1</sub> / <i>c</i>	<i>P</i> 1	<i>C</i> 2/ <i>c</i>	<i>P</i> 2 <sub>1</sub> / <i>c</i>	<i>P</i> 2 <sub>1</sub> / <i>c</i>	<i>P</i> 2 <sub>1</sub> / <i>n</i>	<i>P</i> 2 <sub>1</sub> / <i>n</i>
<i>a</i> /Å	16.0749(1)	12.9050(4)	31.5490(3)	14.3080(3)	14.17049(12)	14.2120(4)	18.99247(14)
<i>b</i> /Å	14.8045(1)	17.3278(5)	10.32800(10)	16.4910(3)	20.24524(15)	16.5650(4)	11.96944(9)
<i>c</i> /Å	19.0391(1)	25.2732(6)	28.8100(3)	18.8500(4)	14.83697(15)	18.0386(6)	20.99341(16)
<i>α</i> /°	90	75.082(2)	90	90	90	90	90
<i>β</i> /°	111.435(1)	84.432(2)	92.977(1)	95.145(1)	91.0848(8)	108.838(3)	91.1551(7)
<i>γ</i> /°	90	87.444(2)	90	90	90	90	90
<i>U</i> /Å <sup>3</sup>	4217.55(5)	5434.1(3)	9374.73(16)	4429.80(15)	4255.74(6)	4019.2(2)	4771.45(6)
<i>Z</i>	4	2	8	4	4	4	4
<i>ρ</i> <sub>calc</sub> /g cm <sup>-3</sup>	1.352	1.406	1.589	1.329	1.230	1.245	1.230
<i>μ</i> /mm <sup>-1</sup>	0.598	0.836	0.341	0.574	1.343	0.560	1.219
<i>F</i> (000)	1796.0	2364.0	4540.0	1852.0	1688.0	1604.0	1884.0
Crystal size/mm <sup>3</sup>	0.36 × 0.31 × 0.19	0.845 × 0.77 × 0.563	0.2 × 0.1 × 0.1	0.2 × 0.1 × 0.1	0.25 × 0.25 × 0.25	0.621 × 0.378 × 0.062	0.236 × 0.157 × 0.048
2θ range for data collection/°	5.7 to 54.97	6.908 to 54.968	7.078 to 54.872	7.294 to 49.404	15.77 to 144.026	6.814 to 54.968	6.214 to 146.89
Index ranges	−20 ≤ <i>h</i> ≤ 20 −19 ≤ <i>k</i> ≤ 19 −24 ≤ <i>l</i> ≤ 24	−16 ≤ <i>h</i> ≤ 16 −22 ≤ <i>k</i> ≤ 17 −32 ≤ <i>l</i> ≤ 32	−40 ≤ <i>h</i> ≤ 40 −13 ≤ <i>k</i> ≤ 13 −37 ≤ <i>l</i> ≤ 37	−16 ≤ <i>h</i> ≤ 16 −19 ≤ <i>k</i> ≤ 19 −22 ≤ <i>l</i> ≤ 22	−17 ≤ <i>h</i> ≤ 15 −24 ≤ <i>k</i> ≤ 19 −18 ≤ <i>l</i> ≤ 18	−13 ≤ <i>h</i> ≤ 18 −21 ≤ <i>k</i> ≤ 20 −23 ≤ <i>l</i> ≤ 22	−23 ≤ <i>h</i> ≤ 22 −14 ≤ <i>k</i> ≤ 14 −24 ≤ <i>l</i> ≤ 26
Reflections collected	95 090	46 948	75 915	41 528	59 289	38 736	67 012
Independent reflections, <i>R</i> <sub>int</sub>	9656, 0.0282	24 033, 0.0376	10 665, 0.0610	7457, 0.0463	8330, 0.0847	9218, 0.0347	9580, 0.0510
Data/restraints/parameters	9656/0/511	24 033/121/1365	10 665/0/701	7457/128/565	8330/0/511	9218/25/502	9580/0/567
Goodness-of-fit on <i>F</i> <sup>2</sup>	1.089	1.023	1.122	1.053	1.017	1.028	1.027
Final <i>R</i> <sub>1</sub> , w <i>R</i> <sub>2</sub> [ <i>I</i> > 2σ( <i>I</i> )]	0.0284, 0.0809	0.0635, 0.1396	0.0503, 0.1086	0.0435, 0.1070	0.0528, 0.1401	0.0472, 0.1128	0.0343, 0.0828
Final <i>R</i> <sub>1</sub> , w <i>R</i> <sub>2</sub> [all data]	0.0370, 0.0830	0.1297, 0.1745	0.0852, 0.1192	0.0555, 0.1154	0.0580, 0.1457	0.0703, 0.1267	0.0400, 0.0860
Largest diff. peak/hole/e Å <sup>-3</sup>	0.60/−0.36	1.19/−0.93	0.51/−0.30	0.52/−0.48	0.68/−0.64	0.88/−0.94	0.39/−0.29

## Computational details

All density functional theory (DFT) calculations were carried out with ORCA (version 4.0.0.2).<sup>45</sup> The geometries were taken from crystallographic refinements, either optimising only the positions of the hydrogen atoms or fully relaxing the geometry. The geometry optimisations used the BP86 density functional,<sup>46</sup> making use of the zeroth order relativistic correction ZORA retaining one-center terms.<sup>47</sup> The scalar-relativistically recontracted versions of Ahlrich's triple-zeta quality basis sets (ZORA-def2-TZVP) were used on all atoms except carbon and hydrogen for which ZORA-def2-SVP basis sets were used.<sup>48</sup> The resolution of the identity (RI) approximation and the auxiliary basis SARC/J were used.<sup>49</sup> The integration accuracy was increased to 7.0, the grid was set to 7 in ORCA nomenclature, and 'tight' SCF criteria were used. The optimisations considered solvent effects through the conductor-like polarisable continuum model, with the solvents as indicated in the experimental part.<sup>50</sup> Dispersion effects were taken into account with Grimme's D3BJ model including Becke–Johnson damping.<sup>51</sup> Mulliken spin populations were inspected to confirm convergence to the targeted electronic structure.

Broken-symmetry DFT calculations used the functionals TPSSH,<sup>52</sup> B3LYP,<sup>53</sup> PBE0,<sup>54</sup> M06L,<sup>55</sup> additionally making use of the chain-of-spheres approximation (RIJCOSX) and using the 'flipspin' feature in ORCA to generate the initial guess for the broken-symmetry solution, with otherwise unchanged calculation setups.<sup>56</sup> The exchange coupling constants were taken directly from the ORCA output, using the definition by Yamaguchi.<sup>57</sup> For the calculation of EPR parameters, it was found that calculations with a different family of basis sets gave superior results. Generally, the IGLO-II basis set was used on all atoms, with CP for Ni and aug-pc-3 for Br,<sup>58</sup> in conjunction with the PBE0 density functional and the RIJCOSX approximation as for the BS-DFT calculations, making use of the AutoAux feature in ORCA. The grid sizes were set to Grid6 and GridX9 in ORCA nomenclature, with increased grids (7) on the Ni ion and all directly bound atoms as well as the nitrogen atoms in the carbene ligand. The spin-orbit mean field operator (SOMF(1X)) was used, and the origin for the g-tensor was taken at the centre of the electronic charge.<sup>59</sup> All tensor orientations, spin densities and molecular orbitals depicted and discussed in the main text and the ESI† are derived from calculations at this level of theory.

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

We thank the Universities of Bath (studentships for WJMB and RCP; Prize Fellowship to VK) and Cardiff (Cardiff University Research Fellowship to ER), EPSRC (MJP, AF) for financial support and the Royal Society for a Newton International

Fellowship for SS. We acknowledge the Institute Laue-Langevin Scientific Coordination Office for the allocation of neutron beam time on VIVALDI (proposal 5-12-276) and Dr Marie-Hélène Lemée-Cailleau for the data collection on 6. We dedicate this paper to Professor Phil Power on the occasion of his 65<sup>th</sup> birthday.

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