



Cite this: *RSC Adv.*, 2018, 8, 32672

# Experimental and computational preference for phosphine regioselectivity and stereoselective tripodal rotation in $\text{HOs}_3(\text{CO})_8(\text{PPh}_3)_2(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})^\dagger$

Shahin A. Begum,<sup>a</sup> Md. Arshad H. Chowdhury,<sup>a</sup> Shishir Ghosh,<sup>id</sup><sup>a</sup>  
 Derek A. Tocher,<sup>id</sup><sup>b</sup> Michael G. Richmond,<sup>id</sup><sup>c</sup> Li Yang,<sup>c</sup> K. I. Hardcastle,<sup>d</sup>  
 Edward Rosenberg<sup>e</sup> and Shariff E. Kabir<sup>\*a</sup>

The site preference for ligand substitution in the benzothiazolate-bridged cluster  $\text{HOs}_3(\text{CO})_{10}(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})$  (**1**) has been investigated using  $\text{PPh}_3$ . **1** reacts with  $\text{PPh}_3$  in the presence of  $\text{Me}_3\text{NO}$  to afford the mono- and bisphosphine substituted clusters  $\text{HOs}_3(\text{CO})_9(\text{PPh}_3)(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})$  (**2**) and  $\text{HOs}_3(\text{CO})_8(\text{PPh}_3)_2(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})$  (**3**), respectively. **2** exists as a pair of non-interconverting isomers where the  $\text{PPh}_3$  ligand is situated at one of the equatorial sites syn to the edge-bridging hydride that shares a common Os–Os bond with the metalated heterocycle. The solid-state structure of the major isomer establishes the  $\text{PPh}_3$  regiochemistry at the N-substituted osmium center. DFT calculations confirm the thermodynamic preference for this particular isomer relative to the minor isomer whose phosphine ligand is located at the adjacent C-metalated osmium center. **2** also reacts with  $\text{PPh}_3$  to give **3**. The locus of the second substitution occurs at one of the two equatorial sites at the  $\text{Os}(\text{CO})_4$  moiety in **2** and gives rise to a pair of fluxional stereoisomers where the new phosphine ligand is scrambled between the two equatorial sites at the  $\text{Os}(\text{CO})_3\text{P}$  moiety. The molecular structure of the major isomer has been determined by X-ray diffraction analysis and found to represent the lowest energy structure of the different stereoisomers computed for  $\text{HOs}_3(\text{CO})_8(\text{PPh}_3)_2(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})$ . The fluxional behavior displayed by **3** has been examined by VT NMR spectroscopy, and DFT calculations provide evidence for stereoselective tripodal rotation at the  $\text{Os}(\text{CO})_3\text{P}$  moiety that serves to equilibrate the second phosphine between the two available equatorial sites.

Received 5th September 2018  
 Accepted 12th September 2018

DOI: 10.1039/c8ra07400c

rsc.li/rsc-advances

## Introduction

The activation and value-added functionalization of benzoannulated heterocycles containing a pyridyl nitrogen by low-valent trimetallic clusters,  $\text{M}_3(\text{CO})_{12-n}(\text{NCMe})_n$  ( $\text{M} = \text{Os}, \text{Ru}; n = 0, 1, 2$ ), have been extensively studied over the past few decades.<sup>1–11</sup> While the early impetus involved industrial considerations with respect to hydrotreatment of nitrogen-contaminated fuel

stocks,<sup>1,3,12,13</sup> current interests remain focused on the ability to use the cluster as a templating agent to help guide the selective functionalization of the bound benzoheterocycle.<sup>6,7</sup> Recently, we have also reported our data on the biological applications of such complexes and their ability to serve as potential bio-organometallic conjugates in enzyme and telomerase inhibition studies, as well as tagging agents for use in protein crystallography.<sup>2,9</sup>

The ligation of benzoheterocycles at trimetallic clusters is readily achieved *via* the direct reaction of a suitably activated cluster with the corresponding benzoheterocycle. For example, the labile cluster  $1,2\text{-Os}_3(\text{CO})_{10}(\text{NCMe})_2$  reacts with quinoline ( $\text{C}_9\text{H}_7\text{N}$ ) to afford isomeric products with the formula  $\text{HOs}_3(\text{CO})_{10}(\mu\text{-}\eta^1, \kappa^1\text{-C}_9\text{H}_6\text{N})$  which differ in the coordination mode adopted by the activated heterocycle.<sup>6a</sup> Here the quinolate residue bridges one of the three metal–metal edges *via* the pyridyl nitrogen and either a metalated aryl C-8 or C-2 carbon atom (Scheme 1). The 1,8 isomer has been extensively examined by us and it undergoes CO loss during controlled thermolysis or photolysis to give the face-capped cluster  $\text{HOs}_3(\text{CO})_9(\mu_3\text{-}\eta^1, \kappa^1\text{-}$

<sup>a</sup>Department of Chemistry, Jahangirnagar University, Savar, Dhaka-1342, Bangladesh. E-mail: skabir\_ju@yahoo.com; Tel: +880-27791099

<sup>b</sup>Department of Chemistry, University College London, 20 Gordon Street, London WC1H 0AJ, UK

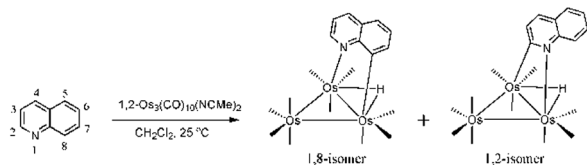
<sup>c</sup>Department of Chemistry, University of North Texas, Denton, TX 76209, USA

<sup>d</sup>Department of Chemistry, Emory University, Atlanta, GA 30322, USA

<sup>e</sup>Department of Chemistry, University of Montana, Missoula, MT 59812, USA

† Electronic supplementary information (ESI) available: Atomic coordinates of all DFT-optimized stationary points, ChemDraw pictures, and spectroscopic data. CCDC 1833188 and 1833187 contain the supplementary crystallographic data for compounds **2** and **3**, respectively. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8ra07400c



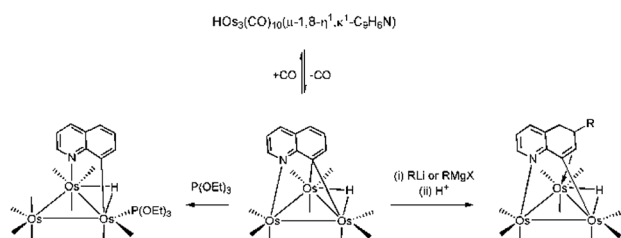


**Scheme 1** Reaction of quinoline with 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> to give the 1,8- and 1,2-metallated isomers of HOs<sub>3</sub>(CO)<sub>10</sub>(μ-η<sup>1</sup>,κ<sup>1</sup>-C<sub>9</sub>H<sub>6</sub>N).

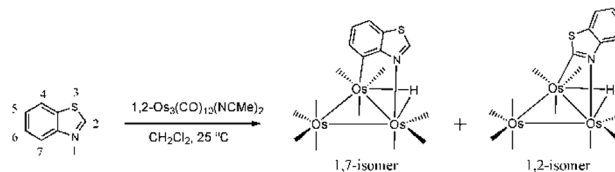
C<sub>9</sub>H<sub>6</sub>N). The nonacarbonyl product reacts with soft pnictogen nucleophiles at an osmium center to yield HOs<sub>3</sub>(CO)<sub>9</sub>P(μ-η<sup>1</sup>,κ<sup>1</sup>-C<sub>9</sub>H<sub>6</sub>N) (where P = phosphine, phosphite) and with hard nucleophiles (RLi and RMgX) at the heterocycle to give HOs<sub>3</sub>(CO)<sub>10</sub>(μ-η<sup>1</sup>,κ<sup>1</sup>-RC<sub>9</sub>H<sub>7</sub>N).<sup>4a,6</sup> The latter product, which may be isolated after protonation, provides the basis for many of our reported heterocyclic functionalizations (Scheme 2).<sup>6c</sup> In comparison to numerous studies involving the 1,8 isomer, the corresponding 1,2 isomer has received relatively little attention.<sup>14</sup>

Metal cluster activation of heterocycles such as benzothiazole is accompanied by added complexity in the form of potential sulfur *versus* nitrogen coordination. We have demonstrated that benzothiazole reacts with 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(NCMe)<sub>2</sub> to furnish the metallated products HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,7-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) and HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) where the metallation occurs with coordination of the nitrogen and carbon atoms (Scheme 3).<sup>6c,15</sup> The benzothiazolate-substituted clusters are formed in comparable yields and may be isolated by column chromatography, albeit with material loss in the case of the 1,2 derivative. This complication in the isolation of the 1,2 isomer, coupled with its limited thermal stability *vis-à-vis* the 1,7 isomer, has hampered meaningful reactivity studies designed to exploit this cluster-activated heterocycle. Systematic reactivity studies are important if we are to gain a better understanding of such systems, especially in light of our earlier observations that have demonstrated that the outcome of such functionalizations is known to be sensitive to the presence of ring substituents and the ligand environment at the metal cluster.

Given the significant influence the heterocyclic auxiliary has on the reactivity in this genre of metal clusters, we have initiated reactivity studies on HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) that will allow us to better understand the observed chemistry as a function of the metallated heterocycle and its control on ligand



**Scheme 2** Decarbonylation in HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,8-C<sub>9</sub>H<sub>6</sub>N) and the different reaction pathways exhibited by HOs<sub>3</sub>(CO)<sub>9</sub>(μ<sub>3</sub>-η<sup>1</sup>,κ<sup>1</sup>-C<sub>9</sub>H<sub>6</sub>N).



**Scheme 3** Reaction of benzothiazole with 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> to give the 1,7- and 1,2-N,C metallated isomers of HOs<sub>3</sub>(CO)<sub>10</sub>(μ-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS).

substitution and value-added functionalizations. The cluster HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,7-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS), whose benzothiazolate bridging ligand has a larger backbone, has received greater attention than its smaller ring orthometalated analogue. Accordingly, we wished to study the reactivity of HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) with added ligand in order to establish the regiochemistry and thermodynamic stability of the resulting products. The effect of the 1,7-N,C ring in HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,7-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) in ligand addition and alkyne insertion reactions has recently been reported by us.<sup>5c,e,16</sup> To our knowledge, only one reactivity report exists for the 1,2-N,C isomer to date and this involves the cyclometalation of thiourea at cluster 1.<sup>17</sup> Herein we present our data on the regiochemical preference for PPh<sub>3</sub> substitution in cluster 1, along with the results of electronic structure calculations that help place the observed outcome in each substitution step in context with the NMR properties and the solid-state structures determined for 2 and 3.

## Results and discussion

### Metalation selectivity in the reaction of benzothiazole with 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub>

Wishing to reconcile the thermodynamic stability of the 1,2- and 1,7-N,C metallated isomers from the reaction of benzothiazole with 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> relative to their reaction yields, along with the relative energies of the corresponding S,C-bridging isomers, we have performed DFT calculations. The input data for 1,7-N,C isomer was taken from our earlier study,<sup>5c</sup> while the 1,2 isomer was modeled using the structure of the thiazolide derivative HOs<sub>3</sub>(CO)<sub>10</sub>(1,2-N,C-C<sub>3</sub>H<sub>2</sub>NS-3-Me) after the coordinated heterocycle was converted to a benzothiazolate ligand.<sup>18</sup> The unknown S,C isomers were modeled by assuming a bridging motif similar to the N,C-ligated clusters. Fig. 1 shows the optimized structures C1–C4 for the isomeric N,C- and S,C-metallated clusters and the energy ordering of these products (plus the release of two MeCN) relative to 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> (A) and benzothiazole (B). The N,C-metallated isomers C1 and C2 are thermodynamically favored compared to the S,C derivatives C3 and C4. The 1,2-N,C isomer C1 is the most stable of the four isomers and it lies 17.3 kcal mol<sup>-1</sup> below the reactants. The 1,7-N,C isomer C2 is 16.1 kcal mol<sup>-1</sup> less stable than C1 and its formation is exergonic by 1.2 kcal mol<sup>-1</sup>. The experimental observation and isolation of these two isomers in comparable yields suggest that these products are produced in a kinetically controlled sequence and their formation is not subject to thermodynamic control. The optimized structures of C1 and C2



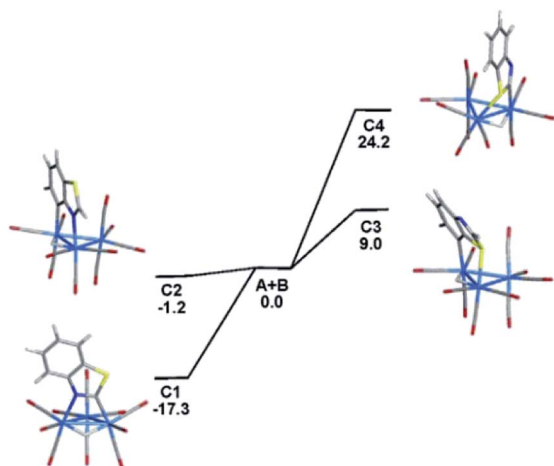
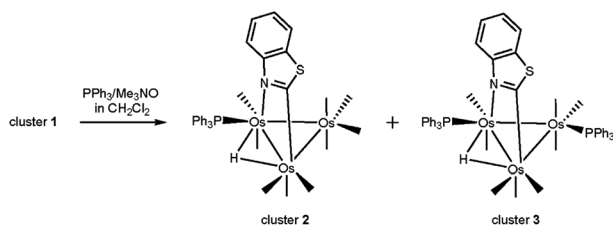


Fig. 1 Optimized metalation products C1–C4 from the reaction of 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> (A) with benzothiazole (B). Energy values ( $\Delta G$ ) are in kcal mol<sup>-1</sup> relative to A and B. The optimized structures for A, B, and free MeCN are not shown.



Scheme 4 Reaction of 1 with PPh<sub>3</sub> to give 2 and 3. Only the major product in each substitution step is shown.

exhibit good agreement with related crystallographically determined structures. Compared to the N,C isomers, both S,C isomers are less stable and lie 9.0 kcal mol<sup>-1</sup> (C3) and 24.2 kcal mol<sup>-1</sup> (C4) higher in energy than the reactants.

Whereas the 1,2-N,C isomer is more stable than the 1,7-N,C species, the reverse trend is found for the S,C isomers. The least stable of these S,C isomers is the 1,2-S,C species (C4). Both C3 and C4 display a highly pyramidalized sulfur atom that results from sulfur coordination at the osmium center, and this perturbation induces a significant loss in the overall aromaticity in the cluster-bound heterocycle. Similar sulfur pyramidalization has been observed in organic alkylation reactions involving sulfur-containing heterocycles<sup>49</sup> and serves as a sufficient driving force that prevents the formation of the two S,C isomers in the reaction of 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> with benzothiazole. Fig. S1† shows the ChemDraw pictures of C1–C4.

### Regioselective PPh<sub>3</sub> addition and DFT evaluation of the different isomers of 2

Treatment of HOs<sub>3</sub>(CO)<sub>10</sub>( $\mu$ -1,2-N,C- $\eta^1$ , $\kappa^1$ -C<sub>7</sub>H<sub>4</sub>NS) (1) with Me<sub>3</sub>NO (one equivalent) at room temperature in the presence of PPh<sub>3</sub> affords the monosubstituted cluster HOs<sub>3</sub>(CO)<sub>9</sub>(PPh<sub>3</sub>)( $\mu$ -1,2-N,C- $\eta^1$ , $\kappa^1$ -C<sub>7</sub>H<sub>4</sub>NS) (2) in crude yields on the order of 80%. Accompanying this reaction is the disubstituted cluster HOs<sub>3</sub>(CO)<sub>8</sub>(PPh<sub>3</sub>)<sub>2</sub>( $\mu$ -1,2-N,C- $\eta^1$ , $\kappa^1$ -C<sub>7</sub>H<sub>4</sub>NS) (3) in 5–10% yield. Control experiments using pure 2 confirm the stepwise nature of the substitution with 3 produced in high yield (>85% crude) when equimolar amounts of 2 and PPh<sub>3</sub> are treated with Me<sub>3</sub>NO (one equivalent). Scheme 4 outlines the course of these reactions where the depicted products of 2 and 3 are based on the crystallographically determined structures (*vide infra*). Both new clusters were isolated by chromatography over silica gel and characterized in solution by IR and NMR spectroscopy, and the solid-state structure of each product was established by single-crystal X-ray diffraction analysis.

The asymmetric unit of 2 contains two independent Os<sub>3</sub> clusters along with a molecule of CH<sub>2</sub>Cl<sub>2</sub> and water. Since the two structures do not show any significant structural differences, we will discuss only one of these crystallographically independent molecules whose structure is depicted in Fig. 2.

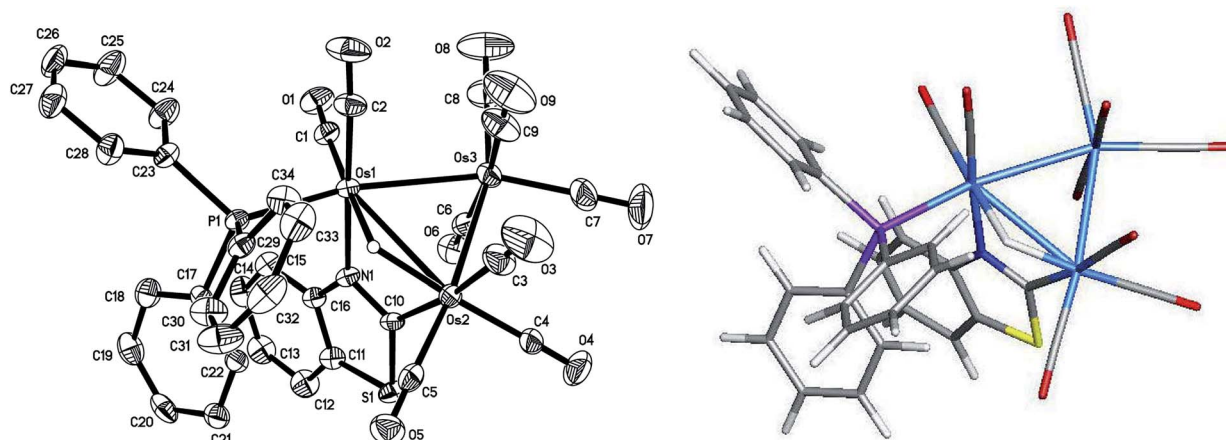


Fig. 2 ORTEP drawing of the molecular structure of 2 (left) showing 50% probability thermal ellipsoids and the DFT-optimized structure of D1 (right). Selected interatomic distances (Å) and angles (°) for 2: Os(1)–Os(2) 2.9843(3), Os(2)–Os(3) 2.8795(3), Os(1)–Os(3) 2.8533(3), Os(1)–N(1) 2.138(4), Os(2)–C(10) 2.086(5), Os(1)–P(1) 2.3888(12), N(1)–C(10) 1.331(6), P(1)–Os(1)–Os(2) 105.08(3), P(1)–Os(1)–Os(3) 163.06(3), P(1)–Os(1)–N(1) 90.49(10), Os(1)–N(1)–C(10) 109.6(3), C(16)–N(1)–C(10) 113.7(4), N(1)–C(10)–Os(2) 116.6(3), C(5)–Os(2)–Os(1) 114.99(13).



Selected bond distances and angles are listed in the figure caption. The molecular core consists of an approximate isosceles triangle of osmium atoms [Os(1)–Os(2) 2.9843(2), Os(2)–Os(3) 2.8795(3) and Os(1)–Os(3) 2.8533(3) Å] ligated by nine carbonyls and a PPh<sub>3</sub> ligand, along with an edge-bridging hydride that shares the same Os(1)–Os(2) vector as the metalated benzothiazolate ligand. The edge-bridged Os–Os bond is the longest of the three metal–metal bonds. The phosphine ligand occupies an equatorial coordination site at the nitrogen-bound osmium and lies cis to the bridging hydride. The Os(1)–P(1) [2.3888(11) Å], Os(1)–N(1) [2.138(4) Å], and Os(2)–C(10) [2.086(5) Å] bond distances involving the ancillary ligands are similar to those reported by us in related tris-osmium clusters.<sup>14,18,20</sup> However, the locus of PPh<sub>3</sub> substitution in **2** is quite different from that displayed in the 4-methylquinolinate analogues HOs<sub>3</sub>(CO)<sub>9</sub>(PPh<sub>3</sub>)[(μ-1,2-η<sup>1</sup>,κ<sup>1</sup>-C<sub>9</sub>H<sub>5</sub>(CH<sub>3</sub>)N] and HOs<sub>3</sub>(CO)<sub>9</sub>[P(OMe)<sub>3</sub>][(μ-1,2-η<sup>1</sup>,κ<sup>1</sup>-C<sub>9</sub>H<sub>5</sub>(CH<sub>3</sub>)N)].<sup>14</sup> The pnictogen ligand in these latter derivatives occupies an equatorial site at the non-metalated osmium atom. The different regiochemistry for phosphine substitution in the current cluster that contains a π-excessive heterocycle *versus* our earlier report on the Os<sub>3</sub> cluster with a π-deficient heterocycle is striking and underscores the importance of the metalated platform in controlling the outcome of the reaction. We examined the bonding in **2** by DFT, and the optimized structure (**D1**) is shown alongside the experimentally determined structure in Fig. 2. The computed structure of **D1** shows good agreement with the solid-state structure.

The <sup>1</sup>H and <sup>31</sup>P NMR spectra recorded for **2** indicate the presence of two clusters with the dominant species present in >95% yield; the amount of the minor species varies but never exceeds 5%. The major product shows a high-field hydride at δ –14.21 (J 11.6) and a <sup>31</sup>P singlet δ 11.8 at room temperature,

while the minor species reveals a hydride doublet and <sup>31</sup>P singlet at δ –15.80 (J 12.0) and δ 1.98, respectively. VT NMR measurements over the temperature range 298–223 K showed no change in the relative intensities of these species, allowing us to rule out a dynamic exchange of the hydride and PPh<sub>3</sub> ligand between the two species on the NMR time scale. This latter premise was further corroborated by <sup>1</sup>H and <sup>31</sup>P EXSY NMR experiments conducted at 253 K.

The identity of the two species in the NMR spectra of **2** and the relative stability of the two products were addressed through electronic structure calculations on the likely monophosphine derivatives of **2**. We restricted our calculations to products where the two edge-bridging ligands share a common Os–Os vector due to their enhanced stability compared to isomers that contain two Os–Os bonds with a single edge-bridging ligand.<sup>5e,21</sup> And concerning the disposition of the PPh<sub>3</sub> group in HOs<sub>3</sub>(CO)<sub>9</sub>(PPh<sub>3</sub>)(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS), we have examined only those isomers that possess an equatorial phosphine since axially disposed phosphine ligands experience an energetic penalty of 10–15 kcal mol<sup>–1</sup> due to unfavorable van der Waals interactions between the phosphine and the pair of axial ligands on the adjacent osmium centers.<sup>22</sup> Not counting species **D1**, five additional structures exist with an equatorial PPh<sub>3</sub> ligand, and these are depicted in Fig. 3. The structure for **D2** is 2.5 kcal mol<sup>–1</sup> less stable than **D1** and the PPh<sub>3</sub> ligand is situated cis to the hydride at the C-metalated osmium atom. Species **D3** and **D4** represent a pair of stereoisomers where the PPh<sub>3</sub> occupies one of the two equatorial sites at the Os(CO)<sub>3</sub>P moiety. The energy difference between **D3** and **D4** is negligible, and both species lie at least 3.5 kcal mol<sup>–1</sup> above the thermodynamically favored species **D1**. Phosphine substitution at the alternative equatorial sites associated with the hydride-bridged Os–Os bond furnishes species **D5** and **D6**. The ancillary PPh<sub>3</sub>

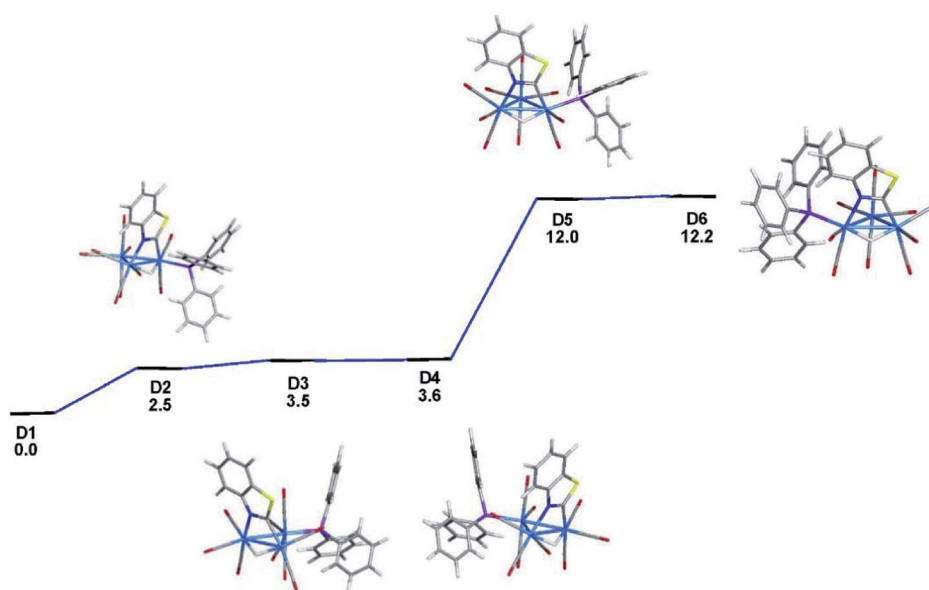


Fig. 3 DFT-optimized structures and energy ordering for the different equatorial-substituted PPh<sub>3</sub> isomers of HOs<sub>3</sub>(CO)<sub>9</sub>(PPh<sub>3</sub>)(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) (**D1**–**D6**). Energy values are ΔG in kcal mol<sup>–1</sup> relative to **D1**. Fig. S2† shows the ChemDraw pictures of species **D1**–**D6**.



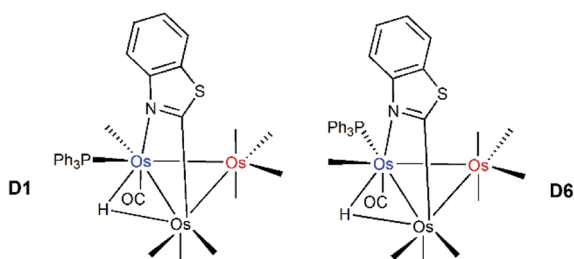
ligand in these two species adopts a distal orientation with respect to the edge-bridging hydride and each species exhibits a nearly linear P–Os–H arrangement of atoms based on the computed bond angles of 167° (**D5**) and 175° (**D6**).

The origin for the computed energy difference in the **D1** and **D6** stereoisomers undoubtedly derives from a combination of steric and electronic effects at each Os<sub>3</sub> product. The observed substitution pattern adopted by the ancillary benzothiazolate, phosphine, and hydride groups largely governs the relative stability of the different isomers **D1–D6**. Adverse interactions emanating from the phosphine and the heterocyclic auxiliary in **D6** *vis-à-vis* **D1** shape the forces that energetically differentiate this pair of stereoisomers. The site of PPh<sub>3</sub> substitution in **D6** places the pnictogen ligand near the benzoannulated portion of the heterocyclic moiety, which promotes a 0.043 Å elongation of the Os–N bond in **D6**; coupled with this perturbation, is a canting of the CO, the one trans to the heterocyclic nitrogen, away from the PPh<sub>3</sub> ligand. The van der Waals interactions serve to reduce the OC–Os–N bond angle from 176° in **D1** to 166° in **D6**. Accompanying this perturbation is an increase in the Os–Os–CO bond angle from 89° (**D1**) to 93° (**D6**) and a 0.1 Å increase in the Os–Os bond distance in **D6**. The defining color labels associated with these structural changes are illustrated in Scheme 5. Analysis of the **D2** and **D5** stereoisomers reveals similar trends at the C-metalated osmium atom.

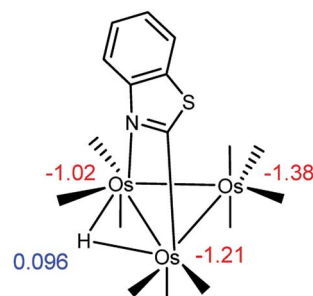
In terms of an operational electronic effect in **D1** and **D6**, the geometrical arrangement of the participant P, Os, and H atoms will influence the net stability of the orbitals involved in the four-centered, four-electron (4c,4e) interaction here. Within this delocalized bonding model, the hydride may be treated as proton that remains associated with the electron-rich Os–Os bond. The “proton” will experience enhanced stabilization if it can interact with additional electron density in an orbital of suitable energy and proper symmetry. The orbital interactions between a hydride(s) and a metal–metal bond in polynuclear systems have been successfully analyzed within a 3c,2e bonding model.<sup>23</sup> And we build on this concept by confirming the thermodynamic preference for products that have the phosphine positioned syn to the bridging hydrogen. Here electron density contributed by the P–Os bond helps stabilize the bridging hydrogen when the phosphine adopts a site syn to the hydride-bridged Os–Os vector. In this scenario, the P–Os–Os atoms display an obtuse angle, a feature underscored by numerous structural studies involving related trimetallic clusters where

the P–Os–Os bond angle ranges from 105–117°.<sup>14,16–18,24</sup> The computed P–Os–Os bond angle in **D1** is 109°. Assuming a fixed cluster polyhedron, expansion of the P–Os–Os bond angle in **D1** effectively permutes the PPh<sub>3</sub> ligand to the other equatorial site at the nitrogen-substituted osmium center. This progression yields **D6** and is accompanied by a 100° increase in the P–Os–Os bond angle. The nearly linear arrangement of the P–Os–Os atoms in **D6** effectively eliminates any through-bond P–Os interaction with the bridging hydrogen that remains tethered to the original Os–Os vector. The stabilizing orbital effect promulgated by the ancillary phosphine when syn disposed to the bridging hydride in **D1** is not unlike many organic reactions where secondary orbital interactions serve to govern the reaction outcome.<sup>25</sup> Syn positioning of the phosphine to the hydride-bridged metal–metal bond in other clusters does not always represent the thermodynamically favored regiochemistry for this pair of ligands.<sup>14</sup> We suspect that in such cases any beneficial orbital effect involving the hydrogen and the electrons in the P–Os bond is counterbalanced by extant steric perturbations within the molecule. The hydride regiochemistry displayed in this genre of metal clusters cannot be generalized with certainty at this time and must be evaluated on a case by case basis.

We have successfully analyzed the course of hydride addition to the metalated heterocycle in a series of heterocyclic-bridged Os<sub>3</sub> clusters earlier by taking into account the charge distribution in the starting material.<sup>7b,11a–c</sup> Accordingly, we have examined the charge distribution in **C1** to help shed light on the isomeric products formed in the reaction that gives cluster **2**. Scheme 6 shows the NPA atomic charges for the osmium atoms in **C1**. The three osmium atoms carry a net negative charge and the nitrogen-substituted osmium is the least electron rich of the three osmium atoms ( $Q = -1.02$ ), followed by the C-metalated osmium center whose  $Q$  was computed as  $-1.21$ . The metal center in the Os(CO)<sub>4</sub> moiety exhibits the greatest negative charge ( $-1.38$ ). The data from the NPA calculations support, inasmuch as charge control is responsible, the formation of **D1** as the major monophosphine product. The presence of **D2** as the other computed component in the reaction mixture is in keeping with PPh<sub>3</sub> association with the second most electro-positive osmium center in **C1**. Finally, the NPA charge for the edge-bridging hydride was positive with a  $Q$  of 0.096.<sup>26</sup>



Scheme 5 Color code used in the discussion of the steric perturbation of the PPh<sub>3</sub> group on the cluster core in **D1** and **D6**.



Scheme 6 Natural population analysis (NPA) charges for the osmiums and bridging hydride in **C1**.



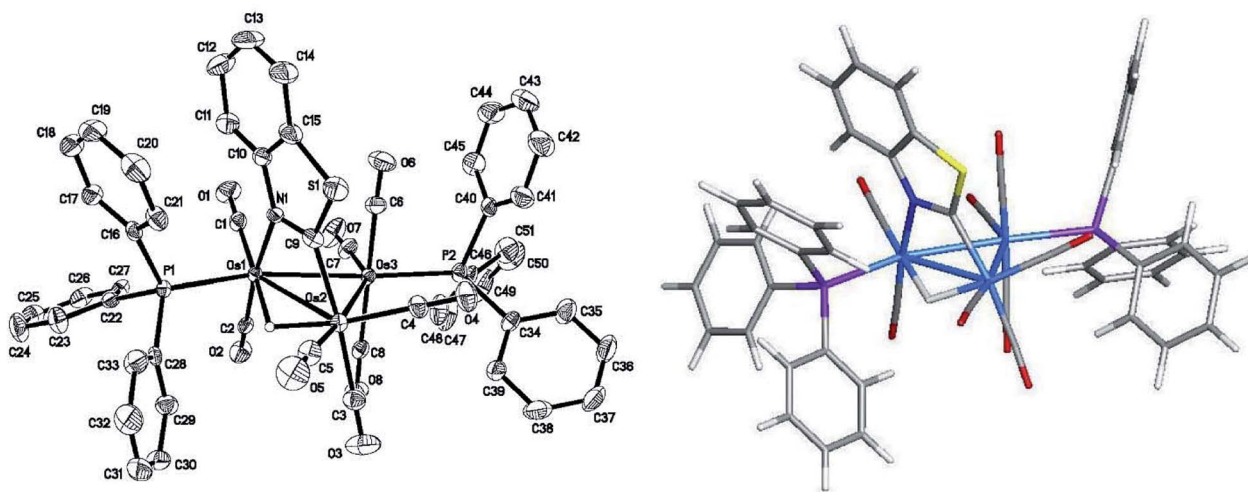


Fig. 4 ORTEP drawing of the molecular structure of **3** (left) showing 50% probability thermal ellipsoids and the DFT-optimized structure of **E1** (right). Selected interatomic distances (Å) and angles (°) for **3**: Os(1)–Os(2) 2.9638(2), Os(2)–Os(3) 2.9066(2), Os(1)–Os(3) 2.8651(2), Os(1)–N(1) 2.143(4), Os(2)–C(9) 2.076(5), Os(3)–P(2) 2.3550(11), Os(1)–P(1) 2.3578(11), N(1)–C(10) 1.408(6), N(1)–C(9) 1.310(6), C(9)–S(1) 1.739(5), Os(2)–Os(1)–Os(3) 59.793(6), Os(1)–Os(2)–Os(3) 58.414(6), Os(1)–Os(3)–Os(2) 61.791(6), P(1)–Os(1)–Os(3) 166.31(3), P(1)–Os(1)–Os(2) 107.16(3), P(1)–Os(1)–N(1) 91.29(10), P(2)–Os(3)–Os(1) 175.57(3), P(2)–Os(3)–Os(2) 116.74(3), N(1)–C(9)–Os(2) 117.3(3), C(9)–Os(2)–Os(1) 66.21(13), C(9)–Os(2)–Os(3) 86.24(12), Os(1)–N(1)–C(9) 108.9(3).

### Formation of $\text{HOs}_3(\text{CO})_8(\text{PPh}_3)_2(\mu\text{-}1,2\text{-N,C-}\eta^1, \kappa^1\text{-C}_7\text{H}_4\text{NS})$ (**3**) and stereoselective tripodal rotation at the $\text{Os}(\text{CO})_3\text{P}$ moiety

The reaction of  $\text{PPh}_3$  with **2** in the presence of  $\text{Me}_3\text{NO}$  affords **3**, which exists as a pair of discernible isomers in solution (298 K) by NMR spectroscopy. The structure of the major isomer was established by X-ray diffraction analysis. Fig. 4 displays the ORTEP diagram of the molecular structure of **3** with selected bond distances and angles listed in the figure caption. **3** consists of a triangular array of osmium atoms [Os(1)–Os(2) 2.9638(2), Os(2)–Os(3) 2.9066(2), and Os(1)–Os(3) 2.8651(2) Å] where the mean Os–Os bond distance is 2.9118 Å. The Os(1)–Os(2) vector is the longest edge and it is bridged by the hydride and benzothiazolate ligands. Eight carbonyls and two  $\text{PPh}_3$  ligands complete the coordination sphere of this cluster. As in **2**, one of the two  $\text{PPh}_3$  ligands is bound to the nitrogen-bound osmium [Os(1)] while the second phosphine [P(2)] is coordinated to the Os(3) atom at an equatorial site syn to the Os(2) center. The mean Os–P bond distance in **3** is 2.3564 Å and not unlike that Os–P distance found in **2** [2.3888(11) Å]. The Os–N [2.143(4) Å] and Os–C(9) [2.076(5) Å] bond distances are unremarkable compared with those bond lengths reported in **1** and related clusters.<sup>4a,14,17,18</sup>

The  $^1\text{H}$  and  $^{31}\text{P}$  NMR spectra of the major species in solution are consistent with the solid-state structure. Here a diagnostic hydride doublet at  $\delta$  –13.74 (J 11.6) confirms the coupling of the edge-bridging hydride to one of the two phosphine groups in the major isomer of **3**. The magnitude of the  $^2J_{\text{PH}}$  is similar to that recorded in **2**, allowing us to confidently assign the coupling to the geminal phosphine defined by the P(1) atom in Fig. 4. The  $^{31}\text{P}\{^1\text{H}\}$  NMR spectrum exhibits two broad singlets at  $\delta$  8.9 and 3.9 for the  $\text{PPh}_3$  ligands. At room temperature, the minor species (<3%) shows a weak hydride at  $\delta$  –15.38 (J 12.5) while the  $^{31}\text{P}$  resonances were not visible. In order to probe the

relationship between the two isomers, we examined **3** by VT NMR spectroscopy over the temperature range 323–223 K. Lowering the temperature to 223 K reveals the presence of four hydride doublets centered at  $\delta$  –13.73, –13.80, –13.90, and –15.45 in the  $^1\text{H}$  NMR spectrum (Fig. S3†). Consistent with this change in the  $^1\text{H}$  NMR spectrum is the observation of eight broad singlets at  $\delta$  2.6, 3.5, 4.2, 4.5, 6.3, 6.8, 8.8, and 10.2 in the  $^{31}\text{P}$  NMR spectrum (Fig. S3 and S4†). Collectively, these data confirm the existence of four isomeric products for **3** whose relationship was established by EXSY measurements at 223 K. The EXSY data confirm the dynamical exchange between the two hydrides representing the minor species ( $\delta$  –13.73 with  $\delta$  –15.45); hydride interconversion between the two species ( $\delta$  –13.80 with  $\delta$  –13.90) associated with the major isomer was also verified. No hydride exchange between the major and minor hydride species was observed. Similar exchange behavior for the pair of  $^{31}\text{P}$  resonances at  $\delta$  10.2 and 4.2 with those at  $\delta$  6.8 and 3.5 for the major species was found. The pairwise exchange of the  $^{31}\text{P}$  resonances at  $\delta$  8.8 and 4.5 with those at  $\delta$  6.6 and 2.6 define the exchanging pairs of resonances in the minor isomeric species.

The identity of the isomers present in the NMR spectra of **3** is best addressed by electronic structure calculations, and here we employed the solid-state structure of **3** as the starting point for our studies. The geometry-optimized structure of species **E1**, which is depicted alongside the X-ray diffraction structure in Fig. 4, exhibits good agreement with the experimental structure. Fig. 5 shows the ground-state energy ordering for the different species based on **3** that possess equatorial phosphines and where one of the  $\text{PPh}_3$  ligands is positioned cis to the edge-bridging hydride. The latter structural feature is required in order to reproduce the  $^2J_{\text{PH}}$  coupling of  $\sim$ 12 Hz in the  $^1\text{H}$  NMR spectra displayed in Fig. S3.† As with cluster **2**, the optimized



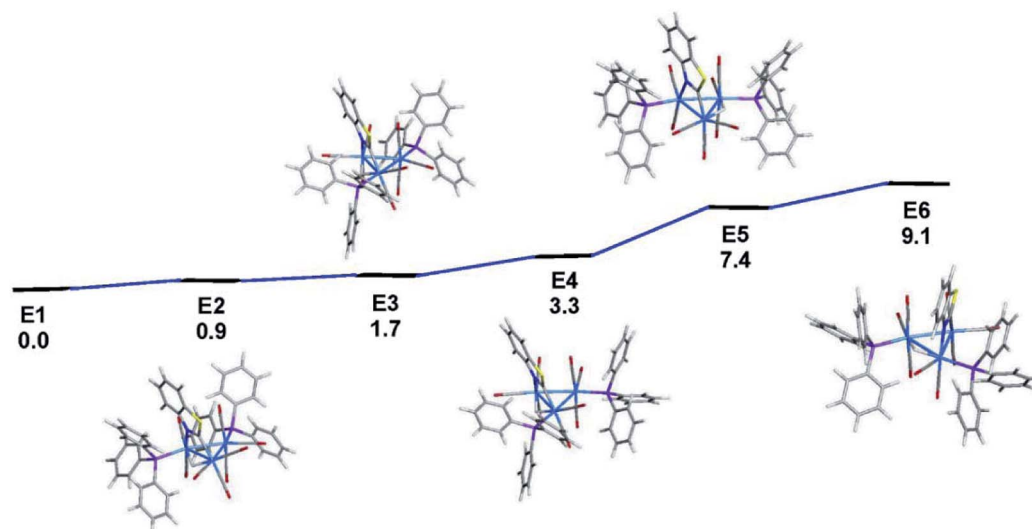
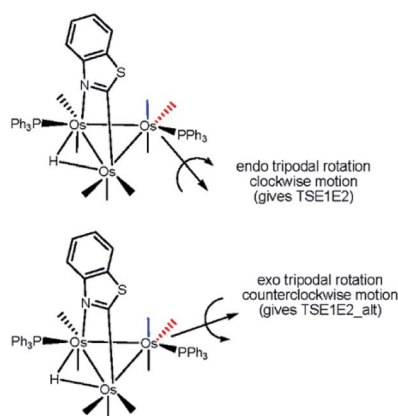


Fig. 5 DFT-optimized structures and energy ordering for selected isomers based on cluster **3**. Energy values are  $\Delta G$  in kcal mol<sup>-1</sup> relative to **E1**. Fig. S5† shows the ChemDraw pictures of **E1**–**E6**.



Scheme 7 Endo and exo tripodal rotation pathways in **E1**.

structure based on the X-ray diffraction structure of **3** (species **E1**) represents the thermodynamically favored isomer. **E2** is a stereoisomer of **E1**, and it lies 0.9 kcal mol<sup>-1</sup> above the latter species. The principal difference between **E1** and **E2** concerns the disposition of the newly added phosphine relative to the two available equatorial sites at the Os(CO)<sub>3</sub>P moiety. In the case of **E1**, the two phosphine groups display a synclinal orientation about the Os–Os bond common to both ligands, while they exhibit an anticlinal arrangement in **E2**.<sup>27</sup> Species **E3** and **E4** are stereoisomers that originate from **D2** and exhibit a substitution pattern similar to that found in species **E1** and **E2**, with synclinal **E3** 1.6 kcal mol<sup>-1</sup> more stable than anticlinal **E4**. Species **E5** and **E6** are isomers of **E1**–**E4**, and they may be viewed as the products of hydride migration from its common locus in **E1** and **E2** to the adjacent Os–Os bond in **E5** and **E6**, respectively. The hydride migration that gives **E5** and **E6** is accompanied by a 7.4 and 9.1 kcal mol<sup>-1</sup> loss in energy, respectively.<sup>28</sup>

The dynamic NMR behavior exhibited by the major component of **3** may be rationalized by an intramolecular exchange

between **E1**  $\rightleftharpoons$  **E2**. The equilibration of these stereoisomers is rapid at room temperature and proceeds *via* a tripodal permutation involving the PPh<sub>3</sub> ligand and two CO groups at the Os(CO)<sub>3</sub>P center. Scheme 7 shows the rotational direction available to the two fragment rotors capable of promoting isomerization. Two of the three rotor substituents in both fragments include the equatorial CO and PPh<sub>3</sub> groups. When the axial CO trans to the benzothiazolate ligand is used as the third rotor group (endo route), a stereoselective clockwise rotation of the fragment rotor was computationally confirmed. In this process the CO *syn* the heterocycle remains unchanged. The participation of the CO in the tripodal rotation proceeds *via* a counterclockwise manner (exo route) and this leads to unfavourable van der Waals interactions between the heterocycle and the two migrating carbonyls (CO and CO). In both tripodal rotations, the PPh<sub>3</sub> group avoids the bridging benzothiazolate ligand by adopting an orientation coincident to the plane defined by the three metals. No evidence for a turnstile rotation involving the Os(CO)<sub>3</sub>P moiety was found, and these data are consistent with our earlier NMR studies that have demonstrated facile tripodal rotation at the Os(CO)<sub>3</sub>L vertices in a variety of heterocyclic-bridged clusters.<sup>8d,29</sup>

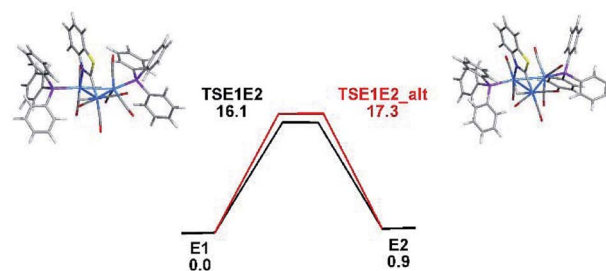


Fig. 6 Free-energy profile for the **E1**  $\rightleftharpoons$  **E2** isomerization through endo (TSE1E2) and exo (TSE1E2<sub>alt</sub>) tripodal rotations. Energy values are in kcal mol<sup>-1</sup> relative to **E1**.



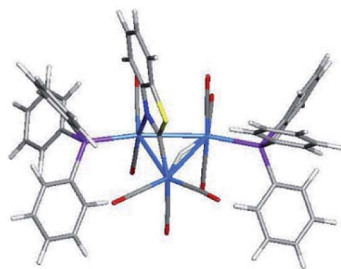


Fig. 7 DFT-optimized structure for TSE1E5.

The optimized transition structures for **TSE1E2** and **TSE1E2\_alt** and the computed barriers for the tripodal rotation that promote the **E1** ⇌ **E2** exchange are displayed in Fig. 6. Both activation barriers are low and in concert with the VT NMR behavior recorded for **3**. The endo tripodal rotation is favored by 1.2 kcal mol<sup>-1</sup> over the exo rotation process. Fig. S6† provides a detailed view of the permutation process as a function of the ancillary ligands at the Os(CO)<sub>3</sub>P moiety.

The dynamical exchange between the minor stereoisomers **E3** ⇌ **E4** follows a similar sequence with the endo tripodal rotation again favored over the exo rotation (Fig. S7†). The computed  $\Delta\Delta G^\ddagger$  for the **TSE3E4** and **TSE3E4\_alt** is 2.0 kcal mol<sup>-1</sup> (Fig. S8†). Here the endo rotation proceeds from **E3** by a counterclockwise rotation of groups associated with the fragment rotor. The change in the rotation direction in **E3** ⇌ **E4** from that found in **E1** ⇌ **E2** is dictated by the synclinal disposition of the phosphines in **E3**.

### Computational analysis of hydride migration in the species

#### **E1** ⇌ **E5** and **E2** ⇌ **E6**

Hydride mobility in metal clusters is a well-established phenomenon, and the dynamical exchange of a hydride(s) between different metal–metal bonds can occur rapidly on the NMR time scale.<sup>29b,30</sup> Since hydride migration in **E1** to the adjacent Os–Os bond in **E5** could account for the NMR behavior displayed by **3**, we investigated this scenario by DFT. Indeed, a suitable transition structure (**TSE1E5**) was found for the concerted migration of hydride through the interior of the cluster core by way of an interstitial-like hydride.<sup>31,32</sup> The optimized structure for **TSE1E5** is depicted in Fig. 7. The barrier computed for this process is 19.4 kcal mol<sup>-1</sup>, confirming that this hydride migration is not competitive to the tripodal rotation associated with **TSE1E2** that lies 3.3 kcal mol<sup>-1</sup> lower in energy.<sup>29a</sup> The migration of the hydride in **E2** to give **E6** was also examined through a series of step-scan calculations. Multiple attempts were made to find a suitable transition structure, but we were unable to successfully model this transformation with a species possessing an acceptable imaginary frequency and reasonable energy.

## Conclusions

DFT calculations on the possible metalated heterocycles from the reaction of benzothiazole and 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> have confirmed the greater stability of the 1,2- and 1,7-N,C

coordination isomers over the corresponding S,C-functionalized species. The reactivity of the 1,2-N,C isomer, HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) (**1**), in PPh<sub>3</sub> substitution has been examined through a combined experimental/computational approach. The high regioselectivity experimentally found in each substitution step site has been successfully analyzed by electronic structure calculations. DFT calculations on the fluxional NMR behavior exhibited by **3** reveal a stereoselective tripodal rotation at the Os(CO)<sub>3</sub>(PPh<sub>3</sub>) moiety where rotation of the fragment rotor proceeds distal to the metalated heterocycle. The relative stabilities of the different isomers of **2** and **3** have been determined by DFT, and the computed species reconciled with those products observed in solution by NMR spectroscopy. The ability to functionalize the benzothiazolate ligand in **1** with soft organic nucleophiles and hydride ligand are planned. These studies will allow us to chart the directing effect that the 1,2-N,C coordination mode of the benzothiazolate group in **1** has, if any, versus our earlier studies based on the 1,7-N,C isomer.

## Experimental

### General and instrumentation

All the reactions were carried out under a nitrogen atmosphere using standard Schlenk techniques unless otherwise stated. Reagent grade solvents were dried by the standard methods and freshly distilled prior to use. The benzothiazole, PPh<sub>3</sub>, and Me<sub>3</sub>NO·2H<sub>2</sub>O used in these studies were purchased from Aldrich Chemical Co.; the former chemicals were used as received. Me<sub>3</sub>NO·2H<sub>2</sub>O was dried by azeotropic distillation using benzene and a Dean–Stark distillation unit. HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) (**1**) was prepared from 1,2-Os<sub>3</sub>(CO)<sub>10</sub>(MeCN)<sub>2</sub> and benzothiazole according to literature procedures.<sup>6c,15</sup> The separation of clusters **2** and **3** were performed by TLC in the air on 0.5 mm silica gel 60 Å F<sub>254</sub> glass plates. Infrared spectra were recorded on a Shimadzu IR Prestige-21 spectrophotometer, and the VT <sup>1</sup>H and <sup>31</sup>P NMR spectra were recorded on a Bruker DPX 400 spectrometer. All NMR chemical shift and coupling constant data are reported in δ units and Hz, respectively. Elemental analyses were performed by the Microanalytical Laboratory of Wazed Miah Research Centre at Jahangirnagar University.

### Reaction of HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) (**1**) with PPh<sub>3</sub>

To a CH<sub>2</sub>Cl<sub>2</sub> solution (20 mL) of HOs<sub>3</sub>(CO)<sub>10</sub>(μ-1,2-N,C-η<sup>1</sup>,κ<sup>1</sup>-C<sub>7</sub>H<sub>4</sub>NS) (**1**) (100 mg, 0.101 mmol) and PPh<sub>3</sub> (27 mg, 0.101 mmol) was added 7 mL of CH<sub>2</sub>Cl<sub>2</sub> containing Me<sub>3</sub>NO (8 mg, 0.106 mmol) dropwise over a period of 30 min. The resultant solution was stirred for another 30 min, during which time the solution color changed from yellow to deep orange. The reaction mixture was then filtered through a short silica column (4 cm) to remove any excess Me<sub>3</sub>NO, and the solvent was removed under reduced pressure. The residue was purified by chromatography over silica gel using a mixture of cyclohexane/CH<sub>2</sub>Cl<sub>2</sub> (7 : 3, v/v) as the eluent. Five bands were isolated, the first of which corresponded to unreacted **1** (12 mg). The second and





third bands afforded  $\text{HOs}_3(\text{CO})_9(\text{PPh}_3)(\mu\text{-}1,2\text{-N,C-}\eta^1,\kappa^1\text{-C}_7\text{H}_4\text{NS})$  (**2**) (55 mg, 41%) and  $\text{HOs}_3(\text{CO})_8(\text{PPh}_3)_2(\mu\text{-}1,2\text{-N,C-}\eta^1,\kappa^1\text{-C}_7\text{H}_4\text{NS})$  (**3**) (15 mg, 10%) as orange crystals after recrystallization from *n*-hexane/ $\text{CH}_2\text{Cl}_2$  at  $-4^\circ\text{C}$ . The contents of the fourth and fifth bands were too small for complete characterization. Data for **2**: anal. calcd for  $\text{C}_{34}\text{H}_{20}\text{NO}_9\text{Os}_3\text{P}_2\text{S}$ : C, 33.47; H, 1.65; N, 1.15. Found: C, 33.21; H, 1.79; N, 1.13%. IR ( $\nu(\text{CO})$ ,  $\text{CH}_2\text{Cl}_2$ ): 2090s, 2051s, 201s, 1998s, 1979sh, 1967m, 1940w  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ,  $25^\circ\text{C}$ ):  $\delta$  7.49 (d, J 6.8, 1H), 7.23 (m, 15H), 6.94 (m, 2H), 6.80 (d, J 7.6, 1H),  $-14.21$  (d, J 11.6, 1H).  $^{31}\text{P}\{^1\text{H}\}$  NMR ( $\text{CDCl}_3$ ,  $25^\circ\text{C}$ ):  $\delta$  11.8 (s). Data for **3**: anal. calcd for  $\text{C}_{51}\text{H}_{35}\text{NO}_8\text{Os}_3\text{P}_2\text{S}\cdot\text{CH}_2\text{Cl}_2$ : C, 40.57; H, 2.42; N, 0.91. Found: C, 41.16; H, 2.48; N, 1.03. IR ( $\nu(\text{CO})$ ,  $\text{CH}_2\text{Cl}_2$ ): 2065m, 2060m, 2023s, 1993s, 1967m, 1952s  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ,  $25^\circ\text{C}$ ):  $\delta$  7.50 (m, 8H), 7.37 (m, 11H), 7.25 (m, 12H), 6.92 (m, 3H),  $-13.74$  (d, J 11.6, 1H).  $^{31}\text{P}\{^1\text{H}\}$  NMR ( $\text{CDCl}_3$ ,  $25^\circ\text{C}$ ):  $\delta$  8.9 (s), 3.9 (s).

### Conversion of **2** to **3**

To a Schlenk tube containing **2** (30 mg, 0.024 mmol) and  $\text{PPh}_3$  (7 mg, 0.027 mmol) in 20 mL of  $\text{CH}_2\text{Cl}_2$  was added a  $\text{CH}_2\text{Cl}_2$  solution (5 mL) of  $\text{Me}_3\text{NO}$  (2 mg, 0.027 mmol) over a period of 15 min with the aid of a dropping funnel. The reaction mixture was then stirred at room temperature for 2 h, after which time TLC analysis confirmed the consumption of the starting cluster and the presence of desired product **3**. Chromatographic workup, followed by recrystallization, afforded **3** (26 mg, 75%).

**Table 1** Crystallographic data and structure refinement details for clusters **2** and **3**

	<b>2</b>	<b>3</b>
CCDC entry no	1833188	1833187
Cryst system	Triclinic	Monoclinic
Space group	$P\bar{1}$	$P\bar{1}$
$a$ , Å	11.0561(2)	18.8799(6)
$b$ , Å	16.0438(3)	15.0410(3)
$c$ , Å	21.7248(5)	18.7314(5)
$\alpha$ , deg	100.9456(19)	90
$\beta$ , deg	92.7106(19)	117.376(4)
$\gamma$ , deg	90.9327(17)	90
$V$ , Å <sup>3</sup>	3777.99(15)	4723.5(3)
Mol formula	$\text{C}_{69}\text{H}_{42}\text{Cl}_2\text{N}_2\text{O}_{19}\text{Os}_6\text{P}_2\text{S}_2$	$\text{C}_{51}\text{H}_{35}\text{NO}_8\text{Os}_3\text{P}_2\text{S}$
$F_w$	2541.20	1454.40
Formula units per cell ( $Z$ )	2	4
$D_{\text{calcd}}$ ( $\text{mg m}^{-3}$ )	2.234	2.045
$\lambda$ (Mo $K\alpha$ ), Å	0.71073	0.71073
$\mu$ ( $\text{mm}^{-1}$ )	10.287	8.218
$F(000)$	2348	2744
Total reflections	61 519	74 577
Independent reflections	14 819	9274
Data/res/paras	14 819/0/936	9274/0/599
GOF on $F^2$	1.076	1.071
$R_1^a$ [ $I \geq 2\sigma(I)$ ]	0.0234	0.0242
$wR_2^b$ (all data)	0.0569	0.0592
$\Delta\rho(\text{max}), \Delta\rho(\text{min})$ ( $\text{e} \text{ \AA}^{-3}$ )	1.79, $-1.52$	2.60, $-1.89$

$$^a R_1 = \sum |F_o| - |F_c| / \sum |F_o|, ^b wR_2 = \{\sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2]\}^{1/2}$$

### X-ray crystallography

Single crystals of **2** and **3** suitable for single crystal X-ray diffraction analysis were grown by slow diffusion of hexane into a  $\text{CH}_2\text{Cl}_2$  solution containing each product. A suitable single crystal of each cluster was mounted on an Agilent Super Nova dual diffractometer (Agilent Technologies Inc., Santa Clara, CA) using a Nylon Loop and the diffraction data were collected at 150(2) K using Mo- $K\alpha$  radiation ( $\lambda = 0.71073$ ). Unit cell determination, data reduction, and absorption corrections were carried out using CrysAlisPro.<sup>33</sup> The structures were solved with the ShelXS<sup>34</sup> structure solution program using Direct Method and refined with the ShelXL<sup>34</sup> refinement package using Least Squares minimization within the Olex2 (ref. 35) graphical user interface. Non-hydrogen atoms were refined anisotropically, and hydrogen atoms (except those directly bonded to metals) were included using a riding model. The bridging hydride ligand in **2** and **3** was found in the difference map and freely refined. Additional details of data collection and structure refinement are given in Table 1.

### Computational methodology and modeling details

All DFT calculations were performed with the Gaussian 09 package of programs<sup>36</sup> using the B3LYP hybrid functional. This functional is comprised of Becke's three-parameter hybrid exchange functional (B3)<sup>37</sup> and the correlation functional of Lee, Yang, and Parr (LYP).<sup>38</sup> Each osmium atom was described with the Stuttgart-Dresden effective core potential and SDD basis set,<sup>39</sup> and the 6-31G(d') basis set<sup>40</sup> was employed for all remaining atoms.

All reported geometries were fully optimized, and analytical Hessian was evaluated at each stationary point to determine whether the geometry was an energy minimum (no negative eigenvalues) or a transition structure (one negative eigenvalue). Unscaled vibrational frequencies were used to make zero-point and thermal corrections to the electronic energies, and the resulting free energies are reported in  $\text{kcal mol}^{-1}$  relative to the specified standard. Intrinsic reaction coordinate (IRC) calculations were performed on all transition-state structures in order to establish the reactant and product species associated with each transition-state structure. The geometry-optimized structures have been drawn with the JIMP2 molecular visualization and manipulation program.<sup>41</sup>

### Conflicts of interest

There are no conflicts to declare.

### Acknowledgements

SEK thanks the Ministry of Education, the Government of the Peoples' Republic of Bangladesh, and the University Grants Commission of Bangladesh for sponsorship and financial support of this work. MGR acknowledges The Robert A. Welch Foundation through Grant B-1093 for generous research support. The DFT calculations were performed at UNT through CASCAM, which is an NSF-supported facility (CHE-1531468).

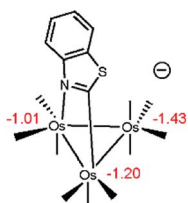


## Notes and references

- 1 (a) A. Eisenstadt, C. M. Giandomenico, M. F. Frederick and R. M. Laine, *Organometallics*, 1985, **4**, 2033; (b) R. H. Fish, T.-J. Kim, J. H. Bushweller, R. K. Rosen and J. W. Dupon, *Organometallics*, 1986, **5**, 2193.
- 2 E. Rosenberg and R. Kumar, *Dalton Trans.*, 2012, **41**, 714.
- 3 S. E. Kabir, D. S. Kolwaite, E. Rosenberg, L. G. Scott, T. McPhillips, R. Duque, M. Day and K. I. Hardcastle, *Organometallics*, 1996, **15**, 1979.
- 4 (a) E. Arcia, D. S. Kolwaite, E. Rosenberg, K. Hardcastle, J. Ciurash, R. Duque, R. Gobetto, L. Milone, D. Osella, M. Botta, W. Dastrú, A. Viale and I. Fiedler, *Organometallics*, 1998, **17**, 415; (b) R. Smith, E. Rosenberg, K. I. Hardcastle, V. Vazquez and J. Roh, *Organometallics*, 1999, **18**, 3519.
- 5 (a) A. K. Raha, S. Ghosh, M. M. Karim, D. A. Tocher, N. Begum, A. Sharmin, E. Rosenberg and S. E. Kabir, *J. Organomet. Chem.*, 2008, **693**, 3613; (b) S. Ghosh, M. N. Uddin, N. Begum, G. M. G. Hossain, K. A. Azam and S. E. Kabir, *J. Chem. Crystallogr.*, 2010, **40**, 572; (c) M. I. Hossain, S. Ghosh, G. Hogarth, G. M. G. Hossain and S. E. Kabir, *J. Organomet. Chem.*, 2011, **696**, 3036; (d) S. Ghosh, M. R. Al-Mamun, G. M. G. Hossain and S. E. Kabir, *Inorg. Chim. Acta*, 2011, **378**, 307; (e) S. A. Begum, M. A. H. Chowdhury, S. Ghosh, D. A. Tocher, M. G. Richmond, E. Rosenberg and S. E. Kabir, *J. Organomet. Chem.*, 2017, **849–850**, 337.
- 6 (a) S. E. Kabir, D. S. Kolwaite, E. Rosenberg, K. Hardcastle, W. Cresswell and J. Grindstaff, *Organometallics*, 1995, **14**, 3611; (b) B. Bergman, R. Holmquist, R. Smith, E. Rosenberg, J. Ciurash, K. Hardcastle and M. Visi, *J. Am. Chem. Soc.*, 1998, **120**, 12818; (c) M. J. Abedin, B. Bergman, R. Holmquist, R. Smith, E. Rosenberg, J. Ciurash, K. Hardcastle, J. Roe, V. Vazquez, C. Roe, S. Kabir, B. Roy, S. Alam and K. A. Azam, *Coord. Chem. Rev.*, 1999, **190–192**, 975.
- 7 (a) E. Rosenberg, S. E. Kabir, M. J. Abedin and K. I. Hardcastle, *Organometallics*, 2004, **23**, 3982; (b) D. G. Musaev, T. Nowroozi-Isfahani, K. Morokuma, J. Abedin, E. Rosenberg and K. I. Hardcastle, *Organometallics*, 2006, **25**, 203; (c) E. Rosenberg, J. Abedin and K. I. Hardcastle, *J. Cluster Sci.*, 2012, **23**, 901.
- 8 (a) A. B. Din, B. Bergman, E. Rosenberg, R. Smith, W. Dastru, R. Gobetto, L. Milone and A. Viale, *Polyhedron*, 1998, **17**, 2975; (b) E. Rosenberg, M. J. Abedin, D. Rokhsana, D. Osella, L. Milone, C. Nervi and J. Fiedler, *Inorg. Chim. Acta*, 2000, **300–302**, 769; (c) S. E. Kabir, K. M. A. Malik, H. S. Mandal, M. A. Mottalib, M. J. Abedin and E. Rosenberg, *Organometallics*, 2002, **21**, 2593; (d) E. Rosenberg, M. J. Abedin, D. Rokhsana, A. Viale, W. Dastru, R. Gobetto, L. Milone and K. Hardcastle, *Inorg. Chim. Acta*, 2002, **334**, 343; (e) S. T. Beatty, B. Bergman, E. Rosenberg, W. Dastru, R. Gobetto, L. Milone and A. Viale, *J. Organomet. Chem.*, 2000, **593–594**, 226.
- 9 (a) E. Rosenberg, F. Spada, K. Sugden, B. Martin, L. Milone, R. Gobetto, A. Viale and J. Fiedler, *J. Organomet. Chem.*, 2003, **668**, 51; (b) D. Colangelo, A. L. Ghiglia, A. R. Ghezzi, M. Ravera, E. Rosenberg, F. Spada and D. Osella, *J. Inorg. Biochem.*, 2005, **99**, 505.
- 10 (a) C. Nervi, R. Gobetto, L. Milone, A. Viale, E. Rosenberg, D. Rokhsana and J. Fiedler, *Chem.–Eur. J.*, 2003, **9**, 5749; (b) E. Rosenberg, D. Rokhsana, C. Nervi, R. Gobetto, L. Milone, A. Viale, J. Fiedler and M. A. Botavina, *Organometallics*, 2004, **23**, 215; (c) E. Rosenberg, F. Spada, K. Sugden, B. Martin, R. Gobetto, L. Milone and A. Viale, *J. Organomet. Chem.*, 2004, **689**, 4729.
- 11 (a) M. A. Mottalib, N. Begum, S. M. T. Abedin, T. Akter, S. E. Kabir, M. A. Miah, D. Rokhsana, E. Rosenberg, G. M. G. Hossain and K. I. Hardcastle, *Organometallics*, 2005, **24**, 4747; (b) D. G. Musaev, T. Nowroozi-Isfahani, K. Morokuma and E. Rosenberg, *Organometallics*, 2005, **24**, 5973; (c) T. Nowroozi-Isfahani, D. G. Musaev, K. Morokuma and E. Rosenberg, *Inorg. Chem.*, 2006, **45**, 4963.
- 12 (a) B. C. Gates, *Catalytic Chemistry*, J. Wiley and Sons, New York, 1992, p. 409; (b) See symposium: *Mechanism of HDS/HDN Reactions*, Prepr. ACS Div. of Pet. Chem. 1988, vol. 33, p. 638; (c) R. H. Fish, in *Aspects of Homogeneous Catalysts*, ed. R. Ugo, Kluwer Academics, Dordrecht, The Netherlands, 1990, pp. 65–83.
- 13 (a) S. D. Gray, D. P. Smith, M. S. Bruck and D. E. Wigley, *J. Am. Chem. Soc.*, 1992, **114**, 5462; (b) S. D. Gray, P. A. Fox, M. S. Kingsborough, M. S. Bruci and D. E. Wigley, *Prepr.–Am. Chem. Soc., Div. Pet. Chem.*, 1993, **38**, 706; (c) I. Horvath and R. D. Adams, *Prog. Inorg. Chem.*, 1985, **33**, 127; (d) A. Eisenstadt, C. M. Giandomenico, M. F. Frederick and R. M. Laine, *Organometallics*, 1985, **4**, 2033; (e) R. M. Laine, *New J. Chem.*, 1987, **11**, 543; (f) R. D. Adams, J. E. Babin, J. T. Tanner and J. A. Wolfe, *J. Am. Chem. Soc.*, 1990, **112**, 3426; (g) S. E. Kabir, M. Day, M. Irving, T. McPhillips, H. Minassian, E. Rosenberg and K. I. Hardcastle, *Organometallics*, 1991, **10**, 3997.
- 14 M. B. Hursthouse, S. E. Kabir, K. M. A. Malik, M. Tesmer and H. Vahrenkamp, *J. Organomet. Chem.*, 1998, **568**, 133.
- 15 N. Begum, M. K. Islam, S. E. Kabir and M. M. Hassan, *J. Bangladesh Acad. Sci.*, 2005, **29**, 145.
- 16 K. M. Uddin, S. Ghosh, A. K. Raha, G. Hogarth, E. Rosenberg, A. Sharmin, K. I. Hardcastle and S. E. Kabir, *J. Organomet. Chem.*, 2010, **695**, 1435.
- 17 N. Begum, A. J. Deeming, M. K. Islam, S. E. Kabir, D. Rokhsana and E. Rosenberg, *J. Organomet. Chem.*, 2004, **689**, 2633.
- 18 K. A. Azam, R. Dilshad, S. E. Kabir, K. Khatoon, L. Nessa, M. M. Rahman, E. Rosenberg, M. B. Hursthouse, K. M. A. Malik and A. J. Deeming, *Dalton Trans.*, 1996, 1731.
- 19 J. A. Joule and K. Mills, *Heterocyclic Chemistry*, Wiley, Hoboken, NJ, 2010.
- 20 K. A. Azam, M. B. Hursthouse, S. A. Hussain, S. E. Kabir, K. M. A. Malik, M. M. Rahman and E. Rosenberg, *J. Organomet. Chem.*, 1998, **559**, 81.



- 21 M. A. H. Chowdhury, S. Rajbangshi, A. Rahaman, L. Yang, V. N. Nesterov, M. G. Richmond, S. M. Mobin and S. E. Kabir, *J. Organomet. Chem.*, 2015, **779**, 21.
- 22 (a) B. F. G. Johnson, J. Lewis, E. Nordlander and P. R. Raithby, *Dalton Trans.*, 1996, 755; (b) X. Zhang, S. Kandala, L. Yang, W. H. Watson, X. Wang, D. A. Hrovat, W. T. Borden and M. G. Richmond, *Organometallics*, 2011, **30**, 1253; (c) M. K. Hossain, S. Rajbangshi, A. Rahaman, M. A. H. Chowdhury, T. A. Siddiquee, S. Ghosh, M. G. Richmond, E. Nordlander, G. Hogarth and S. E. Kabir, *J. Organomet. Chem.*, 2014, **760**, 231.
- 23 (a) R. Bau, R. G. Teller, S. W. Kirtley and T. F. Koetzle, *Acc. Chem. Res.*, 1979, **12**, 176; (b) R. Hoffmann, B. E. R. Schilling, R. Bau, H. D. Kaesz and D. M. P. Mingoss, *J. Am. Chem. Soc.*, 1978, **100**, 6088; (c) S. K. Brayshaw, J. C. Green, N. Hazari and A. S. Weller, *Dalton Trans.*, 2007, 1781.
- 24 (a) R. D. Adams, N. M. Golembeski and J. P. Selegue, *Inorg. Chem.*, 1981, **20**, 1242; (b) R. D. Adams, Z. Dawoodi, D. F. Foust and B. E. Segmuller, *Organometallics*, 1983, **2**, 315; (c) A. J. Deeming, Y. Fuchita, K. Hardcastle, K. Henrick and M. McPartlin, *Dalton Trans.*, 1986, 2259; (d) K. A. Azam, M. B. Hursthouse, S. A. Hussain, S. E. Kabir, K. M. A. Malik, M. Rahman and E. Rosenberg, *J. Organomet. Chem.*, 1998, **559**, 81; (e) M. Day, D. Espitia, K. I. Hardcastle, S. E. Kabir, E. Rosenberg, R. Gobetto, L. Milone and D. Osella, *Organometallics*, 1991, **10**, 3550; (f) S. Monira, S. Afrin, K. A. Azam, M. K. Hossain, D. A. Tocher, S. Ghosh, S. Rajbangshi, S. E. Kabir and G. Hogarth, *J. Organomet. Chem.*, 2015, **799–800**, 281.
- 25 (a) T. A. Albright, J. K. Burdett and M. H. Whangbo, *Orbital Interactions in Chemistry*, Wiley-Interscience, New York, 1985; (b) I. Fleming, *Molecular Orbitals and Organic Chemical Reactions*, Hoboken, New Jersey, 2010.
- 26 Since metal hydrides may be viewed within an amphoteric construct, we have performed NPA calculations on the anionic cluster of **C1** that is formed on deprotonation of the hydride. Here we employed the 6-31+G(d') basis set for the main group atoms in our calculations which were first performed on **C1** as a benchmark. The charges computed for the three metals and the hydride ligand in **C1** using the added diffuse function mimic those *Q* values quoted in Scheme 6. The charges computed for the anionic cluster show little change, as illustrated in the below structure, and reinforce the more important influence the heterocycle plays in directing the ligand substitution chemistry.



- 27 (a) L. C. Cross and W. Klyne, Appendix, Section E., *Pure Appl. Chem.*, 1976, **45**, 11; (b) E. L. Eliel, S. H. Wilen and L. N. Mander, *Stereochemistry of Organic Compounds*, Wiley, New York, 1994.
- 28 The structural isomer containing adjacent equatorial phosphines at the edge-bridged Os–Os bond and cis to the hydride (**E7**, not shown) was also examined computationally and found to be 9.3 kcal mol<sup>-1</sup> less stable than **E1**. The orientation of the two PPh<sub>3</sub> groups in **E7** should furnish a hydride triplet in the <sup>1</sup>H NMR spectrum, allowing us to exclude this species as a participant isomer.
- 29 (a) M. Day, D. Espitia, K. I. Hardcastle, S. E. Kabir, E. Rosenberg, R. Gobetto, L. Milone and D. Osella, *Organometallics*, 1991, **10**, 3550; (b) E. Rosenberg and R. Kumar, *J. Cluster Sci.*, 2014, **25**, 239.
- 30 (a) E. Band and E. L. Muetteries, *Chem. Rev.*, 1978, **78**, 639; (b) B. F. G. Johnson, *Transition Metal Clusters*, Wiley, New York, 1980; (c) E. Rosenberg, *Polyhedron*, 1989, **8**, 383.
- 31 For reports on the hydride fluxionality involving edge-bridge ⇌ interstitial migration, see: (a) T. Eguchi, B. T. Heaton, R. Harding, K. Miyagi, G. Longoni, J. Nahring, N. Nakamura, H. Nakayama, T. A. Pakkanen, J. Pursianen and A. K. Smith, *Dalton Trans.*, 1996, 625; (b) T. Eguchi and B. T. Heaton, *Dalton Trans.*, 1996, 3523.
- 32 For DFT studies where interstitial hydrides have been computed as reactive intermediates (ground-state species) or transition structures in hydride isomerizations, see: (a) M. J. Hossain, S. Rajbangshi, M. M. M. Khan, S. Ghosh, G. Hogarth, E. Rosenberg, K. I. Hardcastle, M. G. Richmond and S. E. Kabir, *J. Organomet. Chem.*, 2014, **767**, 185; (b) M. A. H. Chowdhury, M. R. Haque, S. Ghosh, S. M. Mobin, D. A. Tocher, G. Hogarth, M. G. Richmond, S. E. Kabir and H. W. Roesky, *J. Organomet. Chem.*, 2017, **836–837**, 68; (c) M. M. M. Khan, M. M. Alam, S. Ghosh, A. Rahaman, D. A. Tocher, M. G. Richmond, S. E. Kabir and H. W. Roesky, *J. Organomet. Chem.*, 2017, **843**, 75; (d) M. A. Alvarez, M. E. García, D. García-Vivó, M. A. Ruiz and A. Toyos, *Dalton Trans.*, 2017, **46**, 15317.
- 33 *CrysAlisPro*; Oxford Diffraction, Yarnton, England, 2015.
- 34 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112.
- 35 O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339.
- 36 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev,



- A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision A.02*, Gaussian, Inc., Wallingford CT, 2009.
- 37 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648.
- 38 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785.
- 39 (a) M. Dolg, U. Wedig, H. Stoll and H. Preuss, *J. Chem. Phys.*, 1987, **86**, 866; (b) S. P. Watch and C. W. Bauschlicher, *J. Chem. Phys.*, 1983, **78**, 4597.
- 40 (a) G. A. Petersson, A. Bennett, T. G. Tensfeldt, M. A. Al-Laham, W. A. Shirley and J. Mantzaris, *J. Chem. Phys.*, 1988, **89**, 2193; (b) G. A. Petersson and M. A. Al-Laham, *J. Chem. Phys.*, 1991, **94**, 6081.
- 41 (a) JIMP2, version 0.091, a free program for the visualization and manipulation of molecules, M. B. Hall and R. F. Fenske, *Inorg. Chem.*, 1972, **11**, 768; (b) J. Manson, C. E. Webster and M. B. Hall, Texas A & M University, College Station, TX, 2006: <http://www.chem.tamu.edu/jimp2/index.html>.

