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Unusual $\text{B}_4\text{N}_2\text{Cl}_2$ Ligand in a Ruthenium Pseudo-Triple-Decker Sandwich Complex Displaying Three Reversible Electron Transfer Steps

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Dedicated to Prof. Richard J. Puddephatt

Polydecker sandwich complexes are highly desirable synthetic targets because of their potential applications as advanced materials, due to the considerable interactions between their metal centers.\(^1\) Cyclopentadienyl\(^2\) and benzene\(^3\) have demonstrated early on their ability to coordinate bifacially to transition metals and generate triple-decker sandwich complexes. Although larger oligomers have been identified in the gas phase,\(^4\) the energy of such bridging $\pi$ interactions involving neutral or singly-charged carbon-based ligands is low and oligomers with more than four decks have not been isolated in the condensed phase. Exceptions are the sandwich compounds of the main group metals that associate forming extended structures in the solid-state.\(^5\) In order to circumvent the low tendency of organic ligands to bridge metal centers and build stacked sandwich compounds, annelated cyclopentadienyl derivatives such as indacene\(^6\) and especially pentalene\(^6\)\(^,\)\(^7\) have been used as building blocks for staggered multidecker sandwich compounds. Although promising, these systems have limitations associated to the narrow choice of substitution patterns available, resulting in low solubility of the higher oligomers or in the formation of complex mixtures of isomers. The recent synthesis of hexamethylenalene is a promising development in this area.\(^8\) Heterocyclic rings, particularly those containing boron, have a higher tendency to coordinate bifacially and generate stacked sandwich compounds.\(^9\) The largest multidecker sandwich compounds known to date contain six decks and feature dianionic cyclic ligands with $\text{B}_3\text{C}_3$ and $\text{B}_3\text{C}_2$ frameworks.\(^10\) The only family of polydecker complexes reported so far, [Ni($\text{C}_9\text{B}_9\text{Me}_3$)]\(_n\), displayed conducting properties.\(^11\)

Within the scope of our investigation on heterocyclic cyclopentadienyl analogues we developed the 1,2-diaza-3,5-diboryl ligands \(1\).\(^12\) Herein we report the extension of this project to the pentacenediyl-like dianion in \(3\), which is a very promising $\pi$-bridging ligand for the assembly of polymeric materials.

The precursor \(2\) was obtained as a mixture of cis and trans isomers through condensation of hydrazine and 1,1-bis(phenylchloroboryl)methane\(^12\)\(^b\) in the presence of triethylamine (see Supporting Information). The trans isomer could be separated by crystallization and structurally characterized (Fig. 1).\(^13\) It features a long N-N bond of 1.48 Å, slightly longer than the N-N bonds observed in hydrazine (1.45 Å)\(^14\) and the metal complexes of \(1\) (1.44 – 1.47 Å).\(^12\) The endocyclic B-C and B-N bonds are, at 1.57 and 1.44 Å, longer and slightly shorter, respectively, than the corresponding bonds in the alkali metals salts of the monocyclic analogs.\(^12\)\(^b\) For comparison, the B-N bonds in borazines measure 1.42 – 1.44 Å.\(^15\)

Double deprotonation of \(2\) with K[N(SiMe\(_3\)=)] could be conducted directly or in two successive steps, yielding the orange dipotassium salt \(3\). As expected, both isomers produced the same mono and dianions. The disappearance of the methine protons was noticeable in the $\text{H}^\text{1}$ NMR spectrum of \(3\), and the deprotonation resulted in a 14 ppm upfield shift of the $\text{B}^\text{11}$ NMR signal. Despite its very similar geometry, the 8-$\pi$-electron heterocyclic ligand in \(3\) is

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**Scheme 1.** Synthesis of derivatives 2–4.

**Figure 1.** Molecular structure of trans-2. Selected bond lengths [Å] and angles [°]: N-N 1.480(2), B-N 1.436(2), 1.437(2), B-C(1) 1.571(2), 1.574(2), B-N-B 140.88(14), 109.58(14), N-B-C(1) 109.02(14), B-C-B 101.86(13), B-N-N 109.54(15).

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A crystal structure was determined for 3(tmeda), revealing that the planar B₅N₅C₂ ligand was coordinated by two potassium ions each supporting a TMEDA molecule, in a centrosymmetric arrangement (Fig. 2). The transition from 2 to 3 involves a shortening of the intraannular N-N and B-C bonds and a slight lengthening of the B-N bonds, resembling the transition from cyclopentadiene to cyclopentadienyl. The separation between the two potassium ions measures 5.52 Å and is comparable to the K⋯K separation observed in polymeric cyclopentadienyls (5.52 – 5.85 Å) and the dimeric potassium pentalenediyl derivative (5.43, 5.48 Å).

A computational analysis at the DFT level (see Supporting Information) performed on the model structure 4a, featuring a hydrogen substituted B₅N₅C₂H₅ ligand and Cp groups, gave a molecular geometry in reasonable agreement with the X-ray diffraction data. The frontier MO analysis of 4a shows that substantial mixing of the ligand orbitals with the MOs of the [CpRu]⁺ fragments takes place, resulting in a formal insertion of the metal atoms into the N-N bond. The ruthenium d orbitals interact mainly with the two nitrogen p orbitals which are orientated in the B-N=C plane and perpendicular to it, respectively. The similar oxidative addition of hydrazines to transition metals has recently been reported. Energy decomposition analysis conducted for 4a confirms that the primary, thermodynamic driving force for its formation is the cleavage of the N-N bond, and the subsequent formation of the Ru-N interactions: the total orbital interaction energy is -2400 kJ mol⁻¹. Mayer bond orders obtained for 4a support the description of Ru(2) as η⁶-coordinated whereas coordination of Ru(1) can be best described as η⁵; the largest bond orders are calculated for Ru-N interactions, around 0.45. As expected, the bonding analysis shows no sign of N-N interactions. Hence, the computational data indicates that the atypical linear B-N-B moieties in 4 can be described with formally sp-hybridized nitrogen atoms whose unhybridized p orbitals have an important contribution to the ligand-metal bonding.

Cyclic voltammetry of 4 in THF revealed two reversible reduction steps at -1.30 and -1.99 V vs. SCE and one irreversible oxidation at 0.42 V. The expansion of the electrochemical window using DME confirmed this behaviour and allowed for the observation of an additional, irreversible oxidation at 1.10 V, while the electrochemical range of CH₃Cl₂ enabled the observation of only one reduction and one oxidation step at -1.45 and 0.18 V. All one-electron transfer steps are well separated, indicating significant electron delocalization over the framework. The reversible oxidation likely involves the Ru⁺/Ru²⁺ process while the irreversible electron transfer could be attributed to the oxidation of the Cp ligand, as observed for Cp*Ru (0.55 and 1.25 in CH₂Cl₂). For Cp*Ru₂C₆H₆ (0.11 and 0.40 V in THF, -0.02 and 0.48 in CH₂Cl₂) and
(Cp*Ru)2B14H14 (0.17 and 0.93 V in 8.5:1.5 CH2Cl2/toluene).17a The reduction steps have no parallel in the chemistry of related species containing all-carbon and all-boron ligands and are attributed to processes centred mostly on the B,N,C2- ligand. A theoretical analysis supports this view, as the optimized geometry of [4a]- is C2v symmetric and contains a planar B,N,C2 framework with no apparent N-N interaction. A two-electron oxidation was reported for the related transoid-[Cp(Ru)3C2H4] complex, resulting in cleavage of a C-C bond in the bridging ligand and formation of a flyover dication.17a

In conclusion, a pentalenediyl-like dianion with B,N,C2 framework was isolated as a dipotassium salt,3 which in turn contains a planar B,N,C2 framework with no E4 symmetric and contains a planar B,N,C2 skeleton. A computational analysis showed that this ligand could be considered a distorted form of the bicyclic ligand, brought about by the coordination to the transition metal. The cyclic voltammogram of 4 displayed three reversible, well separated one-electron transfer steps indicating efficient electron delocalization over the framework. Its ligand properties render the highly unusual π-bridging ligand a very promising building block for the design of larger sandwich-like architectures.