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Higher Spin States in Some Low-Energy Bis(Tetramethyl-1,2-diaza-3,5-diboroly) Sandwich Compounds of the First Row Transition Metals: Boraza Analogues of the Metallocenes

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Abstract

The known sandwich compound $[\eta^5-(\text{CH}_2)_3\text{N}_2(\text{BPh})_2\text{CMe}]_2\text{Fe}$ in which adjacent C_2 units are replaced by isoelectronic BN units can be considered as a boraza analogues of ferrocene similar to borazine, $\text{B}_3\text{N}_3\text{H}_6$, considered as a boraza analogue of benzene. In this connection, the related bis(1,2,3,5-tetramethyl-1,2-diaza-3,5-diboroly) derivatives $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ ($\text{M} = \text{Ti}, \text{V}, \text{Cr}, \text{Mn}, \text{Fe}, \text{Co}, \text{Ni}$) for all of the first row transition metals have been optimized using density functional theory for comparison with the isoelectronic tetramethylcyclopentadienyl derivatives $(\text{Me}_4\text{C}_5\text{H})_2\text{M}$. Low-energy sandwich structures having parallel $\text{B}_2\text{N}_2\text{C}$ rings in a *trans* orientation are found for all seven metals. The 1,2-diaza-3,5-diboroly ligand appears to be a weaker field ligand than the isoelectronic cyclopentadienyl ligand as indicated by higher spin ground states for some $(\eta^5-\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ sandwich compounds relative to the corresponding metallocenes $(\eta^5-\text{Me}_4\text{C}_5\text{H})_2\text{M}$. Thus $(\eta^5-\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Cr}$ has a quintet ground state in contrast to the triplet ground state of $(\eta^5-\text{Me}_4\text{C}_5\text{H})_2\text{Cr}$. Similarly, the sextet ground state of $(\eta^5-\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Mn}$ lies ~ 18 kcal/mol below the quartet state in contrast to the doublet ground state of the isoelectronic $(\text{Me}_4\text{C}_5\text{H})_2\text{Mn}$. These sandwich compounds are potentially accessible by reaction of 1,2-diaza-3,5-diborolide anions with metal halides analogous to the synthesis of $[\eta^5-(\text{CH}_2)_3\text{N}_2(\text{BPh})_2\text{CMe}]_2\text{Fe}$.

1. Introduction

Replacement of adjacent pairs of carbon atoms in cyclic hydrocarbons with isoelectronic BN units leads to inorganic boraza analogues of familiar hydrocarbons. For example, borazine or “inorganic benzene,” $B_3N_3H_6$, is the boraza analogue of benzene obtained by complete replacement of the three C_2 units in the six-membered ring of benzene with BN units. Molecular orbital calculations and experimental results for the transmission of substituent effects through the B_3N_3 ring indicate only partial delocalization of the π electrons in borazine as compared with their complete delocalization in benzene [1, 2]. Nevertheless, hexamethylborazine forms a stable chromium tricarbonyl complex ($\eta^6\text{-Me}_6B_3N_3$)Cr(CO) $_3$ (Figure 1) [3, 4] analogous to the well-known benzene chromium tricarbonyl ($\eta\text{-C}_6H_6$)Cr(CO) $_3$ and its derivatives.

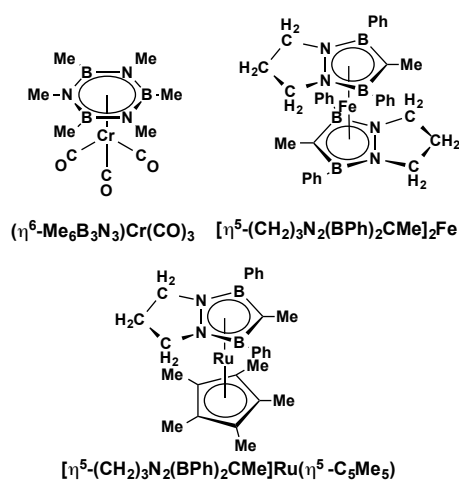


Figure 1. Metal complexes of borazine and diazadiborolyl ligands that have been synthesized.

This paper explores the structures and spin states of the boraza analogues of the metallocenes. An initial step in this direction was taken in 2002 by Fu and coworkers [5]. They used indirect synthetic methods to replace a pair of carbon atoms in one of the cyclopentadienyl rings of ferrocene by an BN pair to give the pentahapto 1,2-azaborolyl-iron derivatives $[\eta^5\text{-C}_3H_3N(tBu)BX]Fe(\eta^5\text{-C}_5H_5)$ ($X = H, F, \text{allyl}, OMe, NMe_2, \text{etc.}$). More extensive substitution of adjacent carbon pairs in ferrocene with BN units to the limit of a $(\eta^5\text{-R}_5B_2N_2C)_2Fe$ derivative was achieved by Roesler and coworkers [6, 7]. They synthesized a diazadiborolyl anion, which was subsequently used to prepare a substituted boraza analogue of ferrocene, namely $[\eta^5\text{-(CH}_2)_3N_2(BPh)_2CMe]_2Fe$ as well as the mixed ruthenium sandwich compound $[\eta^5\text{-(CH}_2)_3N_2(BPh)_2CMe]Ru(\eta^5\text{-C}_5Me_5)$ (Figure 1) [8].

The laboratory availability of the $[\eta^5-(\text{CH}_2)_3\text{N}_2(\text{BPh})_2\text{CMe}]^-$ anion suggests the possibility of synthesizing boraza analogues of the entire series of the first row transition metal metallocenes similar to the ferrocene analogue $[\eta^5-(\text{CH}_2)_3\text{N}_2(\text{BPh})_2\text{CMe}]_2\text{Fe}$. We report here the use of density functional theory to explore the preferred structures and spin states of the boraza metallocene analogues $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ ($\text{M} = \text{Ti}, \text{V}, \text{Cr}, \text{Mn}, \text{Fe}, \text{Co}, \text{Ni}$). The diazadiborole ligand with methyl substituents on the boron and nitrogen atoms was chosen as the simplest reasonable model for the experimental systems.

Experimental studies by Roesler and coworkers on $[\eta^5-(\text{CH}_2)_3\text{N}_2(\text{BPh})_2\text{CMe}]_2\text{Fe}$ [6] as well as the earlier studies by Fu and coworkers [5] showed the boraza analogues of the cyclopentadienyl ligand to be more electron-donating than the cyclopentadienyl ligand itself. In this connection our studies on the $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ metallocene analogues predict a greater preference for higher spin states relative to the $(\eta^5\text{-C}_5\text{H}_5)_2\text{M}$ and $(\eta^5\text{-Me}_4\text{C}_4\text{H})_2\text{M}$ metallocenes. Particularly noteworthy is the energetic preference for a sextet spin state for the manganese derivative $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Mn}$ by nearly 18 kcal/mol. In addition a quintet spin state for the chromium derivative $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Cr}$ is found to be energetically competitive with the triplet spin state, which is the clear ground state for the cyclopentadienyl analogues $(\eta^5\text{-C}_5\text{H}_5)_2\text{Cr}$ [9] and $(\eta^5\text{-Me}_4\text{C}_4\text{H})_2\text{Cr}$.

2. Theoretical Methods

Density functional theory (DFT), including electron correlation effects, is a powerful computational tool that has been successfully used in organotransition metal chemistry [10,11,12,13,14,15,16]. Two DFT methods were used for this work. The first method is the hybrid HF/DFT functional B3LYP* [17,18], which is the modified B3LYP functional [19,20] in order to provide electronic state orderings in good agreement with experiment [21]. The second method is a meta-GGA DFT method M06-L developed by Truhlar's group [22]. The M06-L method is found to predict energy differences close to the experimental values for the transition metal chemistry [23]. We discuss the M06-L results in the text, while the B3LYP* results are listed in the Supporting Information. In fact, the energy orderings predicted by the B3LYP* method are generally consistent with those predicted by the M06-L method.

The double- ζ plus polarization (DZP) basis sets used in this paper for the boron, carbon, and nitrogen atoms add one set of pure spherical harmonic d functions with orbital exponents, $\alpha_d(\text{B}) = 0.70$, $\alpha_d(\text{C}) = 0.75$, and $\alpha_d(\text{N}) = 0.80$, to the standard Huzinaga-Dunning contracted DZ sets and are denoted as (9s5p1d/4s2p1d) [24, 25]. For hydrogen, a set of p polarization functions $\alpha_p(\text{H}) = 0.75$ is added to the Huzinaga-Dunning DZ sets. For the transition metal atoms, our loosely contracted DZP basis set

(14s11p6d/10s8p3d) uses the Wachters' primitive set [26] augmented by two sets of p functions and one set of d functions contracted following Hood, Pitzer, and Schaefer [27].

All computations were carried out using the Gaussian 09 program [28]. For the M06-L functional the (120,974) grid was used for numerical evaluation of the integrals. In the present paper the $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ structures are designated using the format **M-x-n**, where **M** stands for the transition metal, **x** for the spin state, and **n** for the angle in degrees between the H-CB₂N₂ bond in each ring projected onto a plane of the other ring. The spin multiplicities are indicated by **s**, **d**, **t**, **q**, **p**, and **x** for singlets, doublets, triplets, quartets, quintets, and sextets, respectively. Thus, the triplet $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ structure with a *trans* relationship between the two B₂N₂C rings is designated as **Ti-t-180**. The corresponding metallocene $(\eta^5\text{-Me}_4\text{C}_5\text{H})_2\text{M}$ structures are designated as **c-M-x-n**. Thus, the triplet $(\eta^5\text{-Me}_4\text{C}_5\text{H})_2\text{Ti}$ structure with a *cis* relationship between the two cyclopentadienyl rings is designated as **c-Ti-t-0**.

3. Results and Discussion

3.1 $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$

Four low-energy $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ structures (two singlets and two triplets) were found by the DFT methods (Figure 2). Quintet structures were found to lie at least 50 kcal/mol in energy above the lowest energy structure. They thus do not appear to be chemically significant and therefore are not discussed in this paper. In the triplet $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ structure **Ti-t-180** the two B₂N₂C rings are parallel. In the other three structures the two B₂N₂C rings are not parallel, but have dihedral angles of ~30° for the two singlet structures and ~15° for the triplet **Ti-t-144**. Thus the O-Ti-O' angle for **Ti-t-144** is 166.6° whereas the O-Ti-O' angles for **Ti-s-144** and **Ti-s-0** are 150.7° and 152.6°, respectively where O and O' represent the centers of the two rings. Since the B₂N₂C ring is a five-electron donor like the cyclopentadienyl ligand, the titanium atom in the $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ complexes has a 14-electron configuration.

The lowest-energy $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ structure is the C₂ triplet **Ti-t-144** with a rotation angle of 144° between the two B₂N₂C rings (Figure 2). A C_s triplet structure **Ti-t-180** lies only 1.4 kcal/mol in energy above **Ti-t-144**. Structure **Ti-t-180** has a very small 15i cm⁻¹ imaginary vibrational frequency (M06-L) that may be regarded as arising from numerical integration errors. Following the normal mode corresponding to the imaginary vibrational frequency of **Ti-t-180** leads to **Ti-t-144**.

The two singlet $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ti}$ structures **Ti-s-144** (C₂) and **Ti-s-0** (C_s) are significantly higher energy structures, lying 12.3 and 16.0 kcal/mol, respectively, above

Ti-t-144 (M06-L). These two singlet structures have similar geometries, differing only by the rotation of one B_2N_2C ring relative to the other (Figure 2).

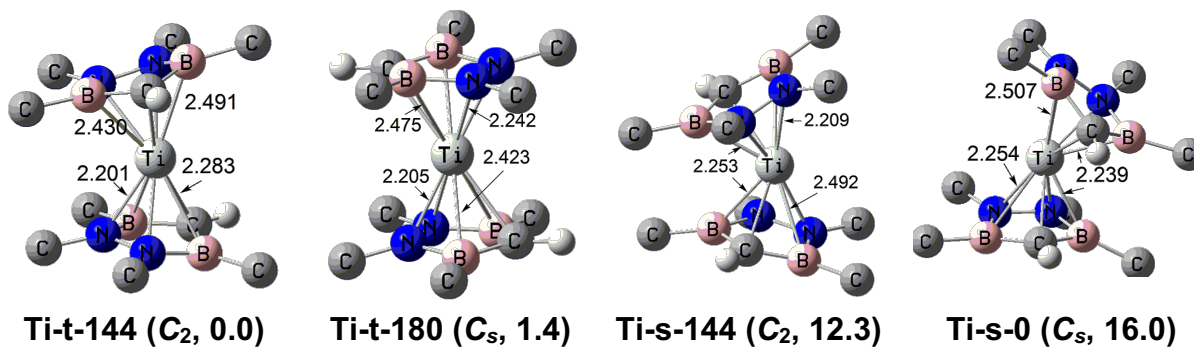


Figure 2. The low-energy $(Me_4B_2N_2CH)_2Ti$ structures using the M06-L method. In Figures 2 to 15, the bond distances are in Å. The relative energies without ZPVE correction (ΔE , in kcal/mol) are given in parentheses. For clarity, the hydrogen atoms in the methyl groups are not shown.

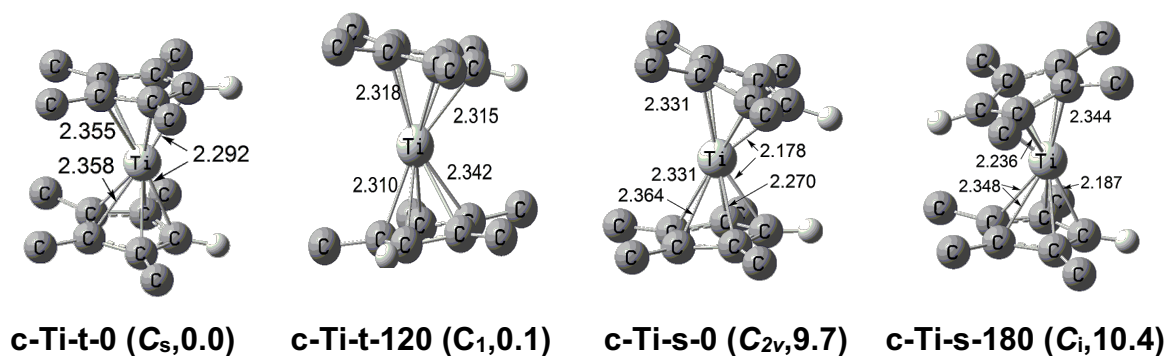


Figure 3. The low-energy $(Me_4C_5H)_2Ti$ structures using the M06-L method.

Four low-energy $(\eta^5-Me_4C_5H)_2Ti$ structures (two triplets and two singlets) were found (Figure 3). The lowest-energy structure is the C_s triplet **c-Ti-t-0**. Another triplet structure **c-Ti-t-120** (C_1) lies only 0.1 kcal/mol in energy above **c-Ti-t-0**. The two singlet $(\eta^5-Me_4C_5H)_2Ti$ structures **c-Ti-s-0** (C_{2v}) and **c-Ti-s-180** (C_i) are of higher energy, lying 9.7 and 10.4 kcal/mol (M06-L), respectively, above **c-Ti-t-0**. In **c-Ti-t-0** the two cyclopentadienyl rings are almost parallel. However, in the other three $(\eta^5-Me_4C_5H)_2Ti$ structures the two cyclopentadienyl rings are not parallel, but have a dihedral angle of $\sim 30^\circ$ for the two singlet structures and $\sim 15^\circ$ for the triplet structure, similar to **Ti-t-144**, **Ti-s-144**, and **Ti-s-0**.

3.2 (Me₄B₂N₂CH)₂V

Only one low-energy quartet (η^5 -Me₄B₂N₂CH)₂V structure **V-q-180** (*C*_{2h}) was found (Figure 4). The corresponding doublet (η^5 -Me₄B₂N₂CH)₂V structure **V-d-180** is a high-energy structure, lying 32.3 kcal/mol in energy above **V-q-180**. Structure **V-q-180** is a genuine minimum adopting the expected parallel sandwich geometry. In **V-q-180** the V–C distances are ~2.33 Å, the V–B distances are ~2.38 Å, while the V–N distances are significantly shorter at 2.16 Å (M06-L). These distances suggest a five-electron donor pentahapto B₂N₂C ring thereby giving the vanadium atom in **V-q-180** a 15-electron configuration.

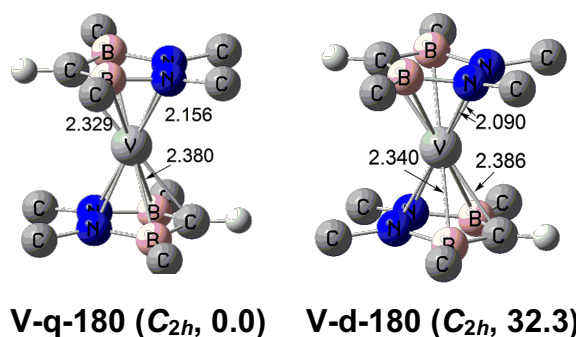


Figure 4. The low-energy (Me₄B₂N₂CH)₂V structures using the M06-L method.

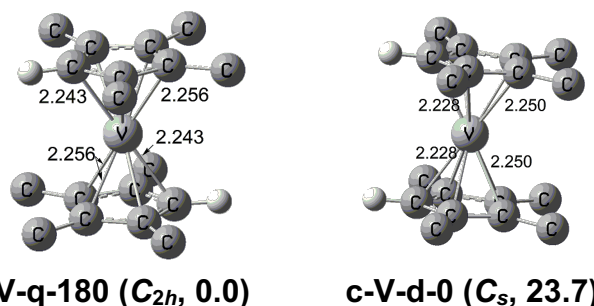


Figure 5. The low-energy (Me₄C₅H)₂V structures using the M06-L method.

Two low-energy (η^5 -Me₄C₅H)₂V structures were found (Figure 5). The quartet structure **c-V-q-180** (*C*_{2h}) is a genuine minimum adopting the expected parallel sandwich geometry. Another minimum, the doublet **c-V-d-0** (*C*_s) is of higher energy, lying 23.7 kcal/mol above **c-V-q-180** (M06-L). In the two (η^5 -Me₄C₅H)₂V structures the V–C distances range from 2.22 to 2.26 Å, which are close to the experimental V–C distances of 2.280 Å in vanadocene, (η^5 -C₅H₅)₂V. [29]

3.3 (Me₄B₂N₂CH)₂Cr

Five low-energy (η^5 -Me₄B₂N₂CH)₂Cr structures (three quintets and two triplets), all with parallel B₂N₂C rings, were found by the DFT methods (Figure 6). The lowest-energy such structure is the C_i quintet **Cr-p-180**. The two C₂ quintet conformers **Cr-p-144** and **Cr-p-36** have energies within 0.8 kcal/mol (M06-L) of **Cr-p-180**. In the three quintet structures the Cr-C distances range from 2.23 to 2.26 Å, the Cr-B distances from 2.32 to 2.50 Å, and the Cr-N distances from 2.14 to 2.40 Å. These interatomic distances indicate two pentahapto B₂N₂C ligands to give each chromium atom a 16-electron configuration. The two triplet (η^5 -Me₄B₂N₂CH)₂Cr structures **Cr-t-180** and **Cr-t-72** are higher energy structures than their quintet isomers, lying 7.9 and 12.0 kcal/mol, respectively, above **Cr-p-180** (M06-L). They also have pentahapto B₂N₂C ligands and thus 16-electron configurations for the chromium atoms.

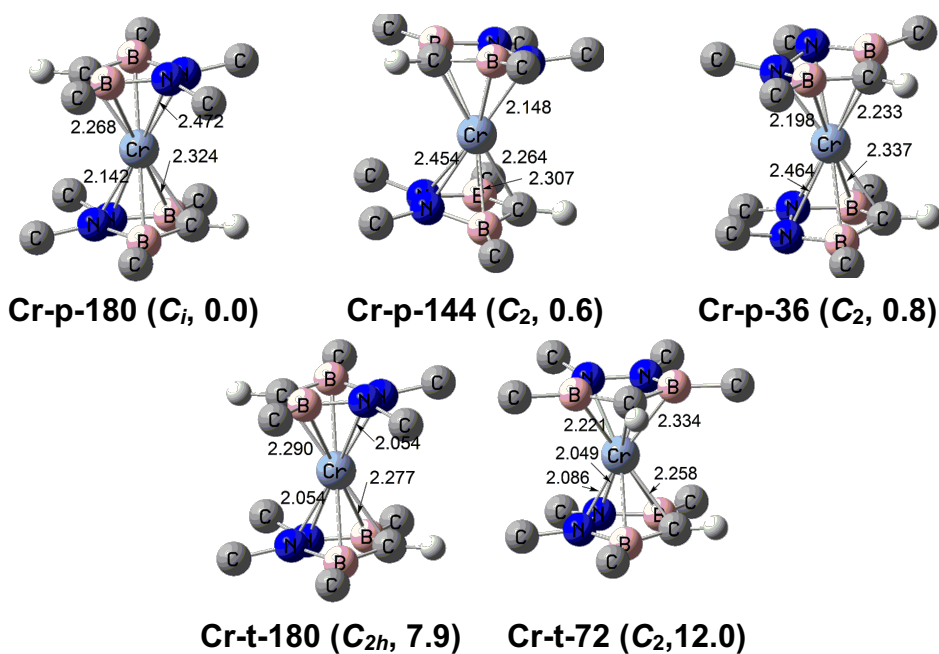


Figure 6. The low-energy (Me₄B₂N₂CH)₂Cr structures using the M06-L method.

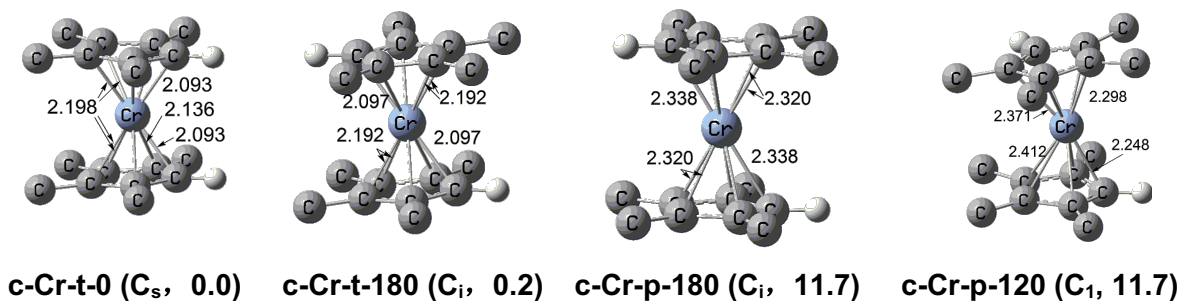


Figure 7. The low-energy (Me₄C₅H)₂Cr structures using the M06-L method.

Four low-energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Cr structures were found (Figure 7). The lowest energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Cr structure is the C_s triplet **c-Cr-t-0**. A triplet structure **c-Cr-t-180** (C_i) lies only 0.2 kcal/mol in energy above **c-Cr-t-0**. The two quintets **c-Cr-p-180** (C_i) and **c-Cr-p-120** (C₁) are of higher energy, lying 11.7 kcal/mol above **c-Cr-t-0**. This is different from the (Me₄B₂N₂CH)₂Cr system for which the quintet structures of lower energies than their triplet isomers. In the two triplet structures the Cr-C distances range from 2.10 to 2.20 Å, which are shorter than those in the quintet structures. The Cr-C distances range from 2.09 to 2.19 Å. This is in reasonable agreement with the experimental Cr-C distances (2.169 Å) in chromocene, ($\eta^5\text{-C}_5\text{H}_5$)₂Cr. [29]

3.4 (Me₄B₂N₂CH)₂Mn

Five low-energy ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Mn structures (one sextet, one doublet, and three quartets) have been optimized using the DFT methods (Figure 8). All five structures are predicted to have the expected parallel sandwich structure. The C₂ sextet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Mn structure **Mn-x-144** is the lowest energy structure. It is a very favorable structure since it lies ~18 kcal/mol in energy below the next lowest energy (Me₄B₂N₂CH)₂Mn structure. Such a high spin sextet ground state is favored by the half-filled d-shell in the d⁵ configuration of the central Mn(II) atom in the sandwich compound (Me₄B₂N₂CH)₂Mn.

The C_i doublet structure **Mn-d-180** lies 21.6 kcal/mol in energy above **Mn-x-144** (M06-L). The three quartet structures **Mn-q-180** (C_i), **Mn-q-36** (C₂), and **Mn-q-144** (C₂) lie 17.9, 18.9, and 24.6 kcal/mol, respectively, above **Mn-x-144** (M06-L). In the five (Me₄B₂N₂CH)₂Mn structures the Mn-C distances range from 2.15 to 2.19 Å, the Mn-B distances range from 2.17 to 2.53 Å, and the Mn-N distances range from 2.01 to 2.51 Å indicating pentahapto B₂N₂C ligands in all cases. Since each Me₄B₂N₂CH ligand is a five-electron donor like the isoelectronic cyclopentadienyl ligand, the manganese atoms in the five structures all have 17-electron configurations. The low energy of the sextet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Mn structure **Mn-x-144** relative to isomers with lower spin states clearly indicates that the B₂N₂C ligand is a weaker field ligand than the cyclopentadienyl or substituted cyclopentadienyl ligands as well as other five-electron cyclic donor ligands forming similar manganese sandwich complexes [30].

Four low-energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Mn structures (one doublet, two sextets, and one quartet) have been optimized using the DFT methods (Figure 9). The lowest energy structure is the doublet **c-Mn-d-0**. Two sextet ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Mn structures **c-Mn-x-90** and **c-Mn-x-180** lie 6.0 and 7.6 kcal/mol, respectively, above **c-Mn-d-0**. The quartet

structure **c-Mn-q-0** (C_{2v}) lies 14.9 kcal/mol above **c-Mn-d-0** (M06-L). In three of the $(\eta^5\text{-Me}_4\text{C}_5\text{H})_2\text{Mn}$ structures, the two cyclopentadienyl rings are almost parallel, while in structure **c-Mn-x-90** the two cyclopentadienyl rings have a dihedral angle of $\sim 30^\circ$. In the doublet **c-Mn-d-0** the Mn-C distances range from 2.05 to 2.13 Å, while in the quartet **c-Mn-q-0** the Mn-C distances range from 2.13 to 2.33 Å. These results are significantly shorter than the experimental Mn-C distances (2.380 Å) in manganocene, $(\eta^5\text{-C}_5\text{H}_5)_2\text{Mn}$. [31]. This difference appears to be a consequence of the sextet spin state of unsubstituted manganocene and the doublet spin state of **c-Mn-d-0**.

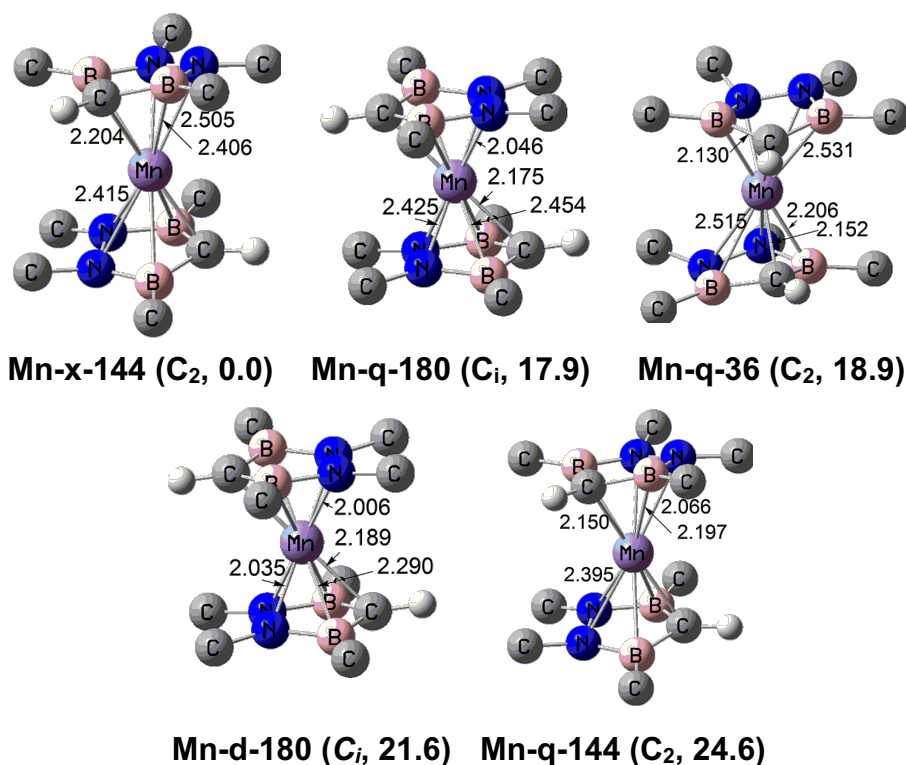


Figure 8. The low-energy $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Mn}$ structures using the M06-L method.

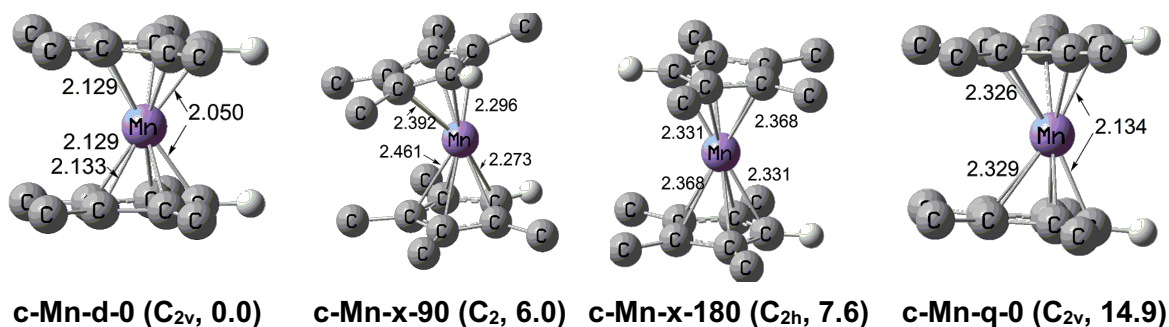


Figure 9. The lowest-energy $(\text{Me}_4\text{C}_5\text{H})_2\text{Mn}$ structures using the M06-L method.

3.5 (Me₄B₂N₂CH)₂Fe

Six low-energy (η^5 -Me₄B₂N₂CH)₂Fe structures (two singlets, one quintet, and three triplets) were found (Figure 10). All six structures are predicted to have the expected sandwich structure with parallel B₂N₂C rings. The global minimum (η^5 -Me₄B₂N₂CH)₂Fe structure is the C_{2h} singlet structure **Fe-s-180**. The other singlet (η^5 -Me₄B₂N₂CH)₂Fe structure **Fe-s-72** lies 3.5 kcal/mol (M06-L) in energy above **Fe-s-180**. (Figure 10), Our predicted Fe-N distances (~1.97 Å), Fe-C distances (~2.15 Å) and Fe-B distances (~2.17 Å) in the singlet (Me₄B₂N₂CH)₂Fe structures **Fe-s-180** and **Fe-s-72** are in very good agreement with the experimental Fe-N, Fe-C, and Fe-B distances of ~1.97, 2.17, and ~2.20 Å, respectively, in [η^5 -(CH₂)₃N₂(BPh)₂CMe]₂Fe, as determined by X-ray crystallography [8]. The B-C, B-N, and N-N bond distances in the B₂N₂C rings are also in very good agreement with corresponding experimental values (Figure 10). X-ray crystallography of [η^5 -(CH₂)₃N₂(BPh)₂CMe]₂Fe shows that the angle between the B₂N₂C-CH₃ bonds is not 180°, but close to the angle between the B₂N₂C-H bonds of structure **Fe-s-72**. This could be a consequence of increased steric hindrance of the phenyl groups in a [η^5 -(CH₂)₃N₂(BPh)₂CMe]₂Fe structure of the type **Fe-s-180** relative to a structure of the type **Fe-s-72**.

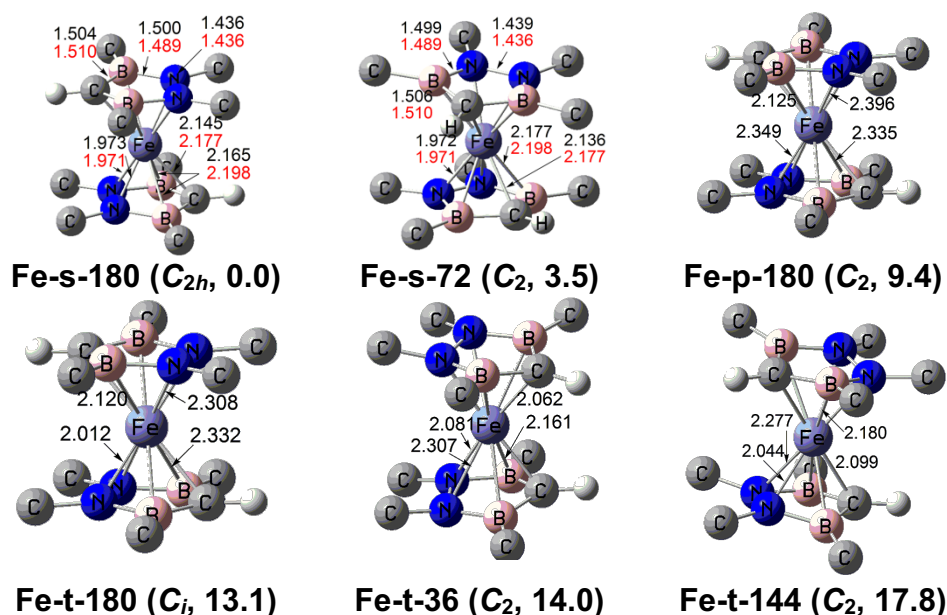


Figure 10. The low-energy (Me₄B₂N₂CH)₂Fe structures using the M06-L method. The lower numbers in red for structures **Fe-s-180** and **Fe-s-72** are the corresponding experimental values for the [η^5 -(CH₂)₃N₂(BPh)₂CMe]₂Fe [8].

The C_{2h} quintet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Fe structure **Fe-p-180** lies 9.4 kcal/mol above **Fe-s-180** (Figure 10). The triplet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Fe structures **Fe-t-180** (C_i), **Fe-t-36** (C_2), and **Fe-t-144** (C_2) are even higher energy structures, lying 13.1, 14.0, and 17.8 kcal/mol (M06-L), respectively, above **Fe-s-180**. In the triplet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Fe structures the Fe-C distances are 2.06 to 2.23 Å, the Fe-B distances are 2.16 to 2.33 Å, and the Fe-N distances are 2.01 to 2.31 Å, suggesting pentahapto $\text{Me}_4\text{B}_2\text{N}_2\text{CH}$ ligands. This gives the iron atoms in even these higher spin state ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Fe structures the favored 18-electron configurations.

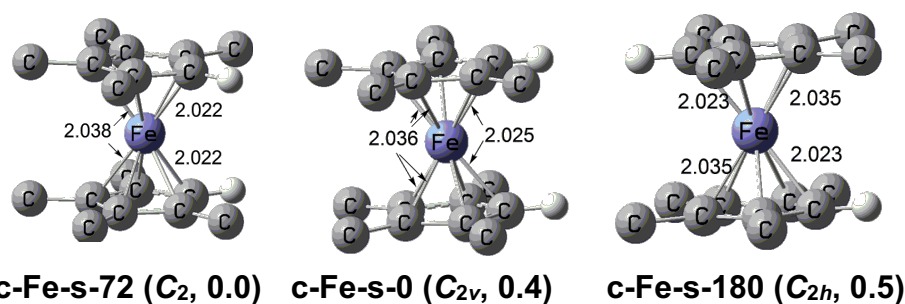


Figure 11. The lowest-energy ($\text{Me}_4\text{C}_5\text{H}$)₂Fe structures using the M06-L method.

Only three low-energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Fe structures were found, all singlets with the favored 18-electron configuration (Figure 11). The triplet and quintet isomers were found to lie least 30 kcal/mol higher in energy and thus are not discussed in this paper. These three low-lying structures are predicted to have the expected parallel sandwich geometry similar to ferrocene. The lowest energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Fe structure **c-Fe-s-72** has C_2 symmetry, while the other isomers **c-Fe-s-0** (C_{2v}) and **c-Fe-s-180** (C_{2h}) have only slightly higher energies of 0.4 and 0.5 kcal/mol, respectively (M06-L). In these three singlet ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Fe structures the Fe-C distances are in a narrow range from 2.02 to 2.03 Å, which are close to the corresponding experimental Fe-C distances (2.064 Å) of the well-known ferrocene, ($\eta^5\text{-C}_5\text{H}_5$)₂Fe [32].

3.6 ($\text{Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Co

Four low-energy ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Co structures (two doublets and two quartets) were found (Figure 12). All four structures adopt the expected parallel sandwich structures. The global minimum is the C_{2h} doublet structure **Co-d-180**. The other doublet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Co structure **Co-d-36** is a C_2 structure lying 1.7 kcal/mol in energy above **Co-d-180**. The quartet ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Co structures **Co-q-180** (C_{2h}) and **Co-q-36** (C_2) are higher energy structures, lying 8.7 and 13.6 kcal/mol (M06-L),

respectively, in energy above **Co-d-180**. In these $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Co}$ structures the Co-C distances range from 2.08 to 2.15 Å, the Co-B distances range from 2.15 to 2.33 Å, and the Co-N distances range from 1.96 to 2.30 Å (M06-L). This indicates pentahapto $\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$ ligands, thereby giving each cobalt atom a 19-electron configuration.

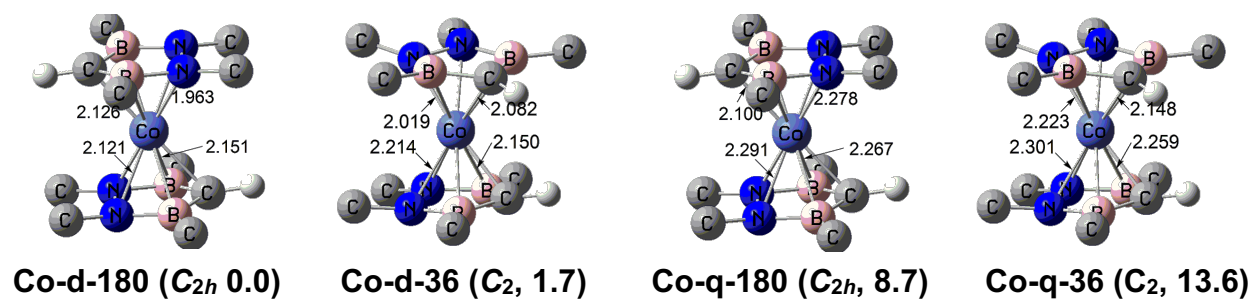


Figure 12. The low-energy $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Co}$ structures using the M06-L method.

Three low-energy $(\eta^5\text{-Me}_4\text{C}_5\text{H})_2\text{Co}$ structures have been located with closely spaced relative energies. The C_{2h} structure **c-Co-d-180** has the lowest energy, but structures **c-Co-d-0** (C_{2v}) and **c-Co-d-36** (C_2) lie only 0.7 and 1.2 kcal/mol, respectively, above **c-Co-d-180**. All three structures are doublet spin state structures, predicted to have the expected parallel sandwich structure (Figure 13). The Co-C distances in these structures range from 2.05 to 2.13 Å (M06-L) in agreement with the corresponding experimental Co-C distances in (2.119 Å) of cobaltocene, $(\eta^5\text{-C}_5\text{H}_5)_2\text{Co}$ [33].

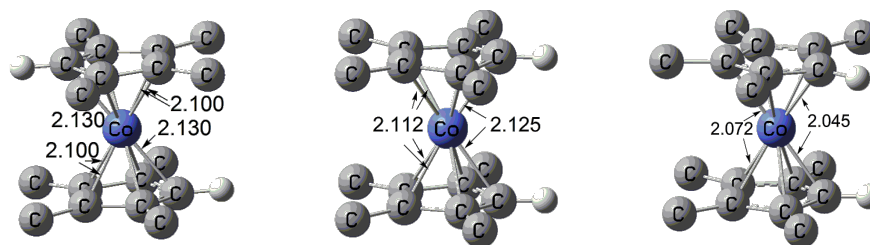


Figure 13. The lowest-energy $(\text{Me}_4\text{C}_5\text{H})_2\text{Co}$ structures using the M06-L method.

3.7 $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ni}$

Only two low-energy $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ni}$ structures **Ni-s-180** (C_{2h}) and **Ni-t-180** (C_{2h}) were found (Figure 14). These structures adopt the expected parallel sandwich geometry and are almost identical in energy with the singlet structure **Ni-s-180** lying 1.2 kcal/mol below the triplet structure **Ni-t-180**. Both $\text{B}_2\text{N}_2\text{C}$ rings in the two $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Ni}$ structures are pentahapto $\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$ ligands, thereby giving the

nickel atoms 20-electron configurations similar to the well-known nickelocene ($\eta^5\text{-C}_5\text{H}_5$)₂Ni.

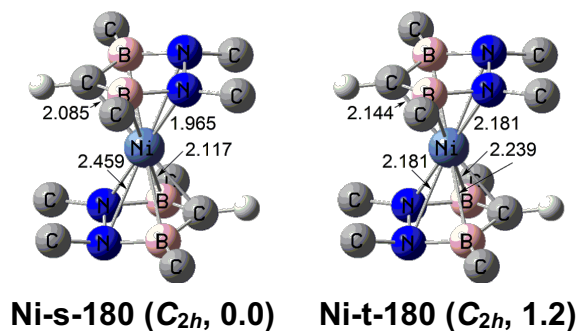


Figure 14. The low-energy ($\text{Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Ni structures using the M06-L method.

Four low-energy ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Ni structures (two singlets and two triplets) were found (Figure 15). All four structures adopt the expected parallel sandwich geometry. The two triplet structures **c-Ni-t-180** and **c-Ni-t-0** have almost the same energies (within 0.05 kcal/mol), whereas the two singlet structures **c-Ni-s-180** (C_i), **c-Ni-s-0** (C_{2v}) have significantly higher energies at 15.2 and 17.4 kcal/mol (M06-L), respectively, above the triplet structure **c-Ni-t-180**. In both triplet structures **c-Ni-t-180** and **c-Ni-t-0**, the Ni-C distances range from 2.16 to 2.17 Å (M06-L) in reasonable agreement with the corresponding experimental Ni-C distances of 2.196 Å in nickelocene, ($\eta^5\text{-C}_5\text{H}_5$)₂Ni [34].

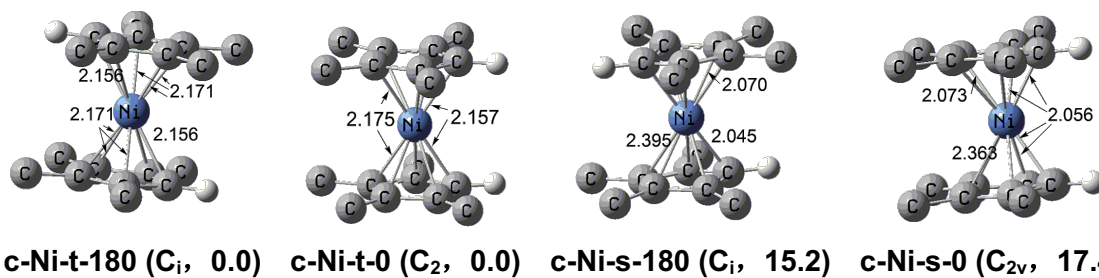
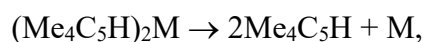


Figure 15. The lowest-energy ($\text{Me}_4\text{C}_5\text{H}$)₂Ni structures using the M06-L method.

3.8 Dissociation Energies

In order to study the stability of complexes under study, we have explored the dissociation energies for the following reactions:



where M = Ti, V, Cr, Mn, Fe, Co, and Ni are the free atoms in their electronic ground states. All of the dissociation energies are found to be substantial (Figure 16). Thus

the DE values for the $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ compounds range from 127 to 193 kcal/mol, while those for the $(\text{Me}_4\text{C}_5\text{H})_2\text{M}$ compounds range from 127 to 204 kcal/mol.

In both cases, the DE values do not monotonously decrease, but have minima at the manganese derivatives $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{Mn}$ and $(\text{Me}_4\text{C}_5\text{H})_2\text{Mn}$. This is similar to the behavior for the experimental mean bond dissociation enthalpies (i.e., half of the total bond dissociation enthalpy) for the metallocenes, which are 88 (V), 68 (Cr), 51 (Mn), 71 (Fe), 64 (Co), 59 (Ni) kcal/mol and thus also have a minimum at manganese.³⁵

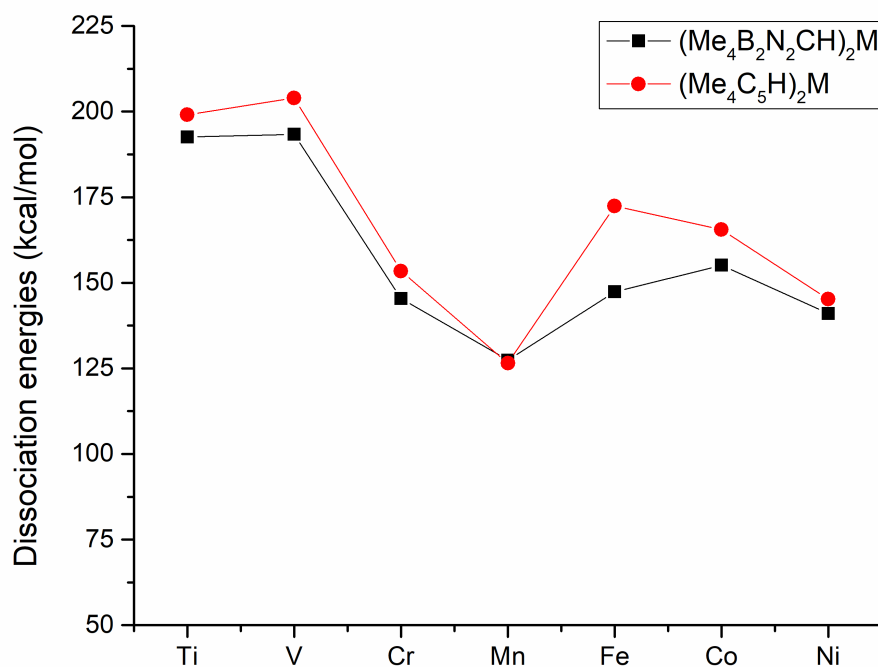


Figure 16. The dissociation energies for the $(\text{Me}_4\text{C}_5\text{H})_2\text{M}$ compounds (M = Ti, V, Cr, Mn, Fe, Co, Ni) using the M06-L method.

4. Conclusion

All seven first-row transition metal atoms form low-energy $(\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ sandwich structures with a *trans* (or nearly *trans* for Mn and Ti) orientation of parallel $\text{B}_2\text{N}_2\text{C}$ rings corresponding to a 180° (or 144° for Mn and Ti) angle between the C–H subunits of the $\text{Me}_4\text{B}_2\text{N}_2\text{CH}$ rings. Furthermore, the π -electrons in these heterocyclic $\text{B}_2\text{N}_2\text{C}$ rings are necessarily more localized than those in the homocyclic $\eta^5\text{-C}_5\text{H}_5$ cyclopentadienyl ligand. Thus the five electrons donated by a neutral $\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$ ligand to a metal atom can at least formally originate from lone pairs on each of the nitrogen atoms and one electron from the carbon atom. The 180° orientation of the $\text{B}_2\text{N}_2\text{C}$ rings in these structures leads to a formal localized octahedral coordination of the

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4 central transition metal forming two M–N bonds and one M–C bond to each ring with
5 two *trans* N–M–N units and one *trans* C–M–C unit.

6
7 This theoretical study shows that this type of octahedral coordination of the two
8 diazadiborolyl rings in sandwich compounds leads to a weaker field ligand than the
9 cyclopentadienyl ligand. This is manifest in higher spin states for the lowest energy
10 ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂M sandwich structures compared with the corresponding metallocenes
11 ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂M. This effect is most noticeable in the manganese derivative
12 ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Mn for which the sextet structure **Mn-x-144** (Figure 5) lies
13 ~18 kcal/mol in energy below the lowest energy quartet structure. The special stability of
14 the half-filled d⁵ shell of manganese(II) is evident here. This contrasts with the compli-
15 cated magnetic behavior of ($\eta^5\text{-C}_5\text{H}_5$)₂Mn, the doublet ground state of ($\eta^5\text{-Me}_5\text{C}_5$)₂Mn
16 found experimentally [30], and the doublet spin state structure of ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Mn
17 predicted in this work to lie ~6 kcal/mol in energy below the lowest energy sextet spin
18 state structure (Figure 9). The chromium derivative ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Cr also has a high-
19 spin quintet ground state in **Cr-p-180** (Figure 4) as compared with the triplet ground state
20 [9] found experimentally for ($\eta^5\text{-C}_5\text{H}_5$)₂Cr and predicted theoretically for
21 ($\eta^5\text{-Me}_4\text{C}_5\text{H}$)₂Cr (Figure 7). Even for the ferrocene analogue ($\eta^5\text{-Me}_4\text{B}_2\text{N}_2\text{CH}$)₂Fe the
22 high-spin quintet structure **Fe-p-180** lies only ~9 kcal/mol in energy above the singlet
23 ground state **Fe-s-180** (Figure 6).

24
25 The only bis(1,2-diaza-3,5-diborolyl)metal sandwich compound that has been
26 synthesized is the substituted ferrocene analogue [$\eta^5\text{-(CH}_2\text{)}_3\text{N}_2\text{(BPh)}_2\text{CMe}$]₂Fe, obtained
27 by Roesler and co-workers from the 1,2-diaza-3,5-diborolyl anion and FeCl₂(thf)₄ [8].
28 Related sandwich compounds of the other first-row transition metals should be accessible
29 by analogous synthetic methods.

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48 There are no conflicts of interest to declare.

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51 **Supporting Information.** Tables S1 to S7: Total energies (*E* in hartree), relative energies
52 (ΔE in kcal/mol), and numbers of imaginary vibrational frequencies (Nimg) for the
53 ($\text{Me}_4\text{B}_2\text{N}_2\text{CH}$)₂M structures; Tables S8 to S35: Atomic coordinates of the optimized
54 structures for the ($\text{Me}_4\text{B}_2\text{N}_2\text{CH}$)₂M complexes.
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Graphical Abstract

Higher Spin States in Some Low-Energy Bis(Tetramethyl-1,2-diaza- 3,5-diboroly) Sandwich Compounds of the First Row Transition Metals: Boraza Analogues of the Metallocenes

Jianlin Chen, Hao Feng,
Yaoming Xie, Schaefer, Henry
and
R. Bruce King

Density functional studies on $(\text{Me}_4\text{B}_2\text{N}_2\text{CH})_2\text{M}$ ($\text{M} = \text{Ti}, \text{V}, \text{Cr}, \text{Mn}, \text{Fe}, \text{Co}, \text{Ni}$) show low-energy sandwich structures for all seven metals. The lowest-energy such chromium and manganese derivatives have higher spin states than the corresponding metallocenes.

