



Kent Academic Repository

Roy, Dipak K., Tröster, Tobias, Fantuzzi, Felipe, Dewhurst, Rian D., Lenczyk, Carsten, Radacki, Krzysztof, Pranckevicius, Conor, Engels, Bernd and Braunschweig, Holger (2021) *Isolation and Reactivity of an Antiaromatic s-Block Metal Compound*. *Angewandte Chemie International Edition*, 60 (7). pp. 3812-3819. ISSN 1433-7851.

Downloaded from

<https://kar.kent.ac.uk/96288/> The University of Kent's Academic Repository KAR

The version of record is available from

<https://doi.org/10.1002/anie.202014557>

This document version

Publisher pdf

DOI for this version

Licence for this version

CC BY (Attribution)

Additional information

Versions of research works

Versions of Record

If this version is the version of record, it is the same as the published version available on the publisher's web site. Cite as the published version.

Author Accepted Manuscripts

If this document is identified as the Author Accepted Manuscript it is the version after peer review but before type setting, copy editing or publisher branding. Cite as Surname, Initial. (Year) 'Title of article'. To be published in *Title of Journal*, Volume and issue numbers [peer-reviewed accepted version]. Available at: DOI or URL (Accessed: date).

Enquiries

If you have questions about this document contact ResearchSupport@kent.ac.uk. Please include the URL of the record in KAR. If you believe that your, or a third party's rights have been compromised through this document please see our [Take Down policy](https://www.kent.ac.uk/guides/kar-the-kent-academic-repository#policies) (available from <https://www.kent.ac.uk/guides/kar-the-kent-academic-repository#policies>).

Antiaromaticity

Isolation and Reactivity of an Antiaromatic s-Block Metal Compound

Dipak Kumar Roy, Tobias Tröster, Felipe Fantuzzi, Rian D. Dewhurst, Carsten Lenczyk, Krzysztof Radacki, Conor Prancevicius, Bernd Engels,* and Holger Braunschweig*

Dedicated to Professor Siegfried Hünig on the occasion of his 100th birthday

Abstract: The concepts of aromaticity and antiaromaticity have a long history, and countless demonstrations of these phenomena have been made with molecules based on elements from the p, d, and f blocks of the periodic table. In contrast, the limited oxidation-state flexibility of the s-block metals has long stood in the way of their participation in sophisticated π -bonding arrangements, and truly antiaromatic systems containing s-block metals are altogether absent or remain poorly defined. Using spectroscopic, structural, and computational techniques, we present herein the synthesis and authentication of a heterocyclic compound containing the alkaline earth metal beryllium that exhibits significant antiaromaticity, and detail its chemical reduction and Lewis-base-coordination chemistry.

Introduction

The concept of antiaromaticity in molecules was presented in the mid-1960s by Breslow and co-workers^[1,2] as a simple counterpoint to aromaticity, that is, aromatic compounds are those in which cyclic delocalization of electrons has a stabilizing effect, while cyclic delocalization imparts destabilization in antiaromatic compounds. The concept has since been

expanded significantly in terms of our understanding of what constitutes antiaromaticity, the techniques available to measure it, and the number of isolated molecules fitting an antiaromatic description.^[3–5] The currently accepted prerequisites for aromaticity and antiaromaticity are that the molecule must be cyclic, planar, and have an unbroken series of orbitals available for π bonding around the ring. From this point, the properties of aromatic and antiaromatic compounds diverge in a number of key ways, including:

- their stability (aromatics are more stable than their acyclic equivalents, while antiaromatics are less stable)
- the number of electrons in their π system ($4n + 2$ π electrons for aromatics and $4n$ π electrons for antiaromatics)
- the magnetic ring current produced by delocalization (diamagnetic for aromatics and paramagnetic for antiaromatics)
- the bond-length equilibration generally found in aromatics is not usually present in antiaromatics

Over the past few decades, the increasing ubiquity of two techniques has allowed more confident identification of antiaromatic species, namely single-crystal X-ray diffraction (for determination of planarity and bond lengths) and computational chemistry approaches, in particular nucleus-independent chemical shift (NICS) methods,^[4,5] which determine the magnetic shielding at the center of a ring or above it. While truly antiaromatic hydrocarbons based on small ring systems are by definition unstable and thus very difficult to prepare and authenticate, the concept of antiaromaticity has found fertile ground in polycyclic π systems and heterocyclic chemistry, as is apparent from the proliferation of heteroatom-containing antiaromatics such as porphyrins and porphyrinoids,^[6–9] boroles,^[10–15] and other boron-containing heterocycles.^[16] Metals of the p, d, and f blocks of the periodic table have also proven able to participate in both aromaticity and antiaromaticity, as predicted theoretically^[17,18] and as confirmed by synthesis and structural authentication, exemplified by the now well-represented families of alumoles,^[19–22] metallabenzenes,^[23–27] and metallacyclopentadienes.^[28–30] The extension of the aromaticity and antiaromaticity concepts to spherical systems,^[31] such as fullerenes,^[32] metallofullerenes,^[33] molecular cages,^[34,35] and heterometallic clusters,^[36,37] is also attracting intense interest. This evidences the substantial importance of these concepts to the rationalization, design, and preparation of novel chemical structures.


In marked contrast to their p, d, and f block congeners, the metals of the s block of the periodic table are generally considered to be inflexible in terms of their oxidation states


[*] Prof. Dr. D. K. Roy, T. Tröster, Dr. F. Fantuzzi, Dr. R. D. Dewhurst, Dr. C. Lenczyk, Dr. K. Radacki, Dr. C. Prancevicius, Prof. Dr. H. Braunschweig
 Institute for Inorganic Chemistry
 Julius-Maximilians-Universität Würzburg
 Am Hubland, 97074 Würzburg (Germany)
 and

Institute for Sustainable Chemistry & Catalysis with Boron
 Julius-Maximilians-Universität Würzburg
 Am Hubland, 97074 Würzburg (Germany)
 E-mail: h.braunschweig@uni-wuerzburg.de

Prof. Dr. D. K. Roy
 Discipline of Chemistry
 Indian Institute of Technology Indore
 Khandwa Road, Simrol, Indore, 453552, M.P. (India)

Dr. F. Fantuzzi, Prof. Dr. B. Engels
 Institute for Physical and Theoretical Chemistry
 Julius-Maximilians-Universität Würzburg
 Emil-Fischer-Strasse 42, 97074 Würzburg (Germany)
 E-mail: bernd.engels@uni-wuerzburg.de

 Supporting information and the ORCID identification number(s) for the author(s) of this article can be found under:
<https://doi.org/10.1002/anie.202014557>.

 © 2020 The Authors. Angewandte Chemie International Edition published by Wiley-VCH GmbH. This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

and have little ability to engage in π -bonding interactions,^[38] which would seemingly disqualify them from forming either aromatic or antiaromatic compounds. However, over the last dozen years, work in s-block chemistry has uncovered surprising oxidation-state flexibility in the lighter Group 2 metals, including formally Mg^I ,^[39,40] Be^0 ,^[41] and Be^I species.^[42] A number of s-block species also exist that possess the six π electrons required for aromaticity, for example base-stabilized 1,3-diaza-2-berylloles and -magnesioles,^[43] although the aromaticity of these species was not discussed in the original articles. One report exists of the synthesis of molecules containing s-block metals as part of 4π -electron rings, namely $[\text{MgC}_4\text{R}^1_2\text{R}^2_2]$ ($\text{R}^1 = \text{SiMe}_3$; $\text{R}^2 = \text{Me, Ph}$),^[44] and these were subsequently calculated to be significantly antiaromatic.^[45] However, these species were not structurally authenticated, preventing the crucial confirmation of their planarity. A complex of one of these magnesioles with the chelating ligand *N,N,N',N'*-tetramethyl-1,2-ethylenediamine (TMEDA) was structurally confirmed,^[44] but the tetracoordinate nature of the Mg atom in this complex suggests that a contiguous π network is not present in this compound. A method to construct an antiaromatic complex of beryllium has also recently been predicted by density functional theory (DFT) calculations,^[46] involving the conceptual combination of a butadiendiyl diradical $[\text{C}_4\text{R}_4]$ fragment with the anionic fragment $[\text{:BeR}]^-$ ($\text{R} =$ anionic substituent), leading to an anionic 4π -electron species. Nevertheless, despite extensive computational design, synthetic effort, and a number of promising results, the goal of preparing an antiaromatic s-block complex remains out of reach.

Our approach to this problem was to conceptually combine the four- π -electron $[\text{C}_4\text{R}_4]$ fragment with the neutral, zero- π -electron fragment $[\text{:BeL}]$ ($\text{L} =$ neutral Lewis donor), which would lead to a neutral species with four π electrons and thus circumvent the potential complications of the negatively charged products predicted theoretically.^[46] As the fragment $[\text{:BeL}]$ is isoelectronic and isolobal with the boron-containing fragment $[\text{:BR}]$ ($\text{R} =$ anionic substituent), present in the diverse family of antiaromatic boroles, we reasoned that this approach had a good chance of success. Interest in the chemistry of carbene-stabilized beryllium species has exploded over the past few years, leading to the discovery of a range of Be^0 , Be^I , and Be^{II} complexes with fascinating properties and reactivity, as well as providing a host of precursors containing stabilized $[\text{:BeL}]$ scaffolds for the construction of compounds with unusual electronic structures.^[41–43,47] The $[\text{:BeL}]$ fragment we chose was $[\text{:Be}(\text{CAAC})]$, bearing a neutral cyclic (alkyl)(amino)carbene (CAAC) donor, inspired by our previous synthesis of the precursor $[(\text{CAAC})\text{BeCl}_2]$ and its use to form a compound in which the formally zerovalent Be atom takes part in strong multiple bonding with carbon, $[\text{Be}(\text{CAAC})_2]$.^[41] Herein, we present the synthesis and isolation of a Lewis-base-stabilized beryllole, a heterocyclic organoberyllium compound, which was subsequently determined by spectroscopic, structural, and computational techniques to have significant antiaromaticity in its *cyclo*- BeC_4 ring. Its facile reactivity, including chemical reduction and the addition of a second Lewis donor, provides stable, non-antiaromatic products.

Results and Discussion

Synthesis and Reactivity of Beryllole 1. Combination of $[(\text{CAAC})\text{BeCl}_2]$ with dilithium tetraphenylbutadiene as its diethyl etherate ($\text{Li}_2[\text{C}_4\text{Ph}_4]\cdot 0.5\text{Et}_2\text{O}$) in benzene at room temperature provided, after stirring and purification, an air-sensitive yellow crystalline solid (yield 38 %) with a ^9Be NMR spectroscopic shift at 22.9 ppm. Multinuclear NMR spectroscopy, high-resolution mass spectrometry, and single-crystal X-ray diffraction techniques unambiguously identified this compound as the CAAC-stabilized beryllole monoadduct $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]$ (**1**) as shown in Figure 1. In order to place the ^9Be NMR signal of **1** in context with comparable compounds, the related $[(\text{CAAC})\text{BeCl}_2]$ showed a ^9Be NMR signal at 12.9 ppm^[41] while that of the only known CAAC complex of a diorganyl beryllium, $[(\text{CAAC})\text{Be}(9,10\text{-anthracenyl})]$, was found at 1.7 ppm.^[47b] Both signals are significantly upfield of that of **1**, suggesting that the Be center in **1** is highly unsaturated, likely due to the low electronegativity of the attached sp^2 carbon atoms of the C_4 ring and the absence of delocalization of the four π electrons in this ring with the Be p orbital.

The solid-state structure of **1** confirmed the tricoordinate nature of the Be center, the effective planarity of the BeC_4 ring (deviations of the BeC_4 atoms from the calculated least-squares plane of this ring: 0.021–0.094 Å), and significant bond-length alternation in the C_4 backbone (inner C–C distance: 1.512(2) Å; outer C–C distances: 1.359(2), 1.358(2) Å). The planarity of the ring and bond-length alternation in the backbone are both key structural attributes required for antiaromaticity, however, an analogous non-aromatic species would also show the same attributes, for example, cyclopentadiene, which in one published solid-state structure has an inner C–C distance of 1.460(1) Å and outer C–C distances of 1.344(1) Å in its butadiene backbone.^[48] Thus, the structural aspects of ring planarity and bond length alternation alone do not allow distinction between antiaromatic/non-aromatic descriptions.

Given the presumed antiaromaticity of beryllole monoadduct **1**, this compound was subjected to reactivity tests that parallel those previously demonstrated for the related family of antiaromatic boroles,^[10–15] namely two-electron reduction and the addition of neutral, two-electron donor groups (Lewis bases). Thereby, stirring **1** in a suspension of diethyl ether and lithium sand, followed by filtration, crystallization, and drying, provided a 50% yield of an air- and thermally sensitive dark-red crystalline solid determined to be the dilithium dietherate $[\text{Li}(\text{OEt}_2)]_2[(\text{CAAC})\text{BeC}_4\text{Ph}_4]$ (**2**, Figure 1) by multinuclear NMR spectroscopy, single-crystal X-ray diffraction, and elemental analysis. The solid-state structure of **2** shows one $[\text{Li}(\text{OEt}_2)]$ fragment bound to each side of the BeC_4 plane, bound somewhat more closely to the carbon atoms than the beryllium atom (avg. Li–C distance: 2.177 Å; avg. Li–Be distance: 2.358 Å). Compound **2** shows distinct bond distance equalization (inner C–C distance: 1.448(3) Å; outer C–C distances: 1.459(2), 1.470(2) Å), in line with the formally aromatic nature of the $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]^{2-}$ heterocycle. However, the π coordination of two $[\text{Li}(\text{OEt}_2)]^+$

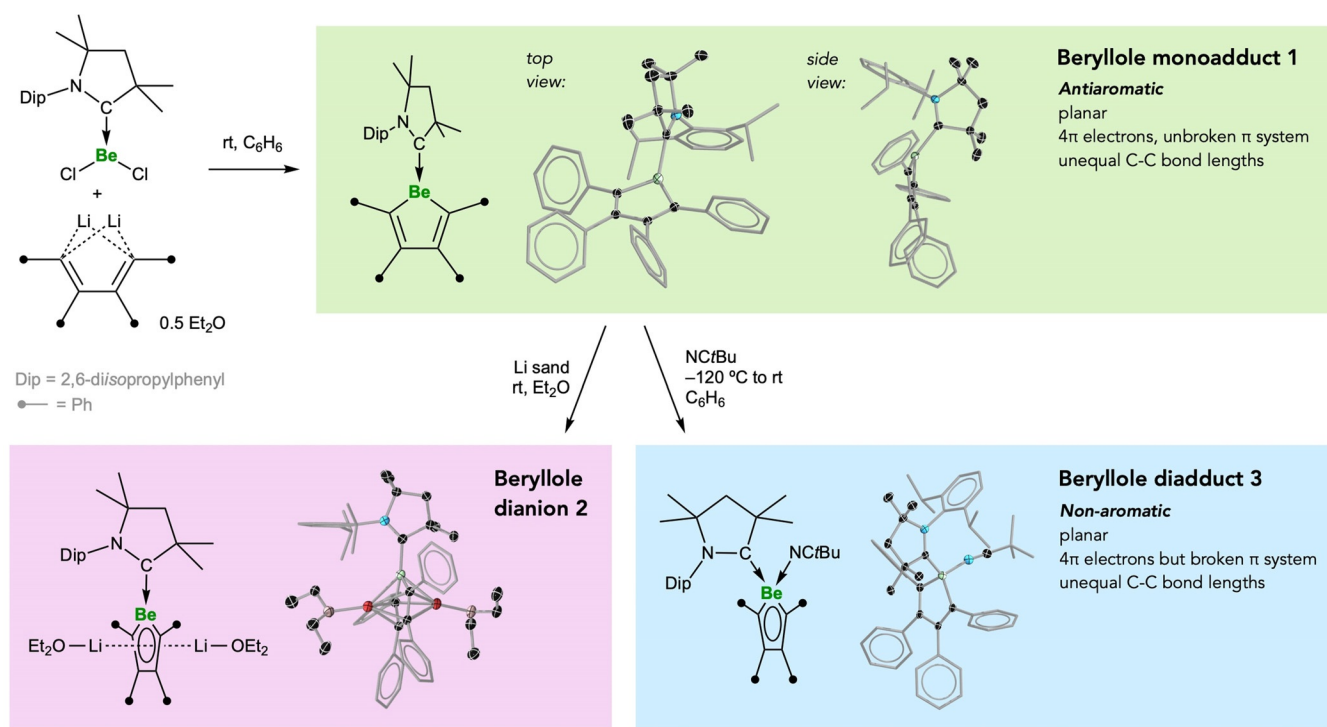


Figure 1. Synthesis and reactivity of the antiaromatic beryllole **1**, and solid-state structures of **1-3**.^[78] Ellipsoids of the crystallographically derived structures are shown at the 50% probability level. All hydrogens and ellipsoids of peripheral groups have been removed for clarity. Atom colors: N blue, Be green, Li red, O pink.

fragments to the ring precludes the description of **2** as a classical aromatic species.

Alternatively, treatment of beryllole monoadduct **1** with pivalonitrile (NCtBu) at -120°C , followed by warming to room temperature, evaporation of solvent and recrystallization, provided a 90% yield of a yellow crystalline solid (**3**) with a ^9Be NMR spectroscopic signal at 4.4 ppm. This signal is significantly upfield of that of **1** (22.9 ppm), reflective of the increase in electron density at Be upon quaternization (i.e. from six to eight valence electrons). The widths at half-height ($\omega_{1/2}$) of reported ^9Be NMR signals have recently been surveyed in relation to coordination number by Buchanan and Plieger,^[49] finding that these values decrease with increasing coordination number at Be due to its increased symmetry. Accordingly, the ^9Be signal of tetracoordinate **3** is significantly less broad ($\omega_{1/2} \approx 141$ Hz) than the very broad signal of tricoordinate **1** ($\omega_{1/2} \approx 506$ Hz). However, this value for **3** is itself very broad for a tetracoordinate Be compound, being only the second such compound with a $\omega_{1/2}$ value above 100 according to the review of Buchanan and Plieger, an effect that could be attributed to its relatively low symmetry. Nevertheless, the ^9Be NMR chemical shifts of **1** and **3** fall within the ranges reported in the aforementioned review for their respective coordination number.

Multinuclear NMR spectroscopy, high-resolution mass spectrometry, and single-crystal X-ray diffraction techniques confirmed this species to be the beryllole diadduct [(tBuCN)-(CAAC)BeC₄Ph₄] (**3**, Figure 1), with a tetracoordinate Be center and strongly alternating C₄ backbone bond distances (inner C–C distance: 1.515(2) Å; outer C–C distances: 1.357-

(2), 1.357(2) Å), these distances being effectively identical to those of **1**. The exo- and endocyclic Be–C distances of **3** are only marginally longer than those of **1** and are presumably the result of the increased coordination number of the Be center.

Computational Determination of the Antiaromaticity of Synthesized Berylloles. Given the structural similarity of the beryllole rings of **1** and **3**, we turned to theory to provide a more concrete determination of the antiaromaticity/non-aromaticity of the compounds. Calculations using DFT, complete active space self-consistent field (CASSCF),^[50] and n-electron valence state second-order perturbation theory (NEVPT2)^[51–53] were performed in order to describe the molecular and electronic structures of the beryllium compounds obtained herein (see SI for details). The nature of the Be–CAAC bond in a truncated model of **1** was investigated using the energy decomposition analysis with natural orbitals for chemical valence (EDA-NOCV) method.^[54,55] The antiaromatic character of the systems was investigated by NICS and aromatic stabilization energy (ASE) calculations,^[56,57] the latter obtained by the thermodynamic analysis of selected homodesmotic reactions.

An informative initial picture of the bonding in these compounds can be obtained from appraisal of the frontier molecular orbitals (FMOs, Figure 2A) of **1**, **2**, and those of a model compound where the [Li(OEt₂)] fragments of **2** are removed, thus leading to the naked dianion [(CAAC)-BeC₄Ph₄]²⁻. As expected, the HOMO of **1** (Figure 2A, left) is very similar to that of the closely related antiaromatic borole species [PhBC₄Ph₄],^[58–60] and is composed of an antisymmetric π orbital with the beryllium atom in the nodal

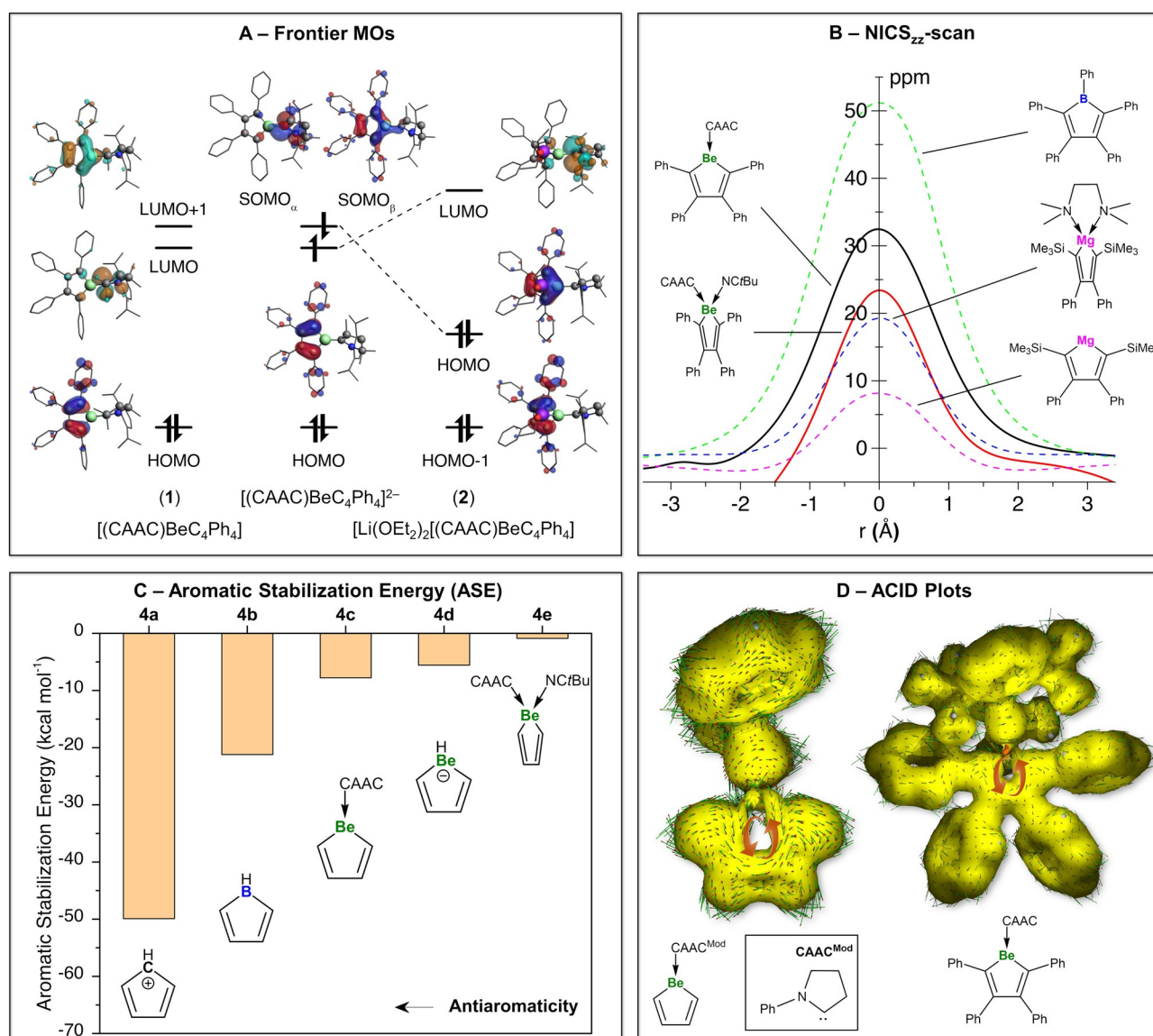


Figure 2. A) Frontier molecular orbitals, B) scan of the zz -components of the nucleus independent chemical shifts (NICS_{zz}-scan), C) aromatic stabilization energies, and D) anisotropy of the induced current density (ACID) plots of selected (anti-/non-)aromatic compounds.

position. A low-lying virtual orbital of π symmetry in which the beryllium atom occupies a non-nodal position, reminiscent of one of the degenerate HOMOs of the triplet cyclopentadienyl cation, C_5H_5^+ (X^3A_2' state of D_{5h} point group symmetry),^[61–63] was also observed. However, while for boroles this orbital is the LUMO, for **1** it is the LUMO + 1, as the LUMO of **1** is predominantly located on the CAAC ligand and can be described as predominantly $\pi^*(\text{CN})$ in character. Thus, substitution of $:\text{BPh}$ with the isolobal $:\text{Be}(\text{CAAC})$ moiety affects the FMOs of **1** mainly by introducing a second low-lying vacant orbital, which, as discussed below, leads to remarkable consequences for the electronic structure of the beryllole dianion.

Computational addition of two electrons to boroles $[\text{R}'\text{BC}_4\text{R}_4]$ results in the formation of borolyd dianions $[\text{R}'\text{BC}_4\text{R}_4]^{2-}$. These 6π -electron systems, although calculated to be aromatic and analogous to the aromatic cyclopenta-

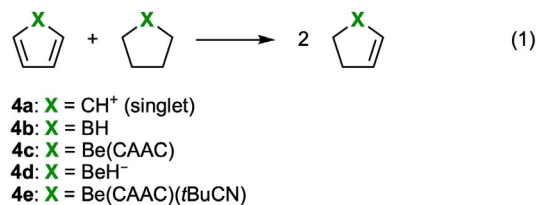
dieryl anion, C_5H_5^- , are unknown in the absence of π -coordinated s-block metals. In contrast, analogous computational addition of two electrons to the CAAC-stabilized beryllole **2** leads to the dianion $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]^{2-}$ (Figure 2 A, middle), which, although possessing six π electrons, is singlet biradicaloid in nature, wherein the low-lying LUMO and LUMO + 1 of **1** are now partially occupied and the C–C bond lengths in the five-membered ring are unequal. This picture is corroborated by DFT in the unrestricted formalism as well as by high-level CASSCF/NEVPT2 calculations, the latter indicating a biradical character of 37.4% for $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]^{2-}$, which is of the same order as that of the recently isolated parent aluminene $[(\text{CAAC})_2\text{AlH}]$,^[64] and a singlet-triplet gap ($\Delta_{\text{T-S}}$) of 3.3 kcal mol⁻¹. Interestingly, the presence of Li atoms bound to the five-membered beryllole ring in compound **2** significantly stabilizes the low-lying π^* orbital associated with the ring, while that of the CAAC ligand is

pushed to higher energies (Figure 2A, right). As a result, in the experimentally realized species **2**, the 6π -electron cloud is stabilized by the addition of two $[\text{Li}(\text{OEt}_2)]^+$ fragments, leading to a closed-shell singlet analogous to an aromatic borole dianion, as well as C–C bond length equalization. Accordingly, the singlet–triplet gap for this system is $42.1 \text{ kcal mol}^{-1}$ as obtained from CASSCF/NEVPT2 calculations.

The aforementioned NICS technique is a useful computational gauge of the magnetic shielding at a given point near a ring system, allowing a measure of aromaticity, antiaromaticity, and non-aromaticity of ring systems.^[4,5] In order to obtain more reliable and comparable data, the so-called zz component of the magnetic shielding tensor (NICS_{zz}) is commonly reported in the literature.^[65,66] The NICS_{zz} values of the experimentally realized compounds **1** and **3** were calculated, along with a number of other relevant molecules for comparison, starting at the ring centroid and at 0.1 \AA steps on an axis perpendicular to the ring plane (in both directions), leading to NICS_{zz} -scan curves.^[67,68] These data, obtained at the $\text{B3LYP}^{[69-72]}/6-311++\text{G}^*_{**[73,74]}$ level of theory, are displayed in Figure 2B. A large negative NICS_{zz} value at a point 1 \AA above and below the ring plane (denoted $\text{NICS}_{zz}(1)$ and $\text{NICS}_{zz}(-1)$) is regarded as a hallmark of aromaticity, while rings with large positive $\text{NICS}_{zz}(1/-1)$ values can be regarded as antiaromatic. Accordingly, the $\text{NICS}_{zz}(1)$ value of the known antiaromatic species pentaphenylborole $[\text{PhBC}_4\text{Ph}_4]$ was calculated to be $+26.1 \text{ ppm}$. Our calculations show that the 4π -electron beryllole monoadduct $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]$ (**1**; $\text{NICS}_{zz}(1)$: $+14.1/+13.5 \text{ ppm}$) is indeed antiaromatic, while the diadduct $[(t\text{BuCN})(\text{CAAC})\text{BeC}_4\text{Ph}_4]$ (**3**; $\text{NICS}_{zz}(1)$: $+5.3/+3.7 \text{ ppm}$), the π network of which is interrupted by the presence of a second donor ligand on the Be atom, is effectively nonaromatic. By comparison, the proposed magnesiole species $[\text{MgC}_4\text{Ph}_2(\text{SiMe}_3)_2]$ ($\text{NICS}_{zz}(1)$: $+0.8/+0.7 \text{ ppm}$) is also nonaromatic, while its structurally authenticated diamine adduct $[(\text{TMEDA})\text{MgC}_4\text{Ph}_2(\text{SiMe}_3)_2]$ ($\text{NICS}_{zz}(1)$: $+6.2/+6.0 \text{ ppm}$) could be regarded as very weakly antiaromatic.^[44]

A number of further points should be noted in order to confirm the antiaromaticity of beryllole monoadduct **1** and distinguish it from beryllole diadduct **3**. The evident perpendicularity of CAAC and BeC_4 planes of **1** (angle between planes: 79.5° in the solid-state structure and 77.3° in the calculated structure) implies the absence of an exocyclic Be–C π interaction and that the CAAC unit acts as a pure σ -donor. This picture is fully corroborated by the EDA–NOCV calculations (see SI for details), which revealed the predominance of a donor–acceptor rather than an electron-sharing Be–C^{CAAC} bond description in **1**, with negligible π interaction. Furthermore, computationally reducing the size of the BeC_4 backbone substituents of **1** causes the tilt of the CAAC unit out of the BeC_4 plane to approach zero (calculated tilt angles for $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]$: 148.0° ; $[(\text{CAAC})\text{BeC}_4\text{Me}_4]$: 160.9° ; $[(^{\text{H}}\text{CAAC})\text{BeC}_4\text{H}_4]$: 179.8° ; $^{\text{H}}\text{CAAC}$ is the hypothetical unsubstituted CAAC ligand pyrrolidin-2-ylidene ($\text{C}_4\text{H}_7\text{N}$), see SI), suggesting that the tilting observed in the solid-state structure is caused by steric, rather than electronic, effects.

In order to test the hypothesis that the presence of four electrons in the BeC_4 π system of **1** is more unfavorable than two analogous molecules containing two π electrons each, we calculated aromatic stabilization energies (ASEs) of a range of related five-membered $\text{C}_4\text{H}_4\text{X}$ rings (Figure 2C). These systems contain four π electrons each, and their ASEs were calculated following the homodesmotic reaction depicted in Equation (1).^[56,57]



The zero-point-energy-corrected electronic energies of the reactions depicted in Equation (1) were calculated using the molecules in their most stable conformations, and thus it is expected that contributions coming from strain effects are suppressed. The ASEs following Equation (1) are thus directly associated with cyclic delocalization/localization effects. Positive ASE values are related to the stability of aromatic compounds due to cyclic delocalization, whereas negative ASE values denote destabilization of antiaromatic species via cyclic localization. The singlet cyclopentadienyl cation C_5H_5^+ (**4a**, $^1\text{A}_1$ state of C_{2v} point group symmetry), which is a low-lying excited state above the $\text{X}^3\text{A}_2'$ ground state,^[64] has the largest destabilization energy ($\text{ASE} = -49.9 \text{ kcal mol}^{-1}$), and therefore is the most antiaromatic $\text{C}_4\text{H}_4\text{X}$ ring among those studied herein. This is followed by the borole ring **4b**, with an ASE value of $-21.2 \text{ kcal mol}^{-1}$. The beryllium-containing systems **4c** and **4d** also present negative ASE values (-7.8 and $-5.6 \text{ kcal mol}^{-1}$, respectively), thus confirming the destabilizing thermodynamic effect and further supporting the antiaromatic description of **1**. Finally, calculations for the reduced model of the beryllole diadduct **4e** found an ASE value of practically zero ($\text{ASE} = -0.9 \text{ kcal mol}^{-1}$), thus confirming the nominal antiaromaticity/non-aromaticity of this species.

Finally, we probed the antiaromaticity of CAAC-stabilized BeC_4 rings by using the anisotropy of the induced current density (ACID) method, as proposed by Herges and co-workers.^[75,76] ACID has been widely used for assessing delocalization, conjugation, and aromaticity in distinct molecular systems.^[76,77] Briefly, by plotting the ACID scalar field isosurfaces together with the current density vectors, it is possible to distinguish between diatropic (clockwise) and paratropic (counterclockwise) π -electron circulation, which are characteristic of aromatic and antiaromatic molecules, respectively. According to the ACID analysis (Figure 2D), BeC_4 rings exhibit counterclockwise, paratropic circulation typical of antiaromatic systems. Furthermore, the heteroatom contribution to the cyclic conjugation is lower in berylloles than boroles (see Figure S15), suggesting that the antiaromatic character of the studied berylloles is less pronounced

than that of boroles. These results are consistent with those found using the aromatic stabilization energy and NICS calculations, supporting our attribution of berylloles as species with distinct but weak antiaromatic character. Future work in our group will include a detailed investigation of the electronic structure of the dilithio beryllole **2** and the electrostructural effects dictating the biradicaloid character of its naked form $[(\text{CAAC})\text{BeC}_4\text{Ph}_4]^{2-}$.

Our results indicate a dramatic decrease in the antiaromaticity in the ring systems $[\text{XC}_4\text{H}_4]$ upon moving left from carbon in the periodic table, that is, from $\text{X} = [\text{HC}]^+$, to $[\text{HB}]$, and to $[(\text{CAAC})\text{Be}]$. The reason for this decrease can be understood by considering the interaction of the π -MOs of the fragments C_4H_4 and X . Figure S16 provides a sketch of the variations obtained for different X fragments while Figure 3

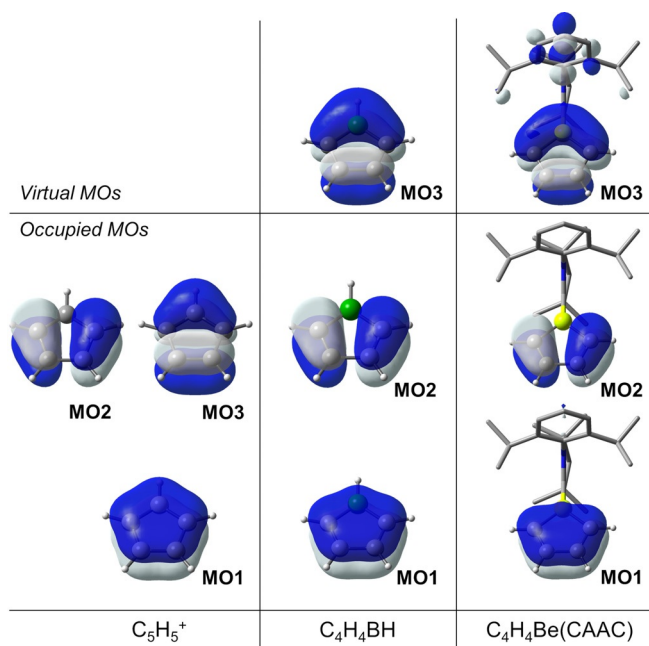


Figure 3. Relevant molecular orbitals calculated for truncated cyclic systems $[\text{XC}_4\text{H}_4]$ ($\text{X} = [\text{HC}]^+$, to $[\text{HB}]$, $[(\text{CAAC})\text{Be}]$), highlighting the extent of participation of the X fragment in the ring π system.

compares the computed MOs of the systems. For clarity we only focus on the MOs of relevance to the XC_4H_4 π system. Upon varying X , the relative energy of the MO in which X lies on the nodal plane (MO2) remains nearly unchanged, but the energy of MO3, to which X contributes, changes its position. For $\text{X} = \text{CH}$ this orbital is degenerate with MO2. The energy of this orbital is higher for $\text{X} = [\text{HB}]$ and $[(\text{CAAC})\text{Be}]$ because the fragment orbital of X increases in energy, thereby lifting the degeneracy. Furthermore, the mixing of the fragment orbitals decreases because the energy difference between the fragment orbitals (i.e. C_4H_4 vs. X) increases. As a consequence, the mixing also decreases so that the fragments become increasingly decoupled. This decoupling is best seen in MO1 (Figure 3), which for $\text{X} = [\text{HC}]^+$ is completely delocalized, but for $\text{X} = [(\text{CAAC})\text{Be}]$ is more localized on the

$[\text{C}_4\text{H}_4]$ fragment. This decoupling in the beryllole system leads to its reduced antiaromaticity.

Conclusion

The structural, thermodynamic, and magnetic considerations detailed above indicate that the beryllole monoadduct **1** has distinct antiaromaticity, setting it apart from the negligible antiaromaticity of the beryllole diadduct **3**. The results herein present the full spectroscopic, structural, and computational authentication of a true antiaromatic species based on the alkaline earth metal beryllium. The results suggest levels of oxidation-state flexibility and participation in sophisticated π systems that are unprecedented for the metals of the s block, and hint at the potentially rich chemistry of molecules combining π systems and alkaline earth metals.

Acknowledgements

Financial support from the Deutsche Forschungsgemeinschaft (H.B.) and the DFG Research Training Group GRK2112, “Molecular Biradicals: Structure, Properties and Reactivity” (B.E.) is gratefully acknowledged. F.F. thanks the Coordenação de Aperfeiçoamento de Pessoal de Nível Superior and the Alexander von Humboldt Foundation for a Capes-Humboldt Research Fellowship. C.P. thanks the NSERC and Alexander von Humboldt Foundation for postdoctoral fellowships. Open access funding enabled and organized by Projekt DEAL.

Conflict of interest

The authors declare no conflict of interest.

Keywords: antiaromaticity · aromaticity · beryllium · heterocycles · s-block metals

- [1] R. Breslow, J. Brown, J. J. Gajewski, *J. Am. Chem. Soc.* **1967**, *89*, 4383–4390.
- [2] R. Breslow, *Acc. Chem. Res.* **1973**, *6*, 393–398.
- [3] P. v. R. Schleyer, H. Jiao, *Pure Appl. Chem.* **1996**, *68*, 209–218.
- [4] P. v. R. Schleyer, C. Maerker, A. Dransfeld, H. Jiao, N. J. R. v. E. Hommes, *J. Am. Chem. Soc.* **1996**, *118*, 6317–6318.
- [5] Z. Chen, C. S. Wannere, C. Corminboeuf, R. Puchta, P. v. R. Schleyer, *Chem. Rev.* **2005**, *105*, 3842–3888.
- [6] Y. M. Sung, J. Oh, W.-Y. Cha, W. Kim, J. M. Lim, M.-C. Yoon, D. Kim, *Chem. Rev.* **2017**, *117*, 2257–2312.
- [7] T. Tanaka, A. Osuka, *Chem. Rev.* **2017**, *117*, 2584–2640.
- [8] B. K. Reddy, A. Basavarajappa, M. D. Ambhore, V. G. Anand, *Chem. Rev.* **2017**, *117*, 3420–3443.
- [9] J. Oh, Y. M. Sung, Y. Hong, D. Kim, *Acc. Chem. Res.* **2018**, *51*, 1349–1358.
- [10] H. Braunschweig, T. Kupfer, *Chem. Commun.* **2011**, *47*, 10903–10914.
- [11] J. H. Barnard, S. Yruegas, K. Huang, C. D. Martin, *Chem. Commun.* **2016**, *52*, 9985–9991.
- [12] L. Ji, S. Griesbeck, T. B. Marder, *Chem. Sci.* **2017**, *8*, 846–863.
- [13] B. Su, R. Kinjo, *Synthesis* **2017**, *49*, 2985–3034.
- [14] W. Zhang, B. Zhang, D. Yu, G. He, *Sci. Bull.* **2017**, *62*, 899–900.

- [15] Y. Su, R. Kinjo, *Chem. Soc. Rev.* **2019**, *48*, 3613–3659.
- [16] Y. Ishida, B. Donnadiou, G. Bertrand, *Proc. Natl. Acad. Sci. USA* **2006**, *103*, 13585–13588.
- [17] D. Y. Zubarev, B. B. Averkiev, H.-J. Zhai, L.-S. Wang, A. I. Boldyrev, *Phys. Chem. Chem. Phys.* **2008**, *10*, 257–267.
- [18] T. R. Galeev, A. I. Boldyrev, *Annu. Rep. Prog. Chem. Sect. C* **2011**, *107*, 124–147.
- [19] T. Agou, T. Wasano, P. Jin, S. Nagase, N. Tokitoh, *Angew. Chem. Int. Ed.* **2013**, *52*, 10031–10034; *Angew. Chem.* **2013**, *125*, 10215–10218.
- [20] T. Wasano, T. Agou, T. Sasamori, N. Tokitoh, *Chem. Commun.* **2014**, *50*, 8148–8150.
- [21] T. Agou, T. Wasano, T. Sasamori, J.-D. Guo, S. Nagase, N. Tokitoh, *Angew. Chem. Int. Ed.* **2015**, *54*, 9568–9571; *Angew. Chem.* **2015**, *127*, 9704–9707.
- [22] Y. Zhang, J. Wei, W.-X. Zhang, Z. Xi, *Inorg. Chem.* **2015**, *54*, 10695–10700.
- [23] J. R. Bleeke, *Chem. Rev.* **2001**, *101*, 1205–1227.
- [24] J. Chen, G. Jia, *Coord. Chem. Rev.* **2013**, *257*, 2491–2521.
- [25] C. Zhu, S. Li, M. Luo, X. Zhou, Y. Niu, M. Lin, J. Zhu, Z. Cao, X. Lu, T. Wen, Z. Xie, P. v. R. Schleyer, H. Xia, *Nat. Chem.* **2013**, *5*, 698–703.
- [26] C. Zhu, H. Xia, *Acc. Chem. Res.* **2018**, *51*, 1691–1700.
- [27] D. Chen, Q. Xie, J. Zhu, *Acc. Chem. Res.* **2019**, *52*, 1449–1460.
- [28] J. Wei, W.-X. Zhang, Z. Xi, *Angew. Chem. Int. Ed.* **2015**, *54*, 5999–6002; *Angew. Chem.* **2015**, *127*, 6097–6100.
- [29] M. Saito, M. Sakaguchi, T. Tajima, K. Ishimura, S. Nagase, M. Hada, *Science* **2010**, *328*, 339–342.
- [30] W. Ma, C. Yu, T. Chen, L. Xu, W.-X. Zhang, Z. Xi, *Chem. Soc. Rev.* **2017**, *46*, 1160–1192.
- [31] A. Hirsch, Z. Chen, H. Jiao, *Angew. Chem. Int. Ed.* **2000**, *39*, 3915–3917; *Angew. Chem.* **2000**, *112*, 4079–4081.
- [32] M. Bühl, A. Hirsch, *Chem. Rev.* **2001**, *101*, 1153–1184.
- [33] M. Garcia-Borràs, S. Osuna, J. M. Luis, M. Swart, M. Solà, *Chem. Soc. Rev.* **2014**, *43*, 5089–5105.
- [34] Y. Ni, T. Y. Gopalakrishna, H. Phan, T. Kim, T. S. Herng, Y. Han, T. Tao, J. Ding, D. Kim, J. Wu, *Nat. Chem.* **2020**, *12*, 242–248.
- [35] A. Muñoz-Castro, *ChemPhysChem* **2020**, *21*, 1384–1387.
- [36] H. Xu, I. A. Popov, N. V. Tkachenko, Z. Wang, A. Muñoz-Castro, A. I. Boldyrev, Z. Sun, *Angew. Chem. Int. Ed.* **2020**, *59*, 17286–17290; *Angew. Chem.* **2020**, *132*, 17439–17443.
- [37] Z.-C. Wang, N. V. Tkachenko, L. Qiao, E. Matito, A. Muñoz-Castro, A. I. Boldyrev, Z.-M. Sun, *Chem. Commun.* **2020**, *56*, 6583–6586.
- [38] L. C. Perera, O. Raymond, W. Henderson, P. J. Brothers, P. G. Plieger, *Coord. Chem. Rev.* **2017**, *352*, 264–290.
- [39] S. P. Green, C. Jones, A. Stasch, *Science* **2007**, *318*, 1754–1757.
- [40] C. Jones, *Nat. Rev. Chem.* **2017**, *1*, 0059.
- [41] M. Arrowsmith, H. Braunschweig, M. A. Celik, T. Dellermann, R. D. Dewhurst, W. C. Ewing, K. Hammond, T. Kramer, I. Krummenacher, J. Mies, K. Radacki, J. K. Schuster, *Nat. Chem.* **2016**, *8*, 890–894.
- [42] G. Wang, J. E. Walley, D. A. Dickey, S. Pan, G. Frenking, R. J. Gilliard, Jr., *J. Am. Chem. Soc.* **2020**, *142*, 4560–4564.
- [43] L. A. Freeman, J. E. Walley, A. D. Obi, G. Wang, D. A. Dickie, A. Molino, D. J. D. Wilson, R. J. Gilliard, Jr., *Inorg. Chem.* **2019**, *58*, 10554–10568.
- [44] J. Wei, L. Liu, M. Zhan, L. Xu, W.-X. Zhang, Z. Xi, *Angew. Chem. Int. Ed.* **2014**, *53*, 5634–5638; *Angew. Chem.* **2014**, *126*, 5740–5744.
- [45] Z. Sun, A. M. Launder, H. F. Schaefer III, *ChemistrySelect* **2017**, *2*, 1442–1453.
- [46] T. E. Field-Theodore, D. J. D. Wilson, J. L. Dutton, *Inorg. Chem.* **2015**, *54*, 8035–8041.
- [47] a) M. R. Buchner, *Chem. Eur. J.* **2019**, *25*, 12018–12036; b) J. K. Schuster, D. K. Roy, C. Lenczyk, J. Mies, H. Braunschweig, *Inorg. Chem.* **2019**, *58*, 2652–2658; c) G. Wang, L. A. Freeman, D. A. Dickie, R. Mokrai, Z. Benkő, R. J. Gilliard, Jr., *Chem. Eur. J.* **2019**, *25*, 4335–4339; d) J. E. Walley, A. D. Obi, G. Breiner, G. Wang, D. A. Dickie, A. Molino, J. L. Dutton, D. J. D. Wilson, R. J. Gilliard, Jr., *Inorg. Chem.* **2019**, *58*, 11118–11126; e) J. E. Walley, Y.-O. Wong, L. A. Freeman, D. A. Dickie, R. J. Gilliard, Jr., *Catalysts* **2019**, *9*, 934; f) J. E. Walley, D. A. Dickie, R. J. Gilliard, Z. *Naturforsch. B* **2020**, *75*, 497–501; g) M. R. Buchner, S. Pan, C. Poggel, N. Spang, M. Müller, G. Frenking, J. Sundermeyer, *Organometallics* **2020**, *39*, 3224–3231.
- [48] T. Haumann, J. Benet-Buchholz, R. Boese, *J. Mol. Struct.* **1996**, *374*, 299–304.
- [49] J. A. Buchanan, P. G. Plieger, *Z. Naturforsch. B* **2020**, *75*, 459–472.
- [50] B. O. Roos, in *Advances in Chemical Physics: Ab Initio Methods in Quantum Chemistry, Part 2, Vol. 69* (Ed.: K. P. Lawley), Wiley, New York, **1987**, pp. 399–445.
- [51] C. Angeli, R. Cimiraglia, J.-P. Malrieu, *Chem. Phys. Lett.* **2001**, *350*, 297–305.
- [52] C. Angeli, R. Cimiraglia, S. Evangelisti, T. Leininger, J.-P. Malrieu, *J. Chem. Phys.* **2001**, *114*, 10252–10264.
- [53] C. Angeli, R. Cimiraglia, J.-P. Malrieu, *J. Chem. Phys.* **2002**, *117*, 9138–9153.
- [54] M. P. Mitoraj, A. Michalak, T. Ziegler, *J. Chem. Theory Comput.* **2009**, *5*, 962–975.
- [55] L. Zhao, M. von Hopffgarten, D. M. Andrada, G. Frenking, *WIREs Comput. Mol. Sci.* **2018**, *8*, e1345.
- [56] P. v. R. Schleyer, H. Jiao, B. Goldfuss, P. K. Freeman, *Angew. Chem. Int. Ed. Engl.* **1995**, *34*, 337–340; *Angew. Chem.* **1995**, *107*, 332–335.
- [57] M. K. Cyrański, P. v. R. Schleyer, T. M. Krygowski, H. Jiao, G. Hohlneicher, *Tetrahedron* **2003**, *59*, 1657–1665.
- [58] J. J. Eisch, J. E. Galle, S. Kozima, *J. Am. Chem. Soc.* **1986**, *108*, 379–385.
- [59] J. Köhler, S. Lindenmeier, I. Fischer, H. Braunschweig, T. Kupfer, D. Gamon, C.-W. Chiu, *J. Raman Spectrosc.* **2010**, *41*, 636–641.
- [60] H. Braunschweig, I. Krummenacher, J. Wahler, *Adv. Organomet. Chem.* **2013**, *61*, 1–53.
- [61] M. Saunders, R. Berger, A. Jaffe, J. M. McBride, J. O'Neill, R. Breslow, J. M. Hoffmann, C. Perchonock, E. Wasserman, R. S. Hutton, V. J. Kuck, *J. Am. Chem. Soc.* **1973**, *95*, 3017–3018.
- [62] E. Wasserman, R. S. Hutton, *Acc. Chem. Res.* **1977**, *10*, 27–32.
- [63] E. P. F. Lee, T. G. Wright, *Phys. Chem. Chem. Phys.* **1999**, *1*, 219–225.
- [64] S. K. Møllerup, Y. Cui, F. Fantuzzi, P. Schmid, J. T. Goettel, G. Bélanger-Chabot, M. Arrowsmith, I. Krummenacher, Q. Ye, V. Engel, B. Engels, H. Braunschweig, *J. Am. Chem. Soc.* **2019**, *141*, 16954–16960.
- [65] H. Fallah-Bagher-Shaidaei, C. S. Wannere, C. Corminboeuf, R. Puchta, P. v. R. Schleyer, *Org. Lett.* **2006**, *8*, 863–866.
- [66] M. D. Wodrich, C. Corminboeuf, S. S. Park, P. v. R. Schleyer, *Chem. Eur. J.* **2007**, *13*, 4582–4593.
- [67] A. Stanger, *J. Org. Chem.* **2006**, *71*, 883–893.
- [68] A. C. Tsepis, *Phys. Chem. Chem. Phys.* **2009**, *11*, 8244.
- [69] S. H. Vosko, L. Wilk, M. Nusair, *Can. J. Phys.* **1980**, *58*, 1200–1211.
- [70] C. Lee, W. Yang, R. G. Parr, *Phys. Rev. B* **1988**, *37*, 785–789.
- [71] A. D. Becke, *J. Chem. Phys.* **1993**, *98*, 5648–5652.
- [72] P. J. Stephens, F. J. Devlin, C. F. Chabalowski, M. J. Frisch, *J. Phys. Chem.* **1994**, *98*, 11623–11627.
- [73] R. Krishnan, J. S. Binkley, R. Seeger, J. A. Pople, *J. Chem. Phys.* **1980**, *72*, 650–654.
- [74] T. Clark, J. Chandrasekhar, G. W. Spitznagel, P. v. R. Schleyer, *J. Comput. Chem.* **1983**, *4*, 294–301.
- [75] R. Herges, D. Geuenich, *J. Phys. Chem. A* **2001**, *105*, 3214–3220.
- [76] D. Geuenich, K. Hess, F. Köhler, R. Herges, *Chem. Rev.* **2005**, *105*, 3758–3772.

- [77] S. Nees, T. Kupfer, A. Hofmann, H. Braunschweig, *Angew. Chem. Int. Ed.* **2020**, *59*, 18809–18815.
- [78] Deposition Numbers 1998824 (1), 1998823 (2), and 1998825 (3) contain the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum

Karlsruhe Access Structures service www.ccdc.cam.ac.uk/structures.

Manuscript received: October 30, 2020
Accepted manuscript online: November 19, 2020
Version of record online: December 27, 2020