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Halogenoborane mediated allene cyclooligomerization[†]

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The halogenoboranes XB(C₆F₅)₂ (X: Cl, Br) react with allene to give a mixture of the cyclotrimer 1,3,5trimethylenecyclohexane (1) and the related halogenoborylated cyclotetramerization products **3a** and **3b**. Alkyl-substituted allenes were catalytically cyclotrimerized metal-free by XB(C₆F₅)₂ (X: Cl, Br) to give *cis,trans*-2,4,6-trialkyl-1,3,5-trimethylenecyclohexanes under mild conditions.

Allenes serve as important organic building blocks.1 The parent allene, propadiene, has been cyclotrimerized at a variety of transition metal catalysts, but this usually gave predominately 1,2,4-trimethylenecyclohexane and only minor amounts of the 1,3,5-trimethylenecyclohexane isomer 1.2,3 Compound 1 had been synthesized in stoichiometric reaction sequences.⁴ We recently developed the metal-free, HB(C₆F₅)₂ catalysed formation of isomerically pure 1 by allene cyclotrimerization under mild conditions.⁵ We also had recently shown that a small series of 1-alkynes could be converted to linear oligomers by treatment with either of the halogenoboranes $XB(C_6F_5)_2$ 2a (X: Cl) or 2b (X: Br). The resulting products contained a halide substituent at one end of the oligoacetylene chain and the $B(C_6F_5)_2$ functional group at the other.⁶ This prompted us to investigate which type of a reaction allene would undergo with either of the reagents 2a and b. We have performed these reactions and here report about their surprising outcome.

We first carried out the reaction of excess allene with $ClB(C_6F_5)_2$ (2a) in d₈-toluene in a Young NMR tube at room temperature. After 7 h reaction time we monitored the NMR features of a mixture that contained the products 1,3,5-trime-thylenecyclohexane (1) and the chloroborylated allene tetramer **3a** in a *ca.* 1 : 1 ratio (both present in *ca.* 8 mol% in the mixture) plus unreacted **2a** (*ca.* 10 mol%) and allene (*ca.* 75 mol%). After 24 h the amount of the pair of allene cyclooligomerization products had almost doubled and there remained only *ca.* 3 mol% of the chloroborane **2a.** That was almost completely consumed after 48 h reaction time at room temperature. The products **1** and **3a** were identified spectroscopically from the mixture [**1**: δ 4.53, 2.70 (¹H)]. Compound **3a** shows the typical olefinic exo-methylene pairs of ¹H NMR resonances at δ 5.08,

4.84 (9-CH₂=) and δ 4.63, 4.48 (4,6-CH₂=) as well as the AXspin system of the diastereotopic 3,7-CH₂ pairs (δ 2.24, 2.00, ${}^{2}J_{\rm HH} = 14.0$ Hz) and the AB pattern of the 5-CH₂ at δ 2.58, 2.54. The 1-CH₂ group at boron and the 8-CH₂ unit give rise to 1 H NMR signals at δ 2.18 and δ 2.26, respectively (atom numbering scheme analogous to Fig. 1). Compound **3a** shows a broad 11 B NMR resonance at δ 71.5, which is typical for a planar tricoordinate Lewis acidic borane in this situation ($\Delta^{19}F_{\rm m,p} =$ 13.2 ppm, for details see the ESI†).

The reaction of allene with $BrB(C_6F_5)_2$ (**2b**) was carried out analogously. The reaction was directly followed by NMR spectroscopy. It resulted in the formation of the products **1** and **3b** in a *ca.* **1** : 2 ratio (4 h, r.t.). Compound **3b** was characterized from the mixture by NMR spectroscopy. It shows similar spectra as its



Fig. 1 A view of the molecular structure of compound 4b. Selected bond lengths (Å) and angles (°): B1-C1 1.634(2) B1-N21 1.661(2) C4-C11 1.323 (2) C6-C12 1.327(2) C9-C10 1.351(9) C9-Br1 1.910(5) B1-C1-C2 124.4(1) C1-C2-C8 108.5(1) C2-C8-C9 116.3(3) C8-C9-C10 124.5(6) C8-C9-Br1 118.4(4).



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chloro-substituted analogue **3a** (see the ESI† for details and the depicted spectra).

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We prepared compound **3a** on a preparative scale. After direct treatment with pyridine we isolated the Lewis adduct **4a** as a white solid in 53% yield (Scheme 1). It was characterized by C, H, N elemental analysis, by spectroscopy and by X-ray diffraction (see the ESI† for details). It showed very similar parameters as the analogous bromine containing compound **4b** (see below).

Treatment of *in situ* prepared **3b** with a slight excess of pyridine gave the adduct **4b**, which we isolated in 72% yield as a white solid on a 100 mg scale. The X-ray crystal structure analysis (Fig. 1) showed the central six-membered ring that was constructed by connecting three allene units. It features the exomethylene C=C double bonds annulated at carbon atoms C4 and C6 and it has the CH₂-CBr=CH₂ unit, derived from the fourth connected allene unit, attached at the ring carbon atom C2. This carbon atom also bears the CH₂-B(C₆F₅)₂ substituent, which has the pyridine donor added to its boron atom. The central six-membered carbocycle of compound **4b** features a distorted chair-like ring conformation.

In solution (CD₂Cl₂) compound **4b** shows the ¹H NMR = CH₂ signals of the symmetry equivalent pair of exo-methylene groups at the ring carbons C4 and C6 and the pair of signals of the C9=CH₂ moiety. The C3/C7 CH₂ hydrogen atoms are pairwise diastereotopic and the C8 and C1 CH₂ groups both show ¹H NMR singlets. Compound **4b** shows a ¹¹B NMR resonance in the typical tetra-coordinated borane range at δ -1.4 (Δ ¹⁹F_{m,p} = 5.6 ppm).

It is probably reasonable to assume that our reaction sequence starts with a halogenoboration reaction⁷ of allene (Scheme 2). This would generate the allylborane 5 that probably undergoes an allylboration⁸ reaction with a second equivalent of allene to form the intermediate 6. Since this contains an allylborane subunit it can undergo another allylboration reaction to give 7. This sequence could in principle be propagated to form a respective series of linear borylated allene oligomers 7, 8 etc., were there not the attractive possibility of these systems to undergo alternative intramolecular allylboration.9 In the case of 7 this would lead to the cyclization product 9. Subsequent XB(C₆F₅)₂ elimination provides an attractive pathway to the observed product 1,3,5-trimethylenecyclohexane (1). This would in principle constitute a cyclotrimerization of allene catalysed by the $XB(C_6F_5)_2$ reagents. However, the intramolecular allylboration of 8, giving the other experimentally observed products



Scheme 2 Reaction pathway for the $XB(C_6F_5)_2$ mediated allene cyclooligomerization.

3, represents a competing stoichiometric reaction branch that eventually removes the $XB(C_6F_5)_2$ reagent from the system (Scheme 2) and thus terminates the catalytic sequence of the formation of **1**.

Compounds **3a,b** are reactive boron Lewis acids. With the bulky ^tBu₃P Lewis base (**10**) they react by ring closure^{10,11} to eventually give the zwitterionic product **12** (Scheme 3). As a typical example, the reaction of the *in situ* generated chloride containing derivative **3a** was carried out with a *ca*. two molar equivalents of ^tBu₃P in toluene (24 h, 60 °C). Workup with pentane and dichloromethane in this case gave a *ca*. 79 : 21 mixture of compound **12** (characterization see below) and the [^tBu₃PH⁺]Cl⁻ phosphonium salt **11a** (³¹P NMR: δ 47.9, ¹J_{PH} ~463 Hz, some of the stoichiometric by-product **11a** was probably lost during the workup procedure).

The reaction between the more reactive borane 3b and ${}^{t}Bu_{3}P$ was carried out similarly (toluene, 24 h, r.t.). Workup in this case gave the pure compound 12 as a white solid, which we isolated in 65% yield. It was characterized by C, H, N elemental analysis, by spectroscopy and by X-ray diffraction (single crystals of compound 12 were obtained from dichloromethane/





Scheme 1 Reaction of allene with the halogenoboranes 2a and 2b.



Scheme 3 Reaction of the products 3 with two molar equiv. of ${}^{t}Bu_{3}P$.

pentane by the diffusion method). The X-ray crystal structure analysis shows the newly formed borataspiro[5,5]undecene framework (Fig. 2) which was selectively formed by boron induced allene tetramerization followed by phosphane induced ring closure reaction. The zwitterionic compound has the bulky ^tBu₃P-substituent attached at the C(9)=C(10) carbon–carbon double bond. The adjacent six-membered ring shows carbon atoms C4 and C6 which serve both as the ring sp²-carbons of the pair of exo-methylene C=CH₂ groups.

In solution (CD₂Cl₂) compound **12** shows the typical ¹H NMR doublet of the ^{*t*}Bu₃P-substituent. It features the ==CH signals of the newly formed internal olefinic moiety at δ 8.21 [¹H: d, ³J_{PH} = 27.8 Hz; ¹³C: δ 177.0 (broad)]. The BCH₂ group shows up as a ¹H NMR signal at δ 1.20 and the C(8)H₂ methylene group as a doublet at δ 2.06 (³J_{PH} = 4.5 Hz). The pair of olefinic exomethylene groups of the adjacent carbocyclic six-membered ring shows typical ¹H NMR signals at δ 4.61/4.41 and the resonances of the pairwise diastereotopic methylene hydrogen atoms at the C3/C7 pair and at C5 (see the ESI† for details).

We also reacted a pair of alkyl-substituted allenes with the halogenoboranes 2 and found a marked dependence of the overall reaction pathway on the substitution pattern. In the two cases investigated the competition between the two previously observed pathways, namely catalytic cyclotrimerization and stoichiometric formation of a tetramer derivative, was shifted to the catalytic side.

We treated *n*-octylallene (13c), which we prepared according to a procedure reported by Ma *et al.*¹² with 10 mol% of ClB(C₆F₅)₂ (2a) in d₈-toluene solution at 60 °C. Workup after 48 h reaction time involving purification by chromatography gave the cyclotrimer 14c as the major product (Scheme 4).



Compound 14c was isolated in 44% yield as a colorless oil. The NMR spectra showed the symmetry features of the cis,trans-2,4,6-trialkyl-substituted 1,3,5-trimethylenecyclohexane diastereomer. It features the ¹H NMR triplets of the 2,6-CH and the 4-CH ring hydrogens in a 2 : 1 intensity ratio. The $1-H_2C=$ unit shows one ¹H NMR resonance, whereas the $3,5-H_2C$ = moieties showed two due to their stereochemically unsymmetrical environment (see the ESI^{\dagger} for further details). The BrB(C₆F₅)₂ borane is an equally efficient metal-free cyclotrimerization catalyst for the n-octylallene 13c. The catalytic reaction was carried out under analogous conditions and gave the product 14c in 46% yield after workup (some minor byproduct was isolated (ca. 20%) but not positively identified as yet). We performed the XB(C₆F₅)₂ catalysed alkylallene cyclotrimerization reaction for a second example: the reaction of n-dodecylallene (13d) with either of the Cl/BrB(C_6F_5)₂ borane (10 mol%, 60 °C, 48 h in d₈-toluene) gave the tri-substituted cyclotrimer **14d** with cis,trans-attachment of the long chained alkyl groups as the major product. We isolated it as a colorless oil in 40% yield from the ClB(C_6F_5)₂ catalysed reaction [54% with BrB(C_6F_5)₂] (see the ESI[†] for the characterization of 14d).

We had previously shown that $HB(C_6F_5)_2$ serves as an efficient metal-free catalyst for the cyclotrimerization of allene to 1,3,5-trimethylenecyclohexane and of cyclohexylallene to cis,trans-2,4,6-tricyclohexyl-1,3,5-trimethylenecyclohexane.5 Our present study shows that the strongly electrophilic halogenoboranes $XB(C_6F_5)_2$ (X: Cl, Br) in principle can induce the same catalytic reaction. Both these strongly electrophilic halogenoboranes serve as catalysts for the cyclotrimerization of longchain *n*-alkylallenes **13c,d** to give the respective cyclotrimers 14c,d as the major products. Ring-closure and elimination is sufficiently effective to close the catalytic cycle with liberation of the respective $XB(C_6F_5)_2$ (X: Cl, Br) catalyst (Scheme 2). However, the reaction seems to be sensitive to sterics in the competition between ring-closure and further allene incorporation. In the case of the unsubstituted parent allene substrate we found a substantially competing further oligomerization step which eventually leads to the formation of the stoichiometric products 3a and 3b, thereby consuming the borane catalyst of the allene cyclotrimerization reaction and, thus, leading to termination of the catalytic reaction in this specific case. Nevertheless, our study has shown that allene oligomerization beyond trimerization is possible with such borane systems. This makes us hopeful that it might be possible to open novel ways of utilization of allenes by a further development of metal-free reactions along the characteristic lines that have shown up in this study.

Fig. 2 A view of the molecular structure of the zwitterionic FLP product 12. Selected bond lengths (Å) and angles (°): B1–C1 1.630(4) B1–C10 1.624(4) C4–C11 1.328(4) C6–C12 1.323(5) C9–C10 1.343(4) C9–P1 1.834(3) B1–C1–C2 113.5(2) C1–C2–C8 108.5(2) C1–B1–C10 107.0(2) C2–C2–C7 108.0(2) C2–C8–C9 114.0(2) C8–C9–C10 121.4(2) C9–C10–B1 125.7(3).



Scheme 4 $XB(C_6F_5)_2$ (X: Cl, Br) catalyzed cyclotrimerization of alkyl substituted allenes.



Scheme 5 Acid catalyzed isomerization reactions of allene-cyclotrimers to mesitylene derivatives.

We have started to use the allene cyclotrimerization products 1 and 14 as the starting materials for the conversion to the respective arene isomers. It is known that 1,3,5-trimethylenecyclohexane is resistant to thermal isomerization, but it was reported that it could be isomerized by treatment with acid.^{2a,13} We repeated this reaction: treatment of 1 with 5 mol% of p-toluene sulfonic acid in toluene solution at r.t. for 1.5 h cleanly converted 1 to mesitylene (Scheme 5). We also treated the *n*-octyl- and cyclohexylallene cyclotrimers $14c^{14}$ and $14e^{5}$ with p-toluene sulfonic acid. Compound 14c was fully converted on a preparative scale with 10 mol% of the acid catalyst during 48 h at r.t. The hexa-substituted arene 15c was isolated as an oil in 78% after workup. It was characterized by ¹H and ¹³C NMR spectroscopy (see the ESI[†] for details). Compound 14e needed slightly more vigorous conditions. It required treatment with 10 mol% of the *p*-toluene sulfonic acid catalyst at 80 °C for 5 h to become converted. This reaction is not overly selective, but we isolated the aromatic isomer 15e in 37% yield from the reaction mixture. The compound was characterized by spectroscopy and by an X-ray crystal structure analysis (see the ESI[†] for details). In principle, these reactions have shown that the metal-free X- $B(C_6F_5)_2$ catalyzed allene cyclotrimerization opens attractive pathways to the synthesis of interesting highly substituted arene products.

Conflicts of interest

There are no conflicts to declare.

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