The enhanced stability of this compound,⁸ probably due to the additional π -type interaction, is clearly demonstrated by the temperature dependence of the 60-MHz nmr spectra shown in Figure 3. Rapid exchange on the nmr time scale does not occur until a temperature of $\sim 70^{\circ}$ is reached, which implies that the exchange process in this system proceeds with a higher activation energy than that observed for any other system studied so far.

Further study of this system should lead to a better understanding of cyclopropyl π -type interactions and the nature of the electron-deficient bridge bond. To this end the exact analysis of the nmr spectrum is continuing and the crystal structure and optical spectra are currently under investigation.

Acknowledgment. We gratefully acknowledge the support of this research by National Science Foundation Grants GP-3908, GP-6762, and GP-8323.

(8) H. H. Jaffé, Z. Elektrochem., 59, 823 (1955).

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Comment on the Thermal Decomposition of Triphenylthallium

Sir:

In a recent interesting publication¹ concerned with thallium in organic synthesis is the statement: "Contrary to the suggestion of Gilman and Jones, we have shown that triphenylthallium does *not* disproportionate to biphenyl and thallium."² In the earlier experiments described by us³ we obtained biphenyl when triphenylthallium dissolved in xylene was refluxed under nitrogen. Also, in two check experiments, a solution of triphenylthallium in xylene was refluxed while a slow stream of dry carbon dioxide was admitted over the surface of the boiling solution. Subsequent to cooling and hydrolysis, there was obtained biphenyl and benzoic acid. It may be relevant to remark that Birch⁴ stated after heating what was presumably only a small quantity of solid triphenylthallium: "When heated it decomposes suddenly to give metallic thallium and a vapor possessing an odor of biphenyl."

We now confirm that biphenyl is indeed formed by thermal decomposition of triphenylthallium under our reported conditions which are more vigorous than those employed in ref 1. Thus, biphenyl is formed by thermal decomposition in xylene of freshly prepared triphenylthallium. These experiments were carried out with newly prepared materials. As in our earlier work the identity of the biphenyl was confirmed by the method of mixture melting points with an authentic specimen, and in the present work also by comparison of infrared spectra.

The following item may have some significance. An interesting and important aspect of the coupling reac-

(3) H. Gilman and R. G. Jones, J. Am. Chem. Soc., 62, 2357 (1940).
(4) S. F. Birch, J. Chem. Soc., 1132 (1934).

tions¹ is that they take place in the presence of excess TlBr (see footnote 4 of ref 1). In the reference to our experiment in which some biphenyl was obtained when phenyllithium was added to thallous chloride at -15° (see ref 3) it is of interest that the TlCl was present in excess because the phenyllithium was added to it slowly. Ordinarily, in the preparation of organothallium compounds, the thallous chloride or bromide is added slowly to the RMgX or RLi compound so that thallous halide is not present in excess. This suggests that a key step in the coupling reaction may be expressed in eq 2 of ref 1.

$$RT1 + 2TlBr \longrightarrow RTlBr_2 + 2Tl$$

There would then be little opportunity for this reaction to take place when TlBr is added to excess RMgX or RLi.

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Synthesis of Novel Substituted (3)-1,7-Dicarbaundecaborate(12) Ions from 1,8-Dicarba-closo-undecaborane(11) and Carbanions

Sir:

Recently, the Lewis acid behavior of 1,8-dicarbacloso-undecaborane(11)¹⁻³ and the properties of some of its Lewis base adducts⁴ have been reported. We now report the synthesis of a group of analogous ions, including two novel two-cage systems.

The previously described $1,8-B_9C_2H_{11}$ carborane reacts with 1 mole of methyllithium in diethyl etherhexane to produce a new $B_9C_2H_{11}CH_3^-$ ion I, which is isolated as its tetramethylammonium salt in 62% yield. Recrystallization from pure acetone yields white airsensitive crystals. The infrared spectrum of a Nujol mull of I exhibits absorptions at 4.01 (s), 7.73 (w), 9.10 (m), 9.89 (w), 10.24 (m), 10.62 (s), 12.12 (w), and 13.79 μ (s). Anal. Calcd for [(CH₃)₄N][B₉C₂H₁₁CH₃]: C, 34.40; H, 12.51; B, 46.41; N, 6.63; equiv wt, 209.8. Found: C, 35.02; H, 11.29; B, 46.78; N, 6.57; mol wt (osmometric in acetonitrile), 207.

The 60-Mc/sec ¹H nmr spectrum of the tetramethylammonium salt of I in deuterioacetone exhibited a sharp singlet of intensity 12 at τ 6.64, a broad singlet of intensity 2 at τ 8.55, and a broad singlet of intensity 3 at τ 9.65. The intensity 12 resonance was assigned to the cation methyl protons, and the intensity 2 and 3 resonances were assigned to the carborane protons and to the cage methyl protons, respectively. The 32.1-Mc/sec ¹¹B nmr spectrum of I is complex, with a high-field doublet of intensity 2, a set of overlapping doublets of intensity 3, a singlet of intensity 1, and a low-field set of overlapping doublets of intensity 3. This spectrum, except for the aforementioned singlet, bears a striking resemblance to the spectrum of the previously reported (3)-1,7-B₉C₂H₁₂⁻ ion,⁵ which can now be obtained in

⁽¹⁾ A. McKillop, L. F. Elsom, and E. C. Taylor, J. Am. Chem. Soc., 90, 2423 (1968).

⁽²⁾ The authors¹ have informed us that this observation was made under the conditions of their coupling reaction, *i.e.*, in refluxing tetrahydrofuran-benzene (1:1), thus indicating that triphenylthallium was not an intermediate in the coupling.

⁽¹⁾ F. N. Tebbe, P. M. Garrett, and M. F. Hawthorne, J. Am. Chem. Soc., 86, 4222 (1964).

⁽²⁾ C. D. Tsai and W. E. Streib, ibid., 88, 4513 (1966).

⁽³⁾ T. E. Berry, F. N. Tebbe, and M. F. Hawthorne, *Tetrahedron Letters*, 715 (1965).

⁽⁴⁾ F. N. Tebbe, P. M. Garrett, and M. F. Hawthorne, J. Am. Chem. Soc., **90**, 869 (1968).



Figure 1. Structure of 1,8-dicarba-closo-undecaborane.

80% yield by the reaction of $1,8-B_9C_2H_{11}$ with 1 mole of NaBH₄ in 1,2-dimethoxyethane.

The 1,8-B₉C₂H₁₁ carborane also reacts with a solution of Li-1,2-B₁₀C₂H₁₀CH₃⁶ in diethyl ether to form a corresponding B₉C₂H₁₁B₁₀C₂H₁₀CH₃⁻ ion II, which can be isolated as its tetramethylammonium salt in 72% yield. Recrystallization from water-ethanol yields white, airstable needles which do not melt to 320°. The infrared spectrum of a Nujol mull of II exhibits absorptions at 3.94 (s), 7.09 (m), 7.82 (w), 8.46 (m), 8.87 (m), 9.09 (m), 9.84 (m), 10.16 (m), 10.57 (s), 11.50 (w), 12.49 (w), and 13.69 μ (s). Anal. Calcd for [(CH₃)₄N][B₉C₂-H₁₁B₁₀C₂H₁₀CH₃]: C, 29.70; H, 9.97; B, 56.47; N, 3.86; equiv wt, 364. Found: C, 30.09; H, 10.09; B, 56.23; N, 3.96; mol wt (cryoscopic in ethanol), 363.

The 60-Mc/sec ¹H nmr spectrum of the tetramethylammonium salt of II in pyridine exhibited a sharp singlet of intensity 12 at τ 6.91, a sharp singlet of intensity 3 at τ 7.89, and a broad singlet of intensity 2 at τ 8.32. The resonances were assigned, as before, to cation methyl, carborane proton, and carborane methyl. The 32.1-Mc/sec ¹¹B nmr spectrum is complex, with the highfield doublet reminiscent of the high-field doublet in I and in the aforementioned (3)-1,7-B₉C₂H₁₂⁻. This doublet has intensity 2:17 relative to the remainder of the resonances which features a large "doublet" very reminiscent of 1,2-B₁₀C₂H₁₀R₂ systems.⁷

The 1,8-B₈C₂H₁₁ carborane also reacts with the similarly formed Li-1,10-B₈C₂H₈C₆H₅ in diethyl etherbenzene to produce an analogous B₉C₂H₁₁B₈C₂H₈C₆H₅ ion III, which is isolated as the tetramethylammonium salt in 52% yield. Recrystallization from acetonewater yields air-stable white crystals. The infrared spectrum of a Nujol mull of III exhibited absorptions at 4.03 (s), 7.79 (w), 8.02 (w), 8.54 (w), 9.11 (m), 9.27 (w), 9.72 (m), 10.19 (m), 10.58 (s), 10.85 (w), 11.54 (w), 12.01 (m), 13.08 (s), 13.58 (w), and 14.35 μ (s). Anal. Calcd for [(CH₃)₄N][B₉C₂H₁₁B₈C₂H₈C₆H₅]: C, 41.88; H, 9.02; B, 45.63; N, 3.49; equiv wt, 402.4. Found: C, 39.88; H, 8.94; B, 45.63; N, 3.42; mol wt (cryoscopic in ethanol), 414.

The 60-Mc/sec ¹H nmr of the tetramethylammonium salt of III exhibits a sharp singlet of intensity 12 at τ



Figure 2. Proposed structure of the $B_9C_2H_{11}B_{10}C_2H_{10}CH_3^-$ ion.



Figure 3. Proposed structure of the $B_9C_2H_{11}B_8C_2H_8C_6H_5^-$ ion.

6.64, a broad singlet of intensity 2 centered at τ 8.70, and two multiplets of total intensity 5 centered at τ 2.45. These resonances were assigned to cation methyl, carborane protons, and phenyl protons, respectively. The 32.1-Mc/sec ¹¹B nmr exhibited the familiar highfield doublet of intensity 2, a multiplet of intensity 3, and a low-field multiplet of intensity 12, which was marked by the characteristic doublet exhibited in 1,10-B₈C₂H₈R₂ systems.⁴

The nmr results indicated that the compounds formed were boron-substituted $(3)-1,7-B_9C_2H_{11}R$ ions. Since there is considerably more complexity in the 32.1-Mc/sec ¹¹B spectrum of I compared with the spectrum of $(3)-1,7-B_9C_2H_{12}$, we tentatively propose that the various R groups are substituted in the chemically equivalent but asymmetric 4 or 8 positions. This would be the point of substitution resulting from nucleophilic attack by the Lewis base at boron atoms 3, 5, 7, or 9 on 1.8-B₉C₂H₁₁ (Figure 1). Since the 60-Mc/sec 1 H spectra of II and III exhibited no other carborane proton resonances other than those expected from a $(3)-1,7-B_9$ - C_2H_{12} ion, it is suggested that the point of substitution on the B_{10} and B_8 cages is C-1 in each case. This led us to propose the structures shown in Figures 2 and 3 for ions II and III.

Further preliminary work suggests that similar ions of this type may be prepared and that transition metal complexes⁸ incorporating this basic unit may be formed.

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(8) M. F. Hawthorne, D. C. Young, T. D. Andrews, D. V. Howe,
R. L. Pilling, A. D. Pitts, M. Reintjes, L. F. Warren, Jr., and P. A.
Wegner, J. Am. Chem. Soc., 90, 879 (1968).
(9) NASA Fellow, 1966-1968.

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⁽⁵⁾ M. F. Hawthorne, D. C. Young, P. M. Garrett, D. A. Owen, S. G. Schwerin, F. N. Tebbe, and P. A. Wegner, J. Am. Chem. Soc., **90**, 862 (1968).

⁽⁶⁾ M. M. Fein, D. Grafstein, J. E. Paustian, I. Bobinski, B. M. Lichstein, N. Mayes, N. N. Schwartz, and M. S. Cohen, *Inorg. Chem.*, 2, 1115 (1963).

⁽⁷⁾ R. L. Pilling, F. N. Tebbe, M. F. Hawthorne, and E. G. Pier, Proc. Chem. Soc., 402 (1964).