Addition of Ethylene or Hydrogen to a Main Group Metal Cluster under Mild Conditions


Abstract: Reaction the tin cluster Sn6(ArH)6 (ArH = C6H5-2,6-(C6H5-2,4,6-Me3)2) with excess ethylene or dihydrogen at 25 °C / 1 atmosphere yielded two new clusters that incorporated ethylene or hydrogen. The reaction with ethylene yielded Sn6(ArH)6(C6H2)3 that contained five ethylene moieties bridging four aryl substituted tin atoms and one tin-tin bond. Reaction with H2 produced a cyclic tin species of formula (SnH)6, which could also be synthesized by the reaction of [(ArH)5Snµ-Cl]2 with DIBAL-H. These reactions represent the first instances of direct reactions of isolable main group clusters with ethylene or hydrogen under mild conditions. The products were characterized in the solid state by X-ray diffraction and IR spectroscopy and by multinuclear NMR and UV-Vis spectroscopy. Density functional theory calculations were performed to explain the reactivity of the cluster.

In 2005 it was shown that a main group molecule could react with dihydrogen at room temperature and atmospheric pressure,[1] Since then a wide variety of main group compounds have been investigated for their reactions with small molecules under mild conditions.[2-5] Such reactions are dependent on the existence of donor and acceptor orbitals of suitable symmetry and modest energy separation.[6] Thus, multiply bonded or unsaturated main group species have commonly been used. For example, heavier group 14 alkyne, Ar3E=EEAr3E, and carbene, =E(=E)2, analogues (E = Ge or Sn; Ar3E = C6H5-2,6-(C6H5-2,4,6-Me3)2) react readily with hydrogen, ethylene and other small molecules under mild conditions.[7-10] Activation of small molecules can also be effected by stable carbenes such as :Ct(Bu)(IPr)2N[11] or frustrated Lewis pairs using a phosphine or related electron donor and B(C6F5)3 as the acceptor.[12] Several reactions have been shown to be reversible, which also has generated widespread interest.[13-15] The reactivity of these main group compounds toward small molecules can resemble that of transition metal complexes, and thus, may have use in catalytic applications.[16]

The reactivity of main group clusters towards small molecules under ambient conditions has remained virtually unexplored.[17] There have been a few theoretical studies on dihydrogen activation by aluminum clusters,[18] and the activation of ammonia-borane by a gallium nitrogen cage compound.[19] Main group molecular clusters are also of interest because the coordination of their constrained atoms may resemble that of atoms at elemental surfaces.[20-22] Herein we report the reactions of the tin cluster Sn6(ArH)6 (ArH = C6H5-2,6-(C6H5-2,4,6-Me3)2) with excess ethylene or dihydrogen to afford the products 1 and 2 as depicted in Scheme 1. The product 1 shows that the initial cluster has absorbed five ethylene molecules. A reaction between the tin cluster and dihydrogen yields the tin hydride 2. Significantly, both reactions involve the loss of the unsubstituted tin atoms in the cluster. The syntheses of 1 and 2 are described as well as their characterization by NMR spectroscopy and X-ray crystallography. Computational investigations of model systems for 1 and 2 are also described.

Compound 1 was synthesized by treating Sn6(ArH)6[20] in THF with ethylene under ambient conditions (Scheme 1). The initially dark purple mixture was stirred for 2.5 days at 25 °C to afford a dark red solution and a metallic precipitate, assumed to be elemental tin. After work-up, compound 1 was isolated in 14 % yield. Colorless crystals were grown from diethyl ether at 6 °C overnight. Compound 1 crystallizes in a triclinic P1 space group with two diethyl ether solvent molecules.[24] Figure 1 depicts the solid state structure of 1 and it can be seen that the initial tin cluster, Sn6(ArH)6, has incorporated five ethylene molecules and the four unsubstituted tin atoms have been eliminated. One C6H5 moiety is disordered over two positions, each with 50 % occupancy and only one of these sites (C7A and C8A) is shown.
Each tin is tetrahedrally coordinated and carries one Ar ligand. Two tins, Sn(1) and Sn(2), are bound also to three carbon atoms from different ethylenes whereas Sn(3) and Sn(4) are bonded to two carbon atoms from bridging ethylenes as well as to each other. Thus there is one intact Sn-Sn bond (i.e., Sn(3)-Sn(4), 1.28549(5) Å), similar in length to the ethylene bridged tin-tin single bond in $\text{Ar}_2\text{Sn}[(\mu\eta^1\eta^1\text{C}_2\text{H}_4)_2\text{SnAr}]_4$, formed by reversible addition of ethylene to the distannylene $\text{Ar}_2\text{Sn}[(\mu\eta^1\eta^1\text{C}_2\text{H}_4)_2\text{SnAr}]_4$. The average Sn-C(ethylene) and ethylene C-C bond lengths are 2.177 Å and 1.527 Å, respectively. These are normal for Sn-C and C-C single bonds.[25,26] Interligand angles at the tin atoms are near 109.5° indicating tetrahedral geometry although disorder at the C7-C8 moiety and Sn-Sn bond cause deviation from the ideal value. The $^1$H and $^{13}$C NMR spectra of compound 1 indicate the presence of symmetry in the molecule in solution as only two unique sets of signals corresponding to $\text{Ar}_2\text{Sn}[(\mu\eta^1\eta^1\text{C}_2\text{H}_4)_2\text{SnAr}]_4$ ligand environments are observed. The $^{119}$Sn NMR spectrum reveals two signals at 336.0 [Sn(3) and Sn(4)] and 1.1 ppm [Sn(1) and Sn(2)], indicating two tin environments whose chemical shifts are consistent with reported $\text{Sn(4)}$ and 1.1 ppm $\text{Sn(1) and Sn(2)}$, indicating two tin environments. The overall tetrameric structure may be contrasted with that of the more sterically encumbered hydrogen bridged dimer $\text{Ar}_2\text{Sn}[(\mu\text{H})_2]\text{SnAr}]_2$ or that of the asymmetric stannylstannylene $\text{Ar}_2\text{PhSn}[(\mu\eta^1\eta^1\text{C}_2\text{H}_4)_2\text{SnAr}]_4$ and related species.[30] The orientation of the hydrogens alternate on either side of the Sn4. The Sn-Sn bond lengths (2.8433(4) and 2.8050(3) Å) are slightly shorter than the Sn-Sn single bonds in the other terminal tin hydrides,[29,30] but within the range of typical Sn-Sn single bonds. There are a few structures in the literature that incorporate a Sn4 ring, but none have Sn-H bonds. The known species have formulas Sn$_2$L$_5$ (L = alkyl ligand) or SnL$_4$ (L$'$ = bidentate ligand).[31-33] The $^1$H NMR spectrum of 2 shows a hydride signal at 3.97 ppm, and the $^{119}$Sn NMR spectrum at ~24.3 ppm, $J_{\text{Sn-H}} = 1590$ Hz.[34]

The reactivity of the tin cluster towards H$_2$ and ethylene was investigated computationally using the PBE0 hybrid functional[55-58] with def2-TZVP basis sets.[38] A model compound Sn$_4$Ph$_4$ was used to lower computational cost. The calculations show that the reaction of dihydrogen with the Sn$_4$Ph$_4$ cluster yields an addition product Sn$_4$H$_2$Ph$_4$ (Figure 3) initially, whose formation is thermodynamically disfavored in the gas phase ($\Delta H = 4$ kJ mol$^{-1}$; $\Delta G = 39$ kJ mol$^{-1}$). The Gibbs energy or activation was found to be 134 kJ mol$^{-1}$, consistent with the fact that the reaction with H$_2$ required mild heating and an excess of hydrogen gas to proceed. A similar addition product was located for the reaction of Sn$_4$Ph$_4$ with ethylene (see Figure 3). However, the gas phase reaction is exothermic and only slightly disfavored by entropy ($\Delta H = -33$ kJ mol$^{-1}$; $\Delta G = -22$ kJ mol$^{-1}$). The Gibbs energy of activation is also significantly smaller, ~66 kJ mol$^{-1}$, consistent with experimental observations.
The frontier orbitals of SnPh₄ have both electron donating and accepting features (see SI), which rationalizes the relatively facile formation of Sn₆H₂Ph₄ and Sn₆(C₃H₄)Ph₄. The modelling of mechanisms for the formation of [Sn(H)Ph₄]₅, Sn₆Ph₄(C₂H₅)₄, and elemental tin is beyond the scope of this work. However, the peculiar structure of 1 with an unreacted Sn-Sn bond prompted us to investigate the possible insertion of a sixth ethylene to Sn₆Ph₄(C₂H₅)₄ to form Sn₆Ph₄(C₂H₅)₅. The structures of ethylene, Sn₆Ph₄(C₂H₅)₂ and Sn₆Ph₄(C₂H₅)₄ were optimized (see SI) and their energies compared, which revealed that the addition of a sixth equivalent of ethylene is thermodynamically favored (ΔG = -12 kJ mol⁻¹) though the reaction could be prevented by kinetic factors. Unfortunately, we could not locate a transition state for the addition of the sixth ethylene molecule, thus the magnitude of the activation barrier remains unknown.

![Figure 3](image)

Figure 3. Optimized structures for the production addition products of Sn₆Ph₄ with H₂ (left) and ethylene (right).

In summary, we have described the syntheses and characterization data for two new insertion products of small molecules to a tin cluster under mild conditions. Further studies on the reactivity of Sn₆[(ArMe₆)₄] towards other small molecules and attempts to prepare Sn₆[(ArMe₆)₄(C₂H₅)₄] are on-going.

**Experimental Section**

All manipulations were carried out under anaerobic and anhydrous conditions by using modified Schlenk line techniques under a dinitrogen atmosphere or in a Vacuum Atmospheres HE-43 drybox. Solvents were dried and stored over sodium. Physical measurements were performed under anaerobic and anhydrous conditions. ¹H, ¹³C[¹H] and ¹¹⁹Sn[¹H] NMR spectra were obtained on a Varian 400 or 600 MHz spectrometers and referenced to known standards. IR spectra were recorded as Nujol mulls between CaF₂ plates on a Perkin-Elmer 1430 Infrared Spectrometer. UV-visible spectra were recorded as dilute toluene solutions in 3.5 mL quartz cuvette using an Olis 17 Modernized Cary 14 UV/Vis/NIR Spectrophotometer. Melting points were determined on a Meltemp II apparatus using glass capillaries sealed with vacuum grease, and are uncorrected. All starting materials were obtained from commercial sources and used as received. Sn₆[(ArMe₆)₄]₅ and [(ArMe₆)Snµ-Cl]₅ were prepared by literature procedures.

**Synthesis of Sn₆[(ArMe₆)₄(C₂H₅)₄](1)**: 0.700 g of Sn₆[(ArMe₆)₄]₅ was dissolved in ca. 40 mL of THF. The flask was flushed with ethylene gas for ca. 45 min at 25 °C and then stirred under an ethylene atmosphere for another 48 h. The initially dark purple solution became dark red with a grey precipitate (assumed to be elemental tin). The mixture was allowed to settle and filtered. The filtrate was evaporated and the residue was dissolved in ca. 20 mL of Et₂O. Filtration followed by reduction in volume to ca. 6 mL and storage at 6 °C overnight yielded colorless crystals of 1 which were dried under vacuum. Yield 0.080 g (14 %). ¹H NMR (25 °C, CD₂Cl₂, 600 MHz): δ = -0.10 (t, 3JH-H = 8.0 Hz, 4H, Sn-CH₂CH₂Sn), 0.11 (m, 3JH-H = 8.0 Hz, 4H, Sn-CH₂CH₂Sn), 0.52 (t, 3JH-H = 24.0 Hz, 8H, Sn-CH₂CH₂Sn), 0.74 (t, 3JH-H = 8.0 Hz, 4H, Sn-CH₂CH₂Sn), 2.04 (s, 24H, Mes-CH₂). ²⁷¹Sn NMR (25 °C, CD₂Cl₂): 2.07 (s, 24H, Mes-CH₂), 2.29 (s, 12H, Mes-CH₂), 2.34 (s, 12H, Mes-CH₂), 6.83, 6.84, 6.87, 6.91, 7.19 ppm (aromatic H, 28H), ¹³C[¹H] NMR (25 °C, CD₂Cl₂, 151 MHz): δ = 6.5, 9.2, 9.5, 13.9, 14.4, 19.8 (CH₂-CH₂), 21.4, 21.6, 21.6, 23.1 (Mes-CH₂), 31.9 (CH₂-CH₂), 128.4, 128.8, 128.9, 129.0, 129.2, 135.1, 135.3, 135.5, 135.7, 135.9, 135.6, 141.9, 142.7, 147.2, 150.2 ppm (aromatic C), three ipso-C signals were not observed, ¹¹⁹Sn[¹H] NMR (25 °C, CD₂Cl₂, 149 MHz): 336.0 (Sn-Sn) and 1.1ppm (Sn-CH₂). Melting point: 152-155 °C (decomp.). IR in Nujol mull (cm⁻¹) with CaF₂ plates: 650, 520, 475, 335 and 315 (Sn-C, stretching and bending). UV-Vis (toluene, nm): 307, 321 and 330.

**Synthesis of Sn₆[(ArMe₆)₄](2)**: Method A. Sn₆[(ArMe₆)₄](0.661g) was dissolved in ca. 25 mL of toluene and warmed to 60 °C. The mixture was stirred under an H₂ atmosphere for 3.5 h and then cooled to room temperature. Stirring was continued overnight without any color change but a significant amount of grey precipitate (elemental tin) was formed. The precipitate was allowed to settle and the solution was filtered. The volume was reduced to ca. 10 mL and 5 mL of THF was added. The mixture was placed in a fridge. Crystals of compound 2 were collected from the mixture which contained mainly unreacted Sn₆[(ArMe₄)₄].

**Method B.** A solution of disobutyl aluminium hydride in hexanes (3.2 mL, 1 M, 3.2 mmol, diluted with 2 mL of hexanes) was added dropwise to the diethyl ether solution of [(ArMe₄)Sn(CH₂)]₅ (0.708 g in ca. 30 mL of diethyl ether) at -78 °C. The solution became orange and was stirred for 2 h. A green precipitate was formed and removed by filtration. Storage of the diethyl ether solution at -30 °C overnight yielded yellow crystals of 2. Yield 0.135 g (21 %). ¹H NMR (25 °C, CD₂Cl₂, 600 MHz): δ = 1.66 (s, 12H, Mes-CH₂), 1.80 (s, 12H, Mes-CH₂), 2.07 (s, 12H, Mes-CH₂), 2.15 (s, 12H, Mes-CH₂), 2.33 (s, 12H, Mes-CH₂), 3.97 (s, 4H, Sn-H), 6.73, 6.77, 6.82, 7.07, 7.10, 7.12 ppm (aromatic H, 28H), ¹³C[¹H] NMR (25 °C, CDCl₃, 151 MHz): δ = 21.5, 21.8, 22.0, 22.4 and 23.7 ppm (Mes-CH₂), 128.4, 128.5, 128.7, 128.8, 128.9, 129.0, 129.3, 135.1, 135.3, 135.5, 135.7, 135.9, 136.0, 141.7, 142.0, 144.5, 150.0 and 150.2 ppm (aromatic C). ¹¹⁹Sn NMR (25 °C, CDCİ₃, 149 MHz): δ = -324.3 (d, 3JH-H = 1590 Hz, Sn-H) [37] Melting point: >300 °C. IR in Nujol mull (cm⁻¹) with CaF₂ plates: 1845 (Sn-H stretching), 730 (Sn-H bending). UV-Vis (toluene, nm): 296 and 308.

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