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Here we provide definitive EPR evidence for the existence of α -triphenylstannylyvinyl radicals in the low temperature O-directed free radical hydrostannation of dialkyl propargylic alcohols with $\text{Ph}_3\text{SnH}/\text{cat. Et}_3\text{B}$ and O_2 in PhMe . Isotropic hyperfine splitting patterns and spectral simulations confirm the assignments made. In the case of the α -triphenylstannylyvinyl radical (Z)-2, an isotopic $^{119/117}\text{Sn}$ hyperfine coupling constant of 9.5 mT (95 G) was measured along with a ^1H hyperfine coupling constant of 1.1 mT.

Although the mechanism of the O-directed free radical hydrostannation of propargylically-oxygenated dialkyl acetylenes with stannanes¹ has been the subject of considerable debate² over the past decade, quite recently this controversy was unambiguously resolved^{1a,3} in favor of the proposals of Hale and Manaviazar,^{1a,4} and Alabugin,⁵ who collectively provided strong experimental support for such reactions proceeding by way of an entirely free radical mechanism that operates under strict O-Sn coordinative control.^{1,4-7} In an effort to further validate these conclusions, we recently undertook an examination of the O-directed hydrostannation¹ of several dialkyl acetylenic alcohols by electron paramagnetic resonance (EPR) spectroscopy, and here we present the first readily interpretable EPR spectra of two α -stannylyvinyl radical intermediates (Z)-2 and (Z)-9, generated from the alkynol 1 and 8 by application of our standard, *chemically-initiated*, $\text{Ph}_3\text{SnH}/\text{cat. Et}_3\text{B}/\text{O}_2$ reaction conditions¹ at high substrate concentration (0.5 M) and low reaction temperature (-10 to -50 °C).

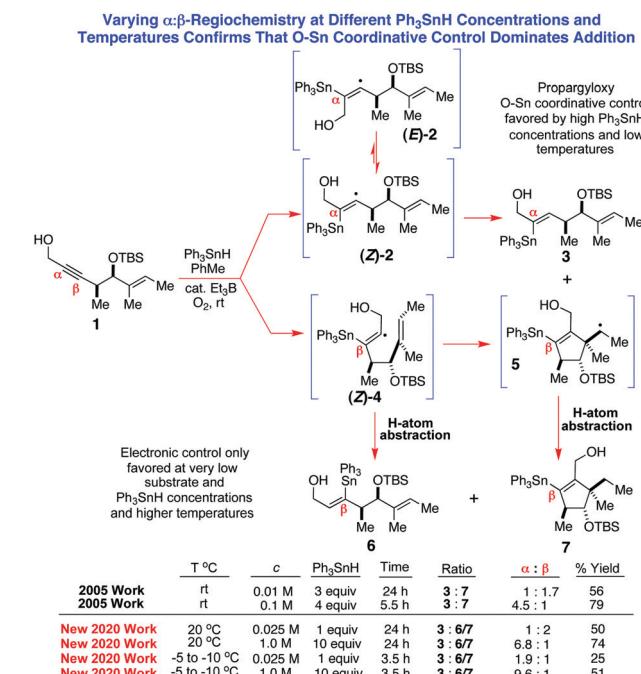
Now with respect to the alkynol 1, our previous mechanistic work^{3,4} had already confirmed that three different radicals 2, 4, and 5 are produced to varying degrees in the room temperature hydrostannation of 1 at different stannane concentrations

EPR evidence for α -triphenylstannylyvinyl radicals in the O-directed hydrostannation of dialkylacetylenes with $\text{Ph}_3\text{SnH}/\text{cat. Et}_3\text{B}/\text{O}_2$ †

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(see Scheme 1). Our finding, also, that the proportion of α -vinylstannane 3 increased steadily as the concentration of stannane increased provided strong mechanistic evidence for these hydrostannations operating under O-Sn coordinative control,^{1,3-7} since a purely electronically-controlled Sn-radical addition mechanism² would give rise to an invariable regiochemical outcome as the stannane concentration increased, provided the reactions were run at identical temperatures.

Indeed, our 2020 work^{3a} very clearly demonstrated the significant influence that temperature could have on the regiochemical outcome of the hydrostannation of 1 with $\text{Ph}_3\text{SnH}/\text{cat. Et}_3\text{B}$, with



P. Dimopoulos, J. George, D. A. Tocher, S. Manaviazar, K. J. Hale *Org. Lett.* **2005**, *7*, 5377.
H. A. Watson, S. Manaviazar, H. G. Steeds, K. J. Hale *Tetrahedron* **2020**, *76*, 131061.

Scheme 1 O-Sn coordinative control with probe 1 is favored by high stannane concentrations and low temperatures.

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the proportion of the α -tin adduct **3** increasing progressively as the reaction temperature decreased, due to lower temperatures promoting a much greater degree of O–Sn coordination between the propargyl O-atom and the stannane, and the newly generated stannyl radical. It was reasoned, therefore, that if the reaction temperature could be decreased further, to -50 °C let us say, this might allow the desired α -vinylstannane **3** to be obtained almost exclusively, and we might thus be able to observe the α -triphenylstannylvinyl radical (*Z*)-**2** as essentially a single regioisomer by EPR. If this proved to be so, then (*Z*)-**2** would, of course, be expected to resonate as an apparent “doublet” with attendant long-range, low-intensity, $^{117,119}\text{Sn}$ hyperfine splittings, although more formally it would be classified as a double doublet.

With this in mind, we initially tried to generate the α -stannylvinyl radical (*Z*)-**2** by photoirradiation (30 W Hg-lamp) of solutions of **1** and Ph_3SnH (1 equiv.) in dry cyclohexane, THF, and PhMe respectively, at room temperature and below (-50 °C), but we were never able to obtain a satisfactory EPR spectrum of (*Z*)-**2**, even when di-*t*-butylperoxide (DTBP) was used as a reaction initiator for some of the runs. In light of these problems, we resolved to generate (*Z*)-**2** by treatment of **1** with Ph_3SnH (1 equiv.), cat. Et_3B (0.1 equiv.) and O_2 in dry THF at -10 °C inside a quartz capillary EPR tube. Under such circumstances, a most encouraging EPR spectrum of predominantly (*Z*)-**2** was obtained, and it is shown in Fig. 1.

It consisted of an apparent “doublet”, due to the newly created vinyl radical coupling to its β -CH. However, because of the rather weak intensity of this signal, we were unable to satisfactorily detect the requisite $^{117,119}\text{Sn}$ hyperfine splittings, because of the high level of background noise. In fact, it was only after we completely abandoned the use of THF and quartz capillary EPR tubes, and we effected the hydrostannation of **1** in PhMe, in a 4 mm EPR tube at -50 °C (223 K), that we were able to finally observe the characteristic “doublet” and accompanying $^{117,119}\text{Sn}$ hyperfine splittings that *unambiguously* confirmed the identity of the α -stannylvinyl radical (*Z*)-**2**. Importantly, the new signal observed (see Fig. 2) once more indicated that we were dealing with predominantly a single Boltzmann conformer/invertomer. However, before we could be absolutely certain

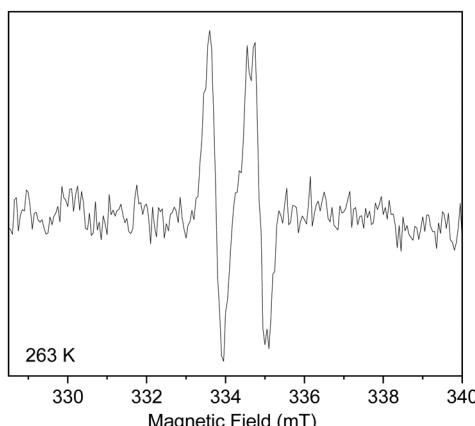


Fig. 1 The EPR spectrum of the α -stannylvinyl radical (*Z*)-**2** generated from the O-directed hydrostannation of a THF solution of **1**, Et_3B (0.1 equiv.), Ph_3SnH (1 equiv.) at 263 K (-10 °C).

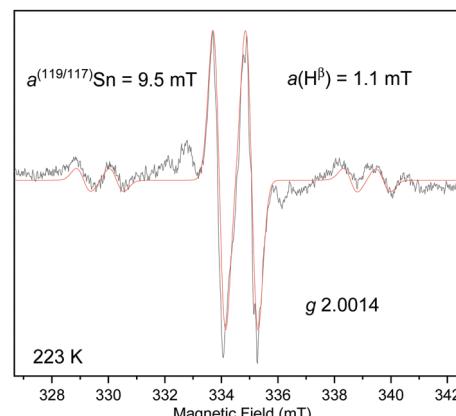


Fig. 2 EPR spectrum of the α -stannylvinyl radical (*Z*)-**2** generated from treatment of **1** with Ph_3SnH (1 equiv.) and Et_3B (0.1 equiv.)/ O_2 in PhMe at 223 K (-50 °C). The simulated signal is overlayed in red.

of our assignment, we felt that we needed to simulate the EPR spectrum of the radical (*Z*)-**2** with the computer programme EasySpin.⁸

After inputting the experimentally determined *g* and *a* values for (*Z*)-**2**, and parameterizing the programme for the presence of C, H and Sn, our simulation of the α -stannylvinyl radical (*Z*)-**2** produced the EPR spectrum shown in Fig. 2 (in red) which has been superimposed upon our experimentally-determined spectrum at 223 K. Clearly there is an excellent correlation, right down to the $^{117,119}\text{Sn}$ hyperfine couplings which, although unresolved, have similar hyperfine splitting values. The aforementioned simulation enabled the *g* value (*g* = 2.0014) and the hyperfine splittings, $a(\text{H}^\beta)$ = 1.1 mT and $a^{(119/117)}\text{Sn}$ = 9.5 mT, to be accurately determined.

Having secured this excellent result in PhMe at -50 °C, we next decided to use near identical experimental conditions to acquire the α -stannylvinyl radical **9** from 3-hexyn-2-ol (**8**). In this instance, the 230 K (-43 °C) EPR spectrum in PhMe gave rise to the anticipated “apparent” triplet (Fig. 3) for the α -stannylvinyl radical (*Z*)-**9**. Following EasySpin analysis⁸ of the radical (*Z*)-**9**, the simulated EPR spectrum in Fig. 3 was obtained; it is again shown in red. It overlaid perfectly with the experimentally-derived spectrum of (*Z*)-**9** at 230 K, whose “triplet” at *g* = 2.0021 had a hyperfine splitting of 0.43 mT (4.3 G) due to the radical coupling with the neighboring $-\text{CH}_2-$ of the Et. Significantly, the simulated spectrum highlighted how the weak intensity of the signal and the strong background noise were both preventing the Sn hyperfine couplings from being seen.

On this basis, we repeated the aforementioned hydrostannation reaction multiple times at progressively lower temperatures on each occasion, in an attempt to reduce the background noise and reveal the $^{117,119}\text{Sn}$ hyperfine splittings.

Although these efforts did not provide the requisite information, they did lead to us observing additional lines of steadily increasing intensity and partial asymmetry as we moved closer to 200 K (see Fig. 4). We believe that these additional signals are due to the rate of inversion of the two interconverting (*E*)- and



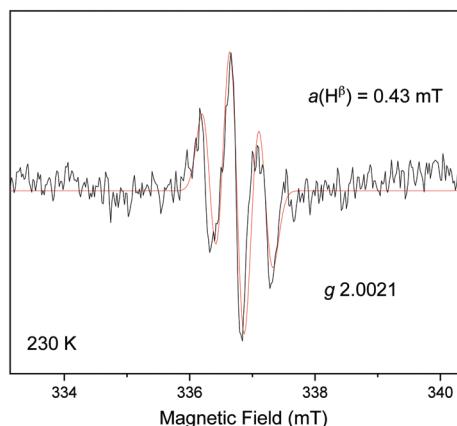
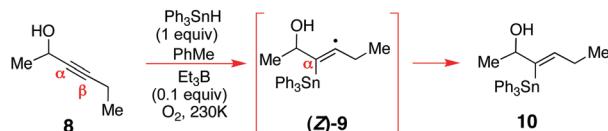


Fig. 3 EPR spectrum of the α -triphenylstannylyvinyl radical (**Z**)-9 generated from 3-hexyn-2-ol (**8**) by hydrostannation with Ph_3SnH (1 equiv.) and Et_3B (0.1 equiv.)/ O_2 in PhMe at 230 K. Simulation in red.

(**Z**)- α -stannylyvinyl radical invertomers being markedly slowed down, to the extent where both conformers of **9** are now observable. According to Galli and Rappoport,⁹ the barriers to inversion of alkyl-bearing σ -vinyl radicals are typically low, and of the order of 1–3 kcal mol^{−1}. It is thus not at all surprising that a decrease in reaction temperature could now be making this process observable on the EPR timescale.

The quite different *g* values observed for these signals would appear to corroborate this proposal, since quite profound structural differences would exist between the two radical conformers. In this connection, the carbon side chains would be positioned *anti*- to one another in (**Z**)-9, while in (**E**)-9 they would be oriented *syn* to each other, and in quite different steric and electronic environments. While this is one possible

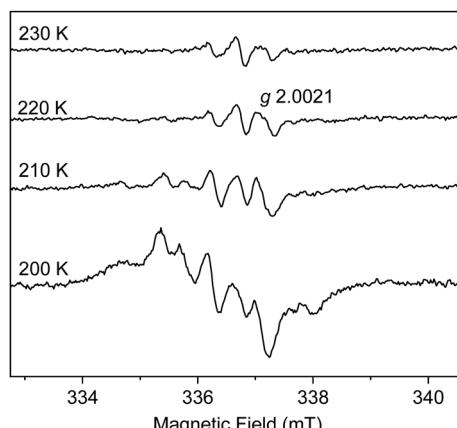
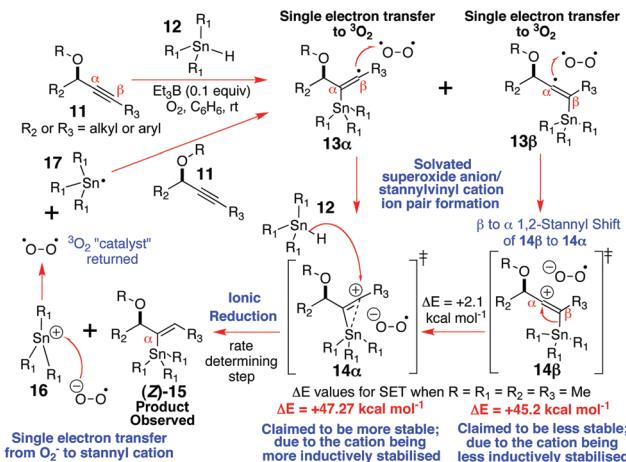


Fig. 4 The multiple EPR spectra obtained from hydrostannation of 3-hexyn-2-ol (**8**) with Ph_3SnH (1 equiv.) and Et_3B (0.1 equiv.) and O_2 in PhMe over a range of temperatures between 200–230 K.

The Invalid Electronically-Controlled Stannylyvinyl Cation Mechanism For The Hydrostannation of Alkyl- and Aryl-Acetylenes with $\text{R}_3\text{SnH}/\text{cat. Et}_3\text{B}/\text{O}_2$



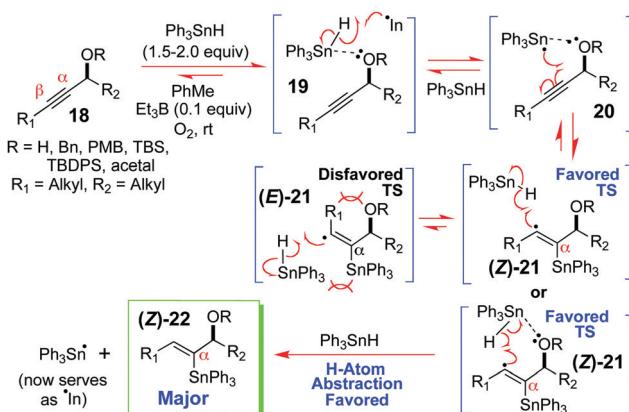
Scheme 2 Organ's stannylyvinyl cation mechanistic proposal.^{2a–d}

explanation of these intriguing effects, equally well, one can offer a rotational isomer rationale of these phenomena.

Given that *continuous wave* EPR spectroscopy cannot detect radical intermediates with lifetimes less than 10^{-6} seconds, and we have been able to continuously acquire these EPR spectra for the α -stannylyvinyl radicals (**Z**)-2 and (**Z**)-9 over timeframes of more than 40 min at -43 to -50 °C, our observations do not align well with Organ's stannylyvinyl cation mechanistic theory^{2a–d} for how such alkyne hydrostannation reactions proceed. According to this proposal (Scheme 2), a non-regiocontrolled stannylyvinyl radical addition occurs to the acetylene.^{2a,b} A fast, near *instantaneous* (10^{-15} s), non-bonded outer-sphere electron-transfer is then suggested to take place from the resulting mixture of α - and β -stannylyvinyl radical intermediates **13** α and **13** β to O_2 , at essentially the diffusion-controlled limit, followed by a cationic equilibration to the all α -stannylyvinyl cation superoxide ion pair **14** α , which is then ionically reduced by O_2^- to **16**. The latter is then reduced by O_2^- to $\text{R}_3\text{Sn}^\bullet$, to allow yet another round of addition.^{2a,d}

The fact that we have been unable to detect the requisite singlet¹⁰ of $g_{\text{av}} = 2.031$ ¹⁰ for the postulated superoxide radical in the EPR spectra of (**Z**)-2 or (**Z**)-9 at -50 °C, and we are able to spectroscopically observe fairly long-lived α -stannylyvinyl radicals, once more calls into question^{1,3,5} the validity of this mechanistic theory,^{2a–d} as does a wide range of other experimental evidence.^{3–5} Clearly the low temperature EPR spectra that we have gathered here for the α -stannylyvinyl radicals (**Z**)-2 and (**Z**)-9, plus Alabugin's^{5a} computational uncovering of a stabilizing $n_{\text{O}} \rightarrow \sigma^*_{\text{Sn-C}}$ interaction of 2.5 kcal mol^{−1} in such radical additions, alongside our own earlier work of Scheme 1,^{3,4} each collectively leave very little room for mechanistic alternatives other than the entirely free radical, O–Sn coordinately-controlled, hydrostannation reaction mechanism of Scheme 3 controlling the outcome of these additions.

Our success in obtaining these EPR spectra for the α -stannylyvinyl radicals (**Z**)-2 and (**Z**)-9 is almost certainly due



Scheme 3 A summary of the primary mechanism of the O-directed free radical hydrostannation of disubstituted alkyl acetylenes.

to both radicals being very strongly stabilized by the powerful C-Sn- σ -bond/SOMO radical hyperconjugative interactions⁵ that can arise. Electronic effects that can involve the C-Sn- σ -bond donating electron density into the partially-filled SOMO of the radical, and the vinyl radical itself delocalising into the empty orbitals associated with the tin. According to Alabugin,⁵ strong hyperconjugative interactions of the former type are typically associated with large stabilization energies of the order of 26–31 kcal mol^{−1}. Our observation of a large ^{117,119}Sn β -hyperfine coupling constant of 9.5 mT (95 Gauss) between the Ph₃Sn unit and the vinylic radical in (Z)-2 seemingly supports^{11,12} the existence of such a computed⁵ stabilizing electronic interaction. It also supports the idea that the stannylvinyl radicals (Z)-2 and (Z)-9 significantly delocalise their unpaired electron into the empty low-lying orbitals associated with the tin. Presumably, the simultaneous manifestation of these two effects leads to the high degree of longevity and persistence¹² that is seen for (Z)-2 and (Z)-9 when they are generated at $−50$ °C; a persistence that diminishes as the radicals are warmed above $−10$ °C, when fast H-atom abstraction occurs from the Ph₃SnH.¹³

How we assigned the bent σ -vinyl radical geometry to (Z)-2 and (Z)-9 was through comparison of their *g* values (2.0014 and 2.0021 respectively) with the *g* values reported for other bent alkyl-bearing σ -vinyl radicals, which are generally below the value of *g* for a free electron (*g* = 2.0023).¹⁴ By way of contrast, linear π -vinyl radicals typically have *g* values that lie well above this number.¹⁴

So, to summarize, we have identified a steady-state set of conditions where rapid stannyl radical addition can proceed

near exclusively at the propargyloxy α -acetylenic carbon, and where the rate of H-atom abstraction by the α -stannylvinyl radical from the stannane is slowed sufficiently to allow these intermediates to be spectroscopically observed by EPR.

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Conflicts of interest

There are no conflicts to declare.

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