Synthesis and Properties of the 5,10,15-Trimesityltruxen-5-yl Radical

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Synthesis and Properties of the 5,10,15-Trimesityltruxen-5-yl Radical

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ABSTRACT

The synthesis of a long-lived, truxene-based radical that is highly delocalized and exhibits a narrow EPR absorption is reported. The radical is stable for multiple hours in a solution exposed to air and remains for months in the solid state under inert gas. Characterization and properties are discussed.

Dynamic nuclear polarization (DNP) is a promising technique to dramatically increase sensitivity in NMR experiments. In a contemporary DNP experiment, the large polarization of a stable radical’s unpaired electron is transferred to proximal nuclei via μ-wave electron–nuclear transitions. Enhancements in nuclear polarization could theoretically reach values approaching the quotient of the gyromagnetic ratios of an electron and the nucleus of interest (γ_e/γ_{1H} ≈ 660; γ_e/γ_{13C} ≈ 2600).

Radicals with narrow EPR absorptions efficiently participate in the solid effect (SE)-DNP mechanism. At present, derivatives of triphenylmethyl (trityl) and 1,3-bisdiphenylene-2-phenylallyl (BDPA) are the only readily available radicals whose EPR absorptions are sufficiently narrow to support SE-DNP. Mixtures of narrow-line radicals whose EPR signals are separated by the Larmor frequency of a nucleus are expected to induce remarkable polarization of that nucleus via the cross effect (CE)-DNP mechanism. CE is a two electron–one nucleus spin-flip process that tends to affect DNP more efficiently than SE at high magnetic fields.

recently demonstrated DNP with a mixture of these radicals achieved \(^{13}\)C-NMR signal enhancements of over 600.\(^6\) The success of C-centered, delocalized radicals in promoting DNP through both SE and CE mechanisms served as motivation for investigating new structures for the generation of novel, narrow-line radicals.

We targeted 5,10,15-triaryltribenzo[a,f,k]trinden-5-yl (1) as a potentially stable radical system. This structure could feature highly delocalized spin density shared primarily by quaternary C-atoms, a condition that could offer narrow EPR linewidths. Structure 1 contains an indeno[1,2-a]fluorene (2) unit, fully \(\pi\)-delocalized derivatives of which have yet to be realized synthetically. A flurry of recent work by Haley, Tobe, and coworkers have yielded stable derivatives of the [2,1-b],\(^9\) [1,2-b],\(^10\) [2,1-a],\(^11\) and [2,1-c]\(^12\) isomers, each of which have fascinating electronic and optical properties.

Truxene-5,10,15-trione (3), envisioned as a synthetic intermediate to 1, was prepared by aldol trimerization\(^13\) of 1,3-indanenedione following a literature procedure\(^14\) (Scheme 1). Addition of mesityllithium to trione 3 afforded triol 4, which was isolated entirely as the syn, \(C_3\)-symmetric stereoisomer. Though it is unclear if the anti diastereomer is formed and removed in purification, it is noteworthy that Eschevarren and coworkers also observe the \(C_3\) tris-adduct exclusively after addition of 9-fluorenlylithium to 3.\(^15\) In an attempt to form either radical 1 or a tribenzotriindene derivative that could be synthetically transformed to 1, triol 4 was treated with SnCl\(_2\) and HCl in CH\(_2\)Cl\(_2\), conditions for reductive aromatization reactions.\(^16\) The result was the formation of a bright green material that could be purified by silica column chromatography.

Solution-phase EPR revealed that the green material is a radical with a narrow absorbance and no clear hyperfine couplings.\(^17\) High resolution ESI-MS (positive mode) spectra of the radical showed a cation with the formula C\(_{54}\)H\(_{72}\)+, corresponding to the ionized form of 5,10,15-trimesityltruxen-5-yl (5). The radical’s IR spectrum shows no discernible O–H vibrations, which are clearly observable in the IR spectrum of triol 4, suggesting that no alcohol groups remain in the radical. This conclusion is corroborated by the radical’s high \(R_f\) value (0.83) on a silica TLC plate with 1:1 CH\(_2\)Cl\(_2\)—hexane as the eluent. Triol 4, like many organic alcohols, remains on the baseline of a TLC plate eluted with this solvent system.
The acidic reduction of 4 must introduce two C–H bonds at the 10- and 15-positions for 5 to be the correct structure of the radical. Accordingly, substituting DCI/D2O for HCl/H2O should yield doubly deuterated radical 5-D,D. This reaction was carried out and ESI-MS (positive mode) analysis of the products showed peaks corresponding to doubly and singly deuterated derivatives of 5 in a 74:26 (D,D to H,D) ratio, indicating 87% total deuteration of the 10- and 15-positions (Figure 1).18 The prevalence of 5-D,D and the absence of species with more than two deuteriums confirms that two C–H/D bonds are formed in the reaction and strongly suggests that 5 is the structure of the non-deuterated radical. The compound may exist as syn and anti diastereomers, depending on the relative configurations of the newly introduced H-atoms.

We propose that the mechanism of the reaction from 4 to 5 may involve a series of cation formation/reduction steps through short-lived indeno[1,2-a]fluorene-containing intermediates, including radical 1 (Scheme 2). These quinoidal species are likely basic19 and may be protonated by HCl to form cations that are stabilized by conjugation to localized benzene rings. The cations then undergo single-electron reduction to radical species and the process repeats until 5 is formed. This mechanistic proposal accounts for yields above 50% for radical 5, whereas a sequence of cation formation–reduction–protonation steps at each methanetriyl group would be expected to afford the diamagnetic trihydride as the major product.

DFT calculations (6-31G*) of radical 5 reveal that spin density is delocalized throughout the truxene core (Figure 2). A large proportion of the radical’s spin density (31%) is located at C(5) and significant spin is shared by alternating sp3-hybridized C atoms extending from C(5). The aromatic ring of the mesityl group lies roughly perpendicular to the truxene system and bears only a small amount of spin density (0.9%). The structure shown is the anti diastereomer, which was found to be less than 0.3 kcal/mol lower in energy than the syn diastereomer. Spin density distribution is almost identical in both configurations (see SI for additional information).

The high-field (140 GHz) EPR spectrum of radical 5 exhibits a narrow absorbance at roughly the same frequency as SAHBDPA (Figure 3). The linewidth at half-height of the radical’s maximum 140 GHz EPR absorption is 1.368 mT, broader than that of SA-BDPA (1.028 mT) and narrower than that of trityl OX063 (1.770 mT). The electron T1 relaxation time of 5, measured through a saturation recovery experiment at 4994.75 mT, was determined to be 7.3 ms (see Figure S7 in SI), an intermediate value compared to BDPA (55.9 ms) and Trityl OX063 (1.28 ms).8
Figure 3. 140 GHz EPR spectrum of radical 5 (bold black, 1 mM in toluene, 80 K), compared to trityl OX063 (red, 1 mM in 60/40 (v/v) glycerol/D$_2$O, 80 K) and SA-BDPA (green, 1 mM in 60:40 (v/v) glycerol/D$_2$O, 80 K). Peak heights are scaled for equal intensity. The SA-BDPA and trityl data shown here are from ref. 8 (see comment in SI).

Figure 4. (A) Cyclic voltammogram of 5 (under N$_2$ in glove box, 0.1 M Bu$_4$NPF$_6$ in CH$_2$Cl$_2$, Pt button (WE), Pt wire (CE), Ag/AgNO$_3$ (RE), scan rate 0.05 V/s. (B) Absorbance spectrum of 5 in CH$_2$Cl$_2$, grey spectrum shows higher wavelength absorption multiplied by a factor of 10.

A kinetic experiment provided insight into the stability of radical 5 in a solution exposed to ambient oxygen. A solution of 5 in CH$_2$Cl$_2$ (4 × 10$^{-5}$ mol/L) was exposed to air and absorption spectra were measured every several hours until only a negligible amount of 5 remained (Figure 5A). Before each measurement, CH$_2$Cl$_2$ was added to a mark on the cuvette to replace the solvent lost by evaporation and to keep the total concentration of the solute roughly consistent. The value for absorbance (Abs) at the peak at 403 nm was used to monitor the disappearance of 5 over time; however, the value does not reach zero, as the oxidized product(s) continue to absorb at this wavelength. The value $Abs^*$, which is equal to $Abs - 0.080$ (0.080 being the extrapolated value for Abs after complete oxidation of 5), clearly follows pseudo-first-order exponential decay kinetics. It is assumed that the rate of diffusion of oxygen into the solution is much faster than the rate of reaction between oxygen and the radical. From the kinetic plots, we determine $k_{obs}$ of the reaction to be roughly ~0.12 h$^{-1}$ and the half-life ($t_{1/2}$) of the radical under these conditions to be roughly ~5.8 h. The ESI-TOF-MS spectrum of the oxidized material after 26.5 h showed a major peak at 711 m/z, which corresponds to [5$^+$O]$^+$ and confirms that oxygen is incorporated into the structure. It is noteworthy that 5 remains for months when properly stored under inert conditions in the solid state.
Figure 5. (A) Absorbance spectra of 5 in CH$_2$Cl$_2$ after exposure to air over specified times. (B) Plots of Abs* (see text) (black circles) and $-\ln$(Abs*/Abs$_0$*) (green squares) over time, indicating that $k_{obs} \approx 1.2$ h and $t_{1/2} \approx 5.8$ h.

In conclusion, we have synthesized a long-lived, truxene-based radical that exhibits an EPR resonance that should be sufficiently narrow to enable DNP by the SE mechanism. Current work is focused on developing derivatives that are both stable in the presence of oxygen and soluble in water/glycerol mixtures amenable to DNP experiments.

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Supporting Information Available. Experimental procedures, full characterization and spectral data for new compounds, Cartesian coordinates for calculated structures. This material is available free of charge via the Internet at http://pubs.acs.org.