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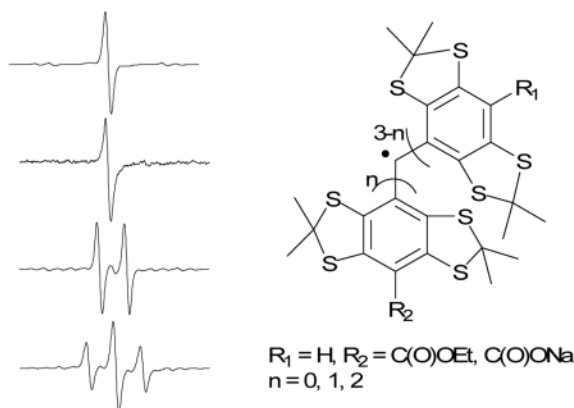
## Reactivity of Molecular Oxygen with Ethoxycarbonyl Derivatives of Tetrathiatriarylmethyl Radicals

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### Abstract



Tetrathiatriarylmethyl (TAM) radicals are commonly used as oximetry probes for electron paramagnetic resonance imaging (EPRI) applications. In this study, the electronic properties and the thermodynamic preferences for O<sub>2</sub> addition to various TAM-type triarylmethyl (trityl) radicals were theoretically investigated. The radicals' stability in the presence of O<sub>2</sub> and biological milieu were also experimentally assessed using EPR spectroscopy. Results show that H substitution on the aromatic ring affects the trityl radical's stability (tricarboxylate salt **1-CO<sub>2</sub>Na** > triester **1-CO<sub>2</sub>Et** > diester **2-CO<sub>2</sub>Et** > monoester **3-CO<sub>2</sub>Et**) and may lead to substitution reactions in cellular systems. We propose that this degradation process involves an arylperoxyl radical which can further decompose to alcohol or quinone products. This study demonstrates how computational chemistry can be used as a tool to rationalize radical stability in the redox environment of biological systems and aid in the future design of more biostable trityl radicals.

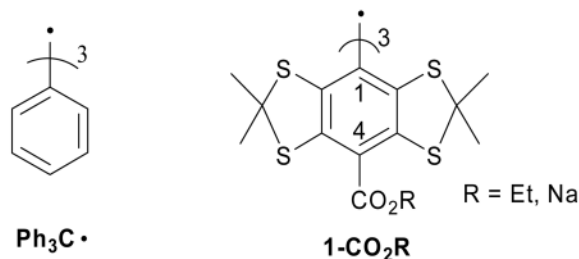
### 1. Introduction

Evaluating the concentration of O<sub>2</sub> as well as O<sub>2</sub>-derived reactive species, including that of superoxide radical anion (O<sub>2</sub><sup>•−</sup>) in *in vivo* and *in vitro* systems, is of utmost importance in the study of numerous pathophysiological processes.<sup>1–4</sup> Over the past decade, major efforts have been made towards the development of electron paramagnetic resonance imaging (EPRI)<sup>5–7</sup>

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of biological samples to provide improved image resolution and quality, and this method has evolved to become an important tool for imaging free radicals in organs<sup>4,8,9</sup> and the entire body of small animals,<sup>10-12</sup> tumors and normal tissues.<sup>13-17</sup>

The development of paramagnetic materials, such as nitroxides,<sup>18</sup> triarylmethyl (trityl) radicals,<sup>19,20</sup> and recently, spin-labeled dendrimers,<sup>21</sup> as probes for EPRI applications is critical for the precise determination of O<sub>2</sub> concentration and detection of superoxide radical anion (O<sub>2</sub><sup>•-</sup>) in biological samples, including cells and tissues.<sup>22-25</sup> Trityl radicals have been the popular choice for EPRI applications due to their stability at physiological pH and long relaxation times which consequently give rise to narrow EPR linewidths. Moreover, trityl radicals offer some advantages over nitroxides, i.e., high analytical resolution at μM concentrations, stability in cells and tissues, and the flexibility to independently determine the concentrations of superoxide radical anion (O<sub>2</sub><sup>•-</sup>) and O<sub>2</sub> by EPR signal loss and line broadening, respectively.<sup>25</sup>

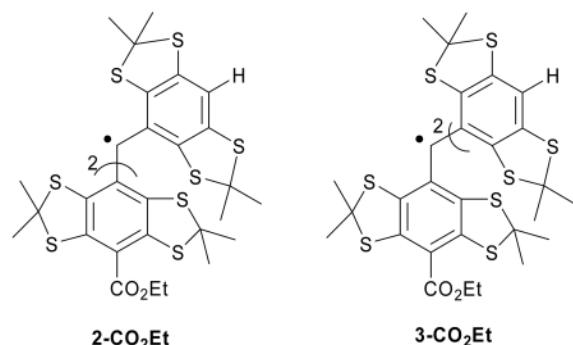


Trityl radicals have a long history in radical chemistry, ever since the initial report by Gomberg in 1900 of triphenylmethyl radical (Ph<sub>3</sub>C<sup>•</sup>).<sup>26</sup> Tetrathiatriarylmethyl (TAM) radicals **1-CO<sub>2</sub>R** belong to a family of trityl radicals which exhibit a very narrow linewidth (<100 mG),<sup>27</sup> even in the presence of biological matrix, rendering this probe ideal for *in vitro* and *in vivo* EPR imaging applications.<sup>23</sup> The perdeuterated analogue of **1-CO<sub>2</sub>Na** has also found applications in magnetic resonance imaging (MRI)<sup>28</sup> and Overhauser-enhanced magnetic resonance imaging (OMRI).<sup>29</sup>

Reddy et al. reported an elegant synthesis of **1-CO<sub>2</sub>Et** and the corresponding salt, **1-CO<sub>2</sub>Na**, and these authors also evaluated the sensitivities of these species to O<sub>2</sub>.<sup>27</sup> However, the underlying mechanism of decay for **1-CO<sub>2</sub>Na** in the presence of O<sub>2</sub> was not reported. In a solid lattice, the reaction of the triphenylmethyl (Ph<sub>3</sub>C<sup>•</sup>) radical with O<sub>2</sub> results in the formation of an equilibrium product, triphenylmethylperoxyl radical, Ph<sub>3</sub>COO<sup>•</sup> (eq. 1).<sup>30,31</sup>



In dichloromethane solution, a second order reaction was observed with Ph<sub>3</sub>C<sup>•</sup> and O<sub>2</sub>, finally yielding benzophenone (Ph<sub>2</sub>C=O) as a major product.<sup>32</sup>



In this paper, we explored the electronic and thermodynamic properties of TAM-type radicals with varying degrees of substitution (**2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et**) and evaluated how these properties can affect radical stability in the presence of O<sub>2</sub> and components of the biological milieu. We complement these experimental efforts with computational chemistry so as to understand and rationalize the radicals' stability.

## II. Results and Discussion

### II. A. EPR Spectroscopic Characterization

Tri- (**1-CO<sub>2</sub>Et**), di- (**2-CO<sub>2</sub>Et**) and mono- (**3-CO<sub>2</sub>Et**) ethoxycarbonyl derivatives of tetrathiatrimethyl (TAM) radical as well as its tri-sodium carboxylate salt (**1-CO<sub>2</sub>Na**) derivative were synthesized and characterized by minor modifications of the method reported by Reddy et al.<sup>27</sup> Figure 1 shows the X-band EPR spectrum for each of the trityl radical species with *g* values that range from 2.0041 to 2.0043, close to that reported for **1-CO<sub>2</sub>Et** and **1-CO<sub>2</sub>Na** of *g* ~ 2.003.<sup>27</sup> The singlet peaks in Figures 1a and 1b are consistent with a tri-substituted trityl radical, while the doublet and triplet spectral features for **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et**, respectively, indicate the presence of a hydrogen hyperfine splitting due to the unsubstituted aromatic ring. The observed hyperfine splitting constant for both **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** trityl radicals of *a*<sub>H,para</sub> = 2.30 G in DMSO is close to our predicted isotropic hyperfine splitting of 2.11 or 2.22 G at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* level of theory in the gas phase, and also similar to that reported experimentally for Ph<sub>3</sub>C· of *a*<sub>H,para</sub> = 2.745 G in toluene solution.<sup>33</sup>

Anaerobic peak-to-peak widths in DMSO for **1-CO<sub>2</sub>Et**, **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** range from  $\Delta B_{pp}$  = 380 to 470 mG and are broader compared to the  $\Delta B_{pp}$  = 111 mG observed for **1-CO<sub>2</sub>Na** in aqueous solution using a modulation amplitude of 0.01 G. Reddy et al.<sup>27</sup> previously reported linewidths for **1-CO<sub>2</sub>Et** and **1-CO<sub>2</sub>Na** of ~270 mG (in pyridine) and 84 mG (in water), respectively, at modulation amplitudes of 0.025-0.086 G. It should be noted, however, that the slight broadening observed in this study may be due to the presence of trace amounts of O<sub>2</sub> since the capillary tubes were only sealed with inert clay after bubbling with Ar. The effect of O<sub>2</sub> on the spectral profile of various trityl radicals was investigated as well. Results show that there is a significant linewidth broadening by about  $\Delta\Delta B_{pp}$  = 0.8-0.9 G for all of the ester-derivatized trityl radicals and about  $\Delta\Delta B_{pp}$  = 0.2 G broadening for **1-CO<sub>2</sub>Na** when O<sub>2</sub> was bubbled through DMSO (or water for **1-CO<sub>2</sub>Na**) solutions. This difference in the  $\Delta\Delta B_{pp}$  between the ester and carboxylated-trityl radicals could be due to the higher solubility of O<sub>2</sub> in DMSO compared to that in water.<sup>34</sup> Purging the O<sub>2</sub>-saturated trityl radical solution with argon gave linewidths similar to that observed originally in the absence of O<sub>2</sub>, thereby demonstrating that the O<sub>2</sub> with trityl radical interaction is reversible.

## II. B. Computational Analysis of Structures, Properties and Decomposition Pathways

The measurement of O<sub>2</sub> in biological systems requires long and continuous monitoring. Hence an investigation of the stability of trityl radicals in the presence of O<sub>2</sub> is critical since O<sub>2</sub> reaction with trityl radicals can lead to the degradation of the radical probe. To gain insight into the stability of these radicals in the presence of O<sub>2</sub>, the electronic as well as the thermodynamic properties of the trityl radical were theoretically investigated. Figure 2 shows the charges and spin ( $\alpha$ - $\beta$ ) populations for various TAM-type compounds as well as for Ph<sub>3</sub>C• using the natural population analysis (NPA) partitioning scheme at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* level. These data show that for TAM-type compounds, there is a 0.62 electron localization at the central carbon which is higher compared to the spin density (population) on the central carbon predicted for Ph<sub>3</sub>C• of 0.56e. However, the spin density of the para-C atoms in TAM-type compounds is in the range of 0.07-0.09e, but higher (0.12e) in Ph<sub>3</sub>C•.

A closer look at the molecular structures of **1-CO<sub>2</sub>Na** and Ph<sub>3</sub>C• showed a significant difference in their respective aromatic ring conformations, i.e., trityl radical **1-CO<sub>2</sub>Na** has a substantial out-of-plane ring twisting compared to that of Ph<sub>3</sub>C• radical.<sup>35</sup> This difference in conformations may have a considerable effect on the electron density distribution and thereby could affect spin relaxation that can result in EPR spectral line broadening.

The preference for O<sub>2</sub> addition to the trityl radical was investigated thermodynamically as to whether the O<sub>2</sub> addition occurs at the central or para-carbon atoms (Scheme 1). Table 1 shows the thermodynamic values for O<sub>2</sub> addition to various trityl radicals to form the peroxy adduct (steps A and B), followed by subsequent H-atom abstraction (steps C and D) from various H-atom donors to form the hydroperoxy adduct at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* level of theory. Results show that, in general, O<sub>2</sub> addition to TAM-type radicals is endothermic (steps A and B in Table 1), and the formation of the peroxy adduct at the aromatic ring C is preferred by 2.5 to 6.5 kcal/mol compared to peroxy adduct formation at the central carbon atom for all of the partially substituted TAM radicals. However, peroxy adduct formation at the central carbon atom is preferred by 1.4 to 2.5 kcal/mol for the fully substituted trityl radicals such as **1-CO<sub>2</sub>Et**, **1-CO<sub>2</sub>H** and **1-CO<sub>2</sub>Na** for the gas-phase calculations. The formation of the peroxy adduct at the central carbon in Ph<sub>3</sub>C• is exothermic by -4.0 kcal/mol, and is preferred by 17.7 kcal/mol relative to O<sub>2</sub> addition to the aromatic ring's carbon atom. Examination of the charge and spin populations for Ph<sub>3</sub>C• and **4-CO<sub>2</sub>Et** shows only a small difference in their respective charge (~0.05e) and spin population (~0.06e) at the central C atom. The spin population on the para C of Ph<sub>3</sub>C•, however, is higher by only 0.03e compared to **4-CO<sub>2</sub>Et** with no difference in their respective atomic charges. Based on the electronic properties predicted for Ph<sub>3</sub>C• and **4-CO<sub>2</sub>Et**, the preference of O<sub>2</sub> addition to the central C of Ph<sub>3</sub>C• appears to be due mainly to a steric effect. Although it has been previously<sup>36,37</sup> shown that O<sub>2</sub> adds preferentially to the carbon centers in delocalized radicals bearing the highest spin density, this behavior does not apply completely for trityl radicals for which steric effects are also important. The effect of solvation on the O<sub>2</sub> addition to **1-CO<sub>2</sub>H** and **1-CO<sub>2</sub>Na** was investigated by employing the polarizable continuum model (PCM) for water as compared to the gas-phase energetics. The reactions for O<sub>2</sub> addition to the aromatic ring shows no significant difference, while O<sub>2</sub> addition reactions to the central carbon atom is preferred by 8~10 kcal/mol in water compared to the gas-phase calculations. We have noted similar solvation effects for peroxy radicals earlier.<sup>38</sup>

Table 1 also shows that the energetics for H-atom abstraction by the peroxy radical adducts from H<sub>2</sub>O, DMSO or MeSH to form the aryl hydroperoxy adduct (steps C and D). Although H-atom abstraction is an endothermic process, abstraction is more preferred with DMSO as an H-atom donor compared to that with H<sub>2</sub>O, and most preferred with MeSH. The overall energetics (steps A + C and B + D) for the formation of hydroperoxy adducts in the various solvents is highly endothermic. For all of the trityl radicals, the formation of the aromatic hydroperoxy adduct (step B + D) is more preferred than the formation of the central carbon

hydroperoxyl adduct (A + C) in the gas phase. The partially substituted trityl radicals are more favored to form aryl hydroperoxyl adducts than the fully substituted trityl radicals. This indicates that the stability of these radicals appears to be correlated with the number of H's on the aryl ring. However, based on the computational results, the **1-CO<sub>2</sub>Na** trityl salt is more preferred to form aryl hydroperoxyl adducts than the fully ester-substituted trityl radicals **1-CO<sub>2</sub>Et**. This is not consistent with the stability study results noted below. Our calculations, of course, are thermodynamic results and can not be used directly to represent kinetic results. Because of the computational expense, searching for the transition state (TS) of the reactions was not practical. Also, transition states for O<sub>2</sub> addition to carbon-center radicals are often difficult to locate and the corresponding wave functions usually suffer from significant spin contamination, thereby rendering the results to be suspect.<sup>38b</sup> Based on Hammond's postulate, the TS should be late on the potential energy surface, and should resemble the product, peroxy radical adduct, or hydroperoxyl adduct. We evaluated the reactions by the energy difference between product and starting material, not the activation barrier which would actually control the reaction process. A possible explanation of the inconsistency between computational and experimental results is that for the ester-substituted trityl radicals, their transition states have a similar correlation with the products and the thermodynamic energies are consistent with their kinetic values, while for the trityl salt, the transition state may have a different correlation with its product. These thermodynamic results are not exclusive, and other pathways for decomposition may be possible.

## II. C. Experimental Stability of Trityl Radicals in Solution: EPR, MS and Product Studies

The stability in solution of various trityl radicals was investigated. Results show that each argon-purged solution of the esterified trityls (**1-CO<sub>2</sub>Et**, **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et**) gave a half-life of ~4 days, while the carboxylate salt, **1-CO<sub>2</sub>Na** did not decay for at least 17 days. This slow decomposition of the esterified trityls could be accounted for by the slow diffusion of O<sub>2</sub> into the solution. However, in the presence of air, the half-life of **3-CO<sub>2</sub>Et** was significantly shorter (~1 day), while the half-lives of **2-CO<sub>2</sub>Et**, **1-CO<sub>2</sub>Et** and **1-CO<sub>2</sub>Na** were the same as in the absence of O<sub>2</sub>. Product analysis of each of the trityl radicals' solutions using electrospray ionization (ESI) and matrix-assisted laser desorption ionization-time of flight (MALDI-ToF) mass spectrometry, before and after exposure to O<sub>2</sub>, was carried out (Figure 3). These analyses reveal that the same spectral pattern was observed for trityl solutions with ESI before and after exposure to O<sub>2</sub> (Table 2). It should be noted, however, that ESI is an atmospheric pressure method, is an oxidizing environment and may lead to the formation of peroxy adducts. In order to differentiate whether the observed mass spectra were due to O<sub>2</sub> addition in the gas or liquid phase, trityl radical solutions were bubbled with O-18 labeled O<sub>2</sub> (<sup>18</sup>O<sub>2</sub>) and the MS were acquired. Unfortunately, the peak for the <sup>18</sup>O-labelled product was too small to be observed and could be due to the reversibility of O-16 O<sub>2</sub> addition to these trityl radicals.

MALDI-ToF spectra for the carboxylate salt radical **1-CO<sub>2</sub>Na** show only a very small peak corresponding to the OH adduct (Figure 3c), while OOH and OH adducts were produced from **1-CO<sub>2</sub>Et**, **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** (Figure 3a-b and Table 2). This result is consistent with the EPR observation that in the presence of O<sub>2</sub>, **1-CO<sub>2</sub>Na** exhibited the greatest stability compared to the three ester derivatives. The stability of these radicals, **1-CO<sub>2</sub>Na** > **1-CO<sub>2</sub>Et** > **2-CO<sub>2</sub>Et** > **3-CO<sub>2</sub>Et**, appears to be correlated with the number of H's on the aryl ring, as we predicted by the energetic calculations in the gas phase. Scheme 2 suggests an explanation consistent with this correlation, and shows the MS fragmentation pattern and the possible origin of the OH adduct as well as the quinone analogue. The scheme shows that the OH adduct could originate from the O-O homolytic cleavage of the peroxy moiety. The formation of an (M+17) ion is evident in analysis of all of the ester trityl radicals **1-CO<sub>2</sub>Et**, **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et**, while a quinone-like product (M+15) was observed only for **3-CO<sub>2</sub>Et**. While **2-CO<sub>2</sub>Et** has one aromatic ring with a H, a species with a mass of (M+15) was not observed and is probably



due to the smaller number of unsubstituted aromatic rings in **2-CO<sub>2</sub>Et** compared to that in **3-CO<sub>2</sub>Et**.

We then performed a product analysis for the decomposed **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** samples, and for which both samples possessed a lack of EPR signals. The products were separated by preparative HPLC, and the major products were analyzed by UV-vis, NMR, and high-resolution MS. Based on the MS results for both purified trityl reaction mixtures, quinone-like products (M+15) and trityl cation or anion (M) species are the major products. Even though these two products can be separated completely by HPLC because of their different polarity, trace quinone-like compound could always be detected from the M residue for both trityl compounds (Figure 4). Based on the MS pattern for the quinone-like compound (M+15) in the M residue, trace (M+16) and (M+17) ions can also be seen from the M residue for both decomposed samples. Furthermore, the relative intensities of (M+16) and (M+17) ions to (M+15) are higher in the M residue of **2-CO<sub>2</sub>Et** than in that of **3-CO<sub>2</sub>Et**. This also supports our previous assumption that the more unsubstituted aromatic rings, as in **3-CO<sub>2</sub>Et** vs **2-CO<sub>2</sub>Et**, will form more quinone-like (M+15) products. The MS results also indicate that the proposed radical pathway mechanism (Scheme 2a) may be a route for the decomposition of the trityl radical. There may also be a redox pathway in which trityl radicals lose their EPR signal by being oxidized or reduced to the cation or anion, respectively (Scheme 2b). (However, we should note that all attempts to obtain negative ion ESI-MS data were unsuccessful, and the MS results are most consistent with the formation of cationic species.) The formed trityl cation (or anion) could then form other products by reactions with nucleophiles (or electrophiles) and/or O<sub>2</sub>. UV-vis spectra of the quinone-like products have  $\lambda_{\text{max}}$  peaks which are shifted to the red to 515 nm compared with their corresponding radicals (493 nm for **2-CO<sub>2</sub>Et** and 481 nm for **3-CO<sub>2</sub>Et**). This red shift is possibly due to the enhanced conjugation between the aromatic rings for the quinone-like products. The trityl cation or anion products, however, show blue shifts of their UV-vis spectra to 486 nm for **2-CO<sub>2</sub>Et** and 473 nm for **3-CO<sub>2</sub>Et**, respectively. NMR analysis was less conclusive as only <sup>1</sup>H NMR could be obtained (see supporting information).

## II. D. Stability of Trityl Radicals in Intracellular Studies

So far, our studies show that the **3-CO<sub>2</sub>Et** trityl radical is the least stable of all of the trityl radicals investigated in the presence of O<sub>2</sub> alone. To test if these theoretical and experimental studies can be used to assess biostability of the trityls, their intracellular stability in rat H9C2 cardiomyocytes was investigated. The ability of these compounds to penetrate into cells as well as their overall biostability were assessed for the four trityl radicals by incubating the compounds in the presence of rat H9C2 cardiomyocytes at various time intervals and subsequent washing of the cells to remove extracellular trityl radicals. After washing, the cells became colored (i.e., green or orange which are the original colors of the trityl radicals) indicating that there is absorption of these compounds inside the cell. EPR data reveal that among all of the trityl radicals used in the study, only the tri-substituted radicals, i.e., **1-CO<sub>2</sub>Na** and **1-CO<sub>2</sub>Et**, exhibited intracellular stability (Figure 5). In spite of the hydrophobic nature of the cell membrane, high permeability to cells was observed for the more polar **1-CO<sub>2</sub>Na** and this may be due to the nature of the carrier solvent used, i.e., 1  $\mu$ L DMSO per mL medium. The **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** trityl radicals, which have at least one H on the aryl-rings, showed only negligible EPR peak intensity after 1 hour of incubation with cells and suggests that these trityl radicals are susceptible to biodegradation in cellular systems.

## II. E. Computational and CV Studies of the Oxidation/Reduction of the Trityl Radicals

In order to gain insights into the nature of the degradation process, the susceptibility of the trityl radicals to oxidation or reduction was investigated by calculating their respective electron affinities (EA) and ionization potentials (IP) as shown in Table 3. The partially substituted

TAM-radicals, such as **2-CO<sub>2</sub>Et**, **3-CO<sub>2</sub>Et** and **4-CO<sub>2</sub>Et**, gave lower EA and IP values compared to those of the fully substituted TAM-radicals (with the exception of the carboxylate salt, **1-CO<sub>2</sub>Na**) indicating that the partially substituted TAM-radicals are harder to reduce and easier to oxidize compared to the fully substituted ones. The IP for the simplest trityl radical Ph<sub>3</sub>C• is intermediate, while the EA is lowest compared to the TAM-type radicals, indicating that the fully unsubstituted trityl radicals are the most difficult to be reduced. We therefore suggest that the loss of EPR signal in the decomposition of the trityl radicals leads to two processes: (1) formation of the trityl cation, which subsequently reacts with nucleophiles, and (2) O<sub>2</sub>-addition to the radical center, which subsequently forms the corresponding quinone product.

Cyclic voltammetry (CV) was performed to obtain the redox potentials for the carboxylate salt **1-CO<sub>2</sub>Na** and triester **1-CO<sub>2</sub>Et** (Table 4). The experimental results showed that the carboxylate salt is easier to oxidize and to reduce than the triester. This is consistent with the computational results using the PCM calculations for the carboxylate salt and with the gas-phase calculations for the triester. Because the trityl radical salt (**1-CO<sub>2</sub>Na**) is more stable than the triester (**1-CO<sub>2</sub>Et**) in organic and biological solutions and **1-CO<sub>2</sub>Na** is more easily oxidized and reduced, the decomposition of the trityl radical is more likely to be a radical process (Scheme 2a). It is possible that the redox reaction proceeds via an equilibrium; recall that the oxidation/reduction of **1-CO<sub>2</sub>Na** is reversible, but is not reversible for **1-CO<sub>2</sub>Et** (Table 4). Also, from the CV data (Table 4), the carboxylate salt **1-CO<sub>2</sub>Na** is easier to oxidize than the **1-CO<sub>2</sub>Et** triester. Therefore, we conjecture that the rate-determining step for decomposition involves the corresponding trityl radical for the **1-CO<sub>2</sub>Na** species, proceeding via peroxy radical formation, but the rate-determining step for the ester radicals, such as **1-CO<sub>2</sub>Et**, may involve the radical, or irreversibly formed cation. Such a scenario would be consistent with the stability of the carboxylate salt **1-CO<sub>2</sub>Na** being greater than the corresponding esters (**1-CO<sub>2</sub>Et**) as well as the greater propensity to oxidize and to reduce the irreversibly formed cation. Such a scenario would be consistent with the stability of the carboxylate salt **1-CO<sub>2</sub>Na**. However, which pathway, Scheme 2a (radical) or 2b (ionic), is the main cause of the decomposition of the trityl radicals is not fully clarified.

## II. F. Stability of Trityl Radicals: Bond Dissociation Energies

If the trityl radicals were decomposing by radical processes, it is important to understand the stability of the carbon-centered radical in this homologous series. A simple hypothesis would be that the more stable trityl radical (Ar<sub>3</sub>C•) would be derived from the parent aryl methane (Ar<sub>3</sub>C-H) derivative that had the lowest C-H bond dissociation energy (BDE). To assess the relative stability of various TAM radicals, the C<sub>central</sub>-H BDE was calculated for the compounds under study (Table 3). The calculations reveal that there is less than 1.5 kcal/mol difference in the BDE's for generating these diverse radicals, and indicates that the unpaired electron on all of the TAM radicals is stabilized, as supported by the spin density (population) results shown in Figure 2. Ph<sub>3</sub>C-H, however, gave the highest BDE value of 80.4 kcal/mol compared to the TAM-type radicals of 77.2-78.1 kcal/mol, thereby indicating that substitution of the aromatic ring has a stabilizing effect on the radical formation and disfavors protonation ( $\Delta E_{\text{acid}}$ ) of their respective carbanion analogues (Table 3). The partially substituted TAM-radicals, such as **2-CO<sub>2</sub>Et**, **3-CO<sub>2</sub>Et** and **4-CO<sub>2</sub>Et**, gave higher BDE and  $\Delta E_{\text{acid}}$  values compared to those of the fully substituted TAM-radicals (with the exception of the carboxylate salt, **1-CO<sub>2</sub>Na**) indicating that the H on the aromatic ring does not stabilize the trityl radical center and may contribute to the instability of the trityl radical.



## IV. Conclusions

Over the years, the use of trityl radicals as probes for electron paramagnetic resonance imaging (EPRI) applications has become increasingly important. An understanding of the degradation pathways of trityl radicals is critical in the design of probes with improved stability. Electronic properties as well as the thermodynamics of O<sub>2</sub> addition to various TAM-type radicals were theoretically investigated. Results show that the presence of hydrogen as a substituent on the aromatic ring affects the stability of the trityl radical by making the intermediate to be more susceptible to O<sub>2</sub> addition at the aryl carbon bearing the H. Addition of O<sub>2</sub> to the aromatic ring is more favored in TAM-type trityl radicals, while addition to the central carbon atom is more preferred in Ph<sub>3</sub>C<sup>•</sup> because of a steric effect. Mass spectrometric analyses reveal the formation of a quinone-type product in partially substituted trityl radicals in the presence of O<sub>2</sub>. The formation of the quinone intermediate after reaction with O<sub>2</sub>, as observed for **3-CO<sub>2</sub>Et**, could be a potential precursor to the formation of glutathione or Michael adducts with other nucleophiles that are present in cells as demonstrated by previous studies<sup>39-43</sup> which showed the addition of certain thiols, such as glutathione, to quinones. Redox chemistry is also evident and suggests the formation of the trityl cation. Cell permeability studies show that fully substituted trityl radicals are biostable compared to the partially substituted ones. Predicted electron affinities and ionization potentials indicate that for TAM-type compounds, the less that the trityl radical is substituted, the easier it is to be oxidized. This study demonstrates how computational chemistry can be used as a tool to assess radical stability in complex systems and aid in the future design of more biostable trityl adducts.

## Experimental Section

Although the preparations of **1-CO<sub>2</sub>Na** and **1-CO<sub>2</sub>Et** have been described in the literature,<sup>27</sup> slight modifications in the reaction conditions were utilized.

**Tris(8-ethoxycarbonyl-2,2,6,6-tetramethylbenzo[1,2-d;4,5-d'] bis[1,3]dithiol-4-yl)methanol (8), bis(8-ethoxycarbonyl)tris-(2,2,6,6-tetramethylbenzo[1,2-d;4,5-d']bis[1,3]dithiol-4-yl)methanol (9), 8-ethoxycarbonyl-tris(2,2,6,6-tetramethylbenzo [1,2-d;4,5-d']bis[1,3]dithiol-4-yl)methanol (10)**

To a stirred solution of TMEDA (250  $\mu$ L, 1.66 mmol) in benzene (1 mL) was added 2.5 M n-BuLi in pentane (0.67 mL, 1.67 mmol) dropwise at 0 °C. After being stirred for 30 min, a solution of **7** (193 mg, 0.2 mmol) in benzene (1 mL) was added to the above solution dropwise at 0 °C. After being stirred at 35-45 °C for 45 min, the resulting solution was added to a solution of diethyl carbonate (1.04 mL, 8.62 mmol) in benzene (1 mL) that was maintained at 0 °C. After the mixture was stirred at 45 °C for 2 h, saturated aqueous KH<sub>2</sub>PO<sub>4</sub> was added. The organic layer was separated, dried over MgSO<sub>4</sub>, and concentrated in vacuo. The red residue was crystallized from MeCN (4 mL) to give 90 mg of a crude orange solid. Flash chromatography on silica gel, using diethyl ether/hexanes (1:20) and then gradually increasing to ethyl acetate/hexanes (1:5) as eluant, provided: 45 mg (25%) of **8** as orange crystals that contained a small amount of inseparable impurities: mp > 280 °C (gradually turned black, dec); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  1.45 (t, 9H), 1.65 (s, 18H), 1.73 (s, 9H), 1.76 (s, 9H), 4.43 (m, 6H), 6.76 (s, 1H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  14.3, 28.6, 29.2, 31.9, 33.8, 60.85, 60.93, 62.3, 84.3, 121.3, 134.0, 139.3, 140.3, 141.4, 141.8, 166.2; IR (CHCl<sub>3</sub>) 3335, 2970, 1699, 1239, 1217, 1018, 728 cm<sup>-1</sup>; MS (ESI, (M+Na)<sup>+</sup>, m/z): 1123.03005 (observed), 1123.025363 (exact).

25 mg (15%) of **9** as orange crystals that contained a small amount of inseparable impurities: mp > 265 °C (gradually turned black, dec); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  1.45 (td, 6H), 1.63-1.82 (m, 36H), 4.42 (m, 4H), 6.68 (s, 1H), 7.18 (s, 1H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  14.2, 14.3, 27.5, 27.7, 29.3, 29.6, 30.1, 30.8, 31.6, 33.0, 34.0, 34.6, 34.9, 60.71, 60.74, 61.2 (d), 61.8, 62.3, 63.2, 63.8, 84.1, 118.5, 121.24, 121.26, 130.6, 134.6, 134.7, 137.0, 137.7, 138.0,

139.1, 139.2, 139.6, 140.2, 140.7, 141.4, 141.7, 141.86, 141.89, 166.22, 166.24; MS (ESI, (M+Na)<sup>+</sup>, m/z): 1051.00782 (observed), 1051.004233 (exact).

15 mg (9%) of **10** as orange crystals that contained a small amount of inseparable impurities: mp > 255 °C (gradually turned black, dec); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.45 (t, 3H), 1.62-1.82 (m, 36H), 4.43 (m, 2H), 6.55 (s, 1H), 7.17 (t, 2H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ 14.3, 27.3, 27.9, 28.3, 29.1, 29.7, 31.6, 33.0, 33.3, 33.6, 34.3, 34.6, 35.0, 60.7, 62.34, 62.39, 63.46, 63.49, 63.54, 63.63, 83.9, 118.3, 118.4, 121.3, 130.7, 131.7, 135.6, 136.9, 137.2, 137.5, 137.6, 137.9, 139.04, 139.06, 139.20, 139.28, 140.8, 141.8, 142.0, 166.3; MS (ESI, (M+Na)<sup>+</sup>, m/z): 978.98481 (observed), 978.983103 (exact).

**Tris(8-ethoxycarbonyl-2,2,6,6-tetramethylbenzo[1,2-d;4,5-d'] bis[1,3]dithiol-4-yl)methyl ester (1-CO<sub>2</sub>Et), bis(8-ethoxycarbonyl)tris (2,2,6,6-tetramethylbenzo[1,2-d;4,5-d']bis[1,3]dithiol-4-yl)- methyl ester (2-CO<sub>2</sub>Et), 8-ethoxycarbonyl-tris(2,2,6,6-tetramethyl benzo[1,2-d;4,5-d']bis[1,3]dithiol-4-yl) methyl ester (3-CO<sub>2</sub>Et), Tris- (8-carboxyl-2,2,6,6-tetramethyl benzo[1,2-d;4,5-d']bis[1,3] dithiol -4-yl) methyl Sodium Salt (1-CO<sub>2</sub>Na)**

To a stirred solution of **8** (11 mg, 0.01 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL) was added BF<sub>3</sub>·Et<sub>2</sub>O (10 μL, 0.08 mmol) dropwise at room temperature. After being stirred for 1 h, the resulting dark green-blue solution was treated with a solution of SnCl<sub>2</sub> (3.2 mg, 0.017 mmol) in THF (0.4 mL). After 10 min, saturated aqueous KH<sub>2</sub>PO<sub>4</sub> was added. The organic layer was separated, dried over Na<sub>2</sub>SO<sub>4</sub>, and concentrated in vacuo to give 10 mg of the crude trityl radical **1-CO<sub>2</sub>Et** as an orange-brown solid: IR 2960, 2924, 2855, 1704, 1490, 1452, 1367, 1234, 1110, 1044 cm<sup>-1</sup>; UV 249, 411, 495 nm; MS (ESI, (M+Na)<sup>+</sup>, m/z): :1106.02081 (observed), 1106.022623 (exact).

To a stirred solution of **9** (10 mg, 0.01 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL) was added BF<sub>3</sub>·Et<sub>2</sub>O (10 μL, 0.08 mmol) dropwise at room temperature. After being stirred for 1 h, the resulting dark green-blue solution was treated with a solution of SnCl<sub>2</sub> (3.2 mg, 0.017 mmol) in THF (0.4 mL). After 10 min, saturated aqueous KH<sub>2</sub>PO<sub>4</sub> was added. The organic layer was separated, dried over Na<sub>2</sub>SO<sub>4</sub>, and concentrated in vacuo to give 10 mg of the crude trityl radical **2-CO<sub>2</sub>Et** as an orange-brown solid: IR 2958, 2923, 2855, 1702, 1488, 1453, 1366, 1237, 1148, 1109, 1043 cm<sup>-1</sup>; UV 247, 405, 493 nm; MS (ESI, (M+Na)<sup>+</sup>, m/z): 1034.00046 (observed), 1034.001493 (exact).

To a stirred solution of **10** (9.5 mg, 0.01 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL) was added BF<sub>3</sub>·Et<sub>2</sub>O (10 μL, 0.08 mmol) dropwise at room temperature. After being stirred for 1 h, the resulting dark green-blue solution was treated with a solution of SnCl<sub>2</sub> (3.2 mg, 0.017 mmol) in THF (0.4 mL). After 10 min, saturated aqueous KH<sub>2</sub>PO<sub>4</sub> was added. The organic layer was separated, dried over Na<sub>2</sub>SO<sub>4</sub>, and concentrated in vacuo to give 9 mg of the crude trityl radical **3-CO<sub>2</sub>Et** as an orange-brown solid: IR 2957, 2923, 2853, 1703, 1605, 1452, 1365, 1239, 1149, 1109, 1020 cm<sup>-1</sup>; UV 239, 449, 481 nm; MS (ESI, (M+Na)<sup>+</sup>, m/z): 961.98135 (observed), 961.980363 (exact).

The crude solid **1-CO<sub>2</sub>Et** was dissolved in dioxane (0.2 mL), and 1M aqueous KOH (0.1 mL) was added. The resultant dark orange-brown solution was heated at 50 °C for 2 h. After being cooled to room temperature, the reaction mixture was diluted with water, and washed twice with ether. The aqueous layer was acidified with 1 M HCl, and the resulting orange brown precipitate was extracted with ether. The ether layer was separated, dried over Na<sub>2</sub>SO<sub>4</sub>, and concentrated in vacuo. The residue was dissolved in 0.1 M aqueous NaOH (0.3 mL), and concentrated in vacuo to give 9 mg of trityl radical **1-CO<sub>2</sub>Na** as a dark green-yellow solid: IR ~3400 (broad), 2923, 1585, 1425, 1260, 1150, 879 cm<sup>-1</sup>. UV 277, 469 nm;

## Computational Methods

Density functional theory<sup>44,45</sup> was applied in this study to determine the optimized geometry of each species.<sup>46-49</sup> All calculations were performed using Gaussian 03<sup>50</sup> at the Ohio Supercomputer Center. Optimized geometries were obtained at the B3LYP/6-31G\* level. Electron spin densities (populations) were obtained from a natural population analysis (NPA) approach using single-point energies evaluated at the B3LYP/6-311+G\*\* level with 6d functions.<sup>51</sup> The effect of solvation on the gas-phase calculations was also investigated using the polarized continuum model (PCM).<sup>52-56</sup> Because the vibrational frequency calculation for each trityl radical could not be finished in more than 14 days of cpu time, we employed the bottom-of-the-well energies to compare the respective radicals.

## EPR Measurements

EPR measurements were carried out on an X band spectrometer with HS resonator at room temperature. General instrument settings are as follows: microwave power, 3 mW; modulation amplitude, 0.03 G; receiver gain  $3.5 \times 10^3$ ; modulation frequency, 6 kHz; scan time, 40 s; time constant, 82 ms. Total volume of all solutions used for EPR measurement was 50  $\mu$ L and was loaded into 50  $\mu$ L micropipettes that were sealed with clay.

## Mass Spectrometric Analysis

**ESI/MS**—Electrospray ionization (ESI) mass spectrometric analyses were performed for **1-CO<sub>2</sub>Et**, **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et**. The trityl solution (~1mM) in tetrahydrofuran (THF) was initially purged with Ar or O<sub>2</sub> gas for 4 min before mixing with THF/CH<sub>3</sub>OH solvent that was previously saturated with NaCl. This combination provided the necessary cationization for charging in the electrospray process for ion generation. Mass analysis was then performed in positive ion detection mode.

**MALDI-TOF/MS**—Matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) was used for **1-CO<sub>2</sub>Na** with the mass spectrometer operated in linear, positive ion mode and with a N<sub>2</sub> laser for laser desorption. Samples were prepared in 0.1% TFA at an approximate concentration of 50 pmol/ $\mu$ L. 2,5-Dihydroxybenzoic acid was prepared as a saturated solutions in 50% ACN/0.1% TFA (in water). Allotments of 1 mL of matrix and 1 mL of ~1 mM aqueous solution of **1-CO<sub>2</sub>Na** (purged with Ar or O<sub>2</sub> gas for 4 min) were thoroughly mixed together; 0.5 mL of this was spotted on the target plate and allowed to dry prior to MS analysis.

## Cell Studies

Rat cardiac H9C2 cells were cultured in DMEM supplemented with 10% fetal bovine serum, 100 U/ml of penicillin, and 100  $\mu$ g/ml of streptomycin in 150 cm<sup>2</sup> tissue culture flasks at 37° C in a humidified atmosphere of 5% CO<sub>2</sub>. The cells were fed every 2-3 days, and subcultured once they reached 80-90% confluence. For determination of penetration of the trityl radicals into cells, H9C2 cells were incubated with the trityl compounds introduced by 10  $\mu$ L (1 mM) in 10:1 DMSO to water solution for various time points. Cells were then collected, and washed twice with PBS prior to EPR measurement.

## Cyclic Voltammetry Measurements

Cyclic voltammetry was performed on a potentiostat and computer-controlled electroanalytical system. Electrochemical measurements were carried out in a 5 ml cell equipped with a gold working electrode (1.6 mm diameter), a platinum-wire auxiliary electrode, and a Ag/AgCl reference electrode. The gold electrode was cleaned before each run by rubbing (circular motion for approximately 10s) on a polishing pad with crystal solution and then washed with water-acetone. Solutions were degassed by bubbling with the argon gas. Background current

(solvent with supporting electrolyte) corrections were done for all measurements. Half-wave potentials were calculated according to the relation  $E_{1/2} = (E_{pa} + E_{pc})/2$ .

Acetonitrile and phosphate buffered saline were used without further purification. Tetrabutylammonium perchlorate (TBAP) and 2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPO) were used as purchase.

All measurements were performed in dry acetonitrile with 0.1 M TBAP as supporting electrolyte for triester trityl radical **1-CO<sub>2</sub>Et**, or in phosphate aqueous solution for carboxylate salt trityl radical **1-CO<sub>2</sub>Na**. The concentration of the investigated compounds was about 1 mM. Voltage was cycled between -2.0 and 0.0 V beginning at 0.0 V, between 0.0 V and 2.0 V beginning at 0.0 V, and -2.0 V and 2.0 V beginning at 0.0 V.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Acknowledgements

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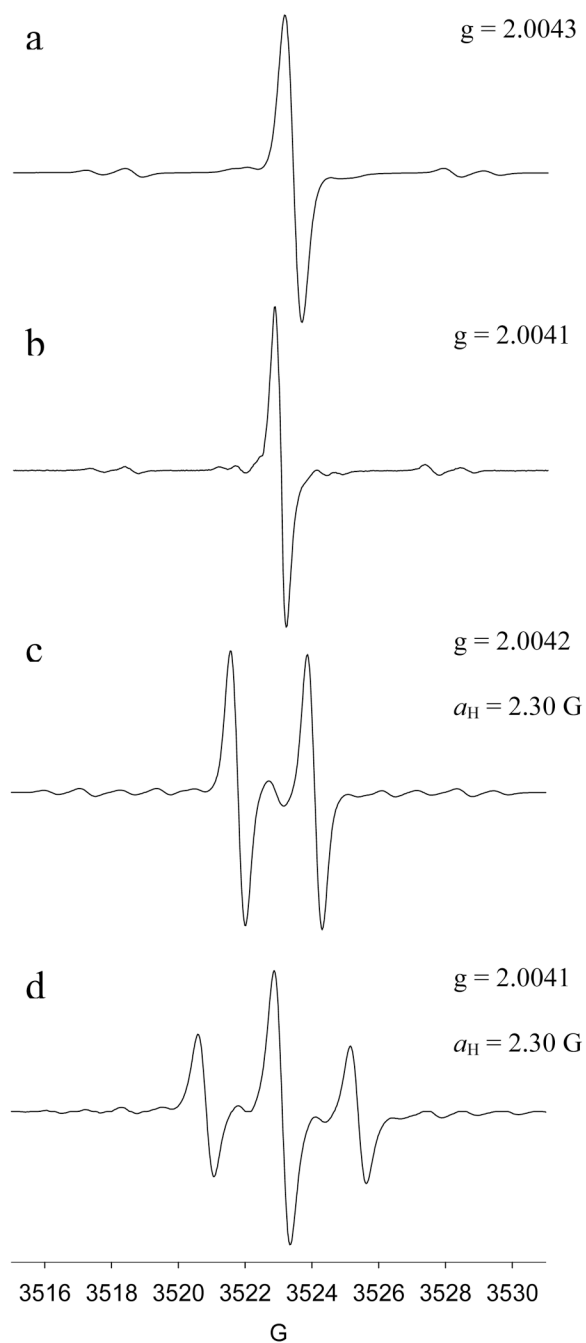
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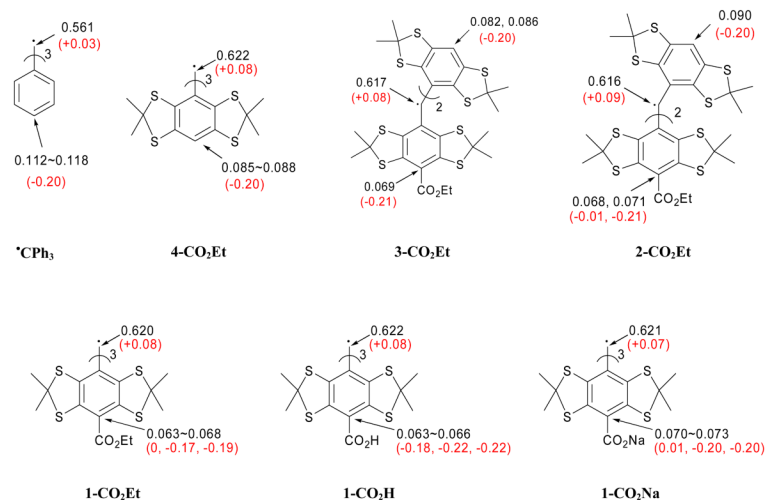
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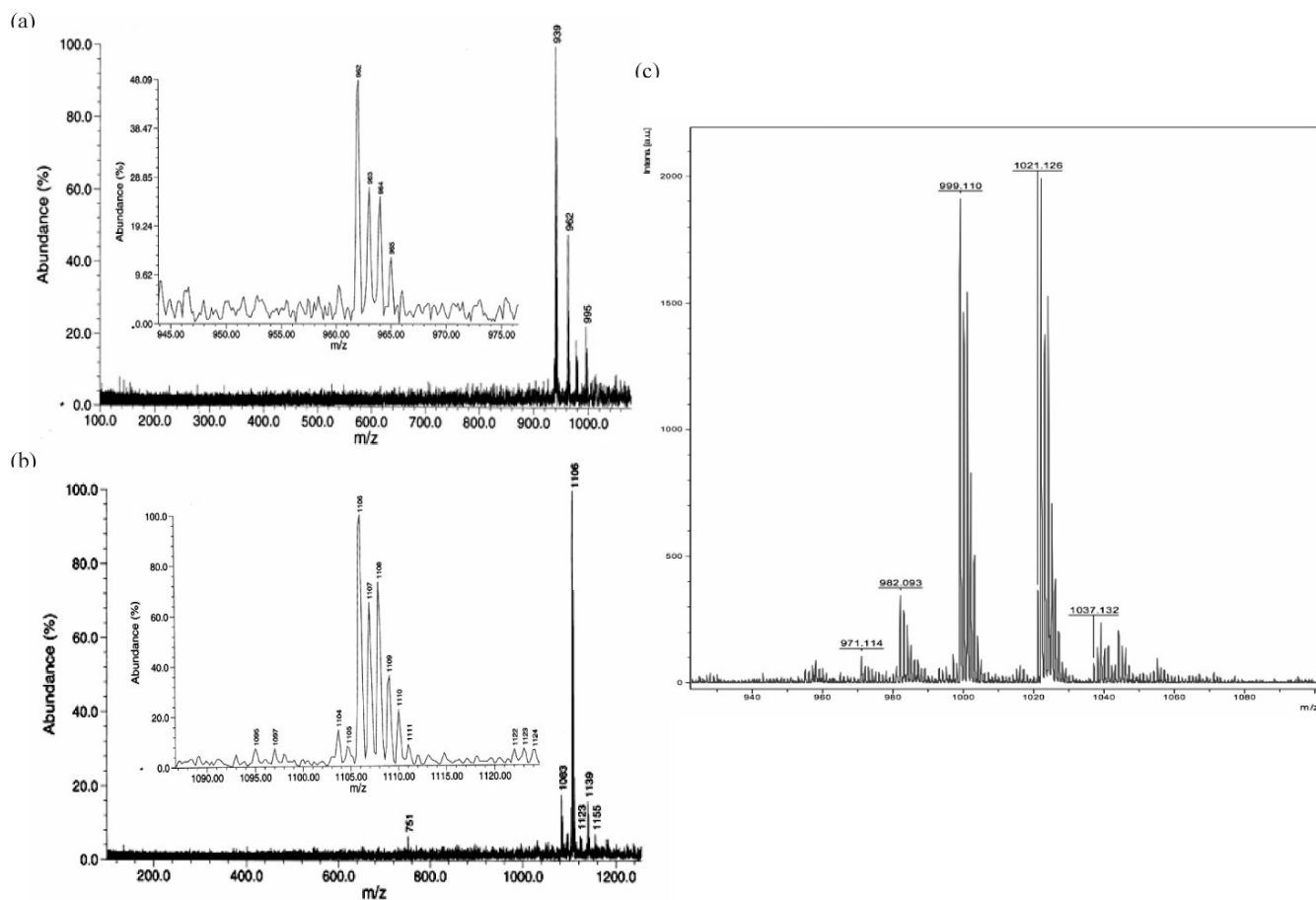




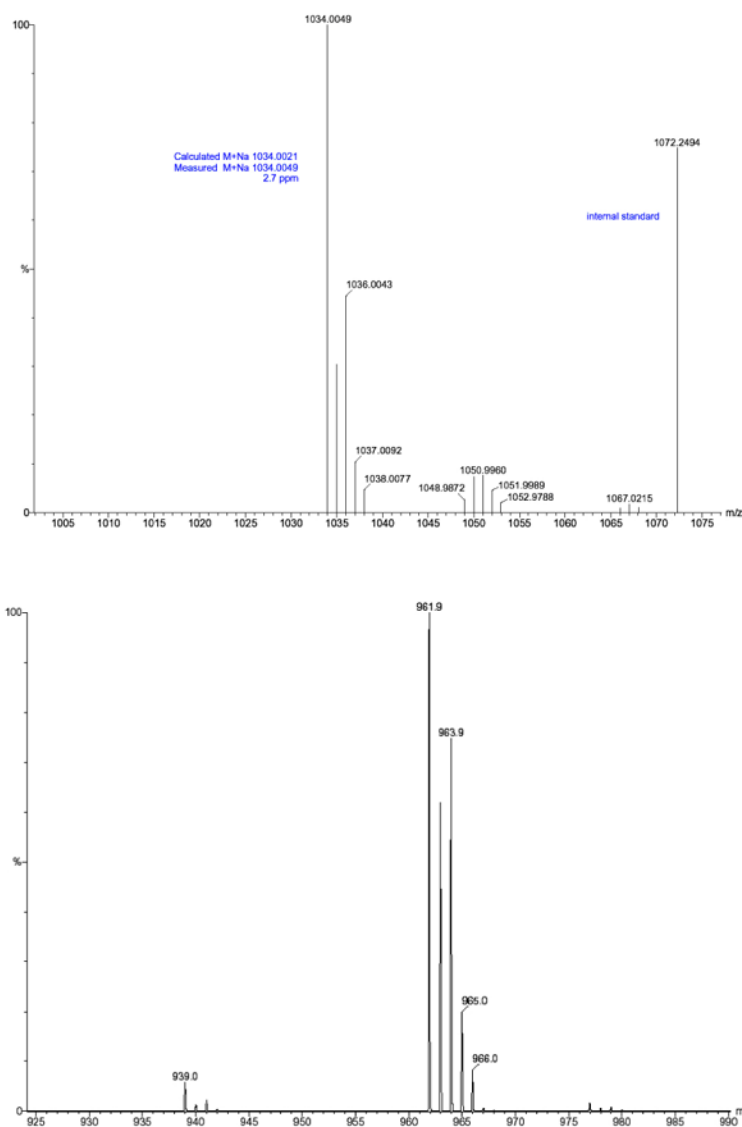
**Figure 1.** CW X-band EPR spectra of various trityl radicals in DMSO under anaerobic condition. (a) **1-CO<sub>2</sub>Et**; (b) **1-CO<sub>2</sub>Na** (in H<sub>2</sub>O); (c) **2-CO<sub>2</sub>Et**; (d) **3-CO<sub>2</sub>Et**. See Experimental Section for the EPR parameters.

**Figure 2.**

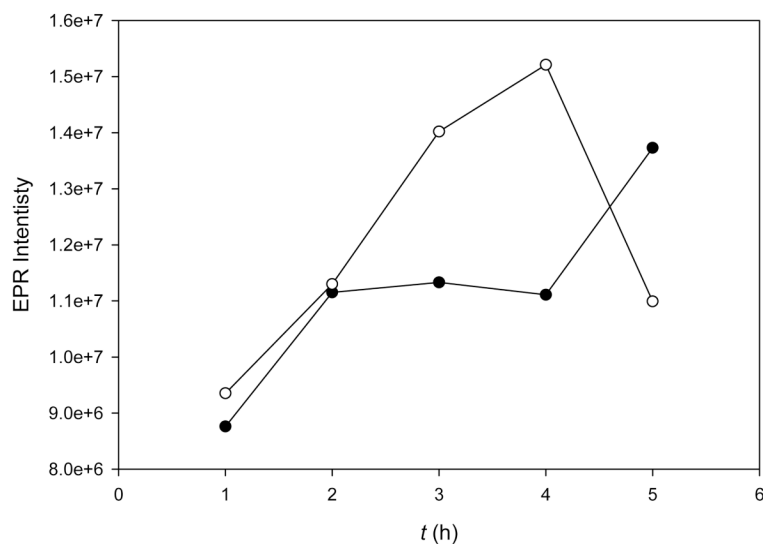
Spin populations ( $\alpha-\beta$ ) and atomic charges (in parentheses) of various trityl radicals at the B3LYP/6-311+G\*\*/B3LYP/6-31G\* level in the gas phase using the natural population analysis partitioning scheme.



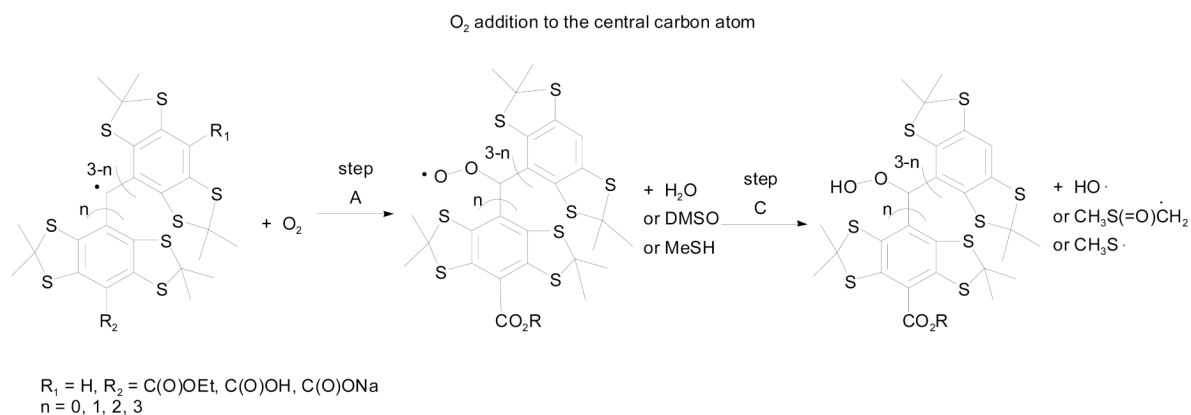
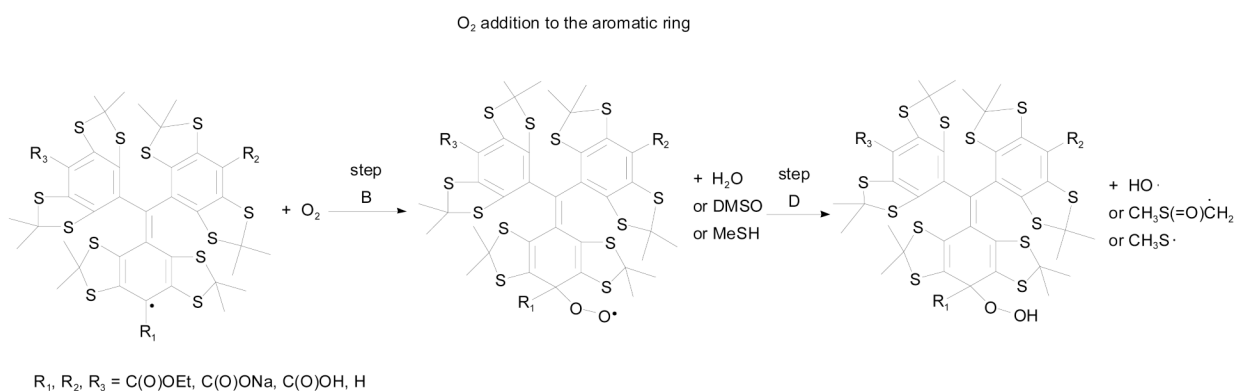
**Figure 3.** Product ESI mass spectra of (a) **3-CO<sub>2</sub>Et** and (b) **1-CO<sub>2</sub>Et**; and MALDI-ToF mass fingerprint of (c) **1-CO<sub>2</sub>Na** upon exposure to O<sub>2</sub>.



**Figure 4.** Product ESI mass spectra of M residues (trityl cation or anion) of (a) **2-CO<sub>2</sub>Et** and (b) **3-CO<sub>2</sub>Et** decomposition products.

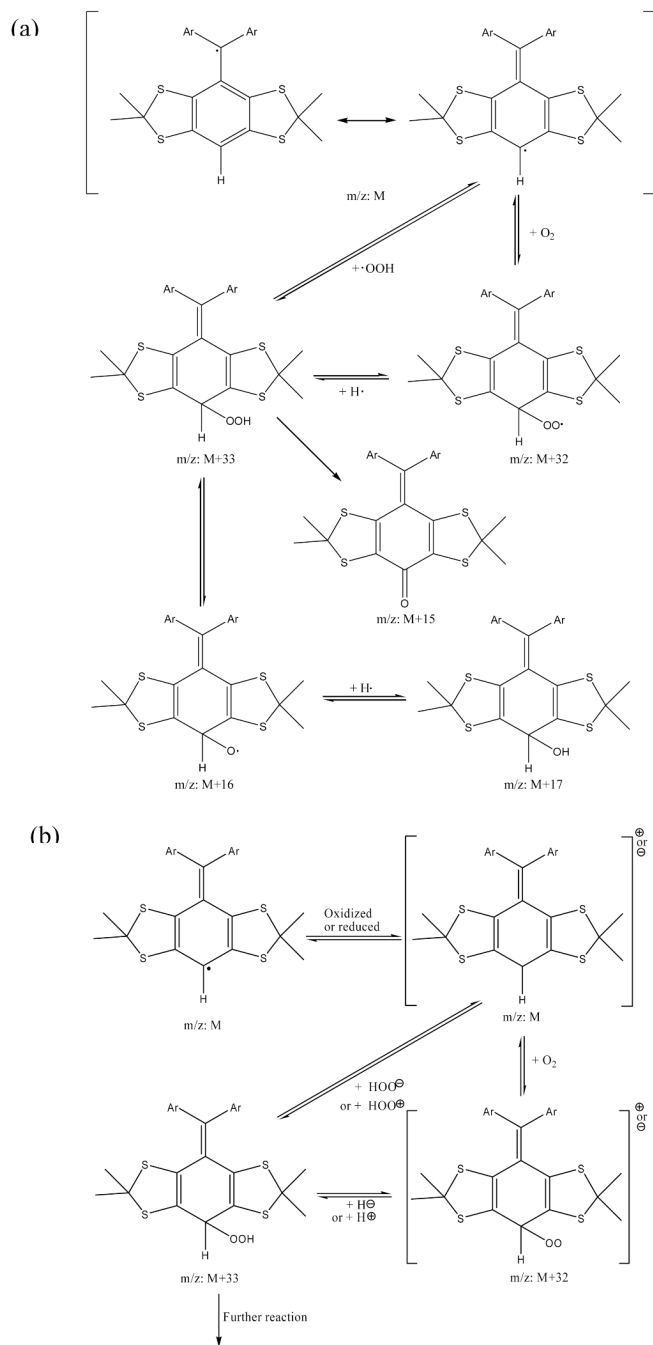


**Figure 5.** EPR signal intensity of **1-CO<sub>2</sub>Na** (○) and **1-CO<sub>2</sub>Et** (●) in rat H9C2 cardiomyocytes after 1, 2, 3, 4 and 5 hours of incubation and washing using phenol-red-free DMEM medium. (The **2-CO<sub>2</sub>Et** and **3-CO<sub>2</sub>Et** trityl radicals did not show an EPR signal after 1 hr of incubation and washing.)

**A****B**

**Scheme 1. Molecular Oxygen Reaction with Trityl Radicals by Addition to the Center Carbon Atom of the Trityl Radical (A) or by Addition to the Aromatic Ring of the Trityl Radical (B)**





**Scheme 2. Proposed mechanism for decomposition of an Unsubstituted Trityl Radical. (a) radical, (b) charged molecule pathways**

**Table 1**  
Predicted Reaction Energies  $\Delta E_0$  (kcal/mol) for the Formation of Various Trityl Radical-OOH Adducts at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* Level in the Gas Phase

Entry <sup>a</sup>	Reaction Energies <sup>b</sup> $\Delta E_0$ (in kcal/mol)													
	Step A	Step B	Step C			Step D			Step A + C			Step B + D		
			H <sub>2</sub> O	DMSO	MeSH	H <sub>2</sub> O	DMSO	MeSH	H <sub>2</sub> O	DMSO	MeSH	H <sub>2</sub> O	DMSO	MeSH
Ph <sub>3</sub> C <sup>+</sup>	-4.0	13.7	35.5	20.7	2.1	36.7	21.9	3.3	31.5	16.7	-1.9	50.4	35.6	17.0
1-CO <sub>2</sub> H	24.2	26.7	36.6	21.9	3.2	34.0	19.3	0.7	60.8	46.1	27.4	60.7	46.0	27.4
1-CO <sub>2</sub> H <sup>c</sup>	15.4	25.3	37.2	20.9	4.1	34.2	17.9	1.1	52.6	36.3	19.5	59.5	43.2	26.4
1-CO <sub>2</sub> Na	22.8	24.2	36.8	22.0	3.4	32.2	17.5	-1.2	59.6	44.8	18.2	56.4	41.7	23.0
1-CO <sub>2</sub> Na <sup>c</sup>	14.3	22.8	37.0	20.7	3.9	34.8	18.5	1.7	51.3	35.0	26.2	57.6	41.3	24.5
1-CO <sub>2</sub> Et	24.0	25.9	38.4	23.7	5.1	34.3	19.5	0.9	62.4	47.7	29.1	60.2	45.4	26.8
2-CO <sub>2</sub> Et	22.5	17.7	38.5	23.8	5.1	35.0	20.3	1.6	61.0	46.3	27.6	52.7	38.0	19.3
3-CO <sub>2</sub> Et	23.9	17.4	35.9	21.2	2.5	35.2	20.5	1.8	59.8	45.1	26.4	52.6	37.9	19.2
4-CO <sub>2</sub> Et	19.7	17.2	37.3	22.5	3.9	35.2	20.4	1.8	57.0	42.2	23.6	52.4	37.6	19.0

<sup>c</sup> In aqueous system, calculated at the B3LYP/6-311+G\*\*/B3LYP/6-31G\* level using the PCM model for water.

Table 2  
ESI and MALDI-ToF Mass Fingerprints of Various Trityl Radicals in the Presence of O<sub>2</sub>

Entry	$M_{obs}$ ( $M_{exact}$ )	$M+Na_{obs}$ ( $M+Na_{exact}$ )	$M+Na+O-H_{obs}$ ( $M+Na+O-H_{exact}$ )	$M+Na+O_{obs}$ ( $M+Na+O_{exact}$ )	$M+Na+OH_{obs}$ ( $M+Na+OH_{exact}$ )	$M+Na+OOH_{obs}$ ( $M+Na+OOH_{exact}$ )
1-CO <sub>2</sub> Et	1083.036 (1083.036)	1106.026 (1106.023)	not observed (1121.010)	1121.998 (1122.018)	1123.054 (1123.026)	1139.066 (1139.021)
2-CO <sub>2</sub> Et	1011.030 (1011.011)	1034.000 (1034.001)	not observed (1048.988)	1049.988 (1049.996)	1050.999 (1051.004)	1067.047 (1067.000)
3-CO <sub>2</sub> Et	938.993 (938.990)	961.981 (961.980)	976.946 (976.967)	977.970 (977.975)	978.937 (978.983)	994.976 (994.978)

<sup>a</sup>1-CO<sub>2</sub>Na: 999.110 (C(ArCO<sub>2</sub>H)<sub>3</sub>), 1021.115 ((ArCO<sub>2</sub>)<sub>2</sub>Na)·C(ArCO<sub>2</sub>H)<sub>2</sub>, 1038.117 ((ArCO<sub>2</sub>)Na)·C(ArCO<sub>2</sub>H)<sub>2</sub>+OH).

**Table 3**  
**Predicted Electron Affinities, Ionization Potentials, BDE's and Energies of Protonation of Various Trityl Radicals at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* Level**

Ar <sub>3</sub> C <sup>•</sup> Radical	Electron Affinity (EA) kcal/mol (eV)	Ionization Potential (IP) kcal/mol (eV)	C <sub>central</sub> -H Bond Dissociation Energy BDE (0K) kcal/mol	ΔE <sub>acid</sub> (0K) kcal/mol
Ph <sub>3</sub> C <sup>•</sup>	32.7 (1.42)	137.8 (5.98)	80.4	362.8
1-CO <sub>2</sub> Na	38.8 (1.68)	118.1 (5.12)	77.1	353.4
1-CO <sub>2</sub> Na (aq) <sup>a</sup>	79.4 (3.44)	109.8 (4.76)	80.2	313.4
1-CO <sub>2</sub> Et	60.7 (2.63)	139.0 (6.03)	77.2	331.7
2-CO <sub>2</sub> Et	57.6 (2.50)	137.2 (5.95)	77.8	335.3
3-CO <sub>2</sub> Et	54.4 (2.36)	136.2 (5.91)	77.9	338.6
4-CO <sub>2</sub> Et	50.3 (2.18)	134.8 (5.85)	78.1	342.9

<sup>a</sup> At the B3LYP/6-311+G\*\*//B3LYP/6-31G\* level using PCM.

**Table 4**  
**Redox Potentials  $E_{1/2}$  of 1-CO<sub>2</sub>Na and 1-CO<sub>2</sub>Et Determined by Cyclic Voltammetry<sup>a</sup> relative to Ag / 4M AgCl**

Compound	Solvent	$E_{1/2}$ (Ox <sub>1</sub> ) (V) ( $\Delta E_p$ <sup>c</sup> [mV])	$E_{1/2}$ (Ox <sub>2</sub> ) (V) ( $\Delta E_p$ [mV])	$E_{1/2}$ (Red <sub>1</sub> ) (V) ( $\Delta E_p$ [mV])
TEMPO	aqueous	0.52 (90)		
TEMPO	CH <sub>3</sub> CN	0.66 (85)		
1-CO <sub>2</sub> Na	aqueous	0.45 (60)	0.83 (50)	-0.65 (60)
1-CO <sub>2</sub> Na	CH <sub>3</sub> CN <sup>b</sup>	0.59 (60)	0.97 (50)	-0.51 (60)
1-CO <sub>2</sub> Et	CH <sub>3</sub> CN	1.42 <sup>d</sup>	0.94 <sup>e</sup>	-1.52 (400)

<sup>a</sup> Scan rate = 100 mV/s.

<sup>b</sup> Using TEMPO to calibrate aqueous solution to acetonitrile.

<sup>c</sup>  $\Delta E_p = |E_{pa} - E_{pc}|$ .

<sup>d</sup> Anodic peak only.

<sup>e</sup> Cathodic peak only.