

Preparation, Structural Characterization, and Thermochemistry of an Isolable 4-Arylphenoxyl Radical

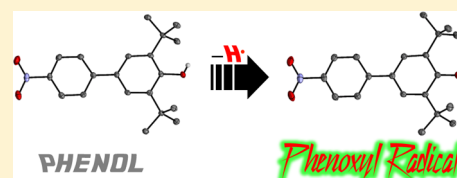
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S Supporting Information

ABSTRACT: The preparation and full characterization of the 4-(nitrophenyl)-phenoxyl radical, 2,6-di-*tert*-butyl-4-(4'-nitrophenyl) phenoxyl radical (^tBu₂NPArO•) is described. This is a rare example of an isolable and crystallographically characterized phenoxyl radical and is the only example in which the parent phenol is also crystallographically well-defined. Analysis of EPR spectra indicates some spin delocalization onto the secondary aromatic ring and nitro group. Equilibrium studies show that the corresponding phenol has an O–H bond dissociation free energy (BDFE) of 77.8 ± 0.5 kcal mol⁻¹ in MeCN (77.5 ± 0.5 kcal mol⁻¹ in toluene). This value is higher than related isolated phenoxyl radicals, making this a useful reagent for hydrogen atom transfer (HAT) studies. Additional thermochemical and spectroscopic parameters are also discussed.



INTRODUCTION

Stable and transient phenoxyl radical species are important in chemical processes spanning a large range of applications. Examples include tyrosine/tyrosyl radical mediated enzymatic electron transfers and hydrogen atom transfers,¹ food preservation (such as butylated hydroxytoluene or BHT),² and fundamental studies of proton-coupled electron transfer (PCET)/hydrogen atom transfer (HAT) reactions.³ While most phenoxyl radicals are transient, sufficiently sterically encumbered phenoxyl radicals can be stable in solution under anaerobic conditions.⁴ The 2,6-di-*tert*-butyl-4-phenylphenoxyl radical (^tBu₂PhArO•), for instance, was prepared by Müller and co-workers in 1959, and isolated in 78–88% purity.⁵ Our laboratory has reported the clean isolation and structural characterization of the 2,4,6-tri-*tert*-butylphenoxyl radical (^tBu₃ArO•)⁶ and the 4,4' coupled dimer of the 2,6-di-*tert*-butyl-4-methoxyphenoxyl radical (^tBu₂MeOArO•).⁷ The latter has a very weak C–C bond and is primarily dissociated in solution.

The O–H bond dissociation free energies (BDFEs) of the 2,6-di-*tert*-butyl-6-R-phenols are significantly modulated by the R substituent. The H atom affinities of the corresponding phenoxyl radicals (described by phenolic O–H BDFEs) range from 73.8 kcal mol⁻¹ for the isolable⁷ R = OMe species to 80.4 kcal mol⁻¹ for the transiently lived⁸ R = NO₂ species (BDFEs in toluene), with the R = *t*Bu and Ph derivatives being the same within error.^{9a} The isolable R = *t*Bu and OMe derivatives have proved to be useful hydrogen atom accepting reagents,¹⁰ complementary due to their different hydrogen atom affinities.^{10a,b} With the goal of preparing an isolable phenoxyl radical with a higher H atom affinity, we report here the preparation, full characterization, and thermochemistry of 2,6-di-*tert*-butyl-4-(4'-nitrophenyl)phenoxyl radical, or ^tBu₂NPArO•.

RESULTS AND DISCUSSION

^tBu₂NPArO• was prepared by treating a benzene solution of 2,6-di-*tert*-butyl-4-(4'-nitrophenyl)phenol, ^tBu₂NPArO-H,¹¹ with aqueous 1 M sodium hydroxide and potassium ferricyanide under anaerobic conditions. After 30 min, removal of the solvent under vacuum, extraction of the dark green material with pentane, and crystallization at –30 °C over 24 h yielded black crystals. These were found to be of high purity from elemental analysis and the ¹H NMR spectra showed only minor diamagnetic impurities (<5%; see the Supporting Information).

High-quality X-ray crystal structures of ^tBu₂NPArO• and its parent phenol were collected for structural comparison (Figure 1, Table 1). This type of direct structural comparison of a phenoxyl radical/phenol has previously not been possible. The parent phenol of the only previously structurally characterized phenoxyl radical, ^tBu₃ArO-H, was found to be disordered over three positions in its crystals.⁶

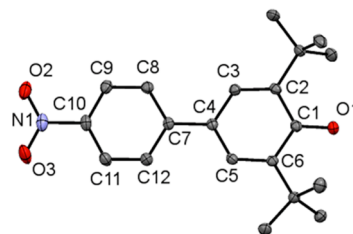


Figure 1. ORTEP drawing of ^tBu₂NPArO• showing 50% probability thermal ellipsoids and labels for select atoms. Hydrogen atoms are omitted for clarity.

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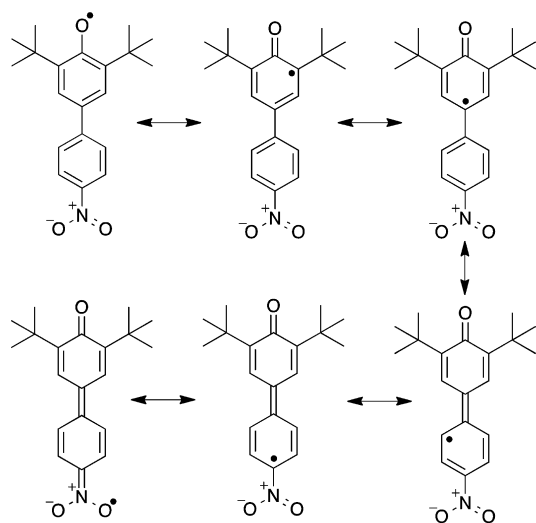
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Table 1. Select Bond Lengths (Å) and Aryl–Aryl Torsion Angles (deg) of ^tBu₂NPArO• and ^tBu₂NPArO-H

	^t Bu ₂ NPArO•	^t Bu ₂ NPArO-H	difference
O1–C1	1.2509(14)	1.3794(12)	–0.1285(18)
C1–C2	1.4699(17)	1.4100(14)	0.0599(22)
C2–C3	1.3696(16)	1.3944(13)	–0.0248(21)
C3–C4	1.4194(16)	1.3928(13)	0.0266(21)
C4–C5	1.4228(17)	1.3964(13)	0.0264(21)
C5–C6	1.3711(16)	1.3941(14)	–0.0230(21)
C6–C1	1.4751(16)	1.4120(14)	0.0631(21)
C4–C7	1.4754(16)	1.4829(13)	–0.0075(21)
C7–C8	1.4069(17)	1.4008(14)	0.0061(22)
C8–C9	1.3833(17)	1.3861(14)	–0.0028(22)
C9–C10	1.3873(18)	1.3839(15)	0.0034(23)
C10–C11	1.3828(19)	1.3842(15)	–0.0014(24)
C11–C12	1.3842(17)	1.3853(14)	–0.0011(22)
C12–C7	1.4114(16)	1.4023(14)	0.0091(21)
C10–N1	1.4272(16)	1.4672(13)	–0.0400(21)
N1–O2	1.2262(16)	1.2312(13)	–0.0050(21)
N1–O3	1.2289(16)	1.2286(13)	–0.0003(21)
avg Ar–Ar torsion angle ^a	17.5(1)	31.9(1)	–14.4(1)

^aAverage aryl–aryl torsion angle (deg) refers to the average dihedral angle measured for C5–C4–C7–C12 and C3–C4–C7–C8.

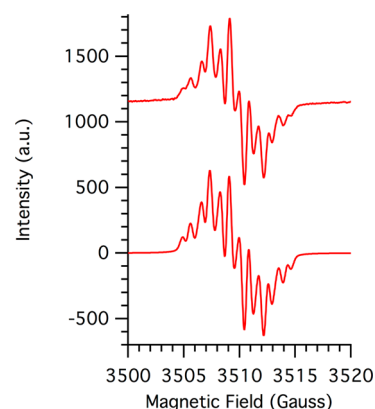
The largest difference between the phenoxyl and phenol structures is in the O1–C1 bond distance, 1.251 vs 1.379 Å. This bond shortening of 0.128 Å is consistent with previous conclusions that phenoxyl radicals have significant ketone character, as suggested by the resonance forms in Scheme 1.^{4,6,12} The changes in the phenolic aromatic bond lengths

Scheme 1. Radical Resonance Forms of ^tBu₂NPArO•

support this model, as the C1–C2 and C1–C6 bond lengths (avg 0.062 Å) more than the C3–C4 and C4–C5 bonds (avg 0.027 Å) while the C2–C3 and C5–C6 bonds shorten (avg –0.024 Å).

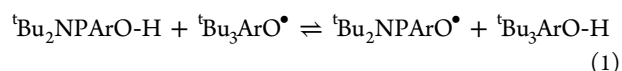
The aryl–aryl linkage is slightly shorter in the radical, by 0.0075(22) Å, suggesting a small quinomethide component (Scheme 1). This is also suggested by the 0.0400(21) shortening of the C10–N1 bond to the nitro group, and the smaller average aryl–aryl torsion angle,¹³ of 17.5° in the radical vs 31.9° observed in the phenol.

The X-band CW EPR spectrum of ^tBu₂NPArO• in toluene displays a multiline pattern centered at $g = 2.007(2)$ that is well modeled by simulation (Figure 2). Hyperfine coupling

**Figure 2.** X-band EPR spectrum of 1 mM ^tBu₂NPArO• in toluene recorded at 25 °C (top) and simulation (bottom).

constants were assigned by comparison to previously reported phenoxyl radical data⁹ and from the structural changes observed in the crystal structure: $a_{3,5}(2H) = 1.80$ G, $a_{8,12}(2H) = 1.61$ G, $a_{9,11}(2H) = 0.74$ G and $a_{NO_2}(1N) = 0.50$ G.¹⁴ The ¹⁴N hyperfine coupling indicates spin density on the nitro group, as depicted in the bottom of Scheme 1. The observed spin density onto the nitro group suggests that the thermochemistry of ^tBu₂NPArO• should be perturbed from that of the unsubstituted ^tBu₂PhArO•.

The O–H BDFE of ^tBu₂NPArO-H, was determined by equilibration with the thermochemically well-established^{3a} ^tBu₃ArO• radical. In either acetonitrile-*d*₃ or toluene-*d*₈, a known concentration of ^tBu₂NPArO-H was combined with several different concentrations of ^tBu₃ArO• (eq 1).



Integration of the ¹H NMR signals of these solutions gave equilibrium concentrations from which equilibrium constants were determined: $K_{\text{eq}}(\text{acetonitrile}) = 0.25 \pm 0.03$, $K_{\text{eq}}(\text{toluene}) = 0.26 \pm 0.03$. Thus, the O–H bond in ^tBu₂NPArO-H is 0.8 ± 0.1 kcal mol^{–1} stronger than that in ^tBu₃ArO-H in both acetonitrile and toluene. Using the known BDFE values of ^tBu₃ArO-H³ and eq 2 gives BDFE(^tBu₂NPArO-H_{MeCN}) = 77.8 ± 0.5 kcal mol^{–1} and BDFE(^tBu₂NPArO-H_{tol}) = 77.5 ± 0.5 kcal mol^{–1}. While this is a small increase, ^tBu₂NPArO• is to our knowledge the thermodynamically strongest isolable, reagent quality organic hydrogen atom abstractor available.

$$\text{BDFE}({}^t\text{Bu}_2\text{NPArO-H}) = \text{BDFE}({}^t\text{Bu}_3\text{ArO-H}) - RT \ln(K_{\text{eq}}) \quad (2)$$

Pedulli and co-workers have previously reported an empirical correlation between the O–H bond strengths of 2,6-*tert*-butyl-substituted phenols with the EPR hyperfine coupling constants, $a_{3,5}$, of the corresponding phenoxyl radicals.^{9a} Figure 3 shows a slightly modified version of this correlation using revised BDFE values.¹⁵ The values for ^tBu₂NPArO• follow this correlation very closely.

Cyclic voltammetry of ^tBu₂NPArO• in acetonitrile with 0.1 M [ⁿBu₄N]PF₆ as a supporting electrolyte displayed a reversible couple with $E_{1/2} = -0.436 \pm 0.010$ V vs Fc^{+/0}. This value is 0.26

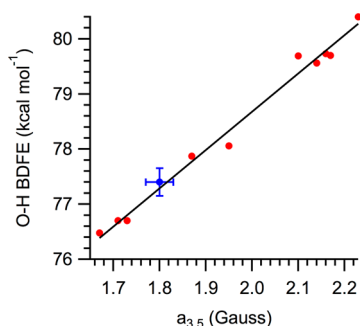


Figure 3. 2,6-^tBu₂-4-X-ArO-H bond dissociation free energies (BDFEs) vs 3,5 hyperfine coupling constant for 2,6-^tBu₂-4-X-ArO[•] radicals. Data in red are part of the correlation reported in ref 9a (revised to use updated BDFE values¹⁵); blue data point is ^tBu₂NPArO(-H).

V less negative than the related ^tBu₃ArO^{0/-} potential of -0.70 V vs Fc^{+/0}.^{3a} This is much larger than the reported potential difference of only 0.045 V between ^tBu₂PhArO[•] and ^tBu₃ArO[•] in 9:1 MeCN/H₂O,¹⁶ illustrating the effect of the nitro substituent on the phenoxyl/phenol thermochemistry.

The cyclic voltammogram of ^tBu₂NPArO-H displayed an irreversible anodic peak centered at 0.975 ± 0.010 V vs Fc^{+/0}. It is presumably irreversible due to loss of the proton from the highly acidic radical cation.

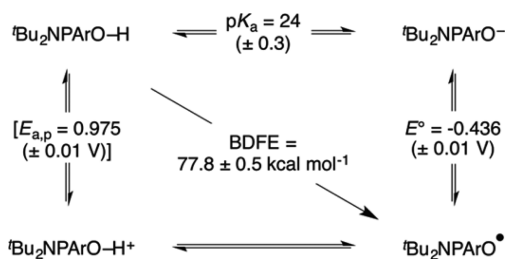
The reduction potential of ^tBu₂NPArO[•] and the BDFE (^tBu₂NPArO-H) imply that in acetonitrile the pK_a of ^tBu₂NPArOH is 24 ± 0.4 by Hess' law (eq 3). Compared to its parent phenylphenol, the (nitrophenyl)phenol has a significantly higher acidity¹⁷ and more positive reduction potential. These are both due to the stabilization of the phenoxide anion by the 4-substituted nitrobenzene group. The higher BDFE of ^tBu₂NPArO-H is due to the shifts in pK_a and E^o not exactly offsetting each other, with the nitro group affecting the pK_a less in free energy terms.¹⁸

$$\text{BDFE} = 1.37\text{p}K_{\text{a}} + 23.06E^{\circ} + C_{\text{G}} \quad (3)$$

These values can be assembled into a “square scheme” that describes the PCET thermochemistry of ^tBu₂NPArO-H (Scheme 2). We have included the irreversible anodic peak potential, E_{a,p}, even though it is not a thermochemical value. Using this value to crudely estimate E^o(^tBu₂NPArO-H^{+/0}) ≅ +0.95 V would imply that the pK_a of the radical cation is about 9 units lower than that of the phenol.

In conclusion, the 4-(nitrophenyl)phenoxyl radical ^tBu₂NPArO[•] is a previously unreported phenoxyl radical that

Scheme 2. Thermochemical “Square Scheme” for ^tBu₂NPArO(-H)^a



^aE_{a,p} is bracketed since it refers to an irreversible anodic peak potential and is not a thermochemical value.

is easily prepared in high purity and reasonable yield. Equilibrium studies show that the ^tBu₂NPArO-H BDFE is modestly stronger than that of its unsubstituted isolable relative, ^tBu₂PhArO[•] (ΔBDFE_{toluene} = 0.8 kcal mol⁻¹). ^tBu₂NPArO-H has the highest reported BDFE of any isolable organic hydrogen atom acceptor: 77.8 ± 0.5 kcal mol⁻¹ in acetonitrile and 77.5 ± 0.5 kcal mol⁻¹ in toluene. The combination of easy isolation of the phenoxyl radical in pure form and its relatively high hydrogen atom affinity should make this a useful reagent for studying hydrogen atom transfer reactions.

EXPERIMENTAL SECTION

Materials. Unless otherwise noted, all chemicals were purchased from commercial sources and used without purification. Toluene was dried using a “Grubb’s type” Seca Solvent System. Acetonitrile was purchased from Burdick & Jackson (low-water brand) and stored in an argon-pressurized glovebox plumbed directly into the glovebox. Toluene-*d*₈ and acetonitrile-*d*₃ were dried over NaK and CaH₂, respectively, and vacuum distilled. ^tBu₃ArO^{•6} and ^tBu₂NPArOH¹⁹ were prepared following literature methods.

Instrumentation. All NMR spectra were collected on 500 MHz spectrometers and chemical shifts referenced to TMS using residual solvent peaks. The reported EPR spectrum was collected using an X-band spectrometer at room temperature in toluene. Simulation of the spectrum was performed using the W9SEPR program.

Synthesis of ^tBu₂NPArO[•]. A 100 mL two neck round-bottom flask was charged with 467 mg (1.43 mmol) of ^tBu₂NPArOH dissolved in ~15 mL of benzene, 5 mL of 1 M NaOH and a stir bar. The flask was fitted with a 180° Schlenk adapter on one neck and a solid addition funnel containing 1.20 g (3.64 mmol) of solid K₃Fe(CN)₆ on the other neck. The biphasic mixture was degassed by 3 sequential freeze-pump-thaw cycles. After degassed, the mixture was frozen and the K₃Fe(CN)₆ was added. The frozen mixture was allowed to thaw at room temperature and left to stir. After 1 h of stirring, the solvents were removed in vacuo and extracted with pentane. Crystals were grown from a saturated pentane solution at -30 °C. Yield: 279 mg, 52%. Anal. Calcd for C₂₀H₂₄NO₃: C, 73.59; H, 7.41; N, 4.29. Found: C, 73.88; H, 7.60; N, 4.34.

ASSOCIATED CONTENT

Supporting Information

NMR and optical spectra, BDFE calculations, electrochemical data, crystallographic information, and an ORTEP of ^tBu₂NPArOH. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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■ REFERENCES

- (1) (a) Styring, S.; Sjöholm, J.; Mamedov, F. *Biochim. Biophys. Acta* **2012**, *1817*, 76. (b) Reece, S. Y.; Nocera, D. G. *Annu. Rev. Biochem.* **2009**, *78*, 673. (c) McEvoy, J. P.; Brudvig, G. W. *Chem. Rev.* **2006**, *106*, 4455. (d) Whittaker, J. W. *Chem. Rev.* **2003**, *103*, 2347.
- (2) Babich, H. *Environ. Res.* **1982**, *29*, 1.
- (3) (a) Warren, J. J.; Tronic, T. T.; Mayer, J. M. *Chem. Rev.* **2010**, *110*, 6961. (b) Weinberg, D. R.; Gagliardi, C. J.; Hull, J. F.; Murphy, C. F.; Kent, C. A.; Westlake, B. C.; Paul, A.; Ess, D. H.; McCafferty, D. G.; Meyer, T. J. *Chem. Rev.* **2012**, *112*, 4016.
- (4) Altwicker, E. R. *Chem. Rev.* **1967**, *67*, 475.
- (5) (a) Müller, E.; Schick, A.; Scheffler, K. *Chem. Ber.* **1959**, *92*, 474. (b) Related 4-tolyl- and 4-anisyl-phenoxy radicals have been reported but not isolated; see ref 4.
- (6) Manner, V. W.; Markle, T. F.; Freudenthal, J. H.; Roth, J. P.; Mayer, J. M. *Chem. Commun.* **2008**, 256.
- (7) Wittman, J. M.; Hayoun, R.; Kaminsky, W.; Coggins, M. K.; Mayer, J. M. *J. Am. Chem. Soc.* **2013**, *135*, 12956.
- (8) Cook, C. D.; Gilmour, N. D. *J. Org. Chem.* **1960**, *25*, 1429.
- (9) (a) Brigati, G.; Lucarini, M.; Mugnaini, V.; Pedulli, G. F. *J. Org. Chem.* **2002**, *67*, 4828. (b) Lucarini, M.; Pedrielli, P.; Pedulli, G. F.; Cabiddu, S.; Fattuoni, C. *J. Org. Chem.* **1996**, *61*, 9259. (c) Lucarini, M.; Pedulli, G. F. *Chem. Soc. Rev.* **2010**, *39*, 2106. (d) Reiker, A.; Scheffler, K. *Justus Liebigs Ann. Chem.* **1965**, 689, 78.
- (10) (a) Waidmann, C. R.; Zhou, X.; Tsai, E. A.; Kaminsky, W.; Hrovat, D. A.; Borden, W. T.; Mayer, J. M. *J. Am. Chem. Soc.* **2009**, *131*, 4729. (b) Warren, J. J.; Mayer, J. M. *Proc. Natl. Acad. Sci. U. S. A.* **2010**, *107*, 5282. (c) Valdez, C. N.; Braten, M. B.; Soria, A.; Gamelin, D. R.; Mayer, J. M. *J. Am. Chem. Soc.* **2013**, *135*, 8492. (d) Schrauben, J. N.; Hayoun, R.; Valdez, C. N.; Braten, M. M.; Fridley, L.; Mayer, J. M. *Science* **2012**, *336*, 1298. (e) Manner, V. W.; Lindsey, A. D.; Mader, E. A.; Harvey, J. N.; Mayer, J. M. *Chem. Sci.* **2012**, *3*, 230. (f) Warren, J. J.; Mayer, J. M. *J. Am. Chem. Soc.* **2011**, *133*, 8544. (g) Mader, E. A.; Manner, V. W.; Markle, T. F.; Wu, A.; Franz, J. A.; Mayer, J. M. *J. Am. Chem. Soc.* **2009**, *131*, 4335.
- (11) Chern, Y. T.; Ju, M. H. *Macromolecules* **2009**, *42*, 169.
- (12) (a) O'Malley, P. J. *J. Phys. Chem. B* **2002**, *106*, 12331. (b) Xie, C.; Lahti, P. M.; George, C. *Org. Lett.* **2000**, *2*, 3417.
- (13) Average torsion angle refers to the average dihedral angle measured for atoms C3–C4–C7–C8 and C5–C4–C7–C12.
- (14) EPR hyperfine coupling constants reported have an assumed error margin of $\sim\pm 0.05$ G.
- (15) The correlation in ref 9a used bond dissociation enthalpies (BDEs), which were determined from equilibration experiments. Δ BDE values were obtained from these equilibrium constants and then converted to BDEs using a gas-phase BDE of ${}^t\text{Bu}_3\text{ArO-H}$. Here, we take their equilibrium constants to be Δ BDFEs and scale them to the reported BDFE of ${}^t\text{Bu}_3\text{ArO-H}$ in benzene.³
- (16) (a) Steuber, F. W.; Dimroth, K. *Chem. Ber.* **1966**, *99*, 258. (b) Values reported in (a): $E_{1/2}({}^t\text{Bu}_2\text{PhArO}^{\bullet-}) = -0.014$ V; $E_{1/2}({}^t\text{Bu}_3\text{ArO}^{\bullet-}) = -0.059$ V (both vs Ag/AgCl; in 9:1 MeCN/ H_2O ; 0.01m $[\text{Me}_4\text{N}^+][\text{OH}^-]$, 0.01m $[\text{Me}_4\text{N}^+][\text{Cl}^-]$).
- (17) The $\text{p}K_{\text{a}}$ of ${}^t\text{Bu}_2\text{PhArO-H}$ is taken to be roughly equal to the $\text{p}K_{\text{a}}$ of ${}^t\text{Bu}_3\text{ArO-H}$ (~ 28 in MeCN; ref 3) since their BDFEs and E° are roughly equivalent.
- (18) C_{G} is a constant that contains the free energy of formation of H^\bullet , free energy of solvation of H^\bullet , as well as the nature of the electrode. In MeCN, $C_{\text{G}} = 54.9$ kcal mol⁻¹. A more detailed description can be found in ref 3.
- (19) Chern, Y. T.; Ju, M. H. *Macromolecules* **2009**, *42*, 169.