

Perylene Diimide as a Precise Graphene-like Superoxide Dismutase Mimetic

Almaz S. Jalilov,[†] Lizanne G. Nilewski,[†] Vladimir Berka,[#] Chenhao Zhang,[†] Andrey A. Yakovenko,^{||} Gang Wu,[#] Thomas A. Kent,^{¶,⊥} Ah-Lim Tsai,^{*,#} and James M. Tour^{*,†,¶,§,||}

[†]Department of Chemistry, [‡]The NanoCarbon Center, [§]Department of Materials Science and NanoEngineering, Rice University, 6100 Main Street, Houston, Texas 77005, United States

[¶]Department of Neurology, Baylor College of Medicine, Houston, Texas 77030, United States

[⊥]Center for Translational Research in Inflammatory Diseases, Michel E. DeBakey VA Medical Center, Houston, Texas 77030, United States

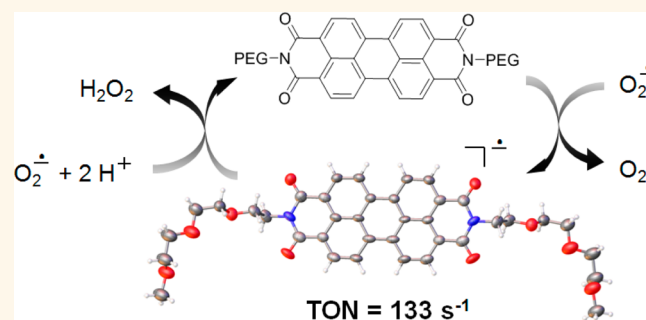
[#]Hematology, Internal Medicine, University of Texas Houston Medical School, Houston, Texas 77030, United States

^{||}Argonne National Laboratory, X-ray Science Division, Advanced Photon Source, Argonne, Illinois 60439, United States

Supporting Information

ABSTRACT: Here we show that the active portion of a graphitic nanoparticle can be mimicked by a perylene diimide (PDI) to explain the otherwise elusive biological and electrocatalytic activity of the nanoparticle construct. Development of molecular analogues that mimic the antioxidant properties of oxidized graphenes, in this case the poly(ethylene glycolated) hydrophilic carbon clusters (PEG-HCCs), will afford important insights into the highly efficient activity of PEG-HCCs and their graphitic analogues. PEGylated perylene diimides (PEG_n-PDI) serve as well-defined molecular analogues of PEG-HCCs and oxidized graphenes in general, and their antioxidant and superoxide dismutase-like (SOD-like) properties were studied. PEG_n-PDIs have two reversible reduction peaks, which are more positive than the oxidation peak of superoxide (O₂^{•-}). This is similar to the reduction peak of the HCCs. Thus, as with PEG-HCCs, PEG_n-PDIs are also strong single-electron oxidants of O₂^{•-}. Furthermore, reduced PEG_n-PDI, PEG_n-PDI^{•-}, in the presence of protons, was shown to reduce O₂^{•-} to H₂O₂ to complete the catalytic cycle in this SOD analogue. The kinetics of the conversion of O₂^{•-} to O₂ and H₂O₂ by PEG₈-PDI was measured using freeze-trap EPR experiments to provide a turnover number of 133 s⁻¹; the similarity in kinetics further supports that PEG₈-PDI is a true SOD mimetic. Finally, PDIs can be used as catalysts in the electrochemical oxygen reduction reaction in water, which proceeds by a two-electron process with the production of H₂O₂, mimicking graphene oxide nanoparticles that are otherwise difficult to study spectroscopically.

KEYWORDS: superoxide dismutase, reactive oxygen species, radical anion, electron paramagnetic resonance, perylene diimide



Superoxide dismutase (SOD) is a metalloenzyme that controls the fluxes of the partially reduced products of molecular O₂ or reactive oxygen species (ROS) during the cycle of molecular O₂ in biological systems.^{1–3} ROS are involved in the progression of aging as well as a variety of acute and chronic diseases such as cancer and vascular diseases.^{4–6} Superoxide is a radical anion (O₂^{•-}) that is the primary ROS produced by direct single-electron reduction of molecular O₂.⁷ Subsequent proton-coupled reduction of O₂^{•-} would lead to the other secondary members of ROS such as hydrogen peroxide (H₂O₂), organic peroxides (ROOH), hydroperoxyl radical (HO₂[•]) and hydroxyl radical (HO[•]).^{8–10} In aprotic solvents, O₂^{•-} is stable and exhibits reversible outer-sphere electron-transfer (ET) behavior. However, in the presence of

protons, O₂^{•-} proceeds quickly to form secondary ROS through consecutive proton-coupled electron transfer (PCET) reactions. SOD possesses the combination of redox-active metal centers and ligand environments that catalytically dismutate 2 O₂^{•-} into O₂ and H₂O₂: 2 O₂^{•-} + 2 H⁺ → O₂ + H₂O₂.³

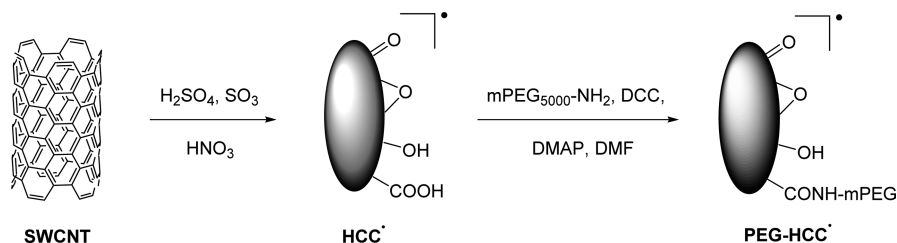
There have been many efforts to synthesize molecular organometallic complexes that can mimic the catalytic cycle of SOD.^{11–13} Over the past two decades, many high-valent metal

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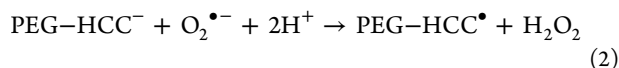
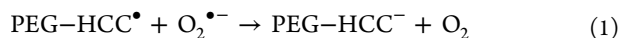
Scheme 1. Synthesis of PEG–HCCs from SWCNTs as the Starting Material



complexes have been studied to gain fundamental insights into the structural, functional and mechanistic aspects of the enzymes and their counterparts.^{14–17} Detailed mechanistic studies have shown that ET occurs through either inner-sphere or outer-sphere mechanisms and the $M-O_2^{\bullet-}$ species is an important reactive intermediate formed via the inner-sphere mechanism.¹⁸ However, another class of effective inhibitors of ROS are antioxidants that are purely organic molecules.^{19,20} Organic molecule ET mechanisms are generally complex and in most cases they occur through a continuum of mixed inner-sphere and outer-sphere mechanisms.^{21,22}

Recently, we developed metal-free carbon nanomaterials, poly(ethylene glycolated) hydrophilic carbon clusters (PEG–HCCs, Scheme 1), which show highly efficient catalytic conversion of $O_2^{\bullet-}$ into O_2 and H_2O_2 (eqs 1 and 2) that rivals that of bovine Cu/Zn SOD, the most efficient native SOD enzyme.²³

Single-walled carbon nanotubes (SWCNTs) are oxidatively disintegrated and split open to form graphene domains that are $3\text{ nm} \times 35\text{ nm}$ with large amounts of oxidized functionalities that make the carbon system extremely electron deficient.²⁴ They have been extensively characterized by transmission electron microscopy (TEM), atomic force microscopy (AFM), X-ray photoelectron spectroscopy (XPS), dynamic light scattering, Raman spectroscopy (confirming the loss of the tubular form through complete diminution of the radial breathing modes), electron paramagnetic resonance (EPR), and further tested for in vitro and in vivo toxicity and biological activity as antioxidants.^{25–32} The HCCs also bear, on average, one unpaired electron per nanoparticle and the nanoparticle radical is air stable.²³ As we have shown previously, HCCs can efficiently oxidize $O_2^{\bullet-}$ into O_2 , one of the two half-reactions processes (eq 1) seen in SOD, and this might also be the rate-determining step of the SOD catalytic cycle.²⁴ This is followed by reduction of $O_2^{\bullet-}$ in the presence of water to form H_2O_2 (eq 2).



In this work, we report poly(ethylene glycolated) perylene diimide (PEG_n–PDI) as an example of small molecular analogues of PEG–HCCs, and more generally well-defined highly oxidized graphene analogues. In our previous work using HCC as an electrocatalyst for the oxygen reduction reaction (ORR),²⁴ we showed that proton transfer steps are much faster than the electron-transfer steps, and here we use the same technique to evaluate the involvement of proton transfer steps during the dismutation of $O_2^{\bullet-}$ using the well-defined carbon core molecule PDI in PEG_n–PDI. The understanding of graphene-based catalysts can benefit from the study of

molecular analogues, and we propose that PDI serves as a simplified model of oxidized graphenes.

RESULTS AND DISCUSSION

PEG_n–PDI and Its Reactions with Superoxide. PDI was identified as an electron deficient molecular analogue of HCCs, and was further linked with two polyethylene glycol (PEG) substituents for increased solubility in solvents such as water, DMF and DMSO. Two different PEG units were used in this study, with the shorter being three and the longer being eight ethylene glycol units, PEG₃–PDI and PEG₈–PDI, respectively (Figure 1). PEG₃–PDI is less soluble in aqueous and organic

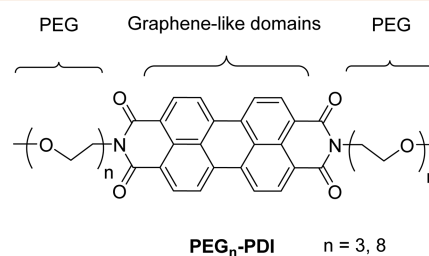


Figure 1. Structural formula of PEG_n–PDI.

solvents; however, it is more suitable for crystallization and characterization in the solid state, while PEG₈–PDI has better solubility therefore better suitability for characterization in the solution state. Note that with PEG–HCCs, 5000 MW PEG was used in order to make the nanoparticles soluble in phosphate buffered saline.³³ Details of the synthetic procedure for preparation of PEG_n–PDIs and structural characterization by NMR and X-ray crystallography can be found in the Supporting Information.

The electrochemical behavior of PDI is well-known and the cyclic voltammetry (CV) shows two reversible redox peaks in both neutral organic and aqueous media.³⁴ In DMSO, PEG₈–PDI gave two redox values (E_1 and E_2) at -0.88 V and -1.12 V versus Fc/Fc^+ (Figure S1). Both redox potentials of PEG₈–PDI are more positive than the redox peak of $O_2^{\bullet-}$ ($E = -1.25\text{ V}$ versus Fc/Fc^+ in DMSO)²⁴ by 0.37 and 0.13 V , respectively. This indicates that single-electron oxidation of $O_2^{\bullet-}$ to O_2 by both PEG₈–PDI and PEG₈–PDI^{•-} are thermodynamically favorable and exothermic processes, which is also the first half-reaction of the SOD catalytic cycle. The HCC carbon core of the PEG–HCC is the electron deficient redox center of the carbon nanoparticles, and it has a broad reduction potential with the initial onset being more positive by 0.60 V than the reduction peak of $O_2^{\bullet-}$ in aqueous media.²⁴ This indicates that PEG₈–PDI has comparable thermodynamic ability to oxidize $O_2^{\bullet-}$ to the oxidized graphene analogue PEG–HCC.

To further assess the reactivity of PEG₈–PDI with $O_2^{\bullet-}$, we monitored the optical spectral changes upon the addition of

$\text{O}_2^{\bullet-}$ to the solution of PEG₈-PDI under inert atmosphere. Figure 2 (in red) shows the optical spectrum of 0.04 mM

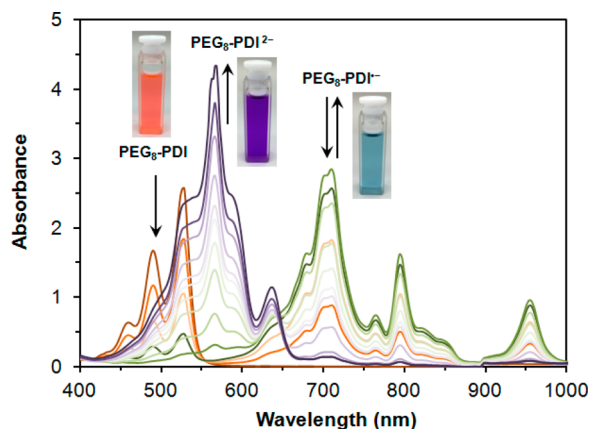
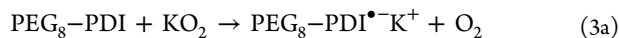


Figure 2. Spectral changes upon treatment of a 0.04 mM DMSO solution of PEG₈-PDI after the addition of a DMSO solution of KO₂ in small increments (0.0114 mM). Red: PEG₈-PDI, greenish-blue: PEG₈-PDI^{•-}, purple: PEG₈-PDI^{2•-}.

PEG₈-PDI (Figure S2) in DMSO with the major band at $\lambda = 525$ nm ($\epsilon = 6.38 \times 10^4$ M⁻¹ cm⁻¹), typical for PDI derivatives in various solvents.³⁵ Upon addition of KO₂ in small increments (0.0114 mM), the solution of PEG₈-PDI shows a gradual change in color from red to greenish-blue, which corresponds to the formation of an equimolar amount of the one-electron reduced PEG₈-PDI^{•-}, with the major band at $\lambda = 710$ nm ($\epsilon = 7.20 \times 10^4$ M⁻¹ cm⁻¹). Optical spectra of PDI radical anions are known and well-documented.^{34,36} The gradual addition of another equivalent of KO₂ results in the further change in absorbance of PEG₈-PDI^{•-} to a purple color, with the major band at $\lambda = 568$ nm ($\epsilon = 1.08 \times 10^5$ M⁻¹ cm⁻¹), which corresponds to the two-electron reduced PEG₈-PDI^{2•-} (Figure 2).

Interestingly, addition of extra amounts of KO₂ to the resultant dark-purple colored solution of PEG₈-PDI^{2•-} does not result in any further changes. This is the evidence that PEG₈-PDI^{2•-} is persistent in the presence of KO₂ and does not exhibit additional reduction steps. The two isosbestic points at $\lambda = 559$ nm and then $\lambda = 648$ nm indicate the two successive single-step chemical processes. Therefore, these optical observations provide evidence that oxidation of $\text{O}_2^{\bullet-}$ to O₂ in DMSO can be achieved by both PEG₈-PDI and PEG₈-PDI^{•-} through the consecutive single electron-transfer mechanisms, according to eq 3.



The other evidence for eq 3 is provided by electrochemical measurements. CV of PEG₈-PDI exhibits two successive redox peaks (Figure 3), which corresponds to two consecutive PEG₈-PDI/PEG₈-PDI^{•-} and PEG₈-PDI^{•-}/PEG₈-PDI^{2•-} couples.

In addition to the color changes of the solution, subsequent addition of incremental amounts of KO₂ to the solution of PEG₈-PDI gives rise to the new CV curve centered at $E = -1.23$ V versus Fc/Fc⁺, as shown in Figure 3. The new peak at $E = -1.23$ V corresponds to the CV of dissolved molecular O₂, as supported by the disappearance of the peak upon bubbling

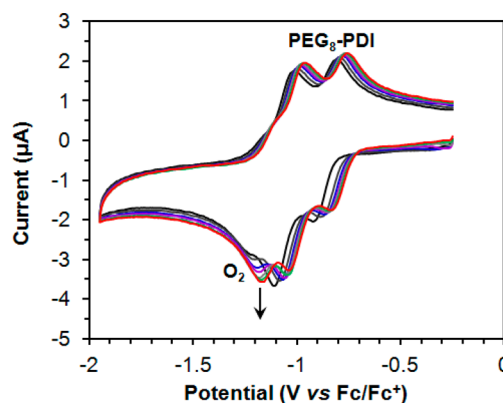


Figure 3. CVs of 10 mM of PEG₈-PDI in DMSO under N₂ after adding incremental amounts of KO₂ (20 μL of KO₂ in DMSO, 0.0114 mM stock) containing of 0.1 M [(*n*-Bu)₄N]ClO₄ as a supporting electrolyte at 298 K with a glassy carbon working electrode and platinum wire as quasi-reference electrode. Scan rate: 50 mV/s. The arrow indicates the newly formed redox peak of the molecular O₂. Color code: KO₂ concentration increased from black to red.

the solution with N₂ for 5 min. Conversely, the peak of dissolved KO₂ in electrolyte without PEG₈-PDI present is persistent for 6 h while bubbling with N₂ (Figure S3).²⁴ These electrochemical tests show that both PEG₈-PDI and PEG₈-PDI^{•-} can oxidize $\text{O}_2^{\bullet-}$ to O₂ through a single electron-transfer reaction (eq 3) and the product, molecular O₂, can be observed electrochemically. Furthermore, the reversible two consecutive 1e⁻ redox peaks of PEG₈-PDI upon addition of KO₂ nicely reveal the underlying mechanism of outer-sphere ET from $\text{O}_2^{\bullet-}$ to PEG₈-PDI and PEG₈-PDI^{•-}, where $\text{O}_2^{\bullet-}$ is a reductant. Also, the symmetric size of the current on the reductive and oxidative transition of PEG₈-PDI in the presence of $\text{O}_2^{\bullet-}$ confirms the simple interpretation of the outer-sphere ET mechanism.³⁷

To examine the SOD-like activity of PEG₈-PDI, and to investigate whether the second-half reaction of SOD proceeds analogous to eq 2, we performed additional experiments to determine the reactivity of PEG₈-PDI^{•-} and PEG₈-PDI^{2•-} with $\text{O}_2^{\bullet-}$ in the presence of protons (HClO₄) in the aprotic environment of DMSO. Both PEG₈-PDI^{•-} and PEG₈-PDI^{2•-} served as reductants of $\text{O}_2^{\bullet-}$ to H₂O₂ in the presence of 2 equiv of H⁺. But upon incremental addition of substoichiometric amounts of HClO₄ in DMSO, the purple solution of PEG₈-PDI^{2•-} in the presence of KO₂ gradually changed color to red (Figure 4). Through analysis of UV/vis spectra taken immediately after the addition of HClO₄, the recovery of the initial PEG₈-PDI was achieved in 70% yield (Figure 4). The loss of the remaining 30% suggests that the PEG₈-PDI, PEG₈-PDI^{•-} or PEG₈-PDI^{2•-} could be thermodynamically unstable in the presence of H₂O₂, which is the product of the second-half of the dismutase according to eqs 2, 4 and 5.

A separate experiment where PEG₈-PDI is treated with H₂O₂ and HClO₄ does not exhibit any changes in the optical spectra (Figure S4), suggesting the persistent nature of PEG₈-PDI under these conditions, thus precluding reaction of PEG₈-PDI with H₂O₂ as a reason for the depressed yield. The fact that PEG₈-PDI^{2•-} rapidly reacts according to eq 4 and immediately reforms stoichiometric amounts of PEG₈-PDI^{•-} as the product (Figure 4), also excludes PEG₈-PDI^{2•-} from being the source for the decreased yields. However, PEG₈-

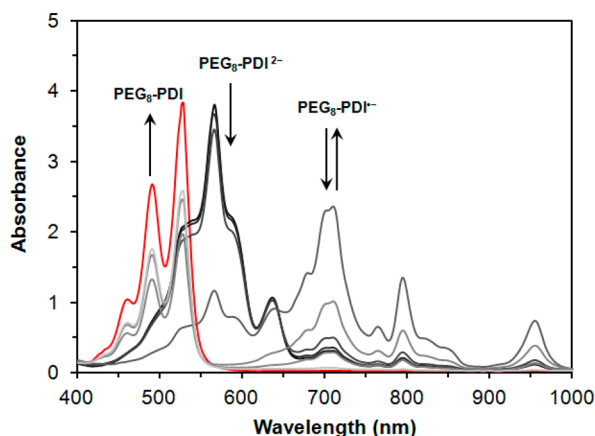
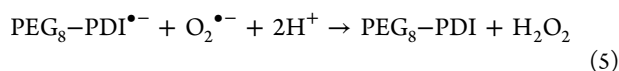
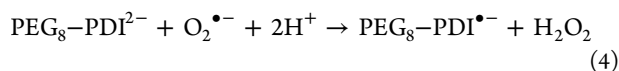


Figure 4. Absorption spectra of PEG₈-PDI²⁻ in DMSO (black curve) upon treatment with KO₂ and HClO₄ indicating the formation of H₂O₂ and regeneration of original neutral PEG₈-PDI (red curve) proceeding via the PEG₈-PDI^{•-} (gray curve).

PDI^{•-} reacts irreversibly with H₂O₂ and gradually forms a product with an optical spectrum uncharacteristic of PEG₈-PDI (Figure S5). This explains the loss of the 30% of the original PEG₈-PDI during the second half of the SOD-mimetic step, where PEG₈-PDI^{•-} reacts with produced H₂O₂ via efficient PCET reaction mechanism as shown in eq 6.^{38,39}



Thus, these optical data provide evidence that PEG₈-PDI mimics the catalytic cycle of SOD.

The kinetics of the reaction between O₂^{•-} and PEG₈-PDI was measured using KO₂ and a direct freeze-trap EPR steady-state kinetics assay rather than the less efficient spin-trapping EPR which also loses kinetic information. 18-Crown-6 was used to increase the solubility of KO₂ in DMSO. In this case excess O₂^{•-} was used to estimate the intrinsic turnover number for O₂^{•-} conversion to O₂ by PEG₈-PDI. In order to slow down

the self-dismutation of superoxide in aqueous media to achieve reasonable initial concentrations of O₂^{•-} in solution, we carried out the kinetic experiments at pH 13 in 50 mM NaOH. PEG₈-PDI is stable at pH 13 as there is no change of optical spectral shape observed by 30 min preincubation. A control EPR sample was prepared using this preincubated PEG₈-PDI to react with 20 mM KO₂ and freeze trapped. The amount of radical determined is basically the same as that of fresh PEG₈-PDI.

Figure 5a shows that the total spin concentration of O₂^{•-} gradually increased with the amount of added KO₂. As expected, the concentration of O₂^{•-} decreased in the presence of PEG₈-PDI. Therefore, the consumption of O₂^{•-} by PEG₈-PDI can be estimated from the difference between the control, which does not contain any PEG₈-PDI, and subsequently recalculated as turnover numbers (defined as number of moles of consumed O₂^{•-} per moles of PEG₈-PDI per second) with an average reaction time of 10 s (Figure 5b). At the saturation behavior of [O₂^{•-}] the highest O₂^{•-} turnover rate was estimated as 133 s⁻¹, which is almost 1000× lower than that of PEG-HCCs.²³ This is not surprising, as it is possible that the redox potential of the two half reactions for PEG₈-PDI are not as optimal as PEG-HCCs²⁴ or the electron transfer kinetic barrier is higher than that of the PEG-HCCs. Overall, the kinetics experiments indicate that PEG₈-PDI behaves similarly to SOD and PEG-HCCs as a “hit-and-run” type reaction, where O₂^{•-} substrate only momentarily stays in the active center, so the rate-limiting step is not the binding but the oxidation/reduction of the O₂^{•-}.

Isolation and Characterization of PEG_n-PDI Radical Anions. PEG₈-PDI^{•-} as a reaction intermediate was additionally characterized by EPR spectroscopy. As with the optical measurements, PEG₈-PDI^{•-} was generated using stoichiometric (1:1) amounts of KO₂ as a reductant under inert atmosphere and it shows an EPR spectrum ($g = 2.0019$) with apparent 18 line hyperfine splitting caused by nitrogens and protons (Figure 6). Simulation of the EPR spectrum reveals the delocalization of unpaired spin over the entire π -conjugated unit of PDI with the following hyperfine splitting: ring protons ($n = 8$): 1.2 G; side chain protons ($n = 4$): 0.6 G; nitrogens ($n = 2$): 0.04 G, line width 0.23 G (Figures S6). The shape of the EPR spectra does not change substantially with temperature, except slight broadening of the lines when the temperature is lowered from 295 to 260 K. This is due to increased viscosity of

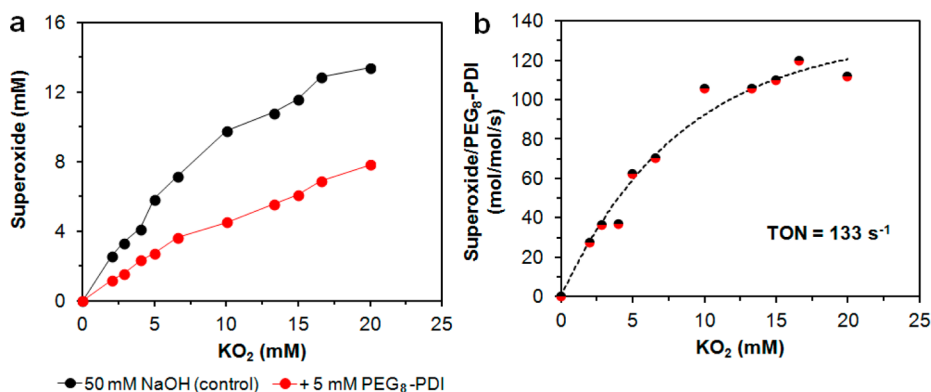


Figure 5. KO₂ experiments in 50 mM NaOH. (a) O₂^{•-} total spin concentration based on double integration of obtained EPR spectra. For details, see Experimental Methods. (b) The turnover number (TON) was calculated by subtracting the amount of remaining O₂^{•-} from the amount of O₂^{•-} in the control, dividing by the amount of PEG₈-PDI and then by 10 s for the reaction time (the average time required for manually freeze-trapping each EPR sample). Two similar experiments performed using 1.5 μM PEG₈-PDI concentration yielded similar kinetic pattern.

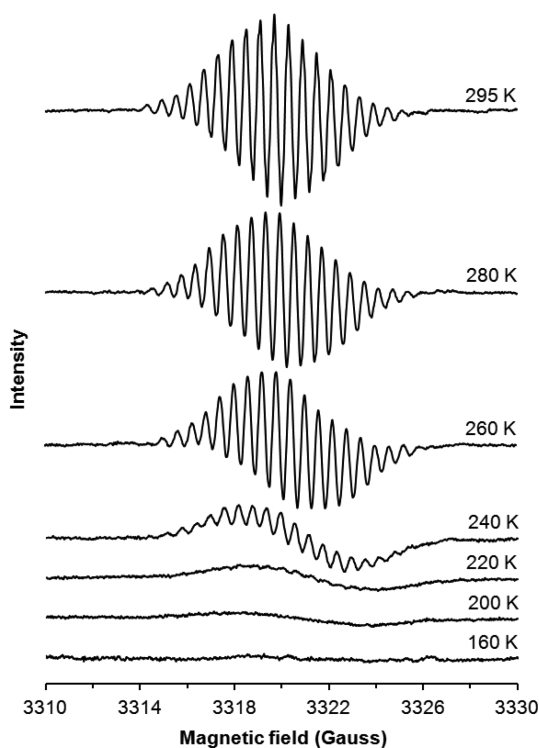
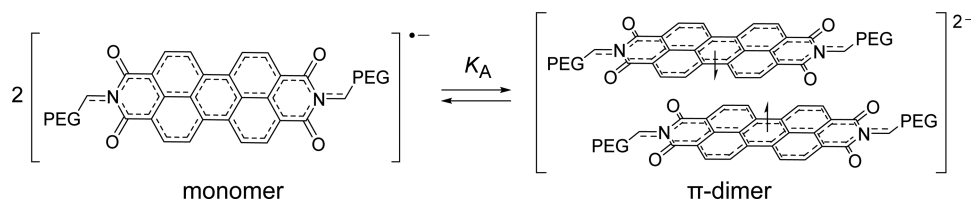


Figure 6. Variable-temperature EPR spectra of $\text{PEG}_8\text{-PDI}^{\bullet-}$ obtained by reduction of $\text{PEG}_8\text{-PDI}$ with equimolar amounts of KO_2 in DMF.

the solvent which could lead to changes in the electron spin distribution and variation in coupling constants (Figure S7). However, further decrease in temperature of the sample results in decrease of intensity and eventually loss of the EPR signal at 160 K as can be seen from Figure 6. π -Dimerization of the stable radical ions to form weakly bonded dimeric species has been frequently considered over the past decade.^{40–44}

Similarly, for $\text{PEG}_8\text{-PDI}^{\bullet-}$ the signal intensity decreases upon cooling the sample with the gradual replacement of the paramagnetic monomer by the diamagnetic π -dimer (Scheme 2). Intensity of signal (I_{EPR}) and concentrations were determined by double integration of the spectra. According to the Curie law, $I_{\text{EPR}} = a \times [M]/T$, where a is a proportionality factor, I_{EPR} is proportional to the concentration of the radical ($\text{PEG}_8\text{-PDI}^{\bullet-}$) in the sample. Assuming that at room temperature the equilibrium (Scheme 2) shifts to the left and only monomeric radical anions exist in solution, and using the proportionality ratio $I_{\text{EPR}}/I_{\text{EPR}}^{298} = [M]/[M]^{298} = \alpha$ where α is a mole fraction of the monomeric radical anion at a particular temperature, the equilibrium concentration of the monomeric radical anion can be estimated at each temperature. By using $K_A = (1 - \alpha)/2[M]^{298}\alpha^2$ with an estimated α , it permits calculation of the equilibrium constants for π -dimerization at each

Scheme 2. Reversible π -Dimerization of $\text{PEG}_n\text{-PDI}^{\bullet-}$



temperature. Thermodynamic parameters for π -dimerization of the monomeric radical anions were calculated by the least-squares procedure from the linear relationship of $\ln(K_A)$ and $1/T$, resulting in the analysis pictured in Figures S8.

The thermodynamic parameters ΔH_D and ΔS_D can be estimated as -7.7 kcal/mol and -17.1 e.u. respectively, which are close to the values of thermodynamic parameters for π -dimerization in most of the extended π -radicals.⁴¹ The strong attraction of two unpaired electrons prevails over electrostatic repulsion of two oppositely charged anions.

Interestingly, a single crystal of the $\text{PEG}_3\text{-PDI}^{\bullet-}$ was obtained using cobaltocene as a reducing agent. A green solution of the $[\text{PEG}_3\text{-PDI}^{\bullet-} \text{CoCp}^+]$ in DMF was layered carefully with dry ethyl ether and left in a freezer at -10 °C for 1 week and small dark-green-colored crystalline needles were collected and subjected to X-ray crystallographic analysis. Crystallographic analysis resulted in infinite π -stacked PDI units of $\text{PEG}_3\text{-PDI}^{\bullet-}$ along the b -axis with close π - π interactions of $3.36(2)$ Å as shown in Figure 7.

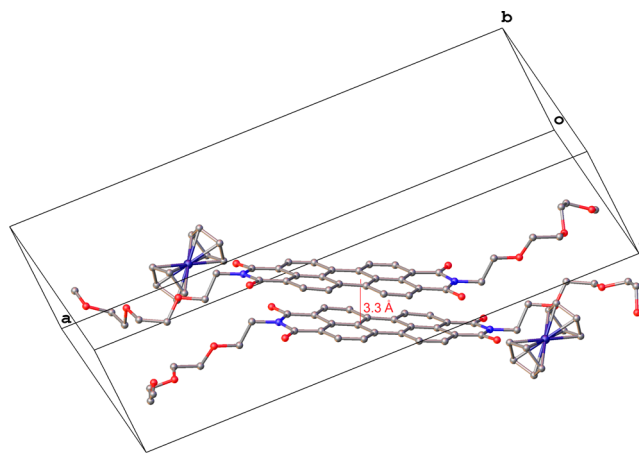


Figure 7. View of the unit cell of the crystal structure of $[\text{PEG}_3\text{-PDI}^{\bullet-} \text{CoCp}^+]$. Solvents and hydrogen atoms are omitted for clarity. Color code: C, gray; N, blue; O, red; Co, purple.

ORR Activity of PDIs. Knowing the properties of PDI within $\text{PEG}_n\text{-PDIs}$ to be efficient electron shuttle agents during the catalytic turnover of $\text{O}_2^{\bullet-}$, and also that PDIs have redox potentials close to that of molecular O_2 , we investigated the electrocatalytic properties of PDIs for their oxygen reduction reaction (ORR) activity. Despite the fact that development of molecular catalysts for electrochemical reduction of O_2 has been an active research field, many of the studied catalysts are high-valent organometallic complexes while examples of pure organic molecules are scarce.^{45–47} A water-insoluble derivative of PDI, bis-2-ethylhexyl PDI (PDI-1, Figure S9) was immobilized on a glassy carbon (GC) working electrode which served as an O_2 -electrode, and it was tested in O_2 -

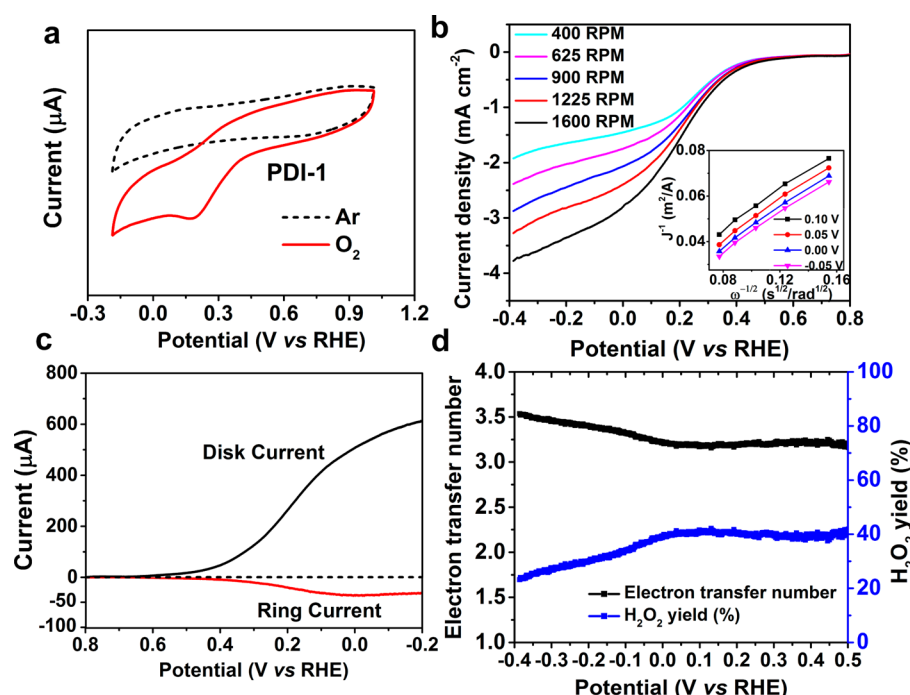
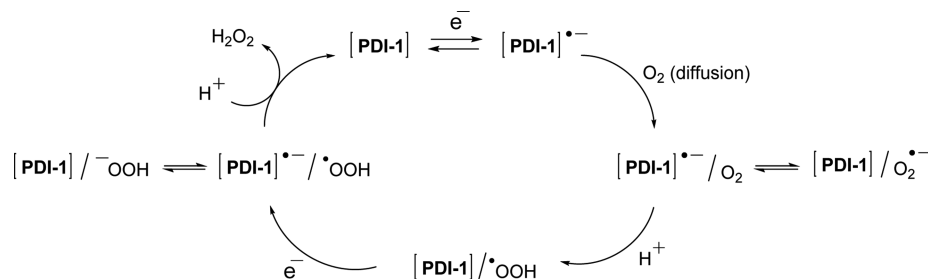


Figure 8. (a) CVs of PDI-1 under 1 atm O_2 or Ar. All scans were collected at 100 mV s^{-1} using a GC working electrode with an area of 0.196 cm^2 . (b) LSV curves of RDEVs of PDI-1 in O_2 -saturated $0.1 \text{ M NaH}_2\text{PO}_4/\text{NaH}_2\text{PO}_4$ buffer solution at pH 7 with different rotating speeds ranging from 400 to 1600 rpm. Inset: Koutecky–Levich plots of PDI-1 showing that $n = 1.9$. (c) RRDEV of PDI-1 in $0.1 \text{ M NaH}_2\text{PO}_4/\text{NaH}_2\text{PO}_4$ buffer solution at pH 7 with a rotation speed 1600 rpm. The disk was scanned from 0.8 to -0.2 V while the ring electrode was held at 1.4 V . (d) The number of electrons transferred and the H_2O_2 yield of PDI-1 during ORR as calculated by ring currents.

Scheme 3. Oxygen Reduction Reaction Mechanism by PDI-1 to form H_2O_2



saturated $0.1 \text{ M NaH}_2\text{PO}_4/\text{NaH}_2\text{PO}_4$ buffer solution at pH 7. The comparison spectra testing under both Ar and O_2 reveal that under Ar there is a very small reduction peak (Figure 8a). In contrast, PDI-1 under O_2 -saturated conditions results in a prominent cathodic ORR peak with an onset potential of $\sim 0.40 \text{ V}$ vs RHE (Figure 8a). Linear sweep voltammetry (LSV) plots were obtained by using a rotating disc electrode (RDE) with varying rotation rates to show the ORR electrocatalytic currents of PDI-1 (Figure 8b). Using the Koutecky–Levich (K–L) equation, the kinetic current density (J_k) was obtained and the electron-transfer number (n , number of electrons exchanged per O_2 molecule for the ORR) was estimated as shown in inset of Figure 8b.⁴⁸ The n value was determined to be 1.9 at a potential range from -0.20 to 0.20 V . This demonstrates that the complete two-electron transfer reduction process of oxygen to H_2O_2 proceeds according to eq 7.



The ORR activity of PDI-1 is also evident from the comparison in electrocatalytic reduction currents of the GC substrate with the PDI-1 immobilized electrode (Figure S10). Higher ORR

currents and larger differences in the onset potential for the PDI-1-covered electrode than for the naked GC electrode reveal that interference of the GC substrate in ORR performance is negligible, in spite of the fact that O_2 can be reduced onto a GC electrode.⁴⁹

These results show that there is a two-electron ORR activity of PDIs in aqueous media, as opposed to the single-electron transfer process in organic solvents. However, the stability of PDI-1 shows a small decline under ORR conditions (Figure S11), likely due to the production of aggressive radicals and H_2O_2 during ORR, which is likely to passivate the electrode surface over extended cycling, similar to the H_2O_2 -mediated decomposition of $\text{PEG}_8\text{-PDI}^{\bullet-}$ discussed above.

The accurate determination of n , using the classic Koutecky–Levich analysis shown above, depends on many factors, including determination of the active surface area. Hence, we performed additional electrochemical tests to support the ORR activity of PDI-1. The yield of H_2O_2 and the potential range of ORR operation were further measured by rotating disc electrode voltammetry (RDEV) and rotating ring-disk electrode voltammetry (RRDEV) as shown in Figures 8c,d.

Significant increase in cathodic disk current was observed starting at 0.4 V, while the disk potential was scanned from 0.8 to -0.2 V at a constant ring electrode potential of 1.4 V. This is consistent with the voltammograms in Figures 8a,b. Symmetric increase in ring current as in that of the disk current (Figure 8c) indicates catalytic formation of H_2O_2 generated at the disk.⁵⁰ Using the differences in the values between disk and ring currents, the value for n and the yield of H_2O_2 were estimated to be 3.0 and 40%, respectively (see Experimental Methods for the details of the calculations). The value for n and the yield of H_2O_2 during the electrocatalytic ORR by PDI-1 indicates that there is nearly equal contribution of 2-electron and 4-electron reductions of O_2 to H_2O_2 and H_2O , respectively.

On the basis of the experimental data shown above, the general PDI-1-catalyzed two-electron-transfer oxygen reduction reaction can be described as an electrochemical-chemical-electrochemical-chemical (ECEC) type mechanism shown in Scheme 3.⁵¹ The initial electron transfer step generates $[\text{PDI-1}]^{\bullet-}$, which complexes with diffused O_2 to form transient single-electron reduced complex $[\text{PDI-1}]/\text{O}_2^{\bullet-}$. That is readily protonated to generate hydroperoxyl radical intermediate, $[\text{PDI-1}]/\bullet\text{OOH}$. $\bullet\text{OOH}$ is a stronger oxidant, therefore formation of $[\text{PDI-1}]/\bullet\text{OOH}$ is followed by the faster second ET-step to form $[\text{PDI-1}]/\text{OOH}$, where the second proton transfer step leads to H_2O_2 as a final product with reformation of $[\text{PDI-1}]$.

As in most electrochemical reactions, according to the mechanistic pathway shown in Scheme 3 and the formation of H_2O_2 as a product during the ORR process, proton transfer steps from reaction intermediate $[\text{PDI-1}]^{\bullet-}/\text{O}_2$ to $[\text{PDI-1}]^{\bullet-}/\bullet\text{OOH}$ are much faster than the heterogeneous electron transfer steps.⁵¹ Furthermore, the two single-electron transfer steps are the rate-determining steps of the ORR process. Finally, in this work we were able to show that under the electrochemical conditions, PDIs can serve as efficient electron shuttles to reduce O_2 to $\text{O}_2^{\bullet-}$ and H_2O_2 .

CONCLUSION

In summary, preparation and characterization of small well-defined conjugated molecular analogues, PEG-PDIs, as a model to elucidate the mechanisms operative in the SOD-mimetic activity of PEG-HCCs, was accomplished. Water-soluble perylene diimide derivatives serve as molecular analogues of PEG-HCCs that mimic the full catalytic cycle of SOD. PEG_{*n*}-PDI was able to oxidize $\text{O}_2^{\bullet-}$, and the PEG_{*n*}-PDI^{•-} intermediate of the catalytic cycle is thoroughly characterized and shown to react with $\text{O}_2^{\bullet-}$ in the presence of protons to form H_2O_2 . Having the redox potential of PEG_{*n*}-PDIs located between the two half reactions, $\text{O}_2^{\bullet-} \rightarrow \text{O}_2 + e^-$ and $\text{O}_2^{\bullet-} + 2 \text{H}^+ + 2 e^- \rightarrow \text{H}_2\text{O}_2$ (-0.16 and $+0.94$ V, respectively, relative to NHE in water), makes them good molecular SOD-mimetics of PEG-HCCs. The freeze quench EPR study substantiated its multiple turnover of $\text{O}_2^{\bullet-}$ in aqueous environment. We show that study of graphene-based catalysts can benefit from the study precise molecular analogues,^{52,53} and PDI serves as a minimal model of HCCs and more generally oxidized graphenes. Furthermore, on the basis of the results from the RDEV and RRDEV, PDI was shown to be a metal-free molecular electrocatalyst for the O_2 reduction reaction with H_2O_2 being produced in 40% yield. Thus, the results indicate that similar to carbon nanomaterials, PDI is an efficient electron shuttle for reactions with O_2 as well as ROS. These results have important implications in the area

of graphene-based materials used in electrocatalytic processes and their potential application in nanoantioxidants.

EXPERIMENTAL METHODS

Materials and Methods. All chemicals were purchased from Sigma-Aldrich and used without further purification unless otherwise stated. Optical spectra were acquired on a Shimadzu 50 Scan UV/vis spectrometer (200–1100 nm) and Cary5000 UV-vis-NIR spectrophotometer (200–3000 nm), using capped quartz cuvettes. NMR data were recorded on Bruker 400–600 MHz spectrometers.

Single-Crystal X-ray Diffraction Analysis. Crystal data, details of data collection and structure refinement parameters for the $[\text{PEG}_3\text{-PDI}^{\bullet-} \text{CoCp}^+]$ are presented in Table S1 and Figure S12. Diffraction data were collected on a Rigaku SCX-Mini diffractometer (Mercury2 CCD) using graphite-monochromated Mo $K\alpha$ radiation ($\lambda = 0.71073$ Å). Integration was performed with CrystalClear-SM Expert 2.0, and the data were corrected for absorption using empirical methods. The structures were solved by direct methods and refined by the full-matrix least-squares technique against F^2 with the anisotropic temperature parameters for all non-hydrogen atoms. All H atoms were geometrically placed and refined in a rigid model approximation. Data reduction and structure refinement calculations were performed using the SHELXTL⁵⁴ program package. The description of the disorder refinement of PEG₄ groups in the PEG₃-PDI^{•-} radical anion is presented in the Supporting Information.

Electrochemistry. The CVs were obtained with a CHI1202 ElectroChemical Analyzer (CHI Instruments) for 10 mL electrolyte solutions (0.1 M $[(n\text{-Bu})_4\text{N}]\text{ClO}_4$ solution in DMSO or PBS buffer, pH 7.4) using a 3-electrode cell. A GC electrode served as working electrode, platinum wires served as a counter electrode and Ag/AgCl as the reference electrode. A platinum wire was used as the pseudoreference electrode in DMSO and ferrocene (Fc) was used as internal potential standard and all potentials are referred to the Fc/Fc⁺ couple. CVs were recorded at a scan rate of 100 mV s⁻¹.

RDE and RRDE experiments were conducted in an electrochemical cell (AUTO LAB PGSTST 302) using a Pine Instrument rotator (model: AFMSRCE) connected to a CH Instruments electrochemical analyzer (model 600D), with an Ag/AgCl reference electrode and a Pt wire counter electrode. A PDI-1 solution ink was prepared by dispersing 4 mg of the PDI-1 into 1 mL of 4:1 DCM/EtOH solvent, and 8 μL of the catalyst solution ink was loaded onto a GC electrode (5 mm in diameter). A constant bubbling by a stream of O_2 in the cell solution was maintained throughout the measurement to ensure continuous O_2 saturation. Measurements were carried out at pH 7 (0.1 M $\text{K}_2\text{HPO}_4/\text{KH}_2\text{PO}_4$ buffer). For RRDE experiments, the electrode rotation speed was 1600 rpm (scan rate, 0.05 V/s; platinum data collected from anodic sweeps), while the ring electrode potential was held at 1.1 V vs a reversible hydrogen electrode (RHE).

The O_2 reduction current increases with increasing rotation rates following the K-L eq 8:

$$J^{-1} = J_K^{-1} + J_L^{-1} \quad (8)$$

where J_K is the potential dependent kinetic current and J_L is the Levich current. J_L is expressed as $0.62nF[\text{O}_2](D_{\text{O}_2})^{2/3}\omega^{1/2}\nu^{-1/6}$, where n is the number of electrons transferred to the substrate, F is the Faraday constant, $[\text{O}_2]$ is the concentration of O_2 in an air-saturated buffer (0.22 mM in case of pH 7) at 25 °C, D_{O_2} is the diffusion coefficient of O_2 ($1.8 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ at pH 7) at 25 °C, ω is the angular velocity of the disc and ν is the kinematic viscosity of the solution ($0.009 \text{ cm}^2 \text{ s}^{-1}$) at 25 °C. Eq 8 can be restated as eq 9; solving for J_L gives eq 10:

$$1/J = 1/J_K + 1/J_L \quad (9)$$

$$J_L = 0.62nF[\text{O}_2](D_{\text{O}_2})^{2/3}\omega^{1/2}\nu^{-1/6} \quad (10)$$

RRDE measurements were carried out to determine the H_2O_2 yield (%) and n , which were calculated by eqs 11 and 12:

$$\text{H}_2\text{O}_2\% = [200i_r/N]/[i_d + i_r/N] \quad (11)$$

$$n = 4i_d/[i_d + i_r/N] \quad (12)$$

where i_d and i_r are the disk and ring currents, respectively. N is the ring current collection efficiency, which was determined to be 25% by the reduction of 10 mM $K_3[Fe(CN)_6]$ in 0.1 M KNO_3 .

Detection of Radicals by EPR Spectroscopy. The EPR spectra of PEG-PDI^{•-} (0.05 mg/mL in DMSO) in a sealed capillary tube at ambient temperature were recorded using the following parameters: center field 3320 G, sweep width 50 G, microwave frequency 9.3 GHz, microwave power 1 mW, modulation frequency 100 kHz, and modulation amplitude 1.0 G. The same sample was remeasured after adding a small amount of KO_2 (1 mg in powder form).

Steady-State Consumption of Superoxide. PEG₈-PDI (5 μ M) in 50 mM NaOH were mixed in ratio 1:5 with increasing amounts of KO_2 (dissolved in DMSO/18-crown-6) for 10 s and then frozen in ethanol/dry Ice (-72 °C) to stop the reaction. The samples were then transfer to LN_2 to preserve the remaining $O_2^{\bullet-}$. EPR spectra were then recorded. To account for background dismutation of $O_2^{\bullet-}$, a sample lacking PEG₈-PDI was measured and its EPR spectra subtracted from sample spectra to obtain the amount of KO_2 decay catalyzed by the PEG₈-PDI.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsnano.6b08211.

Additional graphs and data, single-crystal X-ray diffraction analysis details of [PEG₃-PDI^{•-} CoCp⁺] (PDF)

AUTHOR INFORMATION

Corresponding Authors

*E-mail: ah-lim.tsai@uth.tmc.edu.

*E-mail: tour@rice.edu.

ORCID

James M. Tour: 0000-0002-8479-9328

Notes

The authors declare the following competing financial interest(s): Rice University owns intellectual property on the use of small molecule graphene analogs for use as antioxidants in medicine. The intellectual property has been licensed to Acelerox LLC. J. M. T. and T. A. K. are stockholders in Acelerox, though not an officer or director. Conflicts are managed through disclosures to the Rice University Office of Sponsored Programs and Research Compliance (SPARC) and the Baylor College of Medicine.

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