Perylene Diimide as a Precise Graphene-Like Superoxide Dismutase Mimetic

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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 analogs 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 analogs of PEG–HCCs and oxidized graphenes in general, and
their antioxidant and superoxide dismutase-like (SOD-like) properties were studied. PEGₙ–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, PEGₙ–PDIs are also strong single-electron oxidants of O₂⁻. Furthermore, reduced PEGₙ–PDI, PEGₙ–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 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, \( \text{O}_2^{\cdot-} \) 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 \( 2 \text{O}_2^{\cdot-} \) into \( \text{O}_2 \) and \( \text{H}_2\text{O}_2 \): \( 2 \text{O}_2^{\cdot-} + 2 \text{H}^+ \rightarrow \text{O}_2 + \text{H}_2\text{O}_2. \)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 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 \( \text{M}−\text{O}_2^{\cdot-} \) 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), that show highly efficient catalytic conversion of \( \text{O}_2^{\cdot-} \) into \( \text{O}_2 \) and \( \text{H}_2\text{O}_2 \) (eqs 1 and 2) rivals that of bovine Cu/Zn SOD, the most efficient native SOD enzyme.²³

Scheme 1. Synthesis of PEG-HCCs from SWCNTs as the starting material.
Single-walled carbon nanotubes (SWCNTs) are oxidatively disintegrated and split open to form graphene domains that are 3 nm × 35 nm with large amounts of oxidized functionalities that make the carbon system extremely electron deficient. 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. The HCCs also bear, on average, one unpaired electron per nanoparticle and the nanoparticle radical is air stable. As we have shown previously, HCCs can efficiently oxidize $O_2^-$ 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. This is followed by reduction of $O_2^-$ in the presence of water to form H$_2$O$_2$ (eq 2).

$$\text{PEG−HCC}^+ + O_2^- \rightarrow \text{PEG−HCC}^- + O_2 \quad (1)$$

$$\text{PEG−HCC}^- + O_2^- + 2 H^+ \rightarrow \text{PEG−HCC}^+ + H_2O_2 \quad (2)$$

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), 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 $O_2^-$ 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 analogs, and we propose that PDI serves as a simplified model of oxidized graphenes.
Results and Discussion

**PEGₙ-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 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 PEGₙ−PDIs and structural characterization by NMR and X-ray crystallography can be found in the Supporting Information.

![Figure 1. Structural formula of PEGₙ−PDI.](image)

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 (E₁ and E₂) at −0.88 V and −1.12 V versus Fe/Fe⁺ (Figure
S1). Both redox potentials of PEG₈−PDI are more positive than the redox peak of O₂⁻ (\(E = -1.25\) V \textit{versus} Fe/Fc⁺ in DMSO)²⁴ by 0.37 V and 0.13 V, respectively. This indicates that single-electron oxidation of O₂⁻ to O₂ 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 O₂⁻ in aqueous media.²⁴ This indicates that PEG₈−PDI has comparable thermodynamic ability to oxidize O₂⁻ to the oxidized graphene analogue PEG-HCC.

To further assess the reactivity of PEG₈−PDI with O₂⁻, we monitored the optical spectral changes upon the addition of O₂⁻ to the solution of PEG₈−PDI under inert atmosphere. Figure 2 (in red) shows the optical spectrum of 0.04 mM PEG₈−PDI (Figure S2) in DMSO with the major band at \(\lambda = 525\) nm (\(\varepsilon = 6.38 \times 10^4\) 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 (\(\varepsilon = 7.20 \times 10^4\) 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 (\(\varepsilon = 1.08 \times 10^5\) M⁻¹ cm⁻¹), which corresponds to the two-electron reduced PEG₈−PDI²⁻ (Figure 2).
Figure 2. Spectral changes upon treatment of a 0.04 mM DMSO solution of PEG$_8$–PDI after the addition of a DMSO solution of KO$_2$ in small increments (0.0114 mM). Red: PEG$_8$–PDI, greenish-blue: PEG$_8$–PDI$^+$, purple: PEG$_8$–PDI$_2^-$.

Interestingly, addition of extra amounts of KO$_2$ to the resultant dark-purple colored solution of PEG$_8$–PDI$_2^-$ does not result in any further changes. This is the evidence that PEG$_8$–PDI$_2^-$ is persistent in the presence of KO$_2$ 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 O$_2^-$ to O$_2$ in DMSO can be achieved by both PEG$_8$–PDI and PEG$_8$–PDI$^-$ through the consecutive single electron-transfer mechanisms, according to eq 3.

\[
\text{PEG}_8\text{–PDI} + \text{KO}_2 \rightarrow \text{PEG}_8\text{–PDI}^- \text{K}^+ + \text{O}_2 \quad (3a)
\]
\[
\text{PEG}_8\text{–PDI}^- \text{K}^+ + \text{KO}_2 \rightarrow \text{PEG}_8\text{–PDI}_2^- 2\text{K}^+ + \text{O}_2 \quad (3b)
\]
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−PDI²⁻ couples.

![Figure 3](image_url)

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.

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 \( \text{O}_2 \), as supported by the disappearance of the peak upon bubbling the solution with \( \text{N}_2 \) for 5 min. Conversely, the peak of dissolved \( \text{KO}_2 \) in electrolyte without PEG₈−PDI present is persistent for 6 h while bubbling with \( \text{N}_2 \) (Figure S3).²⁴ These electrochemical tests show that both PEG₈−PDI and PEG₈−PDI⁻ can oxidize \( \text{O}_2⁻ \) to \( \text{O}_2 \) through a single electron-transfer reaction (eq 3) and the product, molecular \( \text{O}_2 \), can be observed electrochemically. Furthermore, the reversible two consecutive 1e⁻ redox peaks of PEG₈−PDI upon addition of \( \text{KO}_2 \) nicely reveal the underlying mechanism of outer-sphere ET from \( \text{O}_2⁻ \) to PEG₈−PDI and PEG₈−PDI⁻, where \( \text{O}_2⁻ \) is a reductant. Also, the symmetric size of the current on the reductive and oxidative transition of PEG₈−PDI in the presence of \( \text{O}_2⁻ \) 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−PDI²⁻ with \( \text{O}_2⁻ \) in the presence of protons (\( \text{HClO}_4 \)) in the aprotic environment of DMSO. Both PEG₈−PDI⁻ and PEG₈−PDI²⁻ served as reductants of \( \text{O}_2⁻ \) to \( \text{H}_2\text{O}_2 \) in the presence of 2 equivalents of \( \text{H}⁺ \). But upon incremental addition of substoichiometric amounts of \( \text{HClO}_4 \) in DMSO, the purple solution of PEG₈−PDI²⁻ in the presence of \( \text{KO}_2 \) gradually changed color to red (Figure 4). Through analysis of UV/Vis spectra taken immediately after the addition of \( \text{HClO}_4 \), 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−PDI²⁻ could be thermodynamically unstable in the presence of \( \text{H}_2\text{O}_2 \), which is the product of the second-half of the dismutase according to eqs 2, 4 and 5.
Figure 4. Absorption spectra of PEG$_8$−PDI$^{2-}$ in DMSO (black curve) upon treatment with KO$_2$ and HClO$_4$ indicating the formation of H$_2$O$_2$ and regeneration of original neutral PEG$_8$−PDI (red curve) proceeding via the PEG$_8$−PDI$^-$ (gray curve).

A separate experiment where PEG$_8$−PDI is treated with H$_2$O$_2$ and HClO$_4$ does not exhibit any changes in the optical spectra (Figure S4), suggesting the persistent nature of PEG$_8$−PDI under these conditions, thus precluding reaction of PEG$_8$−PDI with H$_2$O$_2$ as a reason for the depressed yield. The fact that PEG$_8$−PDI$^{2-}$ rapidly reacts according to eq 4 and immediately reforms stoichiometric amounts of PEG$_8$−PDI$^-$ as the product (Figure 4), also excludes PEG$_8$−PDI$^{2-}$ from being the source for the decreased yields. However, PEG$_8$−PDI$^-$ reacts irreversibly with H$_2$O$_2$ and gradually forms a product with an optical spectrum uncharacteristic of PEG$_8$−PDI (Figure S5). This explains the loss of the 30% of the original PEG$_8$−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

\[
\text{PEG}_8^\cdot \text{PDI}^2^- + \text{O}_2^- + 2 \text{H}^+ \rightarrow \text{PEG}_8^\cdot \text{PDI}^- + \text{H}_2\text{O}_2 \quad (4)
\]
\[
\text{PEG}_8^\cdot \text{PDI}^- + \text{O}_2^- + 2 \text{H}^+ \rightarrow \text{PEG}_8^- \text{PDI} + \text{H}_2\text{O}_2 \quad (5)
\]
\[
\text{PEG}_8^\cdot \text{PDI}^- + \text{H}_2\text{O}_2 \rightarrow \text{PEG}_8^- \text{PDI}^- \text{OH}^- + \text{HO}^* \quad (6)
\]

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 pre-incubation. A control EPR sample was prepared using this pre-incubated 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 5. KO$_2$ experiments in 50 mM NaOH. (a) O$_2^{-}$ 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 O$_2^{-}$ from the amount of O$_2^{-}$ in the control, dividing by the amount of PEG$_8$−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 PEG$_8$−PDI concentration yielded similar kinetic pattern.

Figure 5a shows that the total spin concentration of O$_2^{-}$ gradually increased with the amount of added KO$_2$. As expected, the concentration of O$_2^{-}$ decreased in the presence of PEG$_8$−PDI. Therefore, the consumption of O$_2^{-}$ by PEG$_8$−PDI can be estimated from the difference between the control, which does not contain any PEG$_8$−PDI, and subsequently recalculated as turnover numbers (defined as number of moles of consumed O$_2^{-}$ per moles of PEG$_8$−PDI per second) with an average reaction time of 10 s (Figure 5b). At the saturation behavior of [O$_2^{-}$] the highest O$_2^{-}$ 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 PEG$_8$−PDI are not as optimal as PEG-HCCs$^{24}$ 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 PEG₈-PDI radical anions.** PEG₈–PDI as a reaction intermediate was additionally characterized by EPR spectroscopy. As with the optical measurements, 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; sidechain protons (n = 4): 0.6 G; nitrogens (n = 2): 0.04 G, linewidth 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 K to 260 K. This is due to increased viscosity of 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 last decade.⁴⁰-⁴⁴
Figure 6. The variable-temperature EPR spectra of PEG₈−PDI⁻ obtained by reduction of PEG₈−PDI with equimolar amounts of KO₂ in DMF.

Similarly, for PEG₈−PDI⁻ 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 (IₑPR) and concentrations were determined by double integration of the spectra. According to the Curie law, IₑPR = a × [M] / T, where a is a proportionality factor, IₑSR is proportional to the concentration of the radical (PEG₈−PDI⁻) 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 $I_{EPR}/I_{EPR}^{298} = [M]/[M]^{298} = \alpha$ where $\alpha$ 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 $\alpha$, it permits calculation of the equilibrium constants for $\pi$-dimerization at each temperature. Thermodynamic parameters for $\pi$-dimerization of the monomeric radical anions were calculated by the least square procedure from the linear relationship of $\ln(K_A)$ and $1/T$, resulting in the analysis pictured in Figures S8.

The thermodynamic parameters $\Delta H_D$ and $\Delta S_D$ can be estimated as $-7.7 \text{ kcal/mol}$ and $-17.1 \text{ e.u.}$ respectively, which are close to the values of thermodynamic parameters for $\pi$-dimerization in almost of the extended $\pi$-radicals. The strong attraction of two unpaired electrons prevails over electrostatic repulsion of two oppositely charged anions.

Interestingly, a single crystal of the PEG$_3$-$\text{PDI}^-$ was obtained using cobaltocene as a reducing agent. A green solution of the [PEG$_3$-$\text{PDI}^-$ CoCp$^+$] in DMF was layered carefully with dry ethyl ether and left in a freezer at $-10 \, ^\circ\text{C}$ 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 PEG$_3$−PDI$^-$ 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 [PEG$_3$−PDI$^-$ 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 PEG$_n$−PDIs to be efficient electron shuttle agents during the catalytic turnover of O$_2$$^-$, and also that PDIs have redox potentials close to that of molecular O$_2$, we investigated the electocatalytic properties of PDIs for their oxygen reduction reaction (ORR) activity. Despite the fact that development of molecular catalysts for electrochemical reduction of O$_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.\textsuperscript{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\textsubscript{2}-electrode, and it was tested in O\textsubscript{2}-saturated 0.1 M NaHPO\textsubscript{4}/NaH\textsubscript{2}PO\textsubscript{4} buffer solution at pH 7. The comparison spectra testing under both Ar and O\textsubscript{2} reveal that under Ar there is a very small reduction peak (Figure 8a). In contrast, PDI-1 under O\textsubscript{2}-saturated conditions results in a prominent cathodic ORR peak with an onset potential of ~0.40 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\textsubscript{2} molecule for the ORR) was estimated as shown in inset of Figure 8b.\textsuperscript{48} The $n$ value was determined to be 1.9 at a potential range from −0.20 V to 0.20 V. This demonstrates the complete two-electron transfer reduction process of oxygen to H\textsubscript{2}O\textsubscript{2} proceeds according to eq 7.

$$\text{O}_2 + 2 \text{H}^+ + 2 \text{e}^- \rightarrow \text{H}_2\text{O}_2 \quad (7)$$

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\textsubscript{2} can be reduced onto a GC electrode.\textsuperscript{49}
Figure 8. (a) CVs of PDI-1 under 1 atm O\textsubscript{2} or Ar. All scans were collected at 100 mV s\textsuperscript{-1} using a GC working electrode with an area of 0.196 cm\textsuperscript{2}. (b) LSV curves of RDEVs of PDI-1 in O\textsubscript{2}-saturated 0.1 M NaHPO\textsubscript{4}/NaH\textsubscript{2}PO\textsubscript{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 M NaHPO\textsubscript{4}/NaH\textsubscript{2}PO\textsubscript{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 \( \text{H}_2\text{O}_2 \) yield of PDI-1 during ORR as calculated by ring currents.
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 H₂O₂ during ORR, which is likely to passivate the electrode surface over extended cycling, similar to the H₂O₂-mediated decomposition of PEG₈–PDI⁻ 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 H₂O₂ and the potential range of ORR operation were further measured by rotating disk 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 H₂O₂ generated at the disk.⁵⁰ Using the differences in the values between disk and ring currents, the value for \( n \) and the yield of H₂O₂ were estimated to be 3.0 and 40 %, respectively (see experimental section for the details of the calculations). The value for \( n \) and the yield of H₂O₂ during the electrocatalytic ORR by PDI-1 indicates that there is nearly equal contribution of 2-electron and 4-electron reductions of O₂ to H₂O₂ and H₂O, respectively.

Based on 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 [PDI-1]⁻, which complexes with diffused O₂ to form transient single-
electron reduced complex \([\text{PDI-1}/O_2^-]\). That is readily protonated to generate hydroperoxyl radical intermediate, \([\text{PDI-1}]/'\text{OOH}\). 'OOH is a stronger oxidant, therefore formation of \([\text{PDI-1}]/'\text{OOH}\) 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 re-formation of \([\text{PDI-1}]\).

![Scheme 3](image)

**Scheme 3.** Oxygen reduction reaction mechanism by PDI-1 to form \(H_2O_2\).

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 \([\text{PDI-1}]^-/O_2\) to \([\text{PDI-1}]^-/\text{OOH}\) 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 \(O_2^-\) and \(H_2O_2\).

**Conclusion**

In summary, preparation and characterization of small well-defined conjugated molecular analogs, PEG-PDs, as a model to elucidate the mechanisms operative in the SOD-mimetic reaction.
activity of PEG-HCCs, was accomplished. Water-soluble perylene diimide derivatives serve as molecular analogs of PEG-HCCs that mimic the full catalytic cycle of SOD. PEGₙ-PDI was able to oxidize O₂⁻ and the PEGₙ-PDI⁻ intermediate of the catalytic cycle, is thoroughly characterized and shown to react with O₂⁻ in the presence of protons to form H₂O₂. Having the redox potential of PEGₙ-PDIs located between the two half reactions: O₂⁻ → O₂ + e⁻ and O₂⁻ + 2 H⁺ + 2 e⁻ → H₂O₂ (−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 O₂⁻ in aqueous environment. We show that study of graphene-based catalysts can benefit from the study precise molecular analogs,²²,⁵³ and PDI serves as a minimal model of HCCs and more generally oxidized graphenes. Furthermore, based on the results from the RDEV and RRDEV, PDI was shown to be a metal-free molecular electrocatalyst for the O₂ reduction reaction with H₂O₂ being produced in 40% yield. Thus, the results indicate that similar to carbon nanomaterials, PDI is an efficient electron shuttle for reactions with O₂ 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 nano-antioxidant.

**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 [PEG₃−PDI− CoCp⁺] are presented in Table S1. Diffraction data were collected on a Rigaku SCX-Mini diffractometer (Mercury2 CCD) using graphite-monochromated Mo Kα radiation (λ = 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 SHELXTL 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 (CHInstruments) for 10 mL electrolyte solutions (0.1 M [(n-Bu)₄N]ClO₄ 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\textsubscript{2} in the cell solution was maintained throughout the measurement to ensure continuous O\textsubscript{2} saturation. Measurements were carried out at pH 7 (0.1 M K\textsubscript{2}HPO\textsubscript{4}/KH\textsubscript{2}PO\textsubscript{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\textsubscript{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

\[ 0.62nF[O_{2}](D_{O2})^{2/3} \omega^{1/2} \nu^{-1/6} \]

where \( n \) is the number of electrons transferred to the substrate, \( F \) is the Faraday constant, \( [O_{2}] \) is the concentration of O\textsubscript{2} in an air-saturated buffer (0.22 mM in case of pH 7) at 25 °C, \( D_{O2} \) is the diffusion coefficient of O\textsubscript{2} (1.8 \times 10^{-5} \text{ cm}^{2} \text{ s}^{-1} \text{ at pH 7}) at 25 °C, \( \omega \) is the angular velocity of the disc and \( \nu \) is the kinematic viscosity of the solution (0.009 cm\textsuperscript{2} s\textsuperscript{-1}) at 25 °C. Eq 8 can be restated as eq 9; solving for \( J_{L} \) gives eq 10:

\[ 1/J = 1/J_{K} + 1/J_{L} \quad (9) \]

\[ J_{L} = 0.62nF[O_{2}](D_{O2})^{2/3} \omega^{1/2} \nu^{-1/6} \quad (10) \]

RRDE measurements were carried out to determine the H\textsubscript{2}O\textsubscript{2} yield (%) and \( n \), which were calculated by eqs 11 and 12:

\[ \text{H}_2\text{O}_2\% = \frac{[200 \ i_{d}/N]}{[i_{d} + i_{r}/N]} \quad (11) \]

\[ n = 4 \ i_{d}/[i_{d} + i_{r}/N] \quad (12) \]

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\textsubscript{3}[Fe(CN)\textsubscript{6}] in 0.1 M KNO\textsubscript{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 1.0G. The same sample was re-measured after adding a small amount of KO$_2$ (1 mg in powder form).

**Steady-State Consumption of Superoxide.** PEG$_8$–PDI (5 µM) 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 ºC) to stop the reaction. The samples were then transfer to LN$_2$ to preserve the remaining O$_2$$^•$−. EPR spectra were then recorded. To account for background dismutation of O$_2$$^•$−, a sample lacking PEG$_8$–PDI was measured and its EPR spectra subtracted from sample spectra to obtain the amount of KO$_2$ decay catalyzed by the PEG$_8$–PDI.

**Supporting Information:** additional graphs and data, single-crystal X-ray diffraction analysis details of [PEG$_3$–PDI$^-$ CoCp$^+$]. This material is available free of charge via the Internet at [http://pubs.acs.org](http://pubs.acs.org).

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**Conflicts:** 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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