Conjugated Polymer Energy Level Shifts in Lithium-Ion Battery Electrolytes

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Supporting Information

ABSTRACT: The ionization potentials (IPs) and electron affinities (EAs) of widely used conjugated polymers are evaluated by cyclic voltammetry (CV) in conventional electrochemical and lithium-ion battery media, and also by ultraviolet photoelectron spectroscopy (UPS) in vacuo. By comparing the data obtained in the different systems, it is found that the IPs of the conjugated polymer films determined by conventional CV (IPC) can be correlated with UPS-measured HOMO energy levels (EHUPS) by the relationship EHUPS = (1.14 ± 0.23) × qIPC + (4.62 ± 0.10) eV, where q is the electron charge. It is also found that the EAs of the conjugated polymer films measured via CV in conventional (EAC) and Li⁺ battery (EAB) media can be linearly correlated by the relationship EAB = (1.07 ± 0.13) × EAC + (2.84 ± 0.22) V. The slopes and intercepts of these equations can be correlated with the dielectric constants of the polymer film environments and the redox potentials of the reference electrodes, as modified by the surrounding electrolyte, respectively.

KEYWORDS: lithium-ion battery, conjugated polymer, electron affinity, cyclic voltammetry, electrolyte, organic photovoltaics

INTRODUCTION

Conjugated polymers find use in numerous electronics applications because of their exceptional properties, which include mechanical flexibility, production scalability, and chemically tunable electrical/electronic properties.¹⁻⁴ These materials have been successfully implemented in organic photovoltaic cells,⁵⁻⁸ organic light-emitting diodes,⁹⁻¹² and organic transistors.¹³,¹⁴ Recently, conjugated polymers have also been employed in lithium-ion battery (LIB) applications to replace¹⁵⁻¹⁷ and/or enhance conventional carbon-based electrodes,¹⁸⁻²⁰ and significantly enhanced battery performance has been reported.

The LIB is a widely used energy storage system with applications that range from portable appliances to automobiles.²¹,²² An important LIB figure-of-merit is the specific capacity.²³ One way to increase overall LIB capacity is to replace the ubiquitous carbon anode with one based on Si or Sn, which offer theoretical specific capacities of 4200 and 994 mAh/g, respectively, far surpassing that of graphite, 372 mAh/g.²⁴,²⁵ The principal challenge facing Si and Sn battery anode technologies is managing the up to 320% volumetric expansion/contraction of the anode during Li,M intermetallic formation in the charge/discharge processes.²⁵ Repeated expansion and contraction cause destructive anode fragmentation, exposing fresh anode surfaces to the organic electrolyte. These electrolyte materials readily undergo chemical reduction on the electrode surfaces, yielding passivating solid—electrolyte interface (SEI) films that irreversibly consume the Li⁺ ions essential for energy storage.²⁶⁻²⁹ For these reasons, active research efforts have focused on developing protective materials systems based on conjugated polymers to minimize cycling-induced anode damage.¹⁷⁻²⁰

Note that the electrochemical properties of polymers potentially useful for battery applications must be characterized in depth for the relevant electrochemical environment of use. Specifically, quantifying the energetics of the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) are essential to understanding the aforementioned electrochemical properties. The conjugated polymer LUMO energy is especially important since it relates to charge transport and stability at the anode;¹⁸,¹⁹ and when the anode potential coincides with the polymer LUMO energy, the reduction of Li⁺ to Li occurs and SEI films form.⁵,⁲³,²⁷,²⁸ However, note that conjugated polymer energy levels are typically derived from electrochemical measurements performed in completely different electrolyte environments than...
the commonly used LIB liquid carbonate battery electrolytes. Conjugated polymer thin films are conventionally characterized in moderately polar organic solvents such as acetonitrile with \(\sim 0.1\) M organic salt as the electrolyte. In contrast, LIBs typically utilize high Li\(^+\)PF\(_6\)^\(-\) electrolyte concentrations (\(\sim 1.2\) M) in mixtures of very polar solvents such as ethylene carbonate (EC) + ethyl methyl carbonate (EMC). Therefore, the frontier orbital energies of conjugated polymers of potential interest could, in principal, shift from those measured in conventional and LIB electrochemical media when estimated by cyclic voltammetry (CV).

In this study, we quantify the frontier orbital energetics of a selected series of known and new conjugated polymers in LIB media and in conventional electrochemical media. To quantify HOMO energies without the intrusion of electrochemical medium effects, in vacuo ultraviolet photoelectron spectroscopy (UPS) is also applied since this alternative method is commonly used to characterize the electronic structure of conjugated polymers.\(^{30-32}\) The conjugated polymers characterized in this study are P3HT, poly(3-(2-propoxyethyl)-thiophene) (P(3POET)), poly(3-(2-methoxy-ethoxy)methyl)thiophene) (P(3MEMT)), poly[4,8-bis[(2-ethylhexyl)oxy]-benzo[1,2-b:4,5-b']dithiophene-2,6-diy][3-fluoro-2-[(2-ethylhexyl)carbonyl]thieno[3,4-b']thiophenediyl)] (PTB7), poly[N-9′-heptadecanyl-2,7-carbazole-alt-5,5-(4′,7′-di-2-thienyl-2′,1′,3′-benzothiadiazole)] (PCDTBT), poly[(5,5′-(4,8-bis(thien-2-yl))-2,6-(dihexyl)benzo[1,2-d,4,5-d”]bis[1,2,3]-triazole-alt-(4,8-bis(5-(2-hexyldecyl)thien-2-yl)benzo[1,2,4,5-b’]dithien-2-yl)] (DT003CF), ActiveInk N2200 (Polyera Corp.), and poly(2,7-(N,N′-bis(2-ethylhexyl)-naphthalenediimide)-co-5,5’-(3,3’-bis(2-(2-methoxyethoxy)ethoxy)ethoxy)-2,2’-bithiophene) (P(NDI-tegme2T)). The molecular structures are shown in Scheme 1. We report here that the estimated orbital energies of these conjugated polymers differ substantially with measurement technique, but also that informative predictive correlations exist between energies measured by UPS, conventional CV, and CV in a commonly used battery medium.

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Regarding experimental data, the ionization potentials (IPs), electron affinities (EAs), HOMO energies (\(E_H\)’s), and LUMO

**Scheme 1. Conjugated Polymers Used in This Study**

**Table 1. Cyclic Voltammetry Experimental Parameters and Terminologies**

<table>
<thead>
<tr>
<th>experimental detail</th>
<th>conventional system (CV(_C))</th>
<th>battery system (CV(_B))</th>
</tr>
</thead>
<tbody>
<tr>
<td>working electrode</td>
<td>Pt (0.20 cm(^2))</td>
<td>patterned glass/Cu (0.50 cm(^2))</td>
</tr>
<tr>
<td>reference electrode</td>
<td>Ag/AgCl (aq) in KCl (3.0 M)</td>
<td>Li</td>
</tr>
<tr>
<td>counter electrode</td>
<td>Pt</td>
<td>Li</td>
</tr>
<tr>
<td>electrolyte (conc.)</td>
<td>TBA/PP(_6)^− (0.1 M)</td>
<td>Li/PP(_6)^− (1.2 M)</td>
</tr>
<tr>
<td>solvent</td>
<td>acetonitrile, anhydrous</td>
<td>EC:EMC (3:7)</td>
</tr>
<tr>
<td>scan rate</td>
<td>100 mV/s</td>
<td>10 mV/s</td>
</tr>
<tr>
<td>sweep range</td>
<td>−2300 to 2300 mV</td>
<td>10−3300 mV</td>
</tr>
<tr>
<td>LUMO labeling (CV/absolute scale)</td>
<td>(E_{A_C}/E_{L_C})</td>
<td>(E_{A_B}/E_{L_B})</td>
</tr>
<tr>
<td>HOMO labeling (CV/absolute scale)</td>
<td>(E_{P_C}/E_{H_C})</td>
<td>(E_{P_B}/E_{H_B})</td>
</tr>
</tbody>
</table>
energies ($E_L$) determined by conventional CV ($E_{CV}$), by CV in a battery medium ($E_{CVB}$), and by UPS are first acquired and compared; all energy levels are reported on the corresponding absolute scale. Then, a brief discussion of the origin of the energy level shifts and apparent correlations follows. To aid in comparing data from different measurements, abbreviations and experimental details are summarized in Table 1. Details of experimental procedures and conditions are described in the Supporting Information.

■ RESULTS AND DISCUSSION

Cyclic Voltammetry with Conventional Electrolytes/Electrodes. CV scans in 0.1 M TBA$^+$/PF$_6^-$ in anhydrous acetonitrile. Positive and negative currents represent oxidation and reduction, respectively.

![Figure 1. CV scans of conjugated polymer films in 0.1 M TBA$^+$/PF$_6^-$ in anhydrous acetonitrile. Positive and negative currents represent oxidation and reduction, respectively.](image)

Table 2. Solution-Phase Electrochemical Data

<table>
<thead>
<tr>
<th>polymer</th>
<th>$E_{IPC}$ vs $E_{Fc}$ (V)$^a$</th>
<th>$E_{EAC}$ vs $E_{Fc}$ (V)$^a$</th>
<th>$E_{H,C}$ (eV)$^b$</th>
<th>$E_{L,C}$ (eV)$^c$</th>
<th>literature $E_{IPC}/E_{EA}$ (eV)$^d$</th>
<th>ref</th>
</tr>
</thead>
<tbody>
<tr>
<td>P3HT</td>
<td>0.032</td>
<td>−2.13</td>
<td>5.13</td>
<td>2.97</td>
<td>4.75/2.73</td>
<td>34</td>
</tr>
<tr>
<td>P(3POET)</td>
<td>−0.036</td>
<td>−2.22</td>
<td>5.06</td>
<td>2.88</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>P(3MEMT)</td>
<td>0.075</td>
<td>−1.79</td>
<td>5.13</td>
<td>3.31</td>
<td>5.13/NA</td>
<td>35</td>
</tr>
<tr>
<td>PTB7</td>
<td>0.480</td>
<td>−1.56</td>
<td>5.58</td>
<td>3.54</td>
<td>5.55/3.40</td>
<td>8</td>
</tr>
<tr>
<td>PCDTBT</td>
<td>0.649</td>
<td>−1.59</td>
<td>5.75</td>
<td>3.51</td>
<td>5.65/3.48</td>
<td>36</td>
</tr>
<tr>
<td>DT003CF</td>
<td>0.342</td>
<td>−1.71</td>
<td>5.44</td>
<td>3.39</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>N2200</td>
<td>0.833</td>
<td>−0.86</td>
<td>5.93</td>
<td>4.24</td>
<td>5.97/4.32</td>
<td>37</td>
</tr>
<tr>
<td>P(NDI-tegme2T)</td>
<td>0.076</td>
<td>−0.69</td>
<td>5.18</td>
<td>4.41</td>
<td>NA</td>
<td></td>
</tr>
</tbody>
</table>

$^a$Estimated uncertainty range is 0.1 V. $^b$Calculated from eq 1. $^c$Calculated from eq 2. $^d$Data reprocessed according to eqs 1 and 2.

...polymer oxidation and reduction peaks versus $Fc^+/Fc$, respectively. The 5.1 eV is the commonly used conversion factor to relate the $Fc$ redox couple formal potential ($E_{Fc} = 0.630$ V vs NHE$^{27}$) to an absolute scale ($E_{NHE} = 4.5$ eV). Since there are several variants of eqs 1 and 2, the previously reported orbital energies of P3HT,$^{39}$ P-(3MEMT),$^{40}$ PTB7,$^{8}$ PCDTBT,$^{41}$ and N2200$^{42}$ are recalibrated here to eqs 1 and 2 and are compared to values

dx.doi.org/10.1021/am505416m | ACS Appl. Mater. Interfaces 2014, 6, 19347–19354
determined in the present study in Table 2. Note that the current values generally track those in the literature. These data are also compared to UPS data to assess the validity of eqs 1 and 2, as discussed below. The onsets of the oxidation and reduction peaks along with the corresponding energy levels on the absolute scale from eqs 1 and 2 are summarized in Table 2.

**Cyclic Voltammetry with a Battery Electrolyte/Electrode System.** The first CV scans from a three-electrode system in 1.2 M Li$^{+}$PF$_6^-$/3:7 EC:EMC yield the EA$_B$ (electron affinity in battery medium) values of the conjugated polymers (Figure 2). After a complete scan, all of the present polymer films are visually bleached except N2200 and P(NDI-tegme2T). Note that IPB/E$_{H,B}$ values (ionization potential/HOMO energy in battery medium) could not be obtained here due to competing Cu electrode oxidation (IPB of Cu $\sim$ 3.4 V vs $E_{Li}$). Also, the EA$_B$’s of several polymers could not be observed, presumably because their high-lying LUMOs lie beyond the Li plating potential ($\sim$0.0 V vs $E_{Li}$). Thus, only data for polymers with reproducibly measurable EA$_B$ values are reported. Next, eq 2 was modified by replacing $E_{Fc}$ with the work function of Li ($\Phi_{Li} = 2.49$ eV$^43$), as in eq 3, and polymer $E_{L,B}$ values on an absolute scale estimated. This yields $E_{L,B}$ values

$$E_{L,B} = qEA_B + 2.49 \text{ eV}$$

(LUMO energies in battery medium) close to $E_{L,C}$; however, some discrepancies are apparent, the origins of which are discussed below. CV data and energy differences ($\Delta E_L = E_{L,C} - E_{L,B}$) are summarized in Table 3.

**UPS and CV Correlations.** UPS data (Figure S2, Supporting Information) were used to derive $E_{H,UPS}$ (HOMO energy in vacuo) from the work function ($\Phi$) and the valence band maximum (VBM) values of the conjugated polymers on Cu foil. Here, $\Phi$ is determined from the averaged spectra of four measurements using the difference between the secondary electron cutoff (SECO), the highest binding energy, and the He I source energy (21.22 eV$^{44,45}$). The VBM values of the polymer-coated Cu foils relative to the Fermi level (0.0 eV) were determined from the onset of the lowest binding energy. The IP, or $E_{H,UPS}$, was then calculated from the sum of $\Phi$ and the VBM$^{44,46}$ Note that the UPS-derived $E_{H,UPS}$ values track the $E_{H,C}$’s closely, but some discrepancies ($\Delta E_H = E_{H,C} - E_{H,UPS}$) as large as 0.72 eV are found (Table 4). However, $\Delta E_H$...
is significantly reduced ($\Delta E_{\text{H,cor}}$) when the correction suggested by D’Andrade\textsuperscript{35} (eq 4) is applied instead of eq 1.

$$E_{\text{H,cor}} = (1.4 \pm 0.1) \times q\text{IP}_C + (4.6 \pm 0.08) \text{ eV}$$

(4)

In this model, $E_{\text{H,cor}}$ denotes the corrected $E_{\text{H}}$ value, and the slope of eq 4 reflects analyte image charge and solvation effects on the electrode surface interacting with the surrounding electrolyte, and the intercept of 4.6 is the $\Phi_f$ of ferrocene. D’Andrade argued that the image charges induced by an analyte on a conductive surface under vacuum or in a polar solvent skew the energy level correlation from unity. In general, the greater the image charge, the greater the slope, with the image charge directly proportional to the dipolar strength of the charged analyte. In the present study, a relationship similar to eq 4 (eq 5) is obtained by plotting $\text{IP}_C$ vs $\text{IP}_{\text{UPS}}$ (Figure 3a).

$$E_{\text{H,cor}} = (1.14 \pm 0.23) \times q\text{IP}_C + (4.62 \pm 0.10) \text{ eV}$$

(5)

The principal factors contributing to the disparities in the results of this investigation and D’Andrade’s model are the analyte solvation effects. D’Andrade et al. used small molecule analytes dissolved in the electrolyte for CV measurements, minimizing interanalyte molecule/ion interactions and maximizing electrolyte solvation. In the present study, CV is performed on solid thin films where solvation effects should be far less, but in which interanalyte interactions should be large. However, as D’Andrade et al. note, solvation effects should be small compared to the image charge effects for adsorbed analytes,\textsuperscript{35} contributing to the similarity of eqs 4 and 5. Note that UPS measurements are on analyte films cast on metal electrodes; thus, any disparities between eqs 4 and 5 should not originate from the forms of the UPS samples.

**EA$_C$ – EA$_B$ Correlations.** For the same conjugated polymer, the orbital energies obtained from CV$_C$ and CV$_B$ are clearly different and likely reflect the different reference electrodes employed (Fc$^+$/Fc vs Li$^+$/Li$^-$). Furthermore, the frontier orbital energetics of the conjugated polymers are expected to vary with solvent and electrolyte concentration. If the origin of these differences is ascribed solely to the reference electrode, the analyte energy levels should uniformly shift with the potential differences of the reference electrodes. However, when EA$_C$ and EA$_B$ data are compared, the differences are significantly smaller than the reference electrode potential difference ($E_{\text{FC}} - E_{\text{Li}} = 3.675$ V), and other sources of energy level shifts, such as electrolyte concentration\textsuperscript{47–49} and solvation, must be considered.\textsuperscript{50–52} Classic examples of this effect include the EA shift of Ti$^{3+}$ ($\sim$0.2 V) with increasing electrolyte concentration of 0.1 to 100 mM\textsuperscript{47} and the O$_2$ IP shift of 0.31 V at a Pt electrode on changing the solvent from DMSO to acetonitrile.\textsuperscript{51} In the present case, CV$_C$ and CV$_B$ measurements are conducted in 0.1 M TBA$^+$PF$_{6}^-$/acetonitrile and 1.2 M Li$^+$PF$_{6}^-$/EC:EMC, respectively, corresponding to a large change in electrochemical environment.

Similar to D’Andrade’s approach,\textsuperscript{35} the EA$_C$ and EA$_B$ data for the present polymers are plotted against each other in Figure 3b. A reasonable linear correlation ($R^2 = 0.90$) is observed, indicating that the orbital energetics from one measurement are approximately translatable to the other within the $-2.4$ to $-0.4$ V range. Thus, EAs of conjugated polymers in a battery medium can be estimated (EA$_{\text{rest}}$) from CV$_C$ measurements using eq 6. The parameters in this equation

$$EA_{\text{rest}} = (1.07 \pm 0.13) \times EA_C + (2.84 \pm 0.22) \text{ V}$$

(6)

afford insights into the relevant electrochemical phenomena in that the intercept ($2.84 \pm 0.22$ V) is less than the reference electrode potential difference ($3.675$ V), but the slope is near unity, implying that conjugated polymers measured as thin films on electrodes by CV$_C$ and CV$_B$ experience minimal differential solvation effects. However, note that the slope is greater than unity when UPS and CV$_C$ data are compared. D’Andrade modeled such slopes by the ratio of the dielectric constants of the respective analyte media. The medium in D’Andrade’s model is the neighboring analyte in films for UPS, and electrolyte in CV—hence, the dielectric constant differences are very large. In the present case, the analyte is not likely to be extensively solvated but embedded in a solid film in both CV experiments. Thus, dielectric constant effects in this model should be relatively small and a slope near unity is expected—exactly what eq 6 reveals. Thus, the apparent orbital energy shifts primarily arise from the potential difference between the reference electrodes; however, as noted above, the electrochemical potentials are also affected by the electrolyte concentration,\textsuperscript{47,48} solvation,\textsuperscript{51,52} and image charge effects, which alter the working electrode potential, as well as that of the reference electrode.\textsuperscript{53–55} Indeed, $E_{\text{FC}}$ on the absolute electrochemical scale is 5.1 eV, while $\Phi_{\text{FC}} = 4.6$ eV. Similarly, $E_{\text{Li}}$
on the absolute scale is 1.44 eV, but $\Phi_{\text{Li}} = 2.49$ eV, where the differences likely originate from reference electrode–electrolyte interactions. A similar phenomenon is also observed with the SHE. These results signify that simple addition/subtraction of a single parameter to convert orbital energies in one electrochemical environment to another can introduce significant errors. Note also that some polymers do not strictly follow the linear trend in Figure 3. This phenomenon may arise from differences in the polymer–electrolyte interactions; however, further studies will be required to better define these interactions.

## CONCLUSIONS

The orbital energetics of a series of established conjugated polymers were analyzed by CV in conventional and Li-ion battery solution environments, and by UPS in vacuo. By comparing data obtained from these two regimes, it is seen that the ionization potentials measured by conventional CV can be correlated with the UPS-derived HOMO energies via $E_{\text{HOMO}} = (1.14 \pm 0.23) \times q \Phi_{\text{Li}} + (0.42 \pm 0.01) \text{ (eV)}$ and the electron affinities of conjugated polymers measured by CV in conventional and Li battery environments can be linearly correlated by the relationship $E_{\text{AOMO}} = (1.07 \pm 0.13) \times q \Phi_{\text{Li}} + (2.84 \pm 0.22) \text{ (eV)}$. The slopes and the intercepts of these equations are related to the dielectric constants of the analyte environments and the redox potentials of the reference electrodes as affected by the surrounding electrolyte, respectively. We expect that the correlations and their origins described here will be useful in analyzing/predicting the electronic structures of diverse conjugated polymers being considered for organic electronic and energy storage applications.

## ASSOCIATED CONTENT

2 Supporting Information

More details of polymer synthesis, CV and UPS experimental methods, and data of the conjugated polymers are described. This material is available free of charge via the Internet at http://pubs.acs.org.

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### Author Contributions

*These authors contributed equally to this work.

### Notes

The authors declare no competing financial interest.

## ACKNOWLEDGMENTS

This research was supported in part by the Institute for Sustainability and Energy at Northwestern (ISEN) (C.K.S.) and by the Argonne-Northwestern Solar Energy Research (ANSER) Center, an Energy Frontier Research Center funded by the U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences, under Award Number DE-SC0001059 (B.J.E.). This research was also supported by the Agency of Science, Technology and Research (A*STAR; T.L.D.T.). We acknowledge the Integrated Molecular Structure Education and Research Center (IMSERC) for molecular characterization facilities for NMR spectroscopy supported by Northwestern U., the NSF under grants CHE-0923236 and CHE-9871268, Pfizer, and the State of Illinois. We also thank the Nanoscale Integrated Fabrication, Testing, and Instrument (NIFTI) and Keck Interdisciplinary Surface Science (KECK-II) facilities of Northwestern University's Atomic and Nanoscale Characterization Experimental (NUANCE) Center for UPS experiments, supported by Northwestern U., the State of Illinois, the Keck foundation, NSF-MRSEC, and NSF-NSEC. Finally, we thank Dr. Antonio Facchetti, J. T. Shin, and J. L. Song for helpful discussions.

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