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The Molecular Mechanism of Alternative P450-Catalyzed Metabolism of Environmental Phenolic Endocrine-Disrupting Chemicals

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Abstract

Understanding the bioactivation mechanisms to predict toxic metabolites is critical for risk assessment of phenolic endocrine-disrupting chemicals (EDCs). One mechanism involves ipso-substitution, which may contribute to the total turnover of phenolic EDCs, yet the detailed mechanism and its relationship with other mechanisms are unknown. We used density functional theory to investigate the P450-catalyzed ipso-substitution mechanism of the prominent xenoestrogen bisphenol A. The ipso-substitution proceeds via H-abstraction from bisphenol A by Compound I, followed by essentially barrierless OH-rebound onto the ipso-position forming a quinol, which can spontaneously decompose into the carbocation and hydroquinone. This carbocation can further evolve into the highly estrogenic hydroxylated and dimer-type metabolites. The H-abstraction/OH-rebound reaction mechanism has been verified as a general reaction mode for many other phenolic EDCs, such as bisphenol analogues, alkylphenols and chlorophenols. The identified mechanism enables us to effectively distinguish between type I (eliminating-substituent as anion) and type II (eliminating-substituent as cation) ipso-substitution in various phenolic EDCs. We envision that the identified pathways will be applicable for prediction of metabolites from phenolic EDCs whose fate is affected by this alternative type of P450 reactivity, and accordingly enable the screening of these metabolites for endocrine-disrupting activity.
Introduction

Biotransformation plays a critical role in determining the toxicity of xenobiotics in organisms and has drawn considerable attention as a basis for environmental risk assessment.\(^1,2\) Biotransformation of environmental endocrine-disrupting chemicals (EDCs) is one such example.\(^3,4\) Accurate risk assessment of EDCs requires consideration of bioactivation via biotransformation processes, especially by human cytochrome P450 enzymes (P450), since neglecting these metabolic pathways may lead to undervaluation of their adverse effects on human health, although the metabolism of phenolic chemicals by P450 is minor compared with the glucuronidation pathway under normal circumstances.\(^3,4\) P450 enzymes are a superfamily of monooxygenases distributed through all kingdoms of life, and are responsible for most phase-I biotransformation reactions.\(^5-9\) Some of these conversions produce metabolites that are much more toxic than their parent compounds, an important example being phenolic EDCs.\(^10\) Phenolic EDCs such as bisphenol analogues, alkylphenols and chlorophenols, are ubiquitous in the environment as widely used industrial chemicals, with associated high risk of environmental exposure.\(^10\) Among these, although bisphenol A (BPA) has traditionally been considered a weak environmental xenoestrogen because of its much lower binding affinity to the estrogen receptor than that of estradiol,\(^11\) the biotransformation largely affects the endocrine disrupting activity of BPA.\(^4\)

As shown in Scheme 1, conjugation with the phase II glucuronide enzyme is the predominant metabolic pathway of BPA in humans (more than 90% of all BPA metabolites), which represents a major detoxification pathway;\(^12\) however, BPA is also metabolized by human P450 to form ortho-OH-BPA via hydroxylation of the aromatic ring,\(^12\) to form hydroxycumyl alcohol (HCA), isopropenylphenol (IPP), and hydroquinone (HQ) via an ipso-substitution mechanism,\(^13\) and to form a dimer-type metabolite 4-methyl-2,4-bis(4-hydroxyphenyl)pent-1-ene (MBP) whose
formation may involve IPP reacting with isopropenylphenol radical.\textsuperscript{14,15} \textit{In vitro} assays have shown that HCA can exhibit 100-fold higher estrogen activity than BPA (concentrations of $10^{-5}$ to $10^{-10}$ M),\textsuperscript{13} while MBP can be 1000-fold more potent (concentrations of $10^{-5}$ to $10^{-9}$ M).\textsuperscript{14} Although the \textit{ipso}-substitution pathway of the P450-catalyzed metabolic activation of BPA is most likely a minor pathway under most circumstances, such strong endocrine-disrupting activity of the metabolites makes this pathway important to the overall environmental risk assessment, especially under conditions where glucuronidation is impaired by e.g. other compounds or for genetic or developmental reasons. For example, human fetal livers show little or no glucuronidation\textsuperscript{16} and in contrast to rodents express significant levels of P450 leading to metabolizing many xenobiotic compounds even at the prenatal stage.\textsuperscript{17,18} thus P450-catalyzed metabolic activation is more likely relatively more significant in the fetus.\textsuperscript{3,19,20}

\textbf{Scheme 1. Major Metabolic Pathways of Bisphenol A}
Formation of metabolites via ipso-substitution constitutes about 20% of the competing ortho-OH-BPA formation via traditional aromatic hydroxylation by P450, i.e. ipso-substitution is quantitatively important in competition with the traditional aromatic hydroxylation of phenolic EDCs. Oxidation of diverse p-substituted phenols by rat liver P450 has been found to result in elimination of the substituents, including -NO₂, -CH₂OH, -COCH₃, -COPh, -COOH, -F, -Cl, and –Br. Accordingly, ipso-substitution can be categorized into two types depending on the group eliminated from the quinol intermediate. As shown in Scheme 2, type I ipso-substitution implies that the substituent eliminates as an anion with formation of a quinone, whereas in type II ipso-substitution the eliminating group is a cation, leading to the formation of a hydroquinone. However, during oxidation of 4-nonylphenol, estrone, estradiol etc. by P450, ipso-addition quinol was formed without C-C bond cleavage. Therefore, the ipso-substitution, ipso-addition, as well as the above-mentioned aromatic hydroxylation mechanisms compete under various conditions as relevant pathways, and understanding these mechanisms at the molecular level seems necessary to access the environmental toxicity and fate of phenolic EDCs. However, the active species of P450, the iron(IV)-oxo heme cation radical Compound I (Cpd I), responsible for P450-catalyzed oxidations in all P450 isoenzymes, is short-lived and one of the most potent oxidants in nature, and thus several details of its catalytic action are inaccessible by standard experimental methods. Specially, two possible pathways for P450-catalyzed ipso-substitution via a quinol intermediate should be distinguished; one involving initial formation of a phenoxy radical and the other involving the formation of an epoxide via O-addition.

Analysis of enzyme mechanisms using computational chemistry may identify with semi-quantitative accuracy the electronic structure features governing reactivity. Density functional theory (DFT) has been used to study many P450-catalyzed oxygenation reactions including
hydroxylation of C-H bonds, epoxidation of C=C bonds, oxidation of aromatic rings, oxidation of heteroatoms etc.\textsuperscript{33} The main goal of this work is to show how DFT can be used to elucidate the full molecular mechanism of the P450-dependent metabolism of phenolic EDCs and to identify how and when environmentally related ipso-substitution, and formation of the very estrogenic dimer-type metabolites can occur. BPA was used to obtain the full mechanistic picture because of its prominence in the environment,\textsuperscript{39,40} with rich experimental data of its P450-catalyzed metabolism\textsuperscript{13,14} for validation of the computationally obtained mechanisms. The work was extended to also study the P450-catalyzed biotransformation mechanisms of several other widely-used phenolic EDCs, such as bisphenol analogues, alkylphenols and chlorophenols. The fundamental electronic drivers that govern ipso-addition vs. ipso-substitution and type I vs. type II substitute elimination were identified, directly relevant for screening P450-catalyzed biotransformation of many emerging environmental phenolic EDCs.

\textbf{Scheme 2.} Proposed \textit{ipso}-Substitution Mechanisms of P450-Catalyzed Substituent Elimination\textsuperscript{a}

\begin{itemize}
  \item The reactive position is defined as the \textit{ipso}-position; R represents the elimination substituent.
\end{itemize}
Computational Methodology

**DFT Calculations with Cpd I of P450.** As is common practice, the six-coordinate triradicaloid ferryl complex $\text{Fe}^{4+}O^2-(\text{C}_{20}\text{N}_4\text{H}_{12})\text{SH}^{-1}(\text{SH})^{-1}$ was used to model the enzymatic active site of Cpd I of P450. Cpd I of P450 exists in two close-lying electronic states, a high-spin (HS) quartet state and a low-spin (LS) doublet state. All geometries on both the LS and HS routes were optimized with unrestricted DFT using the B3LYP hybrid density functional in combination with the LAN2DZ basis set on iron and 6–31G on other atoms (denoted BSI). B3LYP was chosen because it can reproduce measured kinetic isotope effects for P450-catalyzed reactions, electron paramagnetic resonance parameters for penta-coordinated heme in P450 enzyme, generate geometries consistent with crystal structures, and show qualitatively accurate relative energies vs. benchmark CASSCF calculations. Intrinsic reaction coordinate (IRC) calculations were performed to verify the rate-determining transition states connecting the reactants and intermediates on the potential energy surface (Figure S1-S22 in the Supporting Information). Please note that the basis-set superposition error (BSSE) has been reported to be very small for reactant complexes of P450-catalyzed oxidation reactions, but they may affect the relative energies of very large vs. small substrates and thus we did not include these minor contributions to the energies in the following as our substrates are similar in size and type.

In order to evaluate broadly the sensitivity of the reaction mechanism toward the choice of density functional, in addition to the B3LYP energies (Table S1 in the Supporting Information), we performed unrestricted single-point calculations with other hybrid, local, and non-hybrid functionals, i.e. TPSSh, B3PW91, BLYP, MPW1PW91, and M06L using the B3LYP/BSI optimized geometries for the P450-catalyzed metabolic mechanisms of BPA (Table S2 in the Supporting Information). The same qualitative picture was obtained with all of the
functionals, and we therefore focused in the following on the B3LYP results. To test the basis set effect on geometry optimization, the molecular species involved in the initial H-abstraction from the phenolic group as well as in the O-addition to the aromatic ring of BPA were optimized at the B3LYP/BSI** level, producing few geometrical and energetic discrepancies as compared with the results obtained at the B3LYP/BSI level (detailed data in Table S3 and Figure S23 in the Supporting Information). Hence the basis set BSI was used for geometry optimizations throughout the remaining work.

Analytical frequency calculations were used to ensure that there was no imaginary frequency for any ground state, and only one imaginary frequency for all transition states. The vibrational frequencies were also used to calculate the zero-point energy (ZPE) and thermal and entropic corrections to the free energy at 298.15 K and 101.325 kPa. More accurate energies were obtained using single-point calculations with the SDD\textsuperscript{59} basis set on iron and the 6–311++G** basis set for all other atoms (denoted BSII). Bulk polarity effects were evaluated by the PCM solvation model\textsuperscript{60} using chlorobenzene with a dielectric constant of 5.6 at the B3LYP/BSI level; this dielectric constant provides a good estimate of the polarization caused by the dipoles of the protein pocket near the axial cysteine.\textsuperscript{61} We also evaluated the bulk polarity effect using the SMD solvation model\textsuperscript{62} for the P450-catalyzed mechanisms of BPA; the H-abstraction and O-addition steps occurred with only slightly higher energies (Table S4 in the Supporting Information). In addition, we evaluated PCM energies using cyclohexane ($\varepsilon=2.0$), 1-bromopropane ($\varepsilon=8.0$), ethanol ($\varepsilon=24.9$), and acetonitrile ($\varepsilon=35.7$). Except for a minor difference in energy for the oxidation of BPA, the same qualitative picture was obtained throughout (Table S5 in the Supporting Information). Dispersion interactions were considered by performing single-point energy calculations with the B3LYP-D3/BSI level since B3LYP itself does not include dispersion by design.\textsuperscript{63} The relative free
energies of the P450 oxidation reactions shown below were estimated by combining B3LYP/BSII
single-point energies with PCM solvation and dispersion corrections, as well as Gibbs free energy
corrections from optimizations at the BSI level, unless pointed out specifically.

The cluster approach of studying the reaction mechanism treats the catalytic active site of the
enzyme by including key surrounding amino acids and treating all these interactions fully quantum
mechanically.\textsuperscript{38} BPA is mainly catalyzed by P450 isoforms 3A4 and 3A5,\textsuperscript{13} and therefore we used
the P450 3A4 crystal structure (PDB code: 1W0G)\textsuperscript{64} to produce a larger model of the active site.
As shown in \textbf{Figure S24} in the Supporting Information, the Cpd I model is the same in the large
and small model, whereas six important second-shell residues, ARG105, ILE301, THR309 and
ALA370 and the peptide chain of ALA305–GLY306 have been included in the large model, with
key central atoms locked in their crystallographic positions to maintain the protein scaffold
packing, steric effects, and hydrogen bond geometries. The large model is charge-neutral and
contains 138 atoms, and the reaction mechanism was investigated for both the HS and LS states.
The geometry optimization, more accurate single-point calculations, evaluation of the bulk polarity
effects, and dispersion interactions were all performed in the same way for both the large and small
models. The results are discussed in detail in the Supporting Information, where all energies are
compiled in \textbf{Tables S1-S31}. Importantly, we conclude that the small and large models are in good
agreement on the preferred pathways (\textbf{Figure S25} in the Supporting Information), probably
because the main energy effects and electronic reorganizations occur near the iron-oxygen moiety.
We thus performed an extended series of calculations based on the small model as discussed below.

\textbf{Reaction Energy Calculations for the Decomposition of Quinol Intermediates.} All
geometries for the decomposition reactions of various \textit{ipso}-addition quinol intermediates from
P450-catalyzed \textit{ipso}-position metabolism were optimized at the B3LYP/6-31G** level in water
solution (\(\varepsilon=78.4\)) with PCM. Then based on the optimized structures, single-point calculations were performed in PCM water solution with D3 dispersion corrections at the B3LYP/6-311++G** level. The reported reaction free energies for decomposition of quinol intermediates are described by PCM//B3LYP/6-311++G** with water solution and D3 dispersion corrections, as well as free energy corrections from B3LYP/6-31G** geometry optimizations.

All calculations of this work were carried out with the Gaussian 09 D.01 program package.65

Results and Discussion

Reaction Mechanisms of P450-Catalyzed Bisphenol A

H-abstraction vs. O-addition. Figure 1 shows two computed competitive reaction mechanisms of BPA catalyzed by P450, one involving initial H-abstraction from the phenolic group, and the other involving initial O-addition to the \(\pi\)-system of the aromatic ring. As is common in P450 reactions,33 both the HS and LS pathways are available due to the near-degenerate states of Cpd I. The reactions start from reactant complexes \((4,2\text{-RC})\), in which the H-atom of the phenolic group of BPA interacts with the iron-oxo moiety of Cpd I. Then, \(4,2\text{-RC}\) may go through H-abstraction transition states \(4,2\text{TS}_H\) with formation of the intermediate complexes \((4,2\text{I}_H)\) involving iron-hydroxo species and the phenoxy radical of BPA. The HS transition state \(4\text{TS}_H\) appears slightly later on the reaction coordinate than its LS counterpart \(2\text{TS}_H\), with BPA–O⋯H and H⋯O–Fe distances of 1.211 vs. 1.207 Å and 1.203 vs. 1.212 Å, respectively. These H-abstraction transition states are characterized by almost linear O⋯H⋯O configurations as well as large imaginary frequencies (HS: \(i1521 \text{ cm}^{-1}\); LS: \(i1569 \text{ cm}^{-1}\)). Cpd I is a potent H-atom abstractor toward the phenolic group, with a H-abstraction barrier of only 0.4/0.3 kcal/mol for the HS/LS state, similar to the minor H-abstraction barriers obtained for the phenolic group of paracetamol29.
and the amino group of anilines\textsuperscript{42}, yet much lower than the H-abstraction barriers obtained from C–H hydroxylation.\textsuperscript{41,52,66} In addition, the formed complex intermediates (\textsuperscript{4,2}I\textsubscript{H}) are stable, with exothermic reaction energies of -8.0/-7.4 kcal/mol for the HS/LS state. Note that dispersion effects lower the H-abstraction barriers by a substantial 2.5 kcal/mol, a magnitude consistent with previous findings for P450 reactions.\textsuperscript{67}

Another possible reaction path starting from \textsuperscript{4,2}RC is the addition of the oxo group of Cpd I onto the unsubstituted aromatic ring of BPA via C-O bond-forming transition states \textsuperscript{4,2}TS\textsubscript{O}, which produce tetrahedral intermediates. As shown in Figure 1, compared with the LS species, TS\textsubscript{O} in the HS state is more advanced (shorter O…C bond) with a higher degree of aromatic activation. The calculated barriers for O-addition at the ortho-position (\textsuperscript{4,2}TS\textsubscript{Oo}) and meta-position (\textsuperscript{4,2}TS\textsubscript{Om}) are 17.5/14.5 and 19.9/17.1 kcal/mol, respectively, on the HS/LS state surfaces. Comparison of the barriers of the H-abstraction and O-addition steps shows clearly that the H-abstraction reaction is much more favorable. Therefore, we focused on the H-abstraction pathway in the following sections.
Figure 1. Free energy profile of BPA catalyzed by Cpd I of P450, along with the optimized geometries of the key reaction species in the HS and LS states. Free energies (kcal/mol) are relative to the quartet reactant complex $^4$RC at the B3LYP/BSII//BSI level including solvation ($\varepsilon=5.6$) and dispersion corrections (no parentheses), and without dispersion (in parentheses). Geometrical parameters (lengths in Å and angles in degrees) are shown as the HS [LS] state.

OH Radical Rebound Mechanism. For the H-abstraction pathway, formation of the intermediate complex ($^4$2IM$_H$) is followed by rebound of the phenoxy radical onto the iron-hydroxo species. This occurs via formation of covalent bonds at the ipso-, ortho- or meta-carbon of the aromatic ring of BPA to yield corresponding addition quinol intermediates IM$_{ipso}$, IM$_{ortho}$ or IM$_{meta}$. As shown in Figure 1, all the rebound steps are essentially barrierless in the LS state, while they proceed with significant barriers of 7.8-17.4 kcal/mol on the HS surface. The rebound reactions at the ipso- and ortho-carbon are exothermic for both the HS and LS pathways, with reaction energies of -31.6/-29.5 and -30.3/-32.7 kcal/mol, respectively, while the rebound reactions
at the meta-carbon are endothermic (+4.1/+8.1 kcal/mol). Importantly, the thermodynamically unfavorable rebound reactions associated with this mechanism can explain the lack of experimental detection of the hydroxylation product of the meta-position during P450-dependent metabolism of BPA. Note that the HS rebound barriers are significantly larger than the initial H-abstraction barriers, implying that $^4$Cpd I is a sluggish oxidant and unlikely to play a key role. Thus, OH recombination with the phenyl ring of BPA only occurs via the LS potential energy surface. Accordingly, OH radical rebound will proceed under thermodynamic control, and the reaction energy difference between formation of IM$_{ortho}$ of -32.7 kcal/mol and IM$_{ipso}$ of -29.5 kcal/mol for the LS state of about 3.2 kcal/mol, favors IM$_{ortho}$ formation but also translates into a lower fraction of IM$_{ipso}$ formed. This is in accordance with the observation that metabolites formed via ipso-substitution constitute approximately 20% of the products of the traditional aromatic hydroxylation pathway of P450.

**Decomposition Reaction of the Quinol Intermediate (IM$_{ipso}$) of BPA.** Hydroquinone, isopropenylphenol (IPP), and hydroxycumyl alcohol (HCA) were detected as metabolites upon C-C bond scission via ipso-substitution in experiments of the P450-catalyzed degradation of BPA, which means that the ipso-metabolism reaction of BPA does not stop in the quinol form. In order to understand the complete mechanistic picture, we need to establish the nature of the quinol intermediate decomposition. As mentioned above, there are two types of substituent elimination from the quinol intermediate. While hydroquinone has been detected in the experiments of oxidation of BPA by P450, quinone is also easily transformed to hydroquinone upon NADPH-induced reduction in rat liver microsomes. Therefore, it is difficult to conclude whether the decomposition of the ipso-addition quinol intermediate (IM$_{ipso}$) proceeds via type I or type II elimination based on the available experimental data.
Table 1. Computed Aqueous-Phase Free Energies (ΔG) (kcal/mol) for the Decomposition Reactions of Quinol of BPA

<table>
<thead>
<tr>
<th>Condition</th>
<th>Elimination Type</th>
<th>ΔG (kcal/mol)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neutralization Type I</td>
<td><img src="image" alt="image" /></td>
<td>82.1</td>
</tr>
<tr>
<td>Neutralization Type II</td>
<td><img src="image" alt="image" /></td>
<td>11.1</td>
</tr>
<tr>
<td>Deprotonation Type I</td>
<td><img src="image" alt="image" /></td>
<td>21.8</td>
</tr>
<tr>
<td>Protonation Type II</td>
<td><img src="image" alt="image" /></td>
<td>-30.7</td>
</tr>
</tbody>
</table>

As shown in Table 1, the heterolytic decomposition of IM_{ipso} may proceed charge-neutrally or after protonation or deprotonation in water solution. The computations suggest that the charge-neutral decompositions of IM_{ipso} leading to a carbocation (type II ipso-substitution) or carbanion intermediate (type I ipso-substitution) have reaction energies of +11.1 kcal/mol and +82.1 kcal/mol, respectively. The decomposition of IM_{ipso} after deprotonation (type I ipso-substitution) is endothermic by +21.8 kcal/mol. Thus, the most feasible pathway is decomposition after protonation with production of the carbocationic intermediate and hydroquinone (type II ipso-substitution) with a reaction energy of -30.7 kcal/mol, which supports that the quinol intermediate generated in the P450 enzyme pocket can readily dissociate from the pocket and decompose in a nonenzymatic environment after protonation.
The carbocationic intermediate can react to produce IPP by fast proton transfer to a hydroxyl ion with a reaction energy of -48.7 kcal/mol, or into HCA by absorbing the hydroxyl ion with a reaction energy of -44.0 kcal/mol (using the same method of calculations as for the decomposition of quinols). This mechanism would explain the puzzling observation that no quinol of BPA has ever been detected as an *ipso*-addition metabolite. From our reaction diagrams, it is an unstable intermediate that quickly collapses to the product.

**MBP Formation.** A dimer-type metabolite MBP has been shown to exhibit the highest estrogen activity among all BPA metabolites, and thus we investigated also the MBP formation mechanism. First, we examined the feasibility of the previously suggested radical pathway of MBP formation; this reaction occurs between the isopropenylphenol radical formed by oxidative cleavage of the carbon–phenyl bond, and IPP, as supported by the disappearance of the mass peak of MBP when a radical scavenger was added to the incubation system. However, as shown in Table S31 in the Supporting Information, the cleavage reactions of the carbon–phenyl bond of BPA and the phenoxy radical of BPA in the enzymatic environment are both highly endothermic, and thus the radical pathway seems unfavorable. According to LC/MS/MS investigation, the metabolite of BPA gave a negative mass peak at [M−H]− 267 in LC/MS and a single daughter ion at m/z 133 on MS/MS analysis, corresponding to MBP and IPP, respectively. Alternatively, the dimer-type structure of MBP triggers cationic polymerization, by which the carbocation reacts with IPP, with both reactants originating from the *ipso*-substitution pathway, initiating polymerization and generation of MBP, as shown in eq 1:

\[
\text{+} \rightarrow \text{+} + \text{OH}^− \xrightarrow{ΔG = -47.7 \text{ kcal/mol}} \text{+} + \text{H}_{2}\text{O} 
\] (1)
The obtained reaction energy of $-47.7$ kcal/mol provides a notable driving force for this cationic polymerization pathway to form MBP (using the same method of calculation as for the decomposition of quinols). The P450-catalyzed *ipso*-substitution suggested above proceeds through the radical pathway involving H-abstraction from BPA to produce a phenoxy radical, which would explain why adding a radical scavenger to the incubation system prevents MBP formation during the experiment.

The Reaction Patterns of P450-Catalyzed *ipso*-Position Metabolism

**Initial Rate-Determining Step for the Production of *ipso*-Addition Quinol Intermediates.**

In order to study the detailed reaction mechanism and to verify the initial rate-determining step for *ipso*-position metabolism, we studied several other widely-used phenolic EDCs distributed in the environment such as bisphenol analogues, alkylphenols and chlorophenols with available *in vitro* or *in vivo* assay data on the P450 metabolism.\(^3,14,15,21,23,71\) As shown in Table 2, these phenolic EDCs include bisphenol F (BPF), bisphenol B (BPB), tetrabromobisphenol A (TBBPA), dimethylbisphenol A (DMBPA), bisphenol AF (BPAF), bisphenol Z (BPZ), 4-n-nonylphenol (NP1), *p*-hydroxybenzoic acid (PHBA), *p*-cresol (PC), and *p*-chlorophenol (PCP). The relative energies of the H-abstraction from the phenolic group as well as O-addition at the aromatic ortho-carbon position on the LS potential energy surface are listed in Table 2. The barriers of H-abstraction (0.4-3.1 kcal/mol) are much lower than that for O-addition (14.2-21.0 kcal/mol) for all phenolic EDCs, i.e. the initial step involves H-abstraction by Cpd I from the phenolic group leading to an intermediate complex consisting of an iron-hydroxo group and a phenoxy radical. Within the intermediate complex, as in the reaction of BPA catalyzed by P450, the OH rebounds onto both the *ipso*- and ortho-carbon to form the hydroxylation intermediates with markedly
exothermic reaction energies (-36.3 to -16.5 kcal/mol). The OH rebound barriers for the HS pathway (4.4-13.5 kcal/mol) are much higher than the initial H-abstraction barriers (see details in Table S9 in the Supporting Information), while the OH rebound on the LS pathway is essentially barrier-free. Therefore, we suggest that the P450-catalyzed ipso-position metabolism of these diverse phenolic EDCs follows the same reaction mode as displayed in Figure 1 of BPA, i.e. via H-abstraction followed by an essentially barrierless OH rebound onto the ipso-carbon to produce the corresponding ipso-addition quinol intermediate mainly via the LS state.

As shown in Table 2, compared with the thermodynamic data on OH rebound onto the ortho-positions, the rebound reactions onto the ipso-positions are 2.3 and 2.6 kcal/mol more favorable for PCP and NP1, respectively, but 0.7-9.1 kcal/mol less favorable for all other phenolic EDCs. Although the driving force for ortho-addition relative to ipso-addition is much larger for PBHA, BPAF and TBBPA, the obtained energy difference of 6-9 kcal/mol may still translate into a lower fraction of the ipso-addition quinol intermediates. Regardless of the external factors, we conclude that the P450-catalyzed ipso-position metabolism competes with ortho-position metabolism in the LS state under thermodynamic control. This is consistent with the experiments, in which ipso-substitution/addition metabolites of all studied phenolic EDCs studied in this work were observed in the presence of P450, such as 4-hexafluorohydroxyisopropilidene-phenol from BPAF, and 2,6-dibromo-4-(2-hydroxypropane-2-yl) phenol from TBBPA, which may be produced by the addition of hydroxyl ion to the carbocations as the ipso-substitution products, as well as 4-nonyl-4-hydroxy-cyclohexa-2,5-dienone produced from 4-NP1 as the ipso-addition product. Until now, there are no reported ratios of ipso-addition vs. ortho-addition products for most phenolic EDCs. However, the calculated energy difference between ipso-addition and ortho-addition can be used as a probe for predicting the relative importance of these two pathways.
Table 2. Relative Free Energies (kcal/mol) for P450-catalyzed ipso-Position Metabolism of Phenolic EDCs via the LS state

<table>
<thead>
<tr>
<th>Phenolic EDCs</th>
<th>2TS_H</th>
<th>2TS_Oortho</th>
<th>2IM_H</th>
<th>2IM_ipso</th>
<th>2IM_ortho</th>
<th>ΔG_gap</th>
</tr>
</thead>
<tbody>
<tr>
<td>BPF</td>
<td>2.1</td>
<td>16.6</td>
<td>-6.6</td>
<td>-32.3</td>
<td>-33.0</td>
<td>0.7</td>
</tr>
<tr>
<td>BPB</td>
<td>1.2</td>
<td>14.2</td>
<td>-6.5</td>
<td>-30.5</td>
<td>-34.1</td>
<td>3.6</td>
</tr>
<tr>
<td>TBBPA</td>
<td>0.4</td>
<td>19.9</td>
<td>-6.7</td>
<td>-30.1</td>
<td>-36.3</td>
<td>6.2</td>
</tr>
<tr>
<td>DMBPA</td>
<td>0.4</td>
<td>15.4</td>
<td>-7.8</td>
<td>-30.3</td>
<td>-33.7</td>
<td>3.4</td>
</tr>
<tr>
<td>BPAF</td>
<td>3.1</td>
<td>19.5</td>
<td>0.6</td>
<td>-17.0</td>
<td>-26.1</td>
<td>9.1</td>
</tr>
<tr>
<td>BPZ</td>
<td>1.8</td>
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</tr>
<tr>
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<tr>
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Decomposition Reaction Mechanisms of Diverse Quinol Intermediates. Experimental work on P450-catalyzed phenolic EDCs has shown that ipso-substitution prior to ipso-addition does not always occur. However, the reason why some phenolic EDCs are stopped at the ipso-addition step is unknown. It is also difficult to determine which type of elimination (type I or type II) occurs during ipso-substitution due to the complex biological redox environment. We focused on the decomposition mechanisms of the diverse ipso-addition quinol intermediates.
derived from the diverse phenolic EDCs described above with the available experimental information, but excluded TBBPA and BPZ, for which our attempts to locate the quinol intermediates after protonation give fragmental type II products directly. The thermodynamic data for the decomposition of quinol intermediates in all possible pathways were evaluated and the most favorable decomposition paths via type II and type I ipso-substitution are shown in Figure 2.

Figure 2. Computed free energies (kcal/mol) for the decomposition reactions of diverse ipso-addition quinols along the favorable pathways: (left) type II substitution with the hydride ion affinity (HIA, kcal/mol) of the formed carbocation; (right) type I substitution. *R represents the elimination substituent.

As shown in Figure 2, for all bisphenol analogues and alkylphenols except for PHBA, the decomposition of the formed ipso-addition quinols after protonation with formation of carbocation and hydroquinone (type II substitution) is the most favorable pathway. The decomposition
reactions for the ipso-addition quinols from PC and NP1 are distinctly endothermic, which is fully in line with experimental observations of the P450-catalyzed conversion of these two alkylphenols showing only ipso-addition quinols were produced without detecting any ipso-substitution products. However, for other bisphenol analogues and alkylphenols, the decomposition of the formed ipso-addition quinols after protonation can proceed, leading to C-C bond cleavage with significant exothermic energies. It is found that the P450-catalyzed ipso-substitution products are obtained from the ipso-addition quinols when the carbon at the benzylic position contains one or more alkyl branches. More alkyl branches stabilize the carbocation via inductive and hyperconjugative effects; this results in the spontaneous decomposition of the formed ipso-addition quinols after protonation. The hydride ion affinity (HIA) can be used for comparing the carbocation stability of dissimilar structures directly, defined according to eq (2):

\[ \text{RH} \rightarrow \text{R}^+ + \text{H}^- \quad \Delta H^0 = \text{HIA} \]  

The HIA obtained at the B3LYP/6-311++G** level using frequency analysis at 298.15 K and 1 atm pressure are listed in the lower left of Figure 2. The experimental HIA is available for CH$_3^+$ (312 kcal/mol), the same as the computed HIA of 312 kcal/mol, which supports the reliability of the computational method. The reaction free energies of decomposition of the quinol intermediates generally increase with increasing HIA of the formed carbocations ($r^2 = 0.95, \Delta G = 1.4 \text{HIA} + 245.5$). This pattern indicates that the HIA values are useful for preliminary evaluation of the decomposition free energies of the ipso-addition quinols produced from bisphenol analogues and alkylphenols with associated formation of a carbocation and a hydroquinone (type II substitution).

For quinol intermediates with electronegative substituents, such as -Cl and -COOH, as shown in Figure 2, there are two possible pathways for substituent elimination from quinol with the formation of an anion and a quinone (type I ipso-substitution): 1) elimination of the substituent
after deprotonation with the formation of an anion and quinone; 2) involving the prior intra-molecular H-arrangement from OH to the electronegative substituents to produce the corresponding inorganic acid and quinone neutrally. The charge-neutral intra-molecular H-arrangement pathway with formation of the inorganic acid and quinone is more favorable for decomposition of quinol intermediates with electronegative substituents; in this case the inorganic acid can dissociate into an anion. The pathway we have obtained for type I ipso-substitution extends the formal definition of type I ipso-substitution in P450 chemistry. In particular, the elimination of -COOH from quinol after deprotonation is not feasible because it is endothermic, while the exothermic elimination of -COOH during the intra-molecular H-arrangement route is favorable. This is in accordance with the experimental observation that PHBA can be subject to ipso-substitution when the reaction is catalyzed by P450.21

Environmental Implications

Identification of EDCs is one of the most important goals of environmental chemical hazard screening, which has come a long way in developing useful test assays and mechanism-based screening techniques.10 Many synthetic compounds released into the environment may be readily transformed, especially by P450 enzymes, into metabolites exhibiting much higher endocrine-disrupting activity than their parent compounds. Knowledge of detailed metabolic mechanisms gives insight into the bioactivation. Accordingly, it is critical in environmental risk assessment to understand metabolic pathways and to have effective tools for predicting the fate of metabolites. Methods that analyze and predict the metabolic fate of molecules thrive within the field of medicinal chemistry,73 but not so much within environmental sciences despite the similarity of involved tools. In medicinal chemistry, many drugs require P450-mediated bioactivation to elicit their pharmacological effect via metabolites that can be characterized in relatively high
concentrations. In contrast, environmental pollutants such as EDCs and their metabolites normally occur in trace amounts while still important at these levels, and thus identification of their biotransformation products seems more difficult, and mechanism-based methods to provide putative metabolites efficiently are of interest. Experimental methods often require expensive equipment, expertise, running costs and time, which may reduce their applicability when screening large libraries of compounds. Thus, there is substantial interest in the development of fast, accurate computational tools that can predict metabolism with higher throughput and lower cost. These computational tools should: (i) predict the site of metabolism and (ii) predict the metabolite structure from these sites.\(^{74}\)

The present work shows how detailed DFT investigations of metabolic pathways can rationalize the formation of metabolites resulting from the P450-catalyzed reactions of diverse environmental phenolic EDCs such as bisphenol analogues, alkylphenols and chlorophenols, thereby achieving these two tasks, as particularly emphasized for one of the prominent phenolic EDCs, BPA. The barrier for the most favorable H-abstraction/OH-rebound mechanism involving both the ipso- and ortho-position hydroxylation is one of the lowest reported barriers, as far as we know. The H-abstraction/OH-rebound reaction with formation of the quinol intermediate seems to be a general reaction mechanism for phenolic EDCs, as shown by studying a diverse group of such compounds in this work. In case of the ipso-addition quinol intermediate, we can distinguish type II vs. type I ipso-substitutions based on thermodynamic data, and ipso-substitution vs. ipso-addition based on the stability of the eliminating carbocation by both qualitative and quantitative analysis. Notably, the formation mechanism of the highly estrogenic metabolites HCA and dimer-type MBP, which arises from oxidation of BPA catalyzed by P450, has been revealed in detail. Our results show that both metabolites originate from a carbocationic intermediate produced in the
**ipso**-substitution pathway. This pathway gives insight into the potentially important bioactivation of many other alternatives to BPA whose metabolic mechanisms remain unidentified, in particular under conditions where P450-catalyzed metabolism is important relative to glucuronidation (e.g. if this pathway is inhibited or genetically or otherwise down-regulated, e.g. in the fetus). However, even when non-P450 pathways dominates by 10-, 100- or even 1000-fold, the **ipso**-position metabolites may still contribute to toxicity due to their correspondingly higher potency.

The hydroxylated metabolites of many emerging phenolic pollutants, such as **OH**-PBDEs and **OH**-PCBs, have been reported to be even stronger EDCs than their precursors\(^ {75,76} \) and based on their similar molecular structures we speculate that they may involve products from the **ipso**-substitution/addition pathway catalyzed by P450, which has thus far largely been neglected. Recently, the biotransformation of sulfonamide antibiotics in the environment has been reported to proceed via the **ipso**-substitution pathway\(^ {77} \). Therefore, **ipso**-substitution seems to be a much more common and, even at low turnover, more important toxification pathway than previously thought for a wide variety of persistent pollutants. Our study has identified the detailed electronic structure changes and transition states probably involved in these processes, as well as provided simple tools for determining the relative importance of these pathways based on thermodynamic considerations that we envision will be valuable for determining the environmental toxicity and fate of emerging phenolic EDCs.

**ASSOCIATED CONTENT**

**Supporting Information.** Full citation for reference 73; Energies for all molecular species; Intrinsic reaction coordinate (IRC) for verifying transition states; Optimized geometries at the
B3LYP/BSI** level of theory; Quantum chemical cluster calculations; Cartesian coordinates of all structures. This material is available free of charge via the Internet at http://pubs.acs.org.

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**Notes**

The authors declare no competing financial interest.

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