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Structural Identification and Kinetic Analysis of the \textit{in Vitro} Products Formed by Reaction of Bisphenol A-3,4-quinone with \(N\)-Acetylcysteine and Glutathione

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Supporting Information

**ABSTRACT:** Bisphenol A (BPA) has received considerable attention as an endocrine disrupting chemical and a possible substrate for genotoxic metabolites. BPA metabolism leads to formation of electrophilic \(o\)-quinones capable of binding to DNA and other endogenous nucleophiles. We have structurally identified the products resulting from the reaction of bisphenol A-3,4-quinone (BPAQ) with \(N\)-acetylcysteine (NAC) and glutathione (GSH). The major and minor isomers are both the result of 1,6-conjugate addition and are produced almost instantly in high yield. Reactions using 1.3 equiv of GSH showed the presence of a bis-glutathionyl adduct which was not observed using higher GSH concentration relative to BPAQ. NAC reactions with BPAQ showed no bis-\(N\)-acetylcysteinyl adducts. Stopped-flow kinetic analysis reveals the 1,6-conjugate additions to be reversible with a forward free energy of activation of 9.2 and 7.8 kcal/mol for the NAC and GSH reactions, respectively. The bimolecular forward rate constant at 19.4 °C was approximately three time faster for GSH compared to NAC, 1547 vs 496 M\(^{-1}\) s\(^{-1}\). The free energy of activation for the reverse reactions were similar, 11.7 and 11.2 kcal/mol for NAC and GSH, respectively. We plan to use this model system to further explore the mechanism of adduct formation between sulfur nucleophiles and \(o\)-quinones and the resulting chemical properties of both NAC and GSH adducts.

**INTRODUCTION**

Bisphenol A (BPA) is a widely used industrial chemical prominent in the manufacture of plastics and epoxy resins.\(^1\) BPA and its metabolites can act as estrogen mimics and are classified as endocrine disrupting chemicals.\(^2\) Evidence for genotoxic effects in vitro have been revealed in studies involving mouse lymphoma,\(^3\) MCF-7\(^4\) and CHO-K1\(^5\) cells. In vivo studies showed structural changes in bone marrow cells of mice.\(^6\)

The major metabolites of BPA include glucuronic acid conjugates and catechols.\(^7\) Catechols can undergo further oxidation to \(o\)-quinones (Scheme 1), known genotoxic compounds capable of binding to the nucleosides of DNA. The mono-\(o\)-quinone of BPA, BPA-3,4-quinone (BPAQ), reacts with deoxyguanosine via 1,4-conjugate addition.\(^8,9\) BPAQ-DNA adducts have been detected \textit{in vitro} and \textit{in vivo} with either BPAQ exposure or metabolically activated BPA.\(^10,11\) BPAQ chemistry has shown to be analogous to estrogen \(o\)-quinones, known genotoxic metabolites.\(^12\) Glutathione (GSH) is a prominent \(o\)-quinone scavenger. Several studies have shown the production BPAQ-GSH adducts \textit{in vitro} as analyzed by LC/MS\(^9,13\) which confirmed the presence of catechol-GSH products but did not indicate the regiochemistry of the conjugate addition, namely 1,4- versus 1,6-addition. Soft sulfur nucleophiles show a preference for 1,6-addition with various \(o\)-quinones, but 1,4-addition products are also observed.\(^14,15\)

Recently we have developed a large-scale synthetic approach to the BPA metabolites, BPAQ, the di-\(o\)-quinone of BPA, the monocatechol of BPA, and the dicatechol of BPA.\(^16\) With this
new synthetic technique at hand, we sought to determine the major product(s) when BPAQ reacts with NAC and GSH. Scheme 2 shows the three possible conjugate addition products when BPAQ reacts with these sulfur nucleophiles. Products 2-SR and 5-SR are the result of 1,6-addition with respect to BPAQ C4 and C3 carbonyls, respectively. Adduct 6-SR would result from 1,4-addition with respect to BPAQ C4 carbonyl.

BPAQ is a relatively stable α,quinone with a deep red color in organic solvents. We recognized that the stability of BPAQ combined with its chromophore properties would make kinetic analysis of the conjugate addition possible. Conjugate addition of sulfur nucleophiles to certain quinones has shown reversible characteristics indicating that sulfur detoxification may lead to other metabolic processes.17−19 Second-order rate constants for both NAC and GSH conjugate addition to various Michael acceptors have been measured and shown to be a function of both steric and electronic factors of the electrophile in addition to the pKs of the thiol nucleophile.20−22

Herein we described the synthesis and full structural characterization of the NAC and GSH adducts of BPAQ. In addition, the measurement of second-order rate constants using stopped-flow techniques and the resulting thermodynamic parameters are presented to establish a better understanding of the reversible mechanism involved with thiol Michael additions to α-quinones.

### EXPERIMENTAL PROCEDURES

**Chemicals and Reagents.** 2-Iodoxybenzoic acid (IBX) was synthesized by the method of Frigerio.23 Caution: IBX is explosive under impact if heated to more than 200 °C. BPAQ was made by IBX oxidation of BPA24 with modifications to product isolation (see below). Deuterated NMR solvents were purchased from Cambridge Isotope (Andover, MA). All other chemicals and solvents purchased from Fisher Scientific Co. (Fair Lawn, NJ) or Aldrich Chemical Co. (Milwaukee, WI) and used as received.

**Instrumentation.** NMR analysis was obtained on a Bruker 400 MHz Avance III spectrometer (Bruker, Billerica, MA, USA). HPLC was conducted on a Waters 2690 Separations Module (Waters Corp. Milford, MA, USA) equipped with a Waters 2487 dual λ absorbance detector using solvent gradients consisting of acetonitrile and 0.5% aqueous formic acid on a YMC, ODS-AQ C18 analytical column (5 μm, 120 A, 250 mm × 4.5 mm, YMC America, Allentown, PA, USA). Flash column purification of compounds was performed on a Biotage Isoler One (Biotage, Charlotte, NC) automated system using Biotage KP-Shere 30 g and SNAP Ultra HP 12 g C18 columns. Stopped-flow analysis was done on TgK Scientific, KinetAsyst system (TgK Scientific, Bradford-on-Avon, UK). Reaction traces were fit to a sum of three exponentials using KaleidaGraph (Synergy Software, Reading, PA, USA). HRMS analysis was conducted at the Nebraska Center for Mass Spectrometry (University of Nebraska-Lincoln, Lincoln, NE). Elemental analysis was obtained at MHW Laboratories (Phoenix, AZ).

**Synthesis of 3-Hydroxy-5-NAC-BPA and 3-Hydroxy-2-NAC-BPA.** BPAQ (0.250 g, 1.10 mmol) was dissolved in 50 mL of a solvent consisting of 3:2 CHCl₃:CH₃OH in a 100 mL round-bottom, equipped with a magnetic stir bar and brought to −15 °C. To this solution was added IBX 0.337 g (1.20 mmol) causing the solution to turn deep red after approximately 15 min. The mixture was stirred at −15 °C for 12 h and transferred to a 250 mL separatory funnel, and the chloroform layer was washed 6 × 50 mL with a 0.1 M phosphate buffer, pH 6.0, to remove the α-iodoxybenzoic acid byproduct. The chloroform solvent was removed via rotovap, and the crude was placed in 50 mL of methanol. HPLC analysis of the crude methanol solution showed >95% BPAQ which was used without further purification. N-Acetylcysteine (235 mg, 1.43 mmol) was dissolved in 5 mL of 0.1 M phosphate buffer, pH 6.2, and added in one portion to the BPAQ solution. The mixture decolorized instantly from bright red to a light yellow color. The mixture was stirred an additional 10 min at rt, an HPLC sample was removed, and all solvents removed via rotovap. The crude mixture was dissolved in 1−2 mL of methanol and placed on a preconditioned Biotage 30 g C18 flash column (see Figure S1 for flash chromatograph with solvent gradient). Each collection tube was analyzed by HPLC on an analytical column, and three fractions were collected: one containing analytical grade 5-NAC, a middle fraction containing a mixture of 5-NAC and 2-NAC, and fraction containing analytical grade 2-NAC. The middle fraction was subject to a second separation using a 12 g C18 SNAP Ultra column, solvents were removed, and solids dried overnight under high vacuum at 50 °C to afford a total of 319 mg (81%) of 3-hydroxy-5-NAC-BPA and 27 mg (7%) of 3-hydroxy-2-NAC-BPA.

**3-Hydroxy-5-NAC-BPA.** Needle white solid; H NMR (DMSO-d₆, 400 MHz): δ 1.52 (s, 3H), 1.84 (s, 3H), 2.95 (dd, J = 13.4 and 9.2 Hz, 1H), 3.32 (dd, J = 13.4 and 4.7 Hz, 1H), 4.28 (m, 1H), 4.63 (d, J = 2.2 Hz, 1H), 6.65 (d, J = 8.7 Hz, 2H), 6.68 (d, J = 2.2 Hz, 1H), 7.00 (d, J = 6.7 Hz, 2H). For 1H spectrum with structure assignment see Supporting Information, p S-4. 13C NMR (DMSO-d₆, 100 MHz): δ 22.39, 30.71, 30.73, 33.76, 41.14, 51.70, 113.65, 114.59, 118.72, 119.69, 127.30, 140.81, 141.90, 142.17, 144.60, 154.95, 169.27, 172.27. For 13C spectrum with structure assignment see S-5. Anal. calcd for C₂₀H₂₃NO₆S: C, 59.25; H, 5.72; N, 3.45; S, 7.91. Found: C, 59.45; H, 5.84; N, 3.30; S, 8.18.

**3-Hydroxy-2-NAC-BPA.** Needle white solid; H NMR (DMSO-d₆, 400 MHz): δ 1.58 (s, 3H), 1.62 (s, 3H), 1.83 (s, 3H), 2.51 (m, 1H, peak obscured by residual DMSO), 2.69 (dd, J = 12.5 and 4.7 Hz, 1H), 4.04 (m, 1H), 6.57 (d, J = 8.5 Hz, 1H), 6.76 (d, J = 8.9 Hz, 1H), 6.79 (d, J = 8.5 Hz, 1H), 6.86 (d, J = 8.9 Hz, 1H), 7.98 (d, J = 7.8 Hz, 9.17 (bs, 3H phenolic OH partially exchanged). For 1H spectrum with structure assignment see S-10. 13C NMR (DMSO-d₆, 100 MHz): δ 22.34, 30.96, 42.80, 52.05, 114.03, 114.44, 116.73, 142.12, 143.05, 143.74, 147.69, 154.26, 169.16, 172.13. For 13C spectrum with structure assignment see S-11. HRMS (ES+): calculated for C₂₀H₂₃NO₆S [M + H]+: 406.1246; found: 406.1253.

**Synthesis of 3-Hydroxy-2,5-diGSH-BPA, 3-Hydroxy-5-GSH-BPA, and 3-Hydroxy-2,5-diGSH-BPA.** A 1.10 mmol solution of BPAQ was made as above except after workup, BPAQ was dissolved in 50 mL of 50:50 methanol:water solvent system. GSH (572 mg, 1.43 mmol) was dissolved in 5 mL of 0.1 M phosphate buffer, pH 6.2, and added in one portion to the BPAQ solution. The mixture was stirred at rt for 15 min, and solvents removed via rotovap. The crude solid was dissolved in 50 mL of water and extracted 3 × 25 mL with ethyl acetate to remove the 3-hydroxy-BPAQ reduction byproduct. The ethyl acetate extract was dried over magnesium sulfate, and solvent removed via rotovap. The aqueous phase was then reduced to 2 mL via rotovap and placed on a pre-conditioned Biotage 30 g C18 SNAP Ultra flash column (see Figure S2 for flash chromatograph with solvent gradient). Each collection tube was analyzed by HPLC on an analytical column, and four fractions containing pure 2,5-diGSH, a mixture of 2,5-diGSH and 5-GSH, pure 5-GSH, and pure 2-GSH were collected.
The fraction containing the 2,5-GSH/5-GSH mixture was subjected to a second separation using a Biotage 30 g C18 SAPN Ultra flash column, solvents were removed, and solids dried overnight under high vacuum at 50 °C to afford a total of 151 mg of 3-hydroxy-2,5-diGSH-BPA (17%), 212 mg of 3-hydroxy-5-GSH-BPA (36%), 43 mg of 3-hydroxy-2-GSH (7%), and 81 mg of 3-GSH-BPA (32%).

### 3-Hydroxy-5-GSH-BPA

The 1H NMR (DMSO-d6, 400 MHz) spectrum showed the following changes: BPAQ concentration 65 μM; GSH concentrations 40, 20, and 10 mM.

1H NMR spectrum:

- δ 1.51 (s, 3H), 1.63 (s, 3H), 1.91 (m, 2H), 2.50 (t, 2H, J = 7.0 Hz), 2.41 (dd, 1H, J = 12.0 and 10.1 Hz), 2.62 (dd, 1H, J = 12.1 and 4.7 Hz), 3.40 (t, 1H, J = 6.4 Hz), 3.65 (m, 2H), 4.16 (m, 1H), 6.59 (d, 2H, J = 8.7 Hz), 6.76 (d, 1H, J = 8.1 Hz), 6.79 (d, 2H, J = 8.7 Hz), 6.86 (d, 1H, J = 8.1 Hz), 8.05 (d, 1H, J = 7.5 Hz), 8.52 (t, 1H, J = 5.6 Hz).

### Results

#### Reaction of BPAQ with NAC and NMR Analysis of Mercapturate Products

When BPAQ was reacted with an aqueous solution of NAC (1.3 equiv), the red BPAQ solution is observed. Reaction traces were monitored at 400 MHz, and different temperatures: BPAQ concentration 65 μM; 1H NMR (D2O, 400 MHz) signal assignments are given in Table 1.

1. **3-Hydroxy-2-GSH-BPA**: Off-white solid; 1H NMR (DMSO-d6, 400 MHz): δ 1.58 (s, 3H), 1.63 (s, 3H), 1.91 (m, 2H), 2.50 (t, 2H, J = 7.0 Hz), 2.41 (dd, 1H, J = 12.0 and 10.1 Hz), 2.62 (dd, 1H, J = 12.1 and 4.7 Hz), 3.40 (t, 1H, J = 6.4 Hz), 3.65 (m, 2H), 4.16 (m, 1H), 6.59 (d, 2H, J = 8.7 Hz), 6.76 (d, 1H, J = 8.1 Hz), 6.79 (d, 2H, J = 8.7 Hz), 6.86 (d, 1H, J = 8.1 Hz), 8.05 (d, 1H, J = 7.5 Hz), 8.52 (t, 1H, J = 5.6 Hz).

2. **3-Hydroxy-5-GSH-BPA**: Off-white solid; 1H NMR (DMSO-d6, 400 MHz): δ 1.57 (s, 3H), 1.63 (s, 3H), 1.91 (m, 2H), 2.51 (t, 2H, J = 7.0 Hz), 2.41 (dd, 1H, J = 12.0 and 10.1 Hz), 2.62 (dd, 1H, J = 12.1 and 4.7 Hz), 3.40 (t, 1H, J = 6.4 Hz), 3.65 (m, 2H), 4.16 (m, 1H), 6.59 (d, 2H, J = 8.7 Hz), 6.76 (d, 1H, J = 8.1 Hz), 6.79 (d, 2H, J = 8.7 Hz), 6.86 (d, 1H, J = 8.1 Hz), 8.05 (d, 1H, J = 7.5 Hz), 8.52 (t, 1H, J = 5.6 Hz).

3. **3-Hydroxy-2,5-diGSH-BPA**: Off-white solid; 1H NMR (DMSO-d6, 400 MHz): δ 1.49 (s, 3H), 1.52 (s, 3H), 2.03–2.23 (m, 6H), 3.16 (dd, 1H, J = 14.5 and 8.2 Hz), 3.34 (dd, 1H, J = 14.4 and 5.0 Hz), 3.72 (m, 2H), 3.86 (m, 2H), 3.95 (m, 2H), 4.03 (dd, 1H, J = 8.2 and 5.9 Hz), 4.35 (dd, 1H, J = 8.0 and 5.0 Hz), 6.66 (d, 2H, J = 8.8 Hz), 6.95 (d, 2H, J = 8.7 Hz).

### HPLC Chromatography

Figure 1 shows the HPLC chromatogram of the crude mixture indicating the presence of two product peaks along with trace amounts of reduced 3-hydroxyBPA (3-OHBPA). The major product (89%) was 5-NAC, and the minor product was 2-NAC (11%).

#### Reaction of BPAQ with GSH and NMR Analysis of Glutathione Adducts

When BPAQ was reacted with an aqueous solution of GSH under similar conditions used with NAC, several significant differences were observed. The chromatogram in Figure 3 shows the presence of four significant peaks when 1.3 equiv of GSH was mixed with BPAQ. The peak corresponding to 3-OHBP, the result of BPAQ reduction, is observed in much larger quantities when compared to the NAC reaction. Moreover, an early eluting peak was identified as the bis-glutathionyl adduct of BPAQ, 2,5-diGSH. Figure 3 also shows reactions done a smaller, 0.5 mM.

Protons H5 and H6 on the catechol ring would show a distinct coupling pattern when compared to 5-SR or 6-SR. Discrimination between 5-NAC and a possible 6-NAC isomer required a combination of both 1-D and 2-D NMR analysis. Heteronuclear (C–H) multiple bond correlation (HMBC) proved to be the most direct 2-D NMR technique in establishing the structure of 5-NAC from 6-NAC. HMBC optimized for three bond correlations would show three cross peaks between the quaternary, saturated carbon connecting the two arene rings to arene protons in structure 5-NAC, while 6-NAC would show only two correlations. Figure 2A shows the portion of the HMBC spectrum where these correlations occur, clearly showing three cross peaks consistent with isomer 5-NAC. Several noteworthy differences in chemical shifts between the 5-NAC and the 2-NAC isomers were also observed. The methylene protons alpha to the sulfur atom on the NAC moiety were shifted downfield relative to NAC on the 5-NAC adduct, while the same signals of the 2-NAC isomer were shifted upfield relative to NAC (Figure 2B). The proton beta to the sulfur atom in the 2-NAC isomer displayed a significant upfield shift relative to NAC, while the 5-NAC shifted upfield slightly. The 5-NAC isomer showed significant through-space NOE correlations between arene proton H6 to both the aforementioned alpha and beta protons. These observations proved useful in establishing the structure of the larger GSH adducts (vida infra).
scale of BPAQ with increasing equivalents of GSH. At 5-fold excess of GSH, little of the di-2,5-GSH adduct or the reduced 3-OHBPA catechol were observed. Only monoadducted adducts were observed for NAC when using 1.3, 2.0, or 5.0 equiv of NAC. For the large-scale reaction using 1.3 equiv of GSH, isolated yields produced 17% 2,5-diGSH, 36% 5-GSH, 7% 2-GSH, and 32% 3-OHBPA. NMR analysis of the 5-GSH and the 2-GSH adducts was similar to the NAC adducts. The NMR properties of the NAC adducts were useful in establishing the location of the two GSH moieties in 2,5-diGSH; specifically, distinguishing between a 2,5-diadduct versus a 2,6- or 5,6-diadduct. First, two resolved sets of methylene protons alpha to the sulfur atom were observed: one shifted downfield and one shifted upfield relative to GSH. This is consistent with a 2,5-disubstitution profile (vida supra, Figure 2B). Also, through-space NOE experiments shows only one of these methylene signals correlates to the lone remaining arene proton on the catechol ring, specifically the methylene protons shifted downfield. Thus, the 2,5-diGSH adduct is the result of two separate 1,6-conjugated additions. No products, mono- or diadducted, were the result of 1,4-addition conjugate addition.

Kinetic Analysis of the BPAQ-NAC Reaction and BPAQ-GSH Reaction. Using the distinct chromophore of the BPAQ, we sought to explore the kinetics of NAC and GSH conjugate addition to BPAQ by following the disappearance of BPAQ at 400 nm. Figure 4A shows the decrease in the absorbance when a 125 μM solution of BPAQ in 50% methanol was mixed with various concentrations of NAC (0.050, 0.10, and 0.125 M; final concentration) in 0.050 M phosphate buffer pH 6.2 and 50% methanol using a stopped-flow spectrophotometer. To simplify the exponential decay, the reaction was made pseudo-first order using a large excess of NAC. A similar process was performed using GSH using more dilute solutions due to the reduced solubility of GSH in 50% methanol when compared to NAC. Fitting software indicates the decay is the sum of three different exponentials, indicating three separate phases. The first phase, which represents 86% of absorbance decrease, is both NAC concentration and temperature dependent. The second phase is NAC concentration independent and temperature dependent. The third phase is both temperature and NAC concentration independent, which suggests it is not part of the reaction process but an artifact of how components were mixed. Reaction with GSH displayed the same three exponential decays.

Analysis of the first phase was carried out by plotting the observed rate constants as a function of NAC or GSH concentration generating liner plots, the slope of which corresponds to the forward rate constants, $k_{\text{f}}$, of the bimolecular conjugate addition (Figure 4B). Nonzero intercepts indicates a reversible process, and the value of these nonzero intercept represents the reverse rate constant, $k_{-1}$. Both NAC reactions and GSH reaction displayed a reversible profile with GSH forward rate constants approximately three times faster than NAC. Each NAC concentration was carried out at four temperatures to obtain Arrhenius plots of both the forward and reverse reactions. Only three different temperatures were conducted with GSH as insolubility limited lower temperatures (Figures S3 and S4). The free energy of activation for the forward reaction was calculated at 9.2 and 7.8 kcal/mol for the NAC and GSH reactions, respectively. These $\Delta G^\ddagger$ values are similar to computational calculations, 10.0 kcal/mol,
that model the reaction of GSH with BPAQ even though the modeled reaction assumed a 1,4-conjugate addition. Error analysis for the line slopes was between 4 and 6%, while the \( y \)-intercepts were between 7 and 10%; the final \( \Delta G^\ddagger \) values reported are ca. \( \pm 0.5 \) kcal/mol. The free energy of activation for the reverse process was similar for both NAC and GSH, 11.7 and 11.2 kcal/mol, respectively (see Figure 4C). The kinetic data of the first phase indicate a reversible 1,6-addition, which is NAC concentration dependent. The second phase observed, Figure 4A, is likely from the proton transfers involved in the tautomization to the final catechol product. These proton transfers would be temperature dependent while independent of the concentration of NAC, which is consistent with our data.

Discussion. If BPAQ is a significant metabolite formed by exposure to BPA, then the most abundant biomarker would likely be mercapturates in the form of the NAC adducts. This work provides a simple and efficient methodology in the synthesis of both the NAC and GSH adducts of BPAQ. Both nucleophiles showed exclusive preference for 1,6-addition in lieu of 1,4-addition. The formation of a bis-glutathionyl adduct with GSH as opposed to NAC, which showed no bis-N-acetylcysetinyl adducts, is of interest in that bis-adduct formation implies differences in the oxidation potential of GSH and NAC adducts. The mechanism for bis-adduct formation likely involves oxidation of the newly formed catechol adduct by the unreacted \( o \)-quinone, BPAQ (Scheme 3). This would explain the concurrent increase of 3-OHBPA observed in the GSH reaction where 2,5-diGSH is produced as compared to the NAC reaction where very little of the 3-OHBPA catechol is observed. This also implies that the 5-GSH adduct has a lower oxidation potential when compared to the 5-NAC adduct. Macedo et al., using cyclic voltammetry, showed GSH adducts formed from the \( o \)-quinones of \( \alpha \)-methyldopamine and \( N \)-methyl-\( \alpha \)-dopamine had lower oxidation potentials (30 to 65 mV), when compared to the corresponding NAC adducts. This suggests that GSH imparts a stronger electron-donating effect when compared to NAC increasing \( k_2 \) in Scheme 3, which facilitates formation of 2,5-diGSH. If \( k_1 \) is greater than \( k_2 \), increasing GSH concentration relative to BPAQ would decrease the rate of 5-GHS-BPAQ formation and subsequent formation of 2,5-diGSH.

Kinetic data show the reversible nature of conjugate addition of both NAC and GSH to BPAQ. The forward rate constant for GSH reaction was significantly faster than NAC reaction. Freeman et al., also using stopped-flow analysis, observed much
faster conjugation adduct of GSH relative to NAC with αβ-un saturated nitro compounds derived from fatty acids and postulated this was a function of thiol acidity.21 GSH, pKa 8.8, is more acidic than NAC, pKa 9.5.20 Thus, at a pH of 6.2, the effective concentration of thiolate ion would be much larger with GSH solutions compared to NAC. Properties of other GSH and NAC adducts have suggested that the sulfur adduction process is reversible. GSH adducts formed between the o-quinone of 4-hydroxyequilenin, which exists in the oxidized o-quinone form, liberated 4-hydroxyequilenin and GSH when exposed to NADPH reduction.31 Isomeric GSH adducts of the o-quinone formed from quercetin oxidation were shown to interconvert when allowed to equilibrate in solution.30 GSH adducts of allyl and benzyl isothiocyanate conjugation were shown to release the isothiocyanate electrophile in vitro.31 GSH conjugation is generally considered a detoxification process used to sequester reactive electrophiles such as o-quinones. However, GSH conjugates formed from various quinones display toxicity in certain tissues, and many of these GSH adducts stem from p- or o-quinones.21−25 Is there a correlation between the reversible nature of quinone conjugate addition and the subsequent toxicity of GSH adducts? The BPAQ adducts synthesized in this work, efficiently made at large scale and generated from a relatively stable quinone with a distinctive chromophore, could serve as a platform for further studies to this effect.

■ ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.chemrestox.7b00239.

Chromatography conditions for the separation of NAC and GSH adducts; 1-D and 2-D NMR spectra of NAC and GSH adducts; Arrhenius plots used to calculate free energies of activation (PDF)

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Notes

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■ ABBREVIATIONS

BPA, bisphenol A; BPAQ, bisphenol A-3,4-quinone; NAC, N-acetylcysteine; GSH, glutathione; IBX, 2-iodobenzoic acid; HMBC, heteronuclear multiple bond correlation; 3-OHBPA, 3-hydroxybisphenol A

■ REFERENCES


