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Article

Photoprotective Properties of Eumelanin: Computational Insights into the Photophysics of a Catechol:Quinone Heterodimer Model System

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Abstract: Melanins are skin-centered molecular structures that block harmful UV radiation from the sun and help protect chromosomal DNA from UV damage. Understanding the photodynamics of the chromophores that make up eumelanin is therefore paramount. This manuscript presents a multi-reference computational study of the mechanisms responsible for the experimentally observed photostability of a melanin-relevant model heterodimer comprising a catechol (C)–benzoquinone (Q) pair. The present results validate a recently proposed photoinduced intermolecular transfer of an H atom from an OH moiety of C to a carbonyl-oxygen atom of the Q. Photoexcitation of the ground state C:Q heterodimer (which has a π-stacked “sandwich” structure) results in population of a locally excited ππ* state (on Q), which develops increasing charge-transfer (biradical) character as it evolves to a “hinged” minimum energy geometry and drives proton transfer (i.e., net H atom transfer) from C to Q. The study provides further insights into excited state decay mechanisms that could contribute to the photostability afforded by the bulk polymeric structure of eumelanin.

Keywords: photophysics; photoprotection; photostability; eumelanin; catechol; benzoquinone; ultraviolet; conical intersection

1. Introduction

The fundamental photochemistry of prototypical organic and biological chromophores is attracting ever more attention [1–3]—driven, in part, by ambitions to advance understanding (and prevention) of photoinduced damage in biomolecules [4–11] and to improve the photoprotection offered by sunscreen molecules [12–15]. Ultraviolet (UV) excitation of any given molecule increases its total energy, typically to values in excess of many of the energy barriers associated with reaction on the ground state potential energy (PE) surface. The excited state molecules formed upon UV absorption may decay in a number of ways that have traditionally been illustrated using a Jablonski diagram. Extremes of these behaviors include:

(i) Photoreaction, by photodissociation, which constitutes the dominant decay mechanism for many small heterocyclic molecules in the gas phase. Much studied examples include phenol [2,4,16–25], pyrrole [26–34] and indole [35–38];

(ii) Photostability, wherein the photoexcited molecule decays back to the ground state, rapidly and with high efficiency, without any permanent chemical transformation. Such non-radiative decay (generically termed internal conversion) is the desired photophysical response for the DNA/RNA nucleobases [4,39–65] and, for example, for derivatives of the p-aminobenzoates, cinnamates, salicylates, anthranilates, camphor, dibenzoyl methanes and/or benzophenones used in commercial sunscreens [13,14]. Internal conversion processes are mediated by conical intersections (CIs) — regions of the PE surfaces where distinct
electronic states become energetically degenerate [66,67]. These points of degeneracy develop into CIs when orthogonal motions are considered, which facilitate non-adiabatic coupling (i.e., the funnelling of population) from the photoexcited state to a lower (e.g., the ground) state.

Sunscreen molecules are chosen on the basis of good photostability. Those that find use in commercial sunscreen products have been chosen/engineered to absorb in the UV-A/B regions of the electromagnetic spectrum and, following photoexcitation, to undergo efficient non-radiative decay back to the ground state—releasing the excess energy as local heating in the formulation of which they are a part. In oxybenzone, for example, \( \pi^* \rightarrow \pi \) excitation of the dominant (in the ground \( S_0 \) state) enol-conformer populates the strongly absorbing \( ^1\pi\pi^* \) state, which relaxes—either directly or via an (optically dark) \( ^1\pi\pi^* \) state—toward its minimum energy keto-configuration (i.e., an intramolecular H atom transfer (HAT) process) and onwards towards the CI with the \( S_0 \) state at non-planar geometries (one ring twists relative to the other about the central aliphatic C–C bond). A reverse H atom transfer process on the \( S_0 \) PES and vibrational energy transfer to the surrounding solvent results in (efficient, but not complete) reformation of the original enol-conformer with a (solvent-dependent) time constant [32,68–72].

Many sunscreen molecules occur naturally on UV-exposed regions of biological systems. In mammals, for example, natural molecular sunscreens are localized in the skin (eumelanin and pheomelanin), sweat glands (urocanic acid) and the cornea of the eye (kynurenines). As with the DNA/RNA nucleobases, natural sunscreen molecules have evolved to cope with exposure to UV radiation, which, in the case of the skin, means protecting the lower epidermis from UV-induced DNA damage. Eumelanin, the most abundant melanin found in humans, provides many beneficial functions including serving as a naturally occurring sunscreen [73–75]. Eumelanin is produced via melanogenesis, wherein tyrosine is oxidized and polymerized, resulting in a heterogeneous pigment composed of cross-linked 5,6-dihydroxyindole (DHI) and 5,6-dihydroxyindole-2-carboxylic acid (DHICA) based polymers. The mechanism(s) underpinning its photostability remain active research topics, however [76–78].

Several studies have sought to address these issues by exploring aspects of the photophysics prevailing within individual molecular components of eumelanin—including DHI [79–82] and DHICA [75,83–85], various monohydroxyindoles [86,87] and catechol [88–93] in the gas and/or solution phase. Building on earlier work by Van Anh and Williams [94], Kohler and co-workers [95] recently reported ultrafast transient absorption studies following UV photoexcitation of non-covalently bonded heterodimers based on ortho-positioned dihydroxyphenol (catechol) and 1,2-benzoquinone groups. For experimental reasons, it was necessary to work with the chemically stable 3,5-di-tert-butyl substituted C and Q molecules. The experimental data provided rather convincing evidence that intermolecular HAT from the (acidic) O–H proton donor on catechol to a carbonyl oxygen (proton acceptor) group on the quinone could constitute another decay pathway and thus another source of photostability following UV photoexcitation. Here we report multi-reference computational studies designed to explore the photophysics of this model catechol (C), 1,2-benzoquinone (Q) (henceforth C:Q) heterodimer (without the tert-butyl substituents, for computational simplicity), which provide mechanistic insight in support of these recent experiments [95].

2. Computational Methods

The ground state minimum energy geometry of the \( \pi \)-stacked configuration of the C:Q heterodimer was optimized using the \( \omega\)B97XD functional of Density Functional Theory [96], coupled to the 6-311+G(d, p) Pople basis set [97]. This functional was chosen since it is capable of describing the long-range correlation effects inherent to charge-separated configurations, as well as the dispersion interactions between the individual chromophores. Optimizations of other plausible side-on, hydrogen-bonded and \( \pi \)-stacked configurations were undertaken but, in all cases, these alternative starting structures converged to the \( \pi \)-stacked configuration shown below.
The present study focused on the net transfer of an H atom, so the tautomer formed by HAT was next optimized. In the closed-shell singlet configuration, this tautomer should represent an unstable high energy point on the diabatic ground state PE surface due to the long-range attractive interaction that encourages reformation of the parent heterodimer. Thus, the corresponding triplet-spin configuration of the C:Q biradical tautomer (which can also be viewed as a semiquinone, or catechoxyl, radical pair) was optimized in order to maintain a long-range minimum energy structure. A similar methodology was used recently to follow the photoinduced evolution of a cyclopropenone-containing enediyne system through to a biradical configuration [98].

Vertical excitation energies and transition dipole moments of the C:Q heterodimer were calculated using complete active space self-consistent field (CASSCF) and complete active space second-order perturbation theory (CASPT2) methods. The CASSCF calculation was state-averaged with the lowest four singlet and the lowest four triplet states and employed an active space of ten electrons in ten orbitals (10/10)—comprising the five highest valence orbitals and five lowest unoccupied molecular orbitals (shown later).

PE profiles associated with the HAT coordinate were then constructed by interpolating the geometries between the ground state minimum of the C:Q heterodimer and the biradical tautomer, using a linear interpolation in internal coordinates (LIIC). Using the CASPT2 method [99] and a cc-pVDZ basis set, PE values at each point along the LIIC were calculated for the lowest four singlet states using a state-averaged CASSCF reference wavefunction. The same (10/10) active space was used, along with an imaginary level shift of 0.5 \(E_H\) to aid convergence and circumvent the involvement of intruder states.

All DFT calculations were calculated using the Gaussian 16 computational package [100], whilst all CASPT2 calculations used the MOLPRO computational package [101].

3. Results and Discussion

An isolated gas-phase multi-reference computational study of the UV photoinduced chemistry of C:Q heterodimers is presented, the results of which support and extend conclusions reached in recent transient absorption studies of this system in a weakly interacting solvent (cyclohexane) [95]. This section is sub-divided into sections addressing the minimum energy structures of the heterodimer and its biradical tautomer, the electronic spectroscopy of the former and then the topography of the PE surfaces sampled following photoexcitation of the heterodimer.

3.1. Minimum Energy Geometries of the Ground State Heterodimer and Its Biradical Tautomer

As a reminder, the \(\omega\)B97XD functional was used in order to achieve an appropriately balanced description of the dominant \(\pi-\pi\) interactions between the C and Q chromophores and the long-range correlation effects. As Figure 1a,b show, the ground state minimum energy geometry of the C:Q heterodimer exhibits a \(\pi\)-stacked configuration. (The Cartesian coordinates of all atoms in this minimum energy structure are provided as Supplementary Materials, as are (harmonic) normal mode wavenumbers for the ground state C:Q heterodimer and the bare C monomer.) Alternative side-on hydrogen bonding could also be expected to provide a strong intermolecular interaction, but the \(\pi\)-stacked ground state configuration shown in Figure 1 offers both \(\pi-\pi\) and hydrogen-bonding, and the stability of this \(\pi\)-stacked structure can be understood by recognizing two stabilizing interactions. One is a \(\pi-\pi\) interaction between bonding \(\pi\) electrons on the catechol moiety and the antibonding \(\pi^*\) orbital localized on the benzoquinone. These orbitals are reasonably well-matched in energy. The second is the inter-chromophore hydrogen-bonding between an O–H donor, local to catechol, and a carbonyl oxygen acceptor, localized on the benzoquinone chromophore. This deduced C:Q heterodimer structure, involving one intra- and one intermolecular H-bond, is fully consistent with that derived by analysis of the Fourier transform infrared spectrum of mixed solutions of (the di-tert-butyl substituted forms of) C and Q in cyclohexane [95].
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Figure 1. (a) Side and (b) top view of the optimized structure of the catechol (C), 1,2-benzoquinone (Q) (C:Q) heterodimer in its ground electronic state. (c) Optimized geometry of the biradical tautomer formed by H atom transfer. Intramolecular and intermolecular H-bonds are indicated by, respectively, dashed purple and green lines.

Figure 1c shows the minimum energy geometry of the biradical formed upon HAT. (The Cartesian coordinates of all atoms in this minimum energy structure are also provided in the Supplementary Materials). The “hinged” structural arrangement of the C and Q chromophores is very different from the π-stacked configuration of the ground state heterodimer, though it again displays one intermolecular and one intramolecular hydrogen bond. The breakdown of the π-stacking upon biradical formation can be understood by recognizing that the lowest energy biradical configuration has ππ* character, wherein the π- and π*-orbitals are localized on, respectively, the C and Q moieties (vide infra). The ensuing electron–electron repulsion destroys the π–π interaction inherent to the ground state parent structure, leaving inter-chromophore H-bonding as the dominant non-covalent interaction in the biradical tautomer.

We note that the experimentally studied C:Q heterodimer contains bulky tert-butyl substituents which may affect the π-stacking. That said, we do not expect this to have a serious impact on the excited state photophysics deduced here, as the tert-butyl group is a σ-perturbing substituent, while the dominant effects observed in the photophysics of C:Q are π-centered.

3.2. The Electronic Spectrum of the C:Q Heterodimer

Table 1 lists the vertical excitation energies (VEEs) to the first few singlet and triplet excited states of the C:Q heterodimer from the π-stacked minimum energy configuration. The active space orbitals used in the CASPT2 computations, shown in Figure 2, may be used along with Table 1 to identify the dominant orbital promotions involved in preparing these various excited states.
calized on the benzoquinone moiety. The S\textsubscript{1} state is best viewed as a locally excited state localized on the carbonyl oxygen atom, to an antibonding

The respective contributions (i.e., the squares of the associated coefficients) shown in parentheses. The entries in the “Character” column show the dominant electron promotions between the active space orbitals shown in Figure 2, with the respective contributions (i.e., the squares of the associated coefficients) shown in parentheses.

| Electronic State | Character | VEE/eV | f    |
|------------------|-----------|--------|------|
| S\textsubscript{1}-S\textsubscript{0} | 6→3 (0.71) | 1.93   | 0.001|
|                  | 6→3 + 6→5 (0.04) |        |      |
| S\textsubscript{2}-S\textsubscript{0} | 6→5 (0.51) | 3.23   | 0.0541|
|                  | 6→4 (0.34) |        |      |
| T\textsubscript{1}-S\textsubscript{0} | 6→5 (0.72) | 1.64   | -    |
|                  | 8→3 (0.04) |        |      |
| T\textsubscript{2}-S\textsubscript{0} | 6→5 (0.73) | 2.25   | -    |
|                  | 8→5 (0.04) |        |      |

Vertical excitation to the S\textsubscript{1} state from the π-stacked ground state minimum energy geometry is dominated by electron promotion from a largely non-bonding (n) orbital, localized on the carbonyl oxygen atom, to an antibonding π\textsuperscript{*} orbital, both of which are localized on the benzoquinone moiety. The S\textsubscript{1} state is best viewed as a locally excited state (i.e., the excitation is concentrated on a common chromophore) with nπ\textsuperscript{*} character, and optically “dark” (i.e., the S\textsubscript{1}–S\textsubscript{0} transition has a low oscillator strength—reflecting the poor spatial overlap of the n and π\textsuperscript{*} orbitals).

The S\textsubscript{2} state is best described by a mixture of two configurations. The 6→5 orbital promotion involves excitation of an electron from the π highest occupied molecular orbital (HOMO) (which is mainly localized on Q but extends over the C moiety also) to the π\textsuperscript{*} lowest unoccupied molecular orbital (LUMO) localized on Q. 6→4 promotion, in contrast, involves excitation from a bonding π orbital, largely localized on the C chromophore, to the Q-localized π\textsuperscript{*} antibonding orbital. Both promotions can be pictured as π\textsuperscript{*}→π transitions: The S\textsubscript{2} state is thus best viewed as having ππ\textsuperscript{*} character but, even in the vertical region, formation of the S\textsubscript{2} state of the heterodimer involves some electron transfer from C to Q—which likely contributes to the high oscillator strength reported in Table 1. For completeness, excitation energies to the first two triplet excited states, both of which are also best described as locally excited ππ\textsuperscript{*} states, are also included in Table 1.
The predicted oscillator strengths and VEEs for the (weak) $S_1 - S_0$ and (strong) $S_2 - S_0$ transitions reported in Table 1 match well with the maxima evident (at $\lambda \sim 595$ nm and $\sim 400$ nm) in the UV absorption spectrum of the ($t$-butyl substituted) C:Q heterodimer in cyclohexane [95], lending further support to our expectation that the tert-butyl substituents have little effect on the electronic properties (and excited state photophysics) of the heterodimer.

3.3. Photophysics of the C:Q Heterodimer

We now consider the possible fate(s) of the C:Q heterodimer following UV photoexcitation and the potential role of such photophysics in explaining the photostability of eumelanin. Motivated by the work of Kohler and co-workers [95], the ensuing discussion focuses on the mechanism of photoinduced HAT. Figure 3a displays PE profiles of the ground and first two excited singlet electronic states along the LIIC between the ground state minimum energy geometry and that of the optimized biradical tautomer formed by HAT (henceforth $Q_{\text{LIIC}}$)—the structures of both of which are reproduced again as insets in Figure 3a. We caution that the use of a LIIC almost inevitably means that the present calculations do not capture the true minimum energy path from reactant to product, but they are expected to identify key topographical features of the PE surfaces under study. The ground state (black) PE profile increases, reaching a maximum at $Q_{\text{LIIC}} \sim 0.5$, beyond which it decreases en route to the biradical tautomer. The electronic wavefunction of the adiabatic ground state switches at $Q_{\text{LIIC}} \sim 0.5$, as illustrated in Figure 3b,c, which illustrates the increasing $\pi\pi^*$ character of the ground state configuration of the biradical tautomer (which correlates diabatically with the $S_2$ state of the parent C:Q heterodimer). The stability of the biradical structure can be understood by considering the change in electronic character. In the $\pi\pi^*$ configuration, the O-atom donor of the pre-existing OH moiety contains a doubly occupied $p$ orbital which, when viewed from the biradical minimum, provides a long-range repulsive interaction in the reverse HAT direction. As Figure 3b,c show, the singly occupied molecular orbitals in the $S_0$ state of the biradical are localized on different chromophores: the $S_0$ state at $Q_{\text{LIIC}} > 0.5$ is best described as a charge-separated (or charge transfer) state.

The $S_1$ state (red in Figure 3a) has $n\pi^*$ character in the vertical region and is bound with respect to initial motion along $Q_{\text{LIIC}}$—reflecting the fact that the $\pi^* \leftarrow n$ transition is localized on the benzoquinone moiety and shows no net driving force for HAT. The $S_2$ state, in contrast, has $\pi\pi^*$ character at the Franck-Condon geometry and shows net reactivity with respect to the “hinge-like” geometry change along $Q_{\text{LIIC}}$. This can be understood by recognizing that the transition involves $\pi$ and $\pi^*$ orbitals that are initially largely localized on a single chromophore but then develop increasing charge-separated character, as shown in Figure 3b,c. The diabatic $^1\pi\pi^*$ state progressively develops charge transfer (CT) character as $Q_{\text{LIIC}} \rightarrow 1$, which is neutralized by proton transfer from the C to the Q moiety. Such photoinduced HATs are also frequently termed proton-coupled electron transfer (PCET) or electron-driven proton transfer (EDPT) processes. Upon increasing $Q_{\text{LIIC}}$ from the Franck-Condon region, the diabatic CT state crosses both the $^1n\pi^*$ state and the ground state. This is the origin of the evolution of the ground state electronic wavefunction described above. As in many related systems [4,102,103], these diabatic crossing points will surely be CIs when motion along orthogonal modes are considered, and represent regions of configuration space where internal conversion between electronic states is favorable (i.e., where population is funneled efficiently to the lower PE surface).

Given the foregoing descriptions of the various electronic states of the C:Q heterodimer, the reported photophysics can be rationalized as follows: Photoexcitation populates the “bright” $^1\pi\pi^*$ state, which is initially largely localized on Q but evolves spontaneously along the coordinate associated with HAT. Internal conversion is likely to occur at both the $S_2/S_1$ and $S_1/S_0$ CIs (see Figure 3a). The former may well lead to some population becoming temporarily trapped in the $^1n\pi^*$ state—as has been proposed in the case of oxybenzone [72]—while non-adiabatic interaction at the latter CI will promote efficient
internal conversion back to the $S_0$ state. Having accessed the $S_0$ PE surface, population may bifurcate to reform the ground state heterodimer (thereby demonstrating photostability) or evolve towards the biradical tautomer and thence to two (potentially harmful) semiquinone free radical species. Thus, the extent to which the C:Q heterodimer offers photoprotection and photostability will be sensitively dependent on the non-adiabatic dynamics prevailing at the $S_1$/S$_0$ CI—which will be sensitive to the detailed topography of the CI and the nuclear momenta within the evolving population. Such details, in turn, are likely to be sensitively dependent upon the natures of any (less benign than tert-butyl) substituents within the C and Q moieties and, in any condensed phase application, to the prevailing solvent [104]. Extrapolating to eumelanin itself, any such competition between reformation of the minimum energy ground state structure and biradical (and thence radical) formation might well be influenced by the extent (or otherwise) to which the system is able to distort away from any structural layering imposed by more extensive $\pi$-stacking between polymer strands.

**Figure 3.** (a) Adiabatic potential energy (PE) profiles of the $S_0$ (black), $S_1$ (red) and $S_2$ (blue) states of the C:Q heterodimer plotted as a function of the linear interpolation in internal coordinates (LIIC) linking the ground state minimum energy geometry (at $Q_{\text{LIIC}} = 0$) with that of the optimized biradical tautomer (at $Q_{\text{LIIC}} = 1.0$). Representations of the evolving structure of the ground state heterodimer and of orbitals 5 and 6 are shown below for $Q_{\text{LIIC}} = (b) 0.6$ and $Q_{\text{LIIC}} = (c) 1.0$ (with, in each case, the square of the coefficient associated with this $\pi\pi^*$ contribution to the $S_0$ state configuration shown in parenthesis).

### 4. General Discussion and Conclusions

This study, which is limited to the isolated heterodimer only, adds to the growing body of computational research aimed at exploring possible excited state decay paths in organic acid-base heterodimers. The present results support earlier suggestions, from analysis of
transient absorption measurements in a weakly interacting solvent [95], that HAT from an OH moiety of a catechol sub-unit to the carbonyl-oxygen atom of a quinone unit arranged in a π-stacked C:Q heterodimer could contribute to the pool of photoprotection mechanisms available to eumelanin upon exposure to UV radiation.

From the photophysical perspective, the π-stacked chromophores in the C:Q heterodimer exhibit similarities and differences with the excited state decay mechanisms identified for the chromophores in double-stranded DNA. UV excitation of a, base pair starts with a π*←π promotion localized on the purine (adenine (A) or guanine (G)), which is dissipated by PCET to the pyrimidine (thymine (T) or cytosine (C)) partner and subsequent coupling via a CI to the S0 state [105–109]. The H atom in these cases is transferred within an H-bonded base pair wherein the individual bases are parts of complementary strands (i.e., an inter-strand HAT process). UV photoinduced intra-strand electron transfer between stacked nucleobases—more reminiscent of the present situation—has been identified also, but the subsequent charge-separation (and ultimate photostability) is again achieved by an inter-strand proton transfer in the resulting radical anion base-pair [110,111].

As noted in the Introduction, eumelanin is a heterogeneous macromolecule, and much remains to be learned both about its exact structure and the mechanisms of the photoprotection it affords. Several studies of intramolecular processes contributing to the decay of excited states of monomers (and oligomers) of various of the proposed key sub-units of eumelanin, like DHI and DHICA, have been reported [75,82,84], along with some studies of their intermolecular interactions with solvent molecules [104]. The present work supports another inter-chromophore excited state decay pathway wherein HAT facilitates non-radiative coupling to, and reformation of, the ground state C:Q heterodimer. But, as experimental studies of (the di-tert-butyl substituted form of) this heterodimer also show, the biradical structure at the asymptote of the HAT coordinate can decompose to two semiquinone radicals [95]. While it is notable from an energetic perspective that absorption of one photon with an energy less than that required to break an O–H bond in bare catechol [88] could result in the formation of two semiquinone radicals, it is unlikely that nature would have adopted eumelanin as a skin pigment if such heterodimers could act as significant light-driven radical generation centers. Clearly, much further work will be needed in order to establish the importance (or otherwise) of the excited state decay pathways identified thus far for small constituent parts to the overall photoprotection afforded by bulk eumelanin.

Supplementary Materials: The following are available online at https://www.mdpi.com/2673-7256/1/1/3/s1, Cartesian coordinates associated with the various optimized structures of C:Q and (harmonic) normal mode wavenumbers for the ground states of bare C and the C:Q heterodimer.

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