

This is a repository copy of *Linking Electronic Relaxation Dynamics and Ionic Photofragmentation Patterns for the Deprotonated UV Filter Benzophenone-4*.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/172665/>

Version: Published Version

Article:

Wong, Natalie, Rankine, Conor orcid.org/0000-0002-7104-847X and Dessent, Caroline Elizabeth Helen orcid.org/0000-0003-4944-0413 (2021) Linking Electronic Relaxation Dynamics and Ionic Photofragmentation Patterns for the Deprotonated UV Filter Benzophenone-4. JOURNAL OF PHYSICAL CHEMISTRY LETTERS. pp. 2831-2836. ISSN 1948-7185

<https://doi.org/10.1021/acs.jpcllett.1c00423>

Reuse

This article is distributed under the terms of the Creative Commons Attribution (CC BY) licence. This licence allows you to distribute, remix, tweak, and build upon the work, even commercially, as long as you credit the authors for the original work. More information and the full terms of the licence here:

<https://creativecommons.org/licenses/>

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.

Linking Electronic Relaxation Dynamics and Ionic Photofragmentation Patterns for the Deprotonated UV Filter Benzophenone-4

Natalie G. K. Wong, Conor D. Rankine, and Caroline E. H. Dessent*



Cite This: *J. Phys. Chem. Lett.* 2021, 12, 2831–2836



Read Online

ACCESS |



Metrics & More

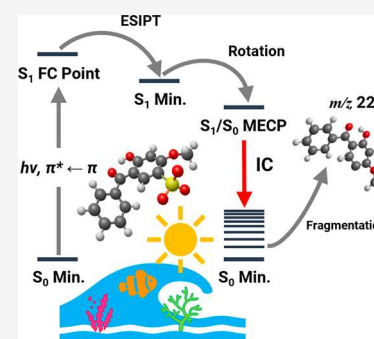


Article Recommendations



Supporting Information

ABSTRACT: Understanding how deprotonation impacts the photophysics of UV filters is critical to better characterize how they behave in key alkaline environments including surface waters and coral reefs. Using anion photodissociation spectroscopy, we have measured the intrinsic absorption electronic spectroscopy (400–214 nm) and numerous accompanying ionic photofragmentation pathways of the benzophenone-4 anion ($[\text{BP4-H}]^-$). Relative ion yield plots reveal the locations of the bright S_1 and S_3 excited states. For the first time for an ionic UV filter, *ab initio* potential energy surfaces are presented to provide new insight into how the photofragment identity maps the relaxation pathways. These calculations reveal that $[\text{BP4-H}]^-$ undergoes excited-state decay consistent with a statistical fragmentation process where the anion breaks down on the ground state after nonradiative relaxation. The broader relevance of the results in providing a basis for interpreting the relaxation dynamics of a wide range of gas-phase ionic systems is discussed.



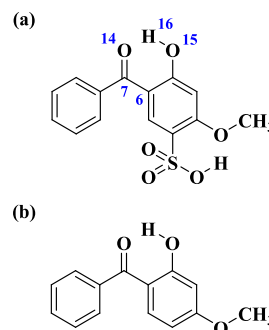
Laser spectroscopy has been increasingly applied over recent years to characterize the intrinsic photophysics of UV filters to provide a more robust understanding of molecular-level sunscreen action.¹ Both solution and gas-phase experiments have been performed, and while the solution phase can constitute an environment closer to that of a commercial sunscreen mixture,^{1–5} gas-phase studies are of particular value in providing data that can readily be interpreted by high-level theory.^{5–9} While several neutral sunscreens have been the subject of gas-phase investigations, protonated and deprotonated analogues have been studied much more sparsely.⁵ These experiments are important given that a number of aquatic environments are alkaline (e.g., surface water and coral reefs),^{10,11} so that the understanding of how deprotonation affects photostability has important environmental implications.

Very recently, laser-interfaced mass spectrometry (LIMS) has been used to probe the photophysics of several ionic sunscreen systems in detail.^{12–16} These studies reveal that protonation and deprotonation can dramatically affect the sunscreen's UV absorption profile. Information on decay dynamics (and hence the intrinsic sunscreen efficiency), however, has only been inferred indirectly in these experiments, through attempting to match the photofragmentation products against the corresponding thermal fragmentation products to elucidate whether excited-state decay is statistical or nonstatistical.^{17,18} This is a general problem for gaseous studies of ionic systems that extends well beyond the specific field of sunscreens,^{17,19–23} since there are currently few

experiments where direct time-resolved measurement of ionic photofragments is possible.^{24,25}

Here, we present the first laser spectroscopy study of benzophenone-4, BP4 (Scheme 1), in its deprotonated form. BP4 is structurally similar to oxybenzone (OB; Scheme 1), which is one of the most widely investigated sunscreens, both experimentally and theoretically.^{5,7,26–29} Studies have revealed

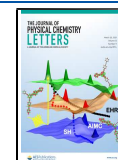
Scheme 1. Molecular Structures of (a) Benzophenone-4 (BP4) and (b) Oxybenzone (OB)



Received: February 5, 2021

Accepted: March 2, 2021

Published: March 15, 2021



that the sunscreen action of OB arises from excited-state intramolecular proton transfer (ESIPT) yielding the *keto* form of neutral oxybenzone, which then undergoes ultrafast formal conversion (IC) from the excited- to ground-state potential energy surface and efficiently thermalizes the excess energy.^{27–29} Notably, for both deprotonated and protonated OB, the observed photofragmentation patterns were interpreted as indicative of nonstatistical excited-state decay, due to disruption of the *keto–enol* moiety.¹⁵ BP4 provides an important analogue to study in this respect, since it contains a strongly acidic sulfonic acid group in addition to the OB *keto–enol* site. Deprotonation of BP4 will therefore produce the sulfonate monoanion, leaving the crucial *keto–enol* site intact for uninterrupted operation of the ultrafast nonradiative relaxation mechanism. Our aim here is to compare the photofragmentation behavior of deprotonated OB and BP4 to investigate whether excited-state decay is in fact nonstatistical and statistical, respectively. For the first time for a deprotonated UV filter, we apply quantum chemical calculations to obtain *ab initio* potential energy surfaces and hence gain direct physical insight into how the photofragment identity maps the nonradiative relaxation channels.

LIMS action spectroscopy was used to record gaseous ion photodepletion and photofragment spectra of $[\text{BP4-H}]^-$ (Section S1).^{12–16} The photodepletion spectrum can be considered to be equivalent to the gaseous absorption spectrum in the limit where radiative decay is absent. Figure 1a displays the photodepletion spectrum of mass-selected $[\text{BP4-H}]^-$ (m/z 307) over the range 3.1–5.8 eV (400–214 nm), displaying strong absorption across the UV. To aid in the discussion of the photofragment production spectra, the key spectral features are labeled I–IV, with bands I and II being the two distinct UVA and UVB absorption bands, peaking at 3.5 and 4.1 eV, respectively. Band III increases gradually in intensity in the UVC between 4.5 and 5.0 eV, and leads into band IV which is a strong, broad feature (onset *ca.* 5.0 eV) that extends further into the UVC.

We next turn to the photofragment ions produced following photoabsorption by $[\text{BP4-H}]^-$. Photofragmentation is extensive, with over 20 photofragments being observed. Figure 1b–i display the action spectra of the most prominent photofragments, with minor photofragments being reported in Section S3. The most intense photofragment ion is observed at m/z 227 (eq 1d), corresponding to loss of neutral SO_3 via a heterolytic cleavage mechanism of the C–S bond of the parent anion. m/z 227 is produced with high intensity across the UVA and lower-energy UVC regions. The other major photodissociation channels of $[\text{BP4-H}]^-$ are given in eqs 1a–1h, with the fragmentation channels discussed further in Section S6. We note that free radical formation is dominant.

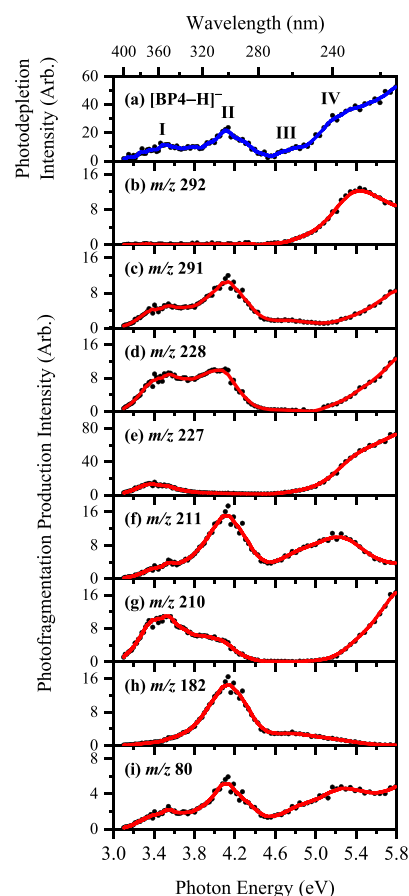
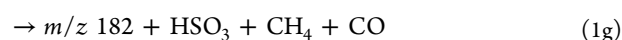
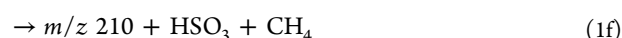
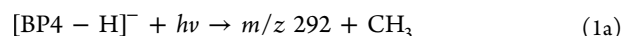


Figure 1. (a) Gas-phase absorption (photodepletion) spectrum of $[\text{BP4-H}]^-$ (m/z 307). (b–i) Photofragment production spectra of the eight major photofragments of $[\text{BP4-H}]^-$: m/z 292, 291, 228, 227, 211, 210, 182, and 80. The solid line is a five-point adjacent average of the data points.



In Figure 1b–i, several distinctive spectral profiles are evident for the various photofragment ions. All the photofragment action spectra, except for those of m/z 292 and 182 fragments, show a prominent peak in the UVA (*ca.* 3.5 eV), corresponding to the photodepletion feature I. A subsequent band, peaking at 4.1 eV, is also evident for the m/z 291, 228, 211, 182, and 80 fragments, in the region of feature II. The growth in production of several of the photofragment ions (m/z 291, 228, 227, 210, and 80) beyond 5.0 eV traces the profile of feature IV (Figure 1a). We note that the vertical detachment energy (VDE) for $[\text{BP4-H}]^-$ is calculated as 5.19 eV, so that most of the spectral range lies below the electron detachment threshold. It is interesting to note that, for the m/z 292 photofragment, production peaks at 5.4 eV, possibly indicating that a dipole-bound excited state is accessed in this region that decays with formation of m/z 292.^{30,31} Section S4 discusses electron detachment further.

Figure 2a presents the relative photofragment ion yields of $[\text{BP4-H}]^-$ as a function of photoexcitation energy, highlighting several maxima that can be attributed to photoexcitation into different electronic states. It is evident that, in both the UVA and low UVC regions, the relative ion yield of the m/z 227 photofragment is *ca.* 50% larger than other photoproduct ions. Conversely, within the range 3.8–5.0 eV, the production of fragment ions m/z 211 and 182 (and to a

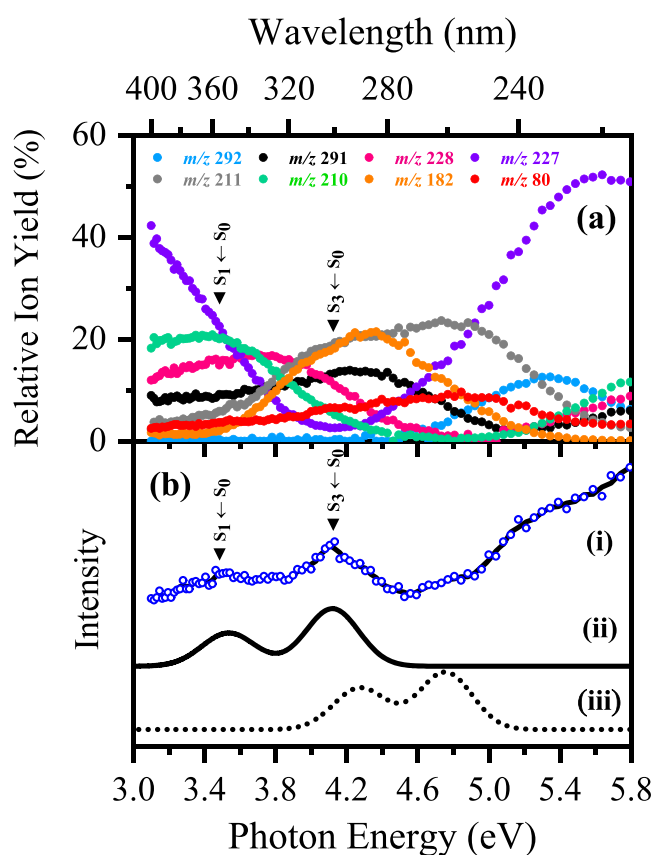


Figure 2. (a) Relative ion yield plot highlighting the eight most intense photofragments of $[\text{BP4-H}]^-$ seen upon laser excitation between 3.1 and 5.8 eV. (b) Gas-phase experimental photodepletion spectrum (i) vs theoretical UV absorption spectra calculated at the (ii) ADC(2)/MP2/ma-def2-SV(P) and (iii) $\omega\text{B97X-D/ma-def2-SV(P)}$ levels. The optically bright $S_1 \leftarrow S_0$ and $S_3 \leftarrow S_0$ $\pi\pi^*$ transitions are indicated.

lesser extent m/z 291) increases significantly in comparison to the remaining ionic photofragments.

To probe the thermal fragmentation pathways of $[\text{BP4-H}]^-$ on its electronic ground state, higher-energy collisional dissociation (HCD) was employed (Table 1, Figure 3, and Section S5).^{14,21} These measurements are essential to identify which ions are secondary products, formed when a precursor species fragments at high internal energy.³² They are also important, as any photofragments not observed in HCD can be identified as purely photochemical products. At relatively low collisional energies (20–42% HCD), the most intense fragment ion is m/z 227, with m/z 291, 228, and 210 also being produced in significant quantities. This indicates that thermal breakdown of the electronic ground state of $[\text{BP4-H}]^-$ is associated with the molecule fragmenting along a number of different pathways. Production of the m/z 227, 228, and 291 ions all decreases at higher energies, concomitant with the m/z 211 fragment increasing. (We note that the m/z 291 fragment persists to higher collisional energies than m/z 227 and 228, indicating higher relative stability.) The HCD results therefore reveal that m/z 211 is a secondary fragment from m/z 227, 228, and 291 at higher internal energy. Similarly, m/z 210 appears to decrease as the m/z 182 ion increases.

Since the m/z 227, 228, 291, and 211 fragments dominate both the UV photofragmentation of $[\text{BP4-H}]^-$ and thermal (HCD) fragmentation, photofragmentation of the anion can

Table 1. Summary of the Ionic Fragments of Deprotonated BP4 (m/z 307) Produced upon UV Laser Photoexcitation and Higher-Energy Collisional Dissociation (HCD) at 40% and 70% HCD Energies (Proposed Structures Are Outlined in Table S1)

Ionic mass fragment (m/z) ^a	Observed in HCD ^b		Observed in UV laser photoexcitation ^b
	40%	70%	
292	√ (xw) ^c	–	√ (m)
291	√ (m)	√ (w)	√ (m)
228	√ (m)	√ (vw)	√ (m)
227	√ (s)	√ (vw)	√ (vs)
211	√ (w)	√ (vs)	√ (m)
210	√ (m)	√ (vw)	√ (m)
182	√ (m)	√ (m)	√ (m)
80	√ (w)	√ (m)	√ (w)

^aDetermined with mass accuracy >0.3 amu. ^bVery strong (vs), strong (s), moderate (m), weak (w), very weak (vw), and extremely weak (xw). ^cHCD fragment m/z 292 is observed to peak at 34% HCD energy, with a relative ion intensity of <2%.

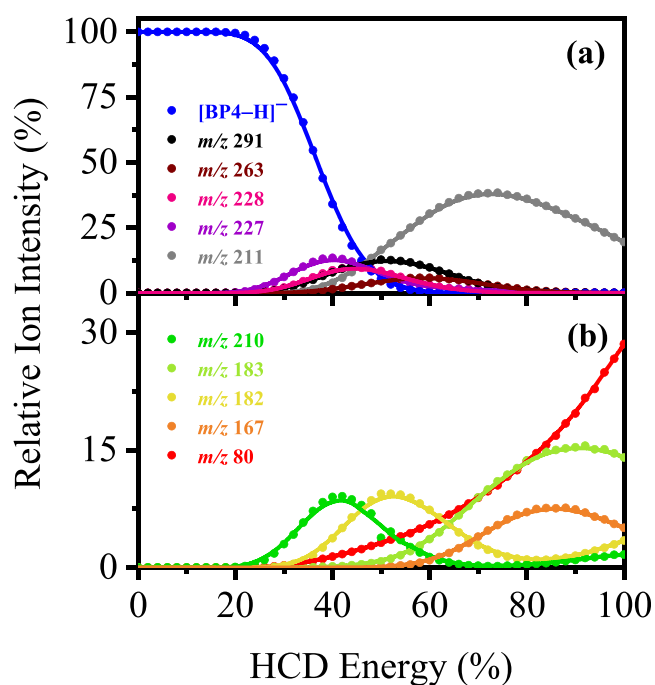


Figure 3. Parent ion dissociation curves for $[\text{BP4-H}]^-$ highlighting its ten most intense thermal fragments between 0% and 100% HCD energy. The curved lines are a five-point adjacent average of such data points and are provided as a viewing guide, to emphasize the profile for each individual fragment.

be categorized as predominantly statistical (ergodic) over the spectral range studied.^{17,18} Section S5 discusses the more minor HCD fragments and branching between the minor fragmentation pathways in more detail.

To explore whether this picture of statistical photofragmentation for $[\text{BP4-H}]^-$ is credible, quantum chemical calculations were performed to characterize the excited-state potential energy surfaces (Section S1). The C_1 -symmetric S_0 minimum-energy geometry of $[\text{BP4-H}]^-$ was located at the $\omega\text{B97X-D}$ level (Table S2), with key excited-state parameters ($\omega\text{B97X-D}$ and ADC(2) levels) summarized in Table 2.

Table 2. Summary of Vertical Excitation Energies, ΔE , Oscillator Strengths, f , and Characters of the $S_n \leftarrow S_0$ ($n = 1, 2, 3$) States As Evaluated at the ω B97X-D/ma-def2-SV(P) and ADC(2)/MP2/ma-def2-SV(P) Levels

State	Char.	ω B97X-D		ADC(2)	
		ΔE (eV)	f	ΔE (eV)	f
S_1	$\pi\pi^*$	4.272	0.256	3.533	0.156
S_2	$n\pi^*$	4.357	0.010	3.701	0.004
S_3	$\pi\pi^*$	4.756	0.365	4.120	0.273

Figure 2b displays the calculated UV absorption spectra of [BP4-H]⁻, along with the experimental photodepletion spectrum. We assign the two bands observed in the UVA/UVB regions of the experimental [BP4-H]⁻ photodepletion spectrum (I and II) as the optically bright $S_1 \leftarrow S_0$ and $S_3 \leftarrow S_0$ $\pi\pi^*$ transitions, respectively. The excellent agreement between the calculated spectra at both the ω B97X-D and ADC(2) levels and the experimental spectrum (Figure 2b) is notable, both in terms of state identities, relative peak positions, and intensities. At the ADC(2) level, quantitative agreement with experiment is obtained ‘out of the box’, whereas, at the ω B97X-D level, the vertical excitation energies of the $S_n \leftarrow S_0$ ($n = 1, 2, 3$) states are characteristically overestimated (ca. 0.7 eV) but in good qualitative agreement.

Based on our understanding of the sister molecule, OB,^{15,29} [BP4-H]⁻ can be expected to relax on the S_1 state via ESIPT. A C_1 -symmetric S_1 minimum-energy geometry for [BP4-H]⁻ was located ca. 4.5 Å Da^{-1/2} from the Franck–Condon point (Table S3). The S_1 minimum-energy geometry is accessed via ESIPT from the Franck–Condon point, with the H₁₆ atom bound to O₁₅ migrating across to O₁₄. ESIPT follows a direct excited-state relaxation coordinate and is consequently expected to occur promptly postphotoexcitation to the S_1 state. Post-ESIPT, [BP4-H]⁻ can access the S_1/S_0 crossing seam at an S_1/S_0 minimum-energy crossing point (MECP). An S_1/S_0 MECP was located ca. 18.2 and 18.0 Å Da^{-1/2} from the Franck–Condon point and S_1 minimum-energy geometry, respectively (Table S4). The S_1/S_0 MECP is accessed via torsion of C₆–C₇ and is characterized by the aromatic rings being rotated into a near-perpendicular conformation, effectively closing the gap between the S_0 and S_1 states.

To map the $S_0 \leftarrow S_1$ IC channel, potential energy surfaces have been constructed between the key geometries via linear interpolation of internal coordinates (LIIC). Independent single-point energy calculations have been carried out at each one of 25 interpolated geometries, respectively, with the calculated potential energy surfaces presented in Figure 4a.

The picture to emerge here is similar to that described for OB by Karsili et al.,²⁹ which is consistent with experiments which identified a subpicosecond lifetime for the IC channel.²⁷ The quantum-chemical calculations reported here are not able to give information on the time scale that the S_1/S_0 crossing seam is accessed (although they could be readily coupled to excited-state dynamics simulations such as nonadiabatic mixed-quantum-classical or trajectory surface-hopping dynamics, to directly obtain this information). However, given the similar potential energy surface morphologies of [BP4-H]⁻ and OB around the key *keto-enol* region, it is reasonable to expect that it is ultrafast (i.e., subpicosecond) and, therefore, able to outcompete other processes efficiently, e.g., excited-state fragmentation, radiative decay, and intersystem crossing. (For ISC; $T_1 \leftarrow S_1$ and $T_2 \leftarrow S_1$ spin–orbit couplings are on the

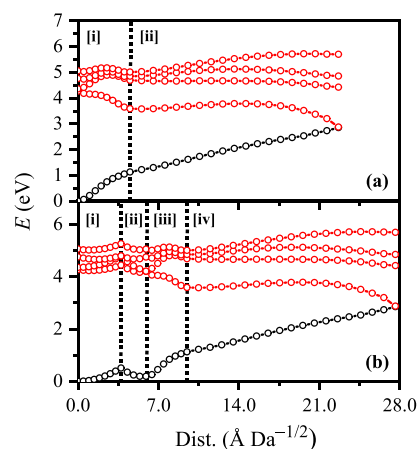


Figure 4. (a) Energies of the S_0 state (black) and excited singlet states (red) between (i) the S_0 and S_1 minimum-energy geometries and (ii) the S_1 minimum-energy geometry and the S_1/S_0 MECP. (b) Energies of the S_0 state (black) and excited singlet states (red) between (i) the S_0 minimum-energy geometry and the S_3/S_2 MECP, (ii) the S_3/S_2 MECP and the S_2/S_1 MECP, (iii) the S_2/S_1 MECP and the S_1 minimum-energy geometry, and (iv) the S_1 minimum-energy geometry and the S_1/S_0 MECP. Points were generated via linear interpolation of internal coordinates (LIIC). Energies were evaluated at the ω B97X-D/ma-def2-SV(P) level.

order of ca. 5–10 cm⁻¹ along the LIIC channel: Section S8.) The calculations are therefore entirely consistent with our deduction from the experimental results of nonradiative relaxation followed by statistical fragmentation on the hot ground state. This leads to ejection of SO₃ as the initial dominant channel, as the C–S bond is the weakest bond in [BP4-H]⁻.³³ Loss of SO₃ is commensurate with production of the m/z 227 fragment, both from excitation at feature I, i.e., the lowest-energy optically bright state, and, crucially, from the HCD production curves (Figure 3).

For the feature II region, which corresponds to excitation into the optically bright S_3 state, the calculations predict decay pathways that appear similar to those outlined for feature I. Figure 4b shows the S_3/S_2 and S_2/S_1 MECPs that have been located (Tables S5–S6), showing that both lie close to (ca. 3.7 and 2.1 Å Da^{-1/2}, respectively), and downhill of, the respective Franck–Condon point. Thus, S_3 excitation is predicted to lead to a prompt $S_1 \leftarrow S_2 \leftarrow S_3$ cascade of population. After arriving on the S_1 state close to the Franck–Condon point, ESIPT and ultrafast $S_0 \leftarrow S_1$ IC will proceed as described above. We speculate that $S_0 \leftarrow S_1$ IC, when the S_1 state is accessed indirectly (from above; i.e., postphotoexcitation into the S_3 state) as opposed to directly (postphotoexcitation to the S_1 state), could be even more efficient, since accessing the S_3/S_2 MECP and the S_2/S_1 MECP directly accesses the proton transfer and torsional coordinates, respectively, that are necessary to subsequently access the S_1/S_0 crossing seam. This could be tested in future work by either excited-state dynamics simulations³⁴ and/or time-resolved experiments.^{24,25}

The differences in fragment production on excitation at features I and II can then be explained as follows. Excitation at feature I (the S_1 state) leads to fission of the C–S bond after nonradiative relaxation (as previously observed for UVB filter 2-phenylbenzimidazole-5-sulfonic acid),¹⁴ producing primarily the m/z 227, 228, and 291 fragments. Excitation at feature II (the S_3 state) will also lead to fission of the C–S bond after nonradiative relaxation and the production of the m/z 227,

228, and 291 fragments. However, as a greater amount of photon energy is pumped into the system (4.1 eV versus 3.5 eV), these fragments possess enough internal energy to undergo secondary fragmentation. The reduction in photofragment intensity can be seen first for m/z 227, then for m/z 228, and finally for m/z 291, exactly mirroring the measured relative stability of these ions from the HCD measurements (Figure 3). (We note that similar arguments can be applied to the m/z 210 and 182 photofragments, where comparison to the HCD data reveals that the m/z 182 ion persists to higher internal energy.) All of these photofragments therefore produce the m/z 211 fragment as a secondary product: indeed, the m/z 211 fragment dominates the medium-high HCD energy range between 42% and 80% HCD energies.

In summary, we have reported the gaseous UV absorption spectrum and photofragmentation profile of $[\text{BP4-H}]^-$ acquired via LIMS. For the first time for an ionic UV filter, *ab initio* potential energy surfaces are presented to provide new insight into the relaxation pathways. The calculations predict that, in the regions of both the optically bright $S_1 \leftarrow S_0$ and $S_3 \leftarrow S_0 \pi\pi^*$ transitions, excited state relaxation will occur via nonradiative decay, associated with a statistical excited state decay process. In the photodissociation experiments, the observed photofragments mirror those observed upon thermal breakdown of the electronic ground state. Importantly, the photon-energy dependent production spectra of the numerous photofragments mirror the fragment production curves in the HCD collisional activation measurements. This is clear evidence of statistical decay, driven by fragmentation on a hot ground state surface, which in turn demonstrates that deprotonated BP4 is behaving like an efficient UV filter. However, the results presented here are of broader importance, as they provide a theoretical basis to support the widely adopted argument linking ionic photofragmentation patterns and decay dynamics that has been used for interpreting the behavior of key gaseous ionic systems including nucleobases and nucleotides.^{17,19–23,35,36}

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.jpcllett.1c00423>.

Experimental and computational methodology; photo-depletion laser power dependence measurements; additional photofragment action spectra; electron detachment yield versus photodepletion yield interpretation; higher-energy collisional dissociation (HCD) production spectra; further discussion of deprotonated benzophenone-4 fragmentation channels; optimized Cartesian coordinate tables; further computational results; schematic structure of deprotonated benzophenone-4 (PDF)

■ AUTHOR INFORMATION

Corresponding Author

Caroline E. H. Dessent – Department of Chemistry, University of York, York YO10 5DD, U.K.; orcid.org/0000-0003-4944-0413; Email: caroline.dessent@york.ac.uk

Authors

Natalie G. K. Wong – Department of Chemistry, University of York, York YO10 5DD, U.K.

Conor D. Rankine – School of Natural and Environmental Sciences, Newcastle University, Newcastle-upon-Tyne NE1 7RU, U.K.

Complete contact information is available at:

<https://pubs.acs.org/doi/10.1021/acs.jpcllett.1c00423>

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was funded through the Leverhulme Trust Research Project Grant RPG-2017-147. We thank the University of York and the Department of Chemistry for provision of funds for the OPO laser system. We are grateful for the computational support from the University of York High Performance Computing service, Viking, and the Research Computing team. The York Centre of Excellence in Mass Spectrometry, used for the higher-energy collisional dissociation (HCD) work, was created thanks to a major capital investment through Science City York, supported by Yorkshire Forward with funds from the Northern Way Initiative, and has more recently received additional support from the EPSRC and BBSRC. C.D.R. thanks the Engineering and Physical Sciences Council (EPSRC) and Newcastle University (Newcastle-upon-Tyne, UK) for funding his research via the award of an EPSRC Doctoral Prize Fellowship (EP/R51309X/1). Finally, we thank Prof. Tolga Karsili for useful discussions on benzophenone-4.

■ REFERENCES

- (1) Holt, E. L.; Stavros, V. G. Applications of Ultrafast Spectroscopy to Sunscreen Development, from First Principles to Complex Mixtures. *Int. Rev. Phys. Chem.* **2019**, *38*, 243–285.
- (2) Chan, C. T.-L.; Ma, C.; Chan, R. C.-T.; Ou, H.-M.; Xie, H.-X.; Wong, A. K.-W.; Wang, M.-L.; Kwok, W.-M. A Long Lasting Sunscreen Controversy of 4-Aminobenzoic Acid and 4-Dimethylaminobenzaldehyde Derivatives Resolved by Ultrafast Spectroscopy Combined with Density Functional Theoretical Study. *Phys. Chem. Chem. Phys.* **2020**, *22*, 8006–8020.
- (3) Zhao, X.; Luo, J.; Liu, Y.; Pandey, P.; Yang, S.; Wei, D.; Han, K. Substitution Dependent Ultrafast Ultraviolet Energy Dissipation Mechanisms of Plant Sunscreens. *J. Phys. Chem. Lett.* **2019**, *10*, 5244–5249.
- (4) Luo, J.; Liu, Y.; Yang, S.; Flourat, A. L.; Allais, F.; Han, K. Ultrafast Barrierless Photoisomerization and Strong Ultraviolet Absorption of Photoproducts in Plant Sunscreens. *J. Phys. Chem. Lett.* **2017**, *8*, 1025–1030.
- (5) Ignasiak, M. T.; Houee-Levin, C.; Kciuk, G.; Marciniak, B.; Pedzinski, T.; Houée-Levin, C.; Kciuk, G.; Marciniak, B.; Pedzinski, T. A Reevaluation of the Photolytic Properties of 2-Hydroxybenzophenone-Based UV Sunscreens: Are Chemical Sunscreens Inoffensive? *ChemPhysChem* **2015**, *16*, 628–633.
- (6) Iida, Y.; Kinoshita, S.; Kenjo, S.; Muramatsu, S.; Inokuchi, Y.; Zhu, C.; Ebata, T. Electronic States and Nonradiative Decay of Cold Gas-Phase Cinnamic Acid Derivatives Studied by Laser Spectroscopy with a Laser-Ablation Technique. *J. Phys. Chem. A* **2020**, *124*, 5580–5589.
- (7) Domingos, S. R.; Schnell, M. Wet Sunscreens in the Gas Phase: Structures of Isolated and Microsolvated Oxybenzone. *J. Phys. Chem. Lett.* **2018**, *9*, 4963–4968.
- (8) Dean, J. C.; Kusaka, R.; Walsh, P. S.; Allais, F.; Zwier, T. S. Plant Sunscreens in the UV-B: Ultraviolet Spectroscopy of Jet-Cooled

Sinapoyl Malate, Sinapic Acid, and Sinapate Ester Derivatives. *J. Am. Chem. Soc.* **2014**, *136*, 14780–14795.

(9) Tan, E. M. M.; Hilbers, M.; Buma, W. J. Excited-State Dynamics of Isolated and Microsolvated Cinnamate- Based UV - B Sunscreens. *J. Phys. Chem. Lett.* **2014**, *5*, 2464–2468.

(10) Jiang, L.-Q.; Carter, B. R.; Feely, R. A.; Lauvset, S. K.; Olsen, A. Surface Ocean PH and Buffer Capacity: Past, Present and Future. *Sci. Rep.* **2019**, *9*, 18624.

(11) Stark, J. S.; Langdon, C. Coral Reef PH Altered in Situ. *Nat. Ecol. Evol.* **2019**, *3*, 1380–1381.

(12) Berenbeim, J. A.; Wong, N. G. K.; Cockett, M. C. R.; Berden, G.; Oomens, J.; Rijs, A. M.; Dessent, C. E. H. Sodium Cationization Can Disrupt the Intramolecular Hydrogen Bond That Mediates the Sunscreen Activity of Oxybenzone. *Phys. Chem. Chem. Phys.* **2020**, *22*, 19522–19531.

(13) Berenbeim, J. A.; Wong, N. G. K.; Cockett, M. C. R.; Berden, G.; Oomens, J.; Rijs, A. M.; Dessent, C. E. H. Unravelling the Keto–Enol Tautomer Dependent Photochemistry and Degradation Pathways of the Protonated UVA Filter Avobenzone. *J. Phys. Chem. A* **2020**, *124*, 2919–2930.

(14) Wong, N. G. K.; Berenbeim, J. A.; Dessent, C. E. H. Direct Observation of Photochemical Free Radical Production from the Sunscreen 2-Phenylbenzimidazole-5-Sulfonic Acid via Laser-Interfaced Mass Spectrometry. *ChemPhotoChem* **2019**, *3*, 1231–1237.

(15) Wong, N. G. K.; Berenbeim, J. A.; Hawkrigde, M.; Matthews, E.; Dessent, C. E. H. Mapping the Intrinsic Absorption Properties and Photodegradation Pathways of the Protonated and Deprotonated Forms of the Sunscreen Oxybenzone. *Phys. Chem. Chem. Phys.* **2019**, *21*, 14311–14321.

(16) Matthews, E.; Dessent, C. E. H. Experiment and Theory Confirm That UV Laser Photodissociation Spectroscopy Can Distinguish Protomers Formed via Electrospray. *Phys. Chem. Chem. Phys.* **2017**, *19*, 17434–17440.

(17) Soorkia, S.; Jouvét, C.; Grégoire, G. UV Photoinduced Dynamics of Conformer-Resolved Aromatic Peptides. *Chem. Rev.* **2020**, *120*, 3296–3327.

(18) Lucas, B.; Barat, M.; Fayeton, J. A.; Jouvét, C.; Çarçalı, P.; Grégoire, G. Statistical versus Non-Statistical Photo-Fragmentation of Protonated GWG Tri-Peptide Induced by UV Excitation. *Chem. Phys.* **2008**, *347*, 324–330.

(19) Noble, J. A.; Marceca, E.; Dedonder, C.; Phasayavan, W.; Féraud, G.; Inceesungvorn, B.; Jouvét, C. Influence of the N Atom Position on the Excited State Photodynamics of Protonated Azaindole. *Phys. Chem. Chem. Phys.* **2020**, *22*, 27280–27289.

(20) Uleanya, K. O.; Cercola, R.; Nikolova, M.; Matthews, E.; Wong, N. G. K.; Dessent, C. E. H. Observation of Enhanced Dissociative Photochemistry in the Non-Native Nucleobase 2-Thiouracil. *Molecules* **2020**, *25*, 3157.

(21) Cercola, R.; Matthews, E.; Dessent, C. E. H. Photoexcitation of Adenosine 5'-Triphosphate Anions in Vacuo: Probing the Influence of Charge State on the UV Photophysics of Adenine. *J. Phys. Chem. B* **2017**, *121*, 5553–5561.

(22) Marcum, J. C.; Halevi, A.; Weber, J. M. Photodamage to Isolated Mononucleotides—Photodissociation Spectra and Fragment Channels. *Phys. Chem. Chem. Phys.* **2009**, *11*, 1740–1751.

(23) Nielsen, S. B.; Andersen, J. U.; Forster, J. S.; Hvelplund, P.; Liu, B.; Pedersen, U. V.; Tomita, S. Photodestruction of Adenosine 5'-Monophosphate (AMP) Nucleotide Ions in Vacuo: Statistical versus Nonstatistical Processes. *Phys. Rev. Lett.* **2003**, *91*, 048302.

(24) Kruppa, S. V.; Bäßler, F.; Klopper, W.; Walg, S. P.; Thiel, W. R.; Diller, R.; Riehn, C. Ultrafast Excited-State Relaxation of a Binuclear Ag(i) Phosphine Complex in Gas Phase and Solution. *Phys. Chem. Chem. Phys.* **2017**, *19*, 22785–22800.

(25) Nolting, D.; Weinkauff, R.; Hertel, I. V.; Schultz, T. Excited-State Relaxation of Protonated Adenine. *ChemPhysChem* **2007**, *8*, 751–755.

(26) Baker, L. A.; Grosvenor, L. C.; Ashfold, M. N. R.; Stavros, V. G. Ultrafast Photophysical Studies of a Multicomponent Sunscreen:

Oxybenzone–Titanium Dioxide Mixtures. *Chem. Phys. Lett.* **2016**, *664*, 39–43.

(27) Baker, L. A.; Horbury, M. D.; Greenough, S. E.; Coulter, P. M.; Karsili, T. N. V.; Roberts, G. M.; Orr-Ewing, A. J.; Ashfold, M. N. R.; Stavros, V. G. Probing the Ultrafast Energy Dissipation Mechanism of the Sunscreen Oxybenzone after UVA Irradiation. *J. Phys. Chem. Lett.* **2015**, *6*, 1363–1368.

(28) Baker, L. A.; Horbury, M. D.; Greenough, S. E.; Ashfold, M. N. R.; Stavros, V. G. Broadband Ultrafast Photoprotection by Oxybenzone across the UVB and UVC Spectral Regions. *Photochem. Photobiol. Sci.* **2015**, *14*, 1814–1820.

(29) Karsili, T. N. V.; Marchetti, B.; Ashfold, M. N. R.; Domcke, W. Ab Initio Study of Potential Ultrafast Internal Conversion Routes in Oxybenzone, Caffeic Acid, and Ferulic Acid: Implications for Sunscreens. *J. Phys. Chem. A* **2014**, *118*, 11999–12010.

(30) Matthews, E.; Dessent, C. E. H. Observation of Near-Threshold Resonances in the Flavin Chromophore Anions Alloxazine and Lumichrome. *J. Phys. Chem. Lett.* **2018**, *9*, 6124–6130.

(31) Harvey, A. J. A.; Yoshikawa, N.; Wang, J.-G.; Dessent, C. E. H. Communication: Evidence for Dipole-Bound Excited States in Gas-Phase I – · MI (M = Na, K, Cs) Anionic Salt Microclusters. *J. Chem. Phys.* **2015**, *143*, 101103.

(32) Cercola, R.; Fischer, K. C.; Sherman, S. L.; Garand, E.; Wong, N. G. K.; Hammerback, L. A.; Lynam, J. M.; Fairlamb, I. J. S.; Dessent, C. E. H. Direct Measurement of the Visible to UV Photodissociation Processes for the PhotoCORM TryptoCORM. *Chem. - Eur. J.* **2020**, *26*, 10297–10306.

(33) Zhang, X. Mass Spectrometric and Theoretical Studies on Dissociation of the CS Bond in the Benzenesulfonic Acid and Benzenesulfonic Acid Anion Series: Homolytic Cleavage vs Heterolytic Cleavage. *J. Mol. Struct.* **2012**, *1028*, 1–6.

(34) Mai, S.; González, L. Molecular Photochemistry: Recent Developments in Theory. *Angew. Chem., Int. Ed.* **2020**, *59*, 16832–16846.

(35) Daly, S.; Porrini, M.; Rosu, F.; Gabelica, V. Electronic Spectroscopy of Isolated DNA Polyanions. *Faraday Discuss.* **2019**, *217*, 361–382.

(36) Broquier, M.; Soorkia, S.; Pino, G.; Dedonder-Lardeux, C.; Jouvét, C.; Grégoire, G. Excited State Dynamics of Cold Protonated Cytosine Tautomers: Characterization of Charge Transfer, Intersystem Crossing, and Internal Conversion Processes. *J. Phys. Chem. A* **2017**, *121*, 6429–6439.