INVESTIGATIVE PHOTOCHEMISTRY: ELUCIDATING THE MECHANISM OF THE PHOTO-FAVORSKII REARRANGEMENT

By

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Abstract

Kenneth F. Stensrud, Ph.D.
Department of Chemistry, April 2008
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The $p$-hydroxyphenacyl chromophore (pHP) is a versatile phototrigger for an assortment of biological nucleofuges, such as $\gamma$-aminobutyric acid (GABA). The photochemistry is trenchant; irradiation discharges the nucleofuge and induces rearrangement of pHP, analogous to the ground state Favorskii rearrangement.

Structure-reactivity relationships were explored with pHP GABA’s. Electron donating groups lowered the release efficiency, $\Phi_{\text{dis}}$, relative to the unsubstituted analog. Electron withdrawing carbonyl groups attenuated $\Phi_{\text{dis}}$. Fluoro moieties modulated the pKa of pHP; protonated pHP manifested higher $\Phi_{\text{dis}}$ than their conjugate base counterparts. The reactive excited states were determined as triplets by Stern-Volmer quenching and laser flash photolysis studies. The triplet lifetimes were $\sim 10^{9}$ s with release rate constants of $10^{7}$ to $10^{8}$ s$^{-1}$. Auxochrome substitutions lengthened the $\lambda_{\text{max}}$ of pHP but lowered $\Phi_{\text{dis}}$. Added perchlorate salts markedly improved $\Phi_{\text{dis}}$ for pHP GABA’s, imputing intermediate ion-radical pairs.

Comprehensive photo-Favorskii mechanisms were posited for protonated and conjugate base forms of pHP GABA.
There have been frequent comments regarding my use of flowery, poetic words instead of those deemed appropriate for scientific writing. Though I have learned to distinguish one from the other while composing this dissertation, I find it truly risible that, at this time, words fail me when appositely expressing my gratitude to my advisor, Professor Richard Givens. Dr. Givens equanimity in dealing with my oft hair-brained ideas and digressions cannot be understated. My motivations to work long hours at solving our research problems originated from his encouragements and well considered proposals. Honestly, I shouldn’t continue to indite a synopsis of his qualities, as they will present as mere platitudes. I can only state with certainty that Dr. Givens will always remain close to my heart for everything he has passed on to me both as a scientist and a man of utmost integrity.

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have persevered without you. Beth, from you I have recognized the existence of angels.

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John Hershberger - we came in together and we will go out together. Through good and bad times, times of confusion and consumption, our experiences will never be forgotten.
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</table>
Glossary of Chemical Compounds Studied

- pHp GABA (101)
- m-F-pHp GABA (122)
- o-F-pHp GABA (123)
- 2,3-diF-pHp GABA (124)
- 2,5-diF-pHp GABA (125)
- 3,5-diF-pHp GABA (126)
- 2,6-diF-pHp GABA (127)
- 2,3,6-triF-pHp GABA (128)
- 2,3,5,6-tetraF-pHp GABA (129)
- m-CF₃-pHp GABA (130)
- m-OCF₃-pHp GABA (131)
- m-NO₂-pHp GABA (132)
- m,m'-diOCH₃-pHp GABA (133)
- m-CH₃-pHp GABA (134)
- o-CH₃-pHp GABA (135)
Introduction

The Favorskii reaction or rearrangement, though discovered over a century ago, continues to be widely exploited by synthetic chemists in the manipulation and assembly of organic molecules, such as Tricycloclavulone and Kelsoene. This assertion is buttressed by the manifestation of 832 references in a SciFinder Scholar® search. The “classical” Favorskii rearrangement involves base-mediated rearrangement of ketones bearing α-substituents with favorable nucleofuge attributes, to produce carboxylic acids, esters, or amides (Equation 1, with KOH as the base). In equation 1, the rearrangement leads to a ring contraction and conversion of the ketone to a carboxylic acid. Further designations can be made depending on whether α-hydrogens exist to the carbonyl moiety (these will be elaborated later).

\[ \text{H} \quad \text{H} \quad \text{O} \quad X \quad \text{KOH} \quad \text{CO}_2\text{H} + \text{X}^- \]

\[ X = \text{Nucleofuge (i.e. Cl, Br, OTs)} \]

Eq. 1

I. Early investigations into the mechanism of Favorskii rearrangement.

Efforts to generate a “working” mechanism to explain classical Favorskii rearrangement have been pervasive and continuous, and have centered on four discrete pathways: a) Aston’s proposition (Equation 2) related a nucleophilic
approach by alkoxide (RO') to the carbonyl moiety (2), followed by nucleofuge (bromide) disjunction to generate an epoxyether (3). Rearrangement through methyl migration proceeded to the Favorskii product (4). An independent study by Stevens and coworkers, however, using a related set of reaction conditions, failed to find the anticipated ester products.

From this and accompanying studies, the transposition of epoxyether intermediates to acids/esters (Aston’s postulate) has been judged as unsuitable for Favorskii-type rearrangements. b) A second hypothesis (Scheme 1) accentuated the formation of a ketene (7), initiated by base induced α-proton abstraction, then abrupt halide departure to form carbene (6), with a supervening “Wolff” rearrangement to

\[
\begin{align*}
\text{Br} & \quad \text{RO'} \quad \text{ROH} \\
\text{1} & \quad \text{2} \\
\text{Hemiketal} & \quad \text{Epoxide} \\
\text{3} & \quad \text{Epoxide} \\
\alpha\text{-hydroxyketal} & \quad \text{Rearr.} \\
\text{4} & \quad \text{R} = \text{H, alkyl}
\end{align*}
\]

Eq. 2
the ketene. Once formed, the ketene was promptly attacked by the nucleophilic alkoxide to produce the Favorskii ester (8). Several related studies evinced the direct formation

**Scheme 1.** Richard’s asserted mechanism of the Favorskii rearrangement.

![Scheme 1](image1)

R = H, alkyl

development of trialkyl acids and esters from ketones,\(^7\) which would directly oppugn a ketene precursor. c) A third proposal suggested an initial unimolecular disjunction of the nucleofuge to afford an \(\alpha\)-ketocarbenium ion (10), followed by tautomerizations 11 and 12, acylium formation from alkyl migration (13) and then rearrangement to the acid/ester\(^{12}\) (14) (Scheme 2). In this mechanism, there appeared to be a marginal role

**Scheme 2.** McPhee’s mechanism of the Favorskii reaction.

![Scheme 2](image2)
for the base, which, in subsequent studies, was shown to be indispensable for the generation of Favorskii products. d) The most widely accepted mechanism was first proposed by Loftfield\textsuperscript{13} and advanced by several researchers.\textsuperscript{14,15,16,17} The reaction sequence has been suggested as stepwise,\textsuperscript{13,18,14} involving base induced deprotonation\textsuperscript{19,14} at the \(\alpha\) position to the carbonyl to form an enolate (16) followed by intramolecular displacement and cyclization via halide disjunction to educe a cyclopropanone intermediate (17) then nucleophilic attack at the carbonyl to induce the signature 1,2-rearrangement product in the furnishing of Favorskii ester (18).

**Scheme 3.** Loftfield’s mechanism of the Favorskii rearrangement.

(Scheme 3). Loftfield demonstrated that the preponderance of products derived only from epoxyether intermediates were generated from the \(\alpha\text{-Cl}\) precursor and posited the rate determining step (RDS) for these analogs as the reversible, initial deprotonation. Dewar,\textsuperscript{20} through computational efforts, averred that a disfavored overlap between the newly formed enolate 20 (Scheme 4) and chloro-bearing carbon would preclude cyclization by an \textit{S}_{\text{N}}\textsubscript{2} process and proposed a dipolar, planar oxyallyl ion 21 formed by chloride departure, followed by rearrangement to the Favorskii ester
This intermediate would be amenable to the formation of $\alpha$-methoxyketone derivatives, which are common by-products in Favorskii rearrangements.

Scheme 4. Dewar’s proposed mechanism of the Favorskii reaction.

A dipolar ion (i.e., a zwitterion) was suggested as a prominent Favorskii intermediate in subsequent studies by Bordwell (vide infra). At this time, both cyclopropanone and oxyallyl intermediates are considered to be practicable intermediates in the course to Favorskii products as neither has been precluded empirically.

II. Bordwell’s contribution to evidence for the Favorskii rearrangement.

A luminary in the study of the Favorskii rearrangement was Bordwell, who, in a series of 11 reports, communicated the results of his all-encompassing efforts towards an improved comprehension of the mechanism: (1) rate parameters (rate determining steps, accelerating/decelerating factors), (2) substituents (structure/reactivity correlations), (3) neighboring groups (anchimeric assistance), (4) nucleofuges, (5) temperature, (6) base concentration and ionic strength, (7) intermediates, (8) competing mechanisms (by-products), and
(9) stereochemistry,\textsuperscript{30} among several others. A conspectus of some noteworthy findings from his research concerning the Favorskii rearrangement is presented.

\textbf{A. Rate, Isotope and Nucleofuge studies.} His seminal publication\textsuperscript{14} scrutinized earlier findings by Loftfield\textsuperscript{13} concerning the disparate product distributions obtained with $\alpha$-Br vs. $\alpha$-Cl ketone precursors. In his studies, Bordwell evaluated isotope and leaving group effects on rates of halogen ion release for protonated and deuterium labeled 2-chloro and 2-bromo-4,4-diphenylcyclohexanones (23-26) (Figure 1) in sodium methoxide/methanol at 0$^\circ$C, common conditions for the Favorskii rearrangement.

\textbf{Figure 1.} Diphenylcyclohexanones examined in the Favorskii reaction.

In choosing di-substitution at the 4 position, he reasoned that the axial orientation of the diphenyl groups would encumber the preferred axial position of the halogen that promotes the formation of epoxyether intermediates (precursors for side reactions) and thus optimize conditions for monitoring any of the aforementioned effects. The series of steps involved in these reactions of interest for 25 are depicted in Scheme 5 with associated steady state expressions.
Scheme 5. The reaction of 2-bromo-4,4’-diphenylcyclohexanone, 25, with CH$_3$O$^-$ to corresponding intermediates. The rates for each step are provided, along with an overall rate expression.

\[
\text{obs} = \frac{k_1 k_2}{(k_1 + k_2)}
\]

Preliminary results from these studies are summarized in Table 1. The lack of a 1$^0$ kinetic isotope effect (KIE) and a considerable rate enhancement with Br$^-$ over Cl$^-$ as the nucleofuge were evident and from these observations, it was asserted that the enolate formation for 25 was formed reversibly and chloride departure was rate determining.

Table 1. Summary of rates of halide release, k; isotope exchange ratios, k$_H$/k$_D$; and rate comparisons, k$_{Br}$/k$_{Cl}$ for reactions of 23-26 under Favorskii conditions. Table adapted from ref. 14.

<table>
<thead>
<tr>
<th>Substrate</th>
<th>k ($10^4$ M$^{-1}$s$^{-1}$)</th>
<th>k$_H$/k$_D$</th>
<th>k$<em>{Br}$/k$</em>{Cl}$</th>
</tr>
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<tbody>
<tr>
<td>α-chlorocyclohexanone</td>
<td>40</td>
<td></td>
<td></td>
</tr>
<tr>
<td>23</td>
<td>0.61</td>
<td>1.05</td>
<td>(23/24)</td>
</tr>
<tr>
<td>24</td>
<td>0.58</td>
<td></td>
<td></td>
</tr>
<tr>
<td>25</td>
<td>71</td>
<td>116</td>
<td>(25/23)</td>
</tr>
<tr>
<td>26</td>
<td>17</td>
<td>4.1</td>
<td></td>
</tr>
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</table>
The near unity \( \frac{k_{H}}{k_{D}} = 1.05 \) for \textbf{23} suggested a rapid, possibly reversible deprotonation prior to the rate determining C-Cl ionization. The relatively low KIE \( \frac{k_{H}}{k_{D}} = 4.1 \) for \textbf{25}, typically 13 for acetone in hydroxide at 0\(^{\circ}\) C,\(^{31}\) indicated the high probability of partial deuterium exchange prior to Br\(^{-}\) departure, but was insignificant when compared to the prodigious leaving group effect \( \frac{k_{Br}}{k_{Cl}} = 116 \) in contribution to the rate determining step, which, from this data, was posited to be proton abstraction for \textbf{25}. The rationalization for differences in the RDS of \textbf{23} and \textbf{25} reside in the relative barriers to the formation of the Favorskii intermediate, for each are expected to be similar, but that a “nucleofuge effect” preponderates; Br\(^{-}\) is a much better nucleofuge than Cl\(^{-}\).

To corroborate this proposal, assays to establish the extent of H-D exchange with \textbf{24} and \textbf{26} and the Favorskii rearrangement product (Table 2) were conducted by \(^1\)H NMR and mass spectrometry (MS) prior to and at one half-life, \( t_{1/2} \sim 3\) h, of the disappearance of reactant. It must be noted that the samples of \textbf{24} and \textbf{26} were not replete with deuterium (<3 D atoms/molecule on average); the rigors of synthesis and prolonged processing times, i.e., isolation and purifying stages, precluded the production of 100% d\(_3\) for either target. The larger amount of deuterons for \textbf{26} vs. \textbf{24} was attributed to truncated processing times; there appears to be a connection between the degree of deuterium insertion and processing time length. In addition, some of the deuterium loss of \textbf{26} at \( t_{1/2} \) was ascribed to processing.

Notably fast H-D exchange occurred during the reaction with \textbf{24}, evidenced by complete loss of deuterium in MS analysis, which revealed that nearly 96% of \textbf{24}
Table 2. Summary of hydrogen-deuterium exchange with 24, 26, and the rearranged Favorovski esters over one half life (~3 h) at 0°C with 0.05 M CH₂Ο⁻. Table adapted from ref. 14.

<table>
<thead>
<tr>
<th>Substrate</th>
<th>Material Analyzed</th>
<th>Deuterons(^a)</th>
<th>NMR(^b)</th>
<th>Mass Spec.(^c)</th>
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<tr>
<td>24</td>
<td>24</td>
<td>~2</td>
<td>1.8</td>
<td>d₀ 3.2</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₁ 24.7</td>
</tr>
<tr>
<td></td>
<td></td>
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<td></td>
<td>d₂ 60.5</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₃ 11.6</td>
</tr>
<tr>
<td>24</td>
<td>24 (recovered at (t_{1/2}))</td>
<td>0</td>
<td>0.3</td>
<td>d₀ 96.7</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₁ 3.3</td>
</tr>
<tr>
<td>24</td>
<td>Rearranged Ester</td>
<td>0</td>
<td>0.05</td>
<td>d₀ 94.6</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₁ 5.4</td>
</tr>
<tr>
<td>26</td>
<td>26</td>
<td>~2.8</td>
<td>2.28</td>
<td>d₀ 0.7</td>
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<td>d₁ 10.9</td>
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<td>d₂ 47.6</td>
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<td></td>
<td>d₃ 40.8</td>
</tr>
<tr>
<td>26</td>
<td>26 (recovered at (t_{1/2}))</td>
<td>1.5</td>
<td>1.68</td>
<td>d₀ 8.0</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₁ 33.1</td>
</tr>
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<td></td>
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<td>d₂ 49.5</td>
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<td></td>
<td></td>
<td></td>
<td>d₃ 9.4</td>
</tr>
<tr>
<td>26</td>
<td>Rearranged Ester</td>
<td>0.7</td>
<td>0.78</td>
<td>d₀ 22.2</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₁ 77.1</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>d₂ 0.7</td>
</tr>
</tbody>
</table>

\(^a\)Deuterium atoms per molecule. \(^b\)The number of deuterium atoms determined by \(^1\)H NMR. \(^c\)Determined by mass spectrometry.

and 95% of the rearranged ester contained no deuterons at \(t_{1/2}\). These observations adduce a rapid pre-equilibrium with 24/23, where \(k_{-1} \gg k_2\) (Scheme 5) and thus \(\alpha\)-proton abstraction as rate determining. This phenomenon is significantly reduced for 26 based on considerable deuterium retention at \(t_{1/2}\) (1.5 deuterons from initial 2.8 deuterons). Bordwell asserted that some of this deuterium loss was not from the reaction but due to; a) processing and b) background H-D exchange that is known to occur with \(\alpha\)-chloro and \(\alpha\)-bromoketones in protic solvents.\(^3^2\) Parallel studies
involving the hydrolysis and reesterification of a partially deuterated, rearranged ester revealed virtually no H-D exchange. Thus, the presence of only 0.05 deuterons per molecule of the rearranged ester corroborates the notion of a swift pre-equilibrium for deprotonation and C-Cl ionization as the RDS.

For 26, based on greater retention of deuterium and, above all, the prodigious $k_{Br}/k_{Cl} \approx 116$ (a typical $k_{Br}/k_{Cl}$ value for $S_N2$ and $S_N1$ reactions is $50^{33}$), it was postulated that $k_2 \gg k_1$ and thus $k_{obs}$ approximated $k_1$. It should be noted that the substantial yield of epoxy ether byproducts from attack of CH$_3$O$^-$ at the carbonyl of unsubstituted $\alpha$-bromocyclohexanone, as demonstrated by Loftfield,$^{19}$ can be explained by competitive side reactions with large $k_{Br}/k_{Cl}$.$^{34}$ The results confirm the proposed $\alpha$-deprotonation as the RDS in the Favorskii process for 26; this also denotes a change in RDS compared to 24. The *nature of the nucleofuge* was emphasized as a pertinent factor in determining the outcomes of Favorskii rearrangement in these studies.

Expeditive H-D exchange obviates a concerted process for the $\alpha$-chlorocyclohexanones 23 and 24. Complementary studies involving 23 and 25, where a methyl or -C$_{20}$H$_{36}$ (cholestan-3-one) had been commuted with one of the phenyl moieties, demonstrated dissimilar $k_{Br}/k_{Cl}$ ratios (35 for -CH$_3$, 52 for cholestan-3-one). If the mechanism was concerted, a relatively constant, sizeable nucleofuge influence would be anticipated. This was inconsistent with the above observations and effectively precluded a concerted process for the Favorskii mechanism for 25 and 26.
As Bordwell anticipated, the 4-substituted-2-halocyclohexanones established the prevalent formation of rearranged, ring contracted esters. For 2-chlorocyclohexanone analogs, the route to these esters was favored whether or not there existed ring substituents. In contrast to this, unsubstituted 2-bromocyclohexanone exhibited an epoxyether as the primary product, resulting from a reaction course through an epoxide intermediate. It is apparent that the 4,4’ moieties effectively prevented the epoxide intermediate from forming, as no epoxyethers were reported. These studies divulged, in addition to the salient nucleofuge impact on the RDS, a structural influence that is pronounced for ring substitutions of α-bromocyclohexanones and that exert direct control over the outcomes of the Favorskii rearrangement.

B. Substituent Effects, Linear Free Energy Relationships (LFER’s), Salt and Media effects, and Further Evidence for a Nucleofuge Effect.

In a supervening report,²³ Bordwell conveyed results from a) temperature studies; b) studies involving adjusting base [CH₃O⁻] and ancillary salts [LiClO₄] (ionic strength) to discern these effects on overall yields and product distributions; c) structure-reactivity linear free energy relationships, i.e., how meta and para arene substituents impact the rates (kₚₒₛ) of nucleofuge release and Favorskii ester formation; d) an expansion on the nucleofuge character of the halogen, i.e., its influence on the mechanism and product outcomes. The model systems for these studies were a series
of 1-chloro-1-phenylpropan-2-ones (28) in CH$_3$OH at 0°C (Scheme 6). The results from these experiments are listed in Table 3.

**Scheme 6.** Disparate reaction pathways for $\alpha$-chlorobenzyl methyl ketone derivatives in CH$_3$O/CH$_3$OH.

![Scheme 6 diagram](image_url)

$R = p$-OCH$_3$, $p$-CH$_3$, $m$-CH$_3$, H, $m$-OCH$_3$, $m$-Cl, $m$-NO$_2$, $p$-NO$_2$

**Table 3.** Results of the reactions of 28 at various [CH$_3$O$^-$] and [LiClO$_4$] in CH$_3$OH at various temperatures. Table adapted from ref. 23.

<table>
<thead>
<tr>
<th>[MeO$^-$], M</th>
<th>T (°C)</th>
<th>Yield of 29 (%)</th>
<th>Yield of 32 (%)</th>
</tr>
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<tbody>
<tr>
<td>Inverse addition$^a$</td>
<td>0</td>
<td>9</td>
<td>68</td>
</tr>
<tr>
<td>0.05</td>
<td>0</td>
<td>13</td>
<td>81</td>
</tr>
<tr>
<td>1.00</td>
<td>0</td>
<td>17</td>
<td>77</td>
</tr>
<tr>
<td>2.00</td>
<td>0</td>
<td>21</td>
<td>70</td>
</tr>
<tr>
<td>0.05</td>
<td>65</td>
<td>37</td>
<td>51</td>
</tr>
<tr>
<td>2.00</td>
<td>65</td>
<td>61</td>
<td>35</td>
</tr>
<tr>
<td>0.05 (1 M LiClO$_4$)</td>
<td>65</td>
<td>57</td>
<td>17</td>
</tr>
<tr>
<td>0.05 (2 M LiClO$_4$)</td>
<td>65</td>
<td>63</td>
<td>19</td>
</tr>
</tbody>
</table>

$^a$Slow addition of 10% excess 0.05 M [CH$_3$O$^-$] to a solution of 28.
The enhanced yield of 29 at higher temperatures, 13% vs. 37% at 0.05 M, was attributed to the calculated 2-3 kcal/mol higher activation energy, $\Delta G^\ddagger$, of 29 compared with hydroxy ketal 32. This was derived from work done by Bordwell that found that $\Delta G^\ddagger$ for 28, based on rates in 0.05 M NaOCH$_3$/CH$_3$OH between 0° and 25°C, was 17.2 kcal/mol and manifested 29 as the major product. $\Delta G^\ddagger$ for the $\alpha$-methyl derivative of 28 (geminal with -Cl) was calculated to be 19.7 kcal/mol under the same conditions with the primary product being 32. Complementary experiments with 29 and 32 in the presence of CH$_3$O$^-$ at high temperatures evinced no reactivity.$^{23}$ Augmented yields of the Favorskii ester 29 as a function of base concentration were shown to be marginal at 0°C but improved at 65°C. Accompanying these observations was an enhancement in the yield of 29, 37% vs. 57%, as a function of increased ionic strength from increasing the LiClO$_4$ concentration, i.e., a sal effect. It was theorized that the Favorskii reaction was impacted to a greater degree by ionic strength adjustments than the corresponding formation of 32 as evidenced by the higher yields of 29.

A parallel study by Skrobek$^{35}$ divulged improved yields of the Favorskii ester from 1-chlorocyclohexyl methyl ketone in the presence of higher concentrations of CH$_3$O$^-$ or LiClO$_4$ and stereochemical differences of trans-1-chloro-2-methylcyclohexyl methyl ketone arose from changes in [CH$_3$O$^-$] or [LiClO$_4$]. No explanations, however, were proffered. Later studies$^{25}$ with 0.0075 M PhCH$_2$COCH$_2$Cl (40, pg 21, vide infra) in CH$_3$O$^-$/CH$_3$OH and 0.1 M LiClO$_4$ at 0°C revealed a 20-30% acceleration in reaction rate and furthermore, a 100% acceleration
in 1.0 M LiClO$_4$. It was hypothesized that LiClO$_4$ imparted a stabilizing effect on the highly ionic transition state through salt-facilitated ionization of the halide. Bordwell theorized this from similar findings with $t$-butylbromide in 90% aqueous acetone and Ph$_2$CHCl in 80% aqueous acetone,$^{36}$ two $S_N1$ model mechanisms. No further studies, however, were conducted by Bordwell to supplement or to elaborate on the *salt effect*. Complementary to the salt effect was a boost in reaction rates for 40 (vide infra) in 0.05 M CH$_3$O$^-$/CH$_3$OH at 0°C as the water content was increased (Table 4).

**Table 4.** Effect of water composition on the reaction rate, $k_2$, of PhCH$_2$COCH$_2$Cl in 0.05 M CH$_3$O$^-$/CH$_3$OH at 0°C. Table adapted from ref. 25.

<table>
<thead>
<tr>
<th>Vol % H$_2$O</th>
<th>$k_2$ (10$^2$, M$^{-1}$s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>2.63</td>
</tr>
<tr>
<td>5</td>
<td>6.23</td>
</tr>
<tr>
<td>10</td>
<td>13.1</td>
</tr>
<tr>
<td>25</td>
<td>50.3</td>
</tr>
<tr>
<td>50</td>
<td>289</td>
</tr>
</tbody>
</table>

A 110-fold acceleration in the reaction rate for 40 occurred as the water concentration increased from 0 to 50% (2.63 to 289 x 10$^2$ M$^{-1}$s$^{-1}$). A plot of the log $k_2$ vs. the Grunwald-Winstein Y,$^{37}$ which represents quantitative measures of the effect of the “ionizing power” of the solvent on the reaction, provided an excellent correlation (slope = 0.647; $R^2 = 0.995$; standard deviation = 0.037). For 28, a 197-fold increase in reaction rate was distinguished by increasing content of water from 0 to 50% (0.22 to 42 x 10$^2$ M$^{-1}$s$^{-1}$) and thus the slope would be expected to be even
greater. The considerable ionizing power of water \((Y = 3.56\) for ionization of \((CH_3)_3C-Cl)\),\(^{37}\) was apparent from data in Table 4. These observations further support the prior contention that heterolysis of the C-Cl possesses a high degree of ionic character in the transition state and accordingly, Cl\(^-\) departure represents the RDS. A media effect was clearly distinguished from these studies.

Several aryl-substituted functionalities were examined for their impacts on observed rates of halide ion release from 28, i.e., to explore whether a linear free energy relationship existed. The data for substituents are revealed in Table 5. A modest surge in yields of 29 was observed for nearly every substituent as the base concentrations were increased from 0.05 to 1.0 M. The influences of temperature and base concentration on product distribution denoted competing reaction pathways, each of which displayed a first order dependence on \([CH_3O^-]\). The impacts of substituents on yield and \(k_{obs}\) were conspicuously related to their electron releasing and withdrawing capacities relative to -H; para-substituted, electron donating groups

**Table 5.** Summary of observed and Favorskii rate constants and product yields in CH\(_3\)OH/CH\(_3\)OH at 0\(^\circ\) for substituted 28 as a function of substituent. Table adapted from ref. 23.

<table>
<thead>
<tr>
<th>Precursor</th>
<th>(k_{obs}/10^3) M(^{-1}) s(^{-1})</th>
<th>(k_{Favorskii}/10^3) M(^{-1}) s(^{-1})</th>
<th>%Yield of 29</th>
<th>%Yield of 29</th>
<th>%Yield of 32</th>
</tr>
</thead>
<tbody>
<tr>
<td>H</td>
<td>3.09</td>
<td>0.4</td>
<td>13</td>
<td>13</td>
<td>81</td>
</tr>
<tr>
<td>p-NO(_2)</td>
<td>5.09</td>
<td>ND</td>
<td>0</td>
<td>-</td>
<td>62</td>
</tr>
<tr>
<td>p-Cl</td>
<td>6.21</td>
<td>0.31</td>
<td>5</td>
<td>9</td>
<td>83</td>
</tr>
<tr>
<td>p-OMe</td>
<td>39.4</td>
<td>26.8</td>
<td>68</td>
<td>65</td>
<td>-</td>
</tr>
<tr>
<td>m-OMe</td>
<td>2.72</td>
<td>0.33</td>
<td>12</td>
<td>25</td>
<td>79</td>
</tr>
<tr>
<td>p-Me</td>
<td>6</td>
<td>1.5</td>
<td>25</td>
<td>36</td>
<td>56</td>
</tr>
<tr>
<td>m-Me</td>
<td>3.23</td>
<td>0.45</td>
<td>14</td>
<td>22</td>
<td>61</td>
</tr>
<tr>
<td>p-F</td>
<td>7.5</td>
<td>0.83</td>
<td>11</td>
<td>16</td>
<td>65</td>
</tr>
</tbody>
</table>
(-OCH$_3$, -CH$_3$) manifested elevated rates for the production of 29 while para-substituted electron withdrawing groups (-F, -Cl and NO$_2$) evinced preferences for 32. Initial attempts to rationalize these phenomena through linear free energy relationships (LFER) led to poor correlations using log $k_{obs}$ and Hammett $\sigma$ substituent constants ($\rho = -1.1$; $R^2 = 0.875$; standard deviation = 0.275); ameliorated correlations were found by employing $\sigma^+$ constants ($\rho = -2.37$; $R^2 = 0.992$; standard deviation = 0.13 (5%)). The negative $\rho$ suggested an accumulation of positive charge (carbocation character) or a loss of electron density (decreased anionic character) in the transition state of the RDS. From this, Bordwell postulated that two key steps in the mechanism leading to 31 were impacted the most by substituents. First, the reversible addition of CH$_3$O$^-$ to 28, the initial step in forming 31, is estimated to have a $\rho \sim 0.8$ based on analogous studies.$^{38}$ Electron withdrawing groups would be expected to accelerate this process; this prediction was confirmed by the present study. Secondly, halogen departure from 30 was calculated to have a $\rho \sim -0.9$. Electron donating groups were expected to accelerate the rate of this process; results from this study were also consistent with this prediction. The electron donating or withdrawing natures of the arene-substituted functional groups were shown to exert a direct control over reaction pathways of 28 in forming 29 and 32.
C. The Ramberg-Backland (R-B) analogy and role of orbital overlap.

To obtain an improved perspective on the step entailing halide ionization, Bordwell examined the Ramberg-Backland (R-B) reaction\(^{39}\) of \(\alpha\)-halosulfones as an alternative model (Scheme 7). The reaction is well understood and characterized by

\textbf{Scheme 7.} A depiction of the Ramberg-Backland reaction.

\[\text{[Diagram of the Ramberg-Backland reaction]}
\]

\(X = \text{Cl, Br}\)

swift, reversible carbanion formation 34 which mediates the rate determining departure of the halide ion to produce an episulfone 35 that readily extrudes SO\(_2\) to form styrene 36. A leaving group effect, \(k_{\text{Br}}/k_{\text{Cl}} > 50\), though smaller than in the Favorskii reaction, nevertheless, suggested considerable C-X cleavage in the transition state of the RDS. A positive \(\rho\) value (+0.8) in 40\% aqueous dioxane was determined from the rate dependence on substituents. Despite significant C-X ionic character, a positive \(\rho\) indicated that electron withdrawing groups should accelerate C-X heterolysis and required the inchoate C-C bond to be nearly completed in the transition state. This finding diverged from the Favorskii reaction; a large negative \(\rho \approx -1.1\) predicted that electron withdrawing groups would decelerate the rate of C-X cleavage\(^{32}\) and the C-C bond formation would be much less developed. Here,
mechanistic disparities in the two reactions became obvious. The negative charge afforded from hydrogen atom abstraction was localized on the carbon of the sulfone due to poor p-d π orbital overlap, whereas the carbanion was primarily a delocalized entity in the Favorskii reaction (Scheme 8); the p-orbital is held preferentially parallel to the contiguous carbonyl π system (I). To educe the appropriate trajectory for a S_N2 cyclopropanation, the carbanion orbitals must contort out of plane (deconjugate) with the carbonyl (II), localizing electron density, i.e., inducing a hybridization change from sp^2 to sp^3 and thus diminishing the feasibility of this route, at least from an

**Scheme 8.** Delocalized (sp^2) vs. localized (sp^3) electron models.

![Scheme 8](image-url)
energetics perspective. Conversely, an arrangement where the C-Cl bond is held parallel to the enolate p orbital (Scheme 9) effectuates partial π overlap of the

**Scheme 9.** Anchimeric π assistance in establishing a cyclopropanone intermediate.

developing p-orbital of the C-Cl center in the transition state, a stabilizing interaction that would promote the formation of cyclopropanone through disrotatory rotation. Energetically, anchimeric π electron participation in halide discharge to effect Favorskii precursors was perceived to be the favored route. To probe this hypothesis, Bordwell determined first order rate constants of bromide release from the base
promoted anions corresponding to three similar α-bromo substrates in 0.05 M CH$_3$O$^-$/CH$_3$OH at 0°C (Scheme 10). A five order of magnitude rate enhancement was evinced by the enolate, 38, compared to the carboxylate, 37. An obvious neighboring group effect, anchimeric π assistance, seemed reasonable to Bordwell, as the only other feature common to both 37 and 38 was resonance delocalization. Bordwell reasoned that if resonance stabilization predominated, 37 would manifest a more rapid rate of bromide release. For 39, as a sulfur enolate is a less important resonance contributor due to the poor p-d orbital overlap, the more localized carbanion displayed further a five order of magnitude increase for bromide release, imputing σ (electrons localized on a sp$^3$ orbital) electron participation. Bordwell inferred from

**Scheme 10.** Rates of bromide release from anionic α-bromo substrates.
these studies that $\pi$ electron participation was a salient feature in the mechanism of Favorskii reaction.

**D. The influence of $\alpha$-substituents.** Earlier, it was revealed by Bordwell that both the nature and locus of the substituent in cyclohexanones strongly influenced product distributions of the Favorskii reaction, i.e. a structure-reactivity relationship was established. Analogous investigations were extended to linear $\alpha$-haloketones, specifically the role of methyl$^{27,28}$ and phenyl$^{26}$ substituents in altering parameters of the Favorskii reaction such as rates of halide departure and product yields. Previous investigations of 1-chloro-3-phenylpropan-2-one 40 (Scheme 11) in 0.05 M

**Scheme 11.** Reactions of $\alpha$-chloro analogs in CH$_3$O$^-$/CH$_3$OH.
CH$_3$O/CH$_3$OH at 0°C exhibited rapid H-D exchange and a $\rho$ of -5.0,$^{23}$ denoting an initial, reversible deprotonation step and suggesting a high degree of C-Cl bond dissociation in the transition state. From this, Bordwell rationalized that $\alpha$-methyl or phenyl substituents (e$^-$ contributors) should stabilize this accumulation of positive charge and augment the rate of halide ionization, $k_2$. He studied the behaviors of 3-chloro-1-phenylbutan-2-one (43) and 1-chloro-1,3-diphenylpropan-2-one (46) under an assortment of conditions and compared results to those obtained for 40.$^{23}$ Results from these examinations are displayed in Table 6. As observed earlier, a direct

<table>
<thead>
<tr>
<th></th>
<th>40</th>
<th>43</th>
<th>46</th>
</tr>
</thead>
<tbody>
<tr>
<td>Relative rates$^a$</td>
<td>1</td>
<td>&gt;250</td>
<td>&gt;330</td>
</tr>
<tr>
<td>Deuterium exchange</td>
<td>~80%</td>
<td>~6%</td>
<td>N$^b$</td>
</tr>
<tr>
<td>$k_{Br}/k_{Cl}$</td>
<td>63</td>
<td>0.9</td>
<td>1.0</td>
</tr>
<tr>
<td>Hammett $\rho$</td>
<td>-5.0</td>
<td>1.4</td>
<td>1.1</td>
</tr>
<tr>
<td>Salt effect$^c$</td>
<td>25-30%</td>
<td>N$^b$</td>
<td>~7%</td>
</tr>
<tr>
<td>Products with:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2 M NaOCH$_3$</td>
<td>100%</td>
<td>100%</td>
<td>39% 47, no 48$^d$</td>
</tr>
<tr>
<td>0.05 M NaOCH$_3$</td>
<td>100%</td>
<td>61% 44, 39% 45</td>
<td>15% 47, 85% 48</td>
</tr>
<tr>
<td>10$^{-5}$ M NaOCH$_3$</td>
<td>100% 41</td>
<td>100% 45</td>
<td>100% 48</td>
</tr>
</tbody>
</table>

$^a$Chloride ionization. $^b$N = negligible. $^c$0.1 M LiClO$_4$. $^d$An unidentified compound was formed.

correlation was found between [OCH$_3^-$] and product composition for 43 and 46; larger quantities of OCH$_3^-$ favored Favorskii product formation. No changes in product distribution as a function of [OCH$_3^-$] for 40 were noticed; the Favorskii ester was the dominant product in all cases. A mechanistic interpretation will be provided later.
Control experiments were performed for validation and found that: 1) No chloride was released from 43 or 46 in the absence of CH₃O⁻ in CH₃OH over 24 h and 2) neither 44 nor 47 reacted in 2 M [CH₃O⁻] for 2 h; this ruled out degradation of these Favorskii esters to other products, which would result in false percent yields to be reported for their formation. Furthermore, it was found that the product ratios for 40, 43 and 46 remained steady at 0°C and 70°C in 0.05 M CH₃O⁻/CH₃OH, 61% and 39% respectively, negating a temperature effect. In the presence of 0.1 M LiClO₄, a salt effect was minimal for 43 as there was essentially no change in yield and for 46 only a 7% increase was reported; a significant salt effect for 40 amounted to a ~25-30% increase in the production of Favorskii esters.

When the Br⁻ analogs were tested under the same conditions, the product ratios were nearly identical. However, the reaction rates for halide detachment in 10 molar excess CH₃O⁻ under the same conditions revealed no variation for 43 and 46 (kBr⁻/kCl⁻ ∼ 0.9 and 1 respectively) but was pronounced for 40 (~63). This phenomenon, in itself, intimates distinct mechanisms for 40 and 43/46 to rearranged products. Corroborating these are the sizeable differences in ρ for 40 (~5.0) compared to 43 (1.4) and 46 (1.1). A substantial buildup of positive charge in the transition state of the RDS for 40 is anticipated and thus C-Cl bond breakage should possess a large degree of ionic character at the transition state. Analogous to its structural isomer (28) the RDS for 40 is hypothesized as chloride ionization. In contrast, positive ρ values for 43 and 46 indicate a buildup of negative charge in the transition state and adduce that the RDS is, in fact, proton abstraction. As Bordwell anticipated,
the α-methyl 43 and phenyl 46 moieties, through stabilization of the positive charge on the chloride-bearing carbon, unambiguously transposed the RDS of the Favorskii reaction to proton abstraction, comparable. From these results and those obtained from related experiments, Bordwell formulated a mechanism to account for the formation of 44 and α-methoxy ketone (45) (Scheme 12).

**Scheme 12.** Discrete mechanistic pathways for α-chlorobenzylpropiophenone 43 to the rearranged ester (44) and α-methoxyketone (45).

![Scheme 12](image)

The RDS, as previously derived, represents $k_1$, or rate of enolate formation. The rearranged ester (44) was engendered from the enolate intermediate (51) through the Favorskii rearrangement. The β-hydroxy allyl chloride (enol) intermediate (52) was assumed to be in rapid equilibrium with 51 and thus permitted solvolysis to occur in forming 45. The relative magnitudes of $k_2$ and $k_3$, along with direction of equilibrium, control the 44 to 45 ratios. When methoxide concentrations are high, enolate [51] is high, $k_2 > k_3$, and the formation of 44 is promoted. Conversely, when
CH₃O⁻ concentrations are reduced, enol [52] is high, k₃ > k₂, and 45 preponderates. Bordwell estimated that the enol to enolate ratio, [52]/[51] ~ 40 at 0.05 M [CH₃O⁻].²⁶ The α-substituted methyl 43 and phenyl 46 groups were vital to this mechanistic change. Structure-reactivity relationships, once again, were indicated as important factors in the Favorskii reaction. Recall that the Favorskii ester is produced solely from 40 irrespective of [CH₃O⁻]. To recapitulate the importance of [CH₃O⁻] in determining product outcomes of the Favorskii reaction, a side by side, detailed mechanistic depiction of the reaction pathways of 43 and its isomer 53 (R = H) is exhibited in Scheme 13. The primary route to 44 for each precursor, after predominant enolate (51ₐ and 51ₐ) formation when concentrations of CH₃O⁻ were higher, was postulated to be through an oxyallyl zwitterion, 54, which was presumed to be in equilibrium with the reactive cyclopropanone. The competing route to 45 was thought to involve the protonated oxyallyl zwitterions (55) when enols (52ₐ and 52ₐ) preponderated at low [CH₃O⁻]. Rearrangement to 44 was precluded from 55, where CH₃O⁻ addition was favored to furnish 45. An additional consideration in determining the pathways to 44 and 45, though not explicitly mentioned, is the interconversion between oxyallyl intermediates 54 and 55, i.e., a Curtin-Hammett condition. Small energy barriers for proton exchange would rapidly interconvert 54 and 55. High [CH₃O⁻] would favor 54, while low concentrations would favor 55.
Scheme 13. Reaction pathways of isomers 53 and 43 to rearranged 44 and \( \alpha \)-methoxy ketone (45).
An extension\textsuperscript{28} of investigations into structure-reactivity relationships of the Favorskii rearrangement was applied to tertiary $\alpha$-haloketones 46, 49, and 53, each containing two substitutions on the $\alpha$-position of the carbonyl (Scheme 14). Control

**Scheme 14.** Reactions of alkyl substituted $\alpha$-chloro ketones with CH$_3$O$^-$.  

![Scheme 14](image)

$X = \text{Cl, Br}$

experiments demonstrated the isomerization of 53 to 43 at $\sim$20% over 6 months at $\sim$ 0\textdegree C; this was accounted for in determining relevant factors pertaining to product formation. Results from these experiments are displayed in Table 7.
Table 7. Summary of product yields and rates of reaction for 46, 49 and 53 as a function of CH$_3$O$^-$ in CH$_3$OH. Table adapted from ref. 28.

<table>
<thead>
<tr>
<th>Substrate</th>
<th>[CH$_3$O$^-$], M</th>
<th>% Yield $\alpha$-methoxyketone 45/48</th>
<th>% Yield of Favorskii ester 44/47</th>
<th>$k$ ($10^3$ M$^{-1}$ s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>46 (Br)</td>
<td>0.05</td>
<td>97</td>
<td>3</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>60</td>
<td>35</td>
<td>-</td>
</tr>
<tr>
<td>46 (Cl)</td>
<td>0.05</td>
<td>-97</td>
<td>-3</td>
<td>-</td>
</tr>
<tr>
<td>49 (Cl)</td>
<td>0.05</td>
<td>-</td>
<td>10</td>
<td>0.016</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>-</td>
<td>45</td>
<td>-</td>
</tr>
<tr>
<td>49 (Br)</td>
<td>1</td>
<td>-</td>
<td>70</td>
<td>0.024</td>
</tr>
<tr>
<td>53$^{ab}$ (Cl)</td>
<td>0.05</td>
<td>30</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

$^a$The yield of Favorskii ester is 100% in 2 M NaOCH$_3$. $^b$At 10$^{-3}$ M, the $\alpha$-methoxyketone was the sole product.

An improved yield of Favorskii ester as the [OCH$_3^-$] increased was manifested. This observation seemed to bolster the plausibility of the previously posited enol/enolate mechanism in accounting for product distributions, i.e., a greater quantity of OCH$_3^-$ (greater basicity) would shift the enol/enolate towards the enolate that promotes the rearrangement process. The lack of a halogen effect, $k_{\text{Br}}/k_{\text{Cl}} \sim 1$, for 49 was similar to previously encountered examples and signifies rate determining proton abstraction. The rate for proton abstraction of PhCH(Cl)COCH$_3$, 28 (vide supra), was accelerated 35 fold compared to 49; this phenomenon implies an *encumbering (possibly steric)* effect exerted by the methyl group in retarding the rate of proton abstraction. The RDS changes from proton abstraction to halide departure. A steric component to the Favorskii mechanism is further advanced when one compares the deprotonation rates of 53 with 46; the former disclosed a 35 fold acceleration over the latter. The RDS for 46 was postulated as proton abstraction based on analogy with previous work.$^{25}$
E. Evidence for a dipolar (oxyallyl) intermediate in the Favorskii reaction.

Bordwell inadvertently provided evidence for the formation of Dewar’s\textsuperscript{20} proposed dipolar ion in studies\textsuperscript{29} involving 1-chloro-1,1-diphenylpropanone (56) and 3-chloro-1,1-diphenylpropanone (60) in CH\textsubscript{3}O/CH\textsubscript{3}OH at 0°C; his specific aims were to probe the effects phenyl substituents on the Favorskii reaction.

Scheme 15. Product distributions from the reaction of α-chloro ketones with low concentrations of CH\textsubscript{3}O\textsuperscript{−}.

At 0.05 M CH\textsubscript{3}O\textsuperscript{−}, both precursors gave the Favorskii ester (57) in approximately 97% yield. At lower concentrations of base, however, an array of products was obtained (Scheme 15), along with, perhaps most interestingly, 59. Noteworthy, were the identical constituents that materialized from each precursor under Favorskii conditions, although the compositions of each product set were
discrete. It was reasoned that the emergence of 59 stemmed from low [CH₃O⁻], permitting the intramolecular cyclization of the enolate to engender an indanone, 61, that thenceforth, would react with O₂ in the presence of CH₃O⁻ (Scheme 16). This

Scheme 16. Proposed mechanism for the reaction of α-chloroketone, 56, with low [CH₃O⁻] to form the methyl ester, 59.

proposed sequence was supported by an independent study from Smith⁴⁰ that demonstrated an “unknown acid, melting point 131°C” from reaction of 61 with cold sodium hydroxide. In a supplementary study, a pure sample of 61 promptly transformed to 59 upon oxidation (O₂) in a methanolic solution of methoxide. The rate of halide ionization for 56 was estimated to be enhanced 625 fold compared to its mono-phenyl analog; this unequivocally demonstrated a stabilizing effect by the phenyl moiety. In addition, only about 11% deuterium exchange occurred with 56.
The RDS for 56, therefore, was posited to be deprotonation. A “dipolar-like” transition state was advanced, resulting from halide ion departure from the enolate intermediate. It was reasoned that the second phenyl group would effectively disperse the positive charge accumulation at the C-Cl center through resonance. A parallel phenomenon, though not as marked, was observed for 60, where the rate of halide ionization was augmented 25 fold vs. its unsubstituted analog 40. Additionally, $k_{Br}/k_{Cl}$ was found to be ~4, supporting enolate formation as being rate determining. After extensive, congruent investigations using analogs of 56 and 60, Bordwell maintained that an oxyallyl intermediate (62) was formed (Scheme 17).

**Scheme 17.** Suggested dipolar (oxyallyl) mechanism to indanone (61).
Hoffmann$^{41}$ independently determined energetic preferences (22 kcal/mol) for dipolar ions at equilibrium relative to cyclopropanone. However, Dewar$^{42}$ and Liberles$^{43}$ asserted cyclopropanones to be more stable by 78 and 83 kcal/mol, respectively.

**F. Studies in Stereochemistry.** Stereochemical investigations of the classical Favorskii reaction have been carried out by several groups,$^6$ most notably Stork and Borowitz,$^{18}$ who discerned the stereospecific formation of esters from non-epimerizable ketones (Scheme 18). They suggested an operating, inversion or $S_N2$

**Scheme 18.** Stork-Borowitz proposal for stereochemical inversion of precursors to Favorskii products.

process. Solvent polarity, however, was determined to direct the stereochemical outcome; this theory was advanced by House$^{44}$ et. al., who’s studies divulged the formation of inversion products only in non-polar media.
Bordwell\textsuperscript{30} also explored Favorskii reactions involving stereochemical conversions, specifically 2-bromo-4-methyl-4-phenylcyclohexanone (63) and the \textit{cis/trans} isomers of 2-bromo-5-methyl-5-phenylcyclohexanones (66) under Favorskii conditions. The reactions of 63 and 66 to generate corresponding products are outlined in Scheme 19 and results presented in Table 8.

\textbf{Scheme 19.} Reactions of $\alpha$-bromo ketones 63 and cis/trans diastereomers 66 with CH$_3$O$^-$ in CH$_3$OH.

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|}
\hline
Substituent & [NaOCH$_3$], M$^a$ & $\%$ ester & $\%$ hydroxyketal \\
\hline
63          & $\sim 10^{-3}$ & 40 (1:3, 64:65) & 36 \\
            & 2            & 55$^a$         & 37 \\
66          & 0.05         & 9 (57:6:14:23, 64:65:67:68) & ND$^b$ \\
            & 1            & 22$^b$         & ND \\
\hline
\end{tabular}
\caption{Results of reactions of 63 and 66 with various [CH$_3$O$^-$] in CH$_3$OH at 0°C.}
\footnotesize{Table adapted from ref. 30.}
\end{table}

$^a$Product ratio not provided. $^b$Percentage not given.
For both precursors, a pronounced increase in Favorskii ester yield was seen to accompany concentrated base solutions, a common occurrence for several α-haloketones previously encountered. The relatively enhanced ester yields for 63 vs. 66 was evidence for a 1,3-diaxial effect, specifically in impeding the halide from adopting the axial position, a favored orientation for epoxyether (α-methoxyketone precursor) formation. This effect was not revealed for 66. When the reaction of 63 was conducted at low [CH$_3$O$^-$], a 40% ester yield was obtained with relative inverse ratios of 1:3 for 64 and 65. Reactions of 66 (either isomers) afforded 64, 65, and two other esters 67 and 68 with relative ratios of 57:6:14:23, respectively, in 0.05 M CH$_3$O$^-$ in methanol, that changed to 45:7:9:39 for 64, 65, 67, and 68, respectively, in 2 M [CH$_3$O$^-$] in methanol, and 26:24:28:22, respectively, when equilibrated (>150 h reflux). It appeared that greater base concentrations fostered the formation of 68 over 64. It was postulated that for 63, an internal S$_N$2 reaction (Loftfield’s mechanism$^{13}$) would educe an inverted product if the halide was equatorial and retention if axial. The principal ester for 63 was found to be 64, advancing the notion that retention had occurred (presumed disjunction of an axial halide). This is not manifested with 66 (cis), where 65 would have been anticipated to be major product according to Loftfield’s mechanism, but, in fact, demonstrated a preponderance of 64. The uniform product distribution for cis/trans 66 denoted that an epimerization at the halide locus may have been operative. To furnish an S$_N$2 reaction, cis/trans 66 would have to first equilibrate and then generate the ester via the equatorial halide; an inversion step required an equatorial halide orientation for 66. Consequently,
Bordwell proposed that the dipolar ion intermediate was a feasible precursor to the Favorskii ester (Scheme 20). Upon the formation of the dipolar ion, 69, a disrotatory ring closure was effected, producing cyclopropanones 70 and 71. The stereochemistry of the products was decided following halide detachment, which adequately justified the product distributions above. Low \([\text{CH}_3\text{O}^-]\) advanced the formation of 65 over 64 but the converse at 2 M concentration. Bordwell speculated that this phenomenon was due to the more rapid formation of 71 over 70; low concentrations of \(\text{CH}_3\text{O}^-\) permitted equilibrium to be achieved between the two cyclopropanone intermediates. The product ratios were, thus, controlled by the equilibrium position and rate

**Scheme 20.** Bordwell’s suggested reaction pathways of 63 to diastereomers 64 and 65 in \(\text{CH}_3\text{O}^-\), directed by the oxyallyl intermediate (69).
constants of reactions of cyclopropanone or corresponding hemi methylketal with CH$_3$O$^-$. The product distributions were directed by the rates in which 70 and 71 were produced. The formation of 4 esters from cis/trans 66 was reasoned to occur from two cyclopropanone intermediates, 66a and 66b (Equation 3), each of which reacted with CH$_3$O$^-$ in CH$_3$OH to bear two products; a) 66a, which manifested the phenyl and carbonyl groups trans to each other, reacted with CH$_3$O$^-$ to produce 64 and 67 and b) 66b, which engendered 65 and 68 under the same conditions. Increasing the [CH$_3$O$^-$] from 0.05 M to 2.0 M seemingly improved the yields of 66b over 66a, since the ratios of 65 and 68 to 64 and 67 increased. A similar explication as for the product ratios from 63 was advanced, i.e., a zwitterion intermediate first formed that underwent disrotatory ring closure in two modes to make the two cyclopropanones. In the end, however, Bordwell conceded that further studies were required to better comprehend the stereochemical inversions and retentions that were exhibited by the product distributions.

Eq. 3

![Chemical structure diagram](image)
III. Computational Investigations. Andres\textsuperscript{45} and coworkers recently communicated results from extensive computational investigations using \textit{ab initio} calculations at the Hartree-Fock level of theory with 6-31G* and 6+31G* basis sets and electron correlations at the MP2/6-31G* level based DFT, BLYP/6-31G* of the Favorskii reaction mechanism. Hydroxide ion (HO\textsuperscript{−}) mediated rearrangement of \(\alpha\)-chlorocyclobutane to cyclopropanecarboxylic acid was utilized as a model system. Two mechanisms were accentuated in the investigations; 1) the signature Favorskii route with cyclopropanone intermediate and 2) semibenzilic path, which has been coined the “quasi”-Favorskii reaction and will be elaborated on later, in the gas and aqueous phase. For each mechanism, a series of reactant complexes (RC), transition states (TS), intermediates (I), and corresponding precursor (R) and product (P) entities were determined along with relative energies. These were termed stationary points. The semibenzilic acid route showed 6 stationary points (Scheme 21).
Scheme 21. The proposed semibenziolic mechanism of conversion of α-chlorocyclobutanone to cyclopropane carboxylic acid through calculated stationary points. Reproduced with permission from the ACS, ref. 45.
The relative energies for stationary points corresponding to the semibenzilic acid rearrangement are tabulated for gas and aqueous phases in Table 9.

**Table 9.** Relative energies (kcal/mol) of stationary points associated with the semibenzilic rearrangement mechanism, in the gas and aqueous phases, as determined by various computational methods. ND, not determined. Table adapted from ref. 45.

<table>
<thead>
<tr>
<th></th>
<th>Gas Phase</th>
<th>Aqueous Phase</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>HF/6-31G*</td>
<td>HF/6+31G*</td>
</tr>
<tr>
<td>R</td>
<td>0.00</td>
<td>0.00</td>
</tr>
<tr>
<td>RC</td>
<td>ND</td>
<td>-19.07</td>
</tr>
<tr>
<td>TS1A</td>
<td>ND</td>
<td>-17.58</td>
</tr>
<tr>
<td>I1A</td>
<td>-63.97</td>
<td>-44.86</td>
</tr>
</tbody>
</table>

In the gas phase, nucleophilic approach by HO\(^-\) was calculated to be, for the most part, barrierless; the initial interaction was manifest by strong hydrogen bonding between the anionic HO\(^-\) and \(\alpha\)-hydrogen atoms to the carbonyl, leading to a stabilized RC complex. The RC complex was thought to ready HO\(^-\) for nucleophilic attack on the carbonyl, in assembling the intermediate IA, through a transition state, TS1A; the energy barrier to IA ranged from 1.5 (HF/6-31G*) to 8.6 (BLYP/6-31G*) kcal/mol; this approach was calculated to occur on the very flat region of a potential energy surface, PES, exhibited in Figure 2.
Figure 2. A potential energy illustration of the semibenzilic rearrangement process, revealing relative energies (in kcal/mol) of stationary points on the reactive surface that were calculated at the HF/6-31G* level. The broken line indicates the aqueous phase; the bold line, gas phase. Reproduced by permission of the ACS, ref. 45.

The rearrangement step (TS2A) displayed a diverse range of activation energies, 17.6 kcal/mol (HF/6+31G*) to 2.0 kcal/mol (BLYP/6-31G*); this was the obvious rate determining step for the reaction, as collectively, these energies were higher than those calculated for TS1A..

In an aqueous environment, the PES was quite disparate from the gas phase for the semibenzilic acid rearrangement. Reliable relative energies originated only from HF/6-31G* level computations. The rate determining step changes to TS1A (recall that it was TS2A in the gas phase from HF/6+31G* and MP2/6-31G* calculations) by nucleophilic attack by HO⁻ at the carbonyl; the activation barrier for this was 5.1 kcal/mol. An extensive solvation of HO⁻ by water was designated as the primary source of this observation and further impacted the relative energies of I1A.
Scheme 22. The Favorskii rearrangement mechanism of $\alpha$-chlorocyclobutanone to cyclopropane carboxylic acid through calculated stationary points. Reproduced with permission from the ACS, ref. 45.
and TS2A that were destabilized 34.4 and 22.0 kcal/mol, respectively. An additional solvation effect was manifest on the energy barrier for the rearrangement (TS2A) step, which dropped from 15.4 kcal/mol in the gas phase to 3.0 kcal/mol in solution.

A total of 8 stationary points were determined for the Favorskii rearrangement mechanism (Scheme 22). The relative energies for associated stationary points in the gas and aqueous phases are presented in Table 10.

Table 10. Relative energies (kcal/mol) of stationary points associated with the cyclopropanone rearrangement mechanism in the gas and aqueous phases, as determined by the connoted computational methods. ND, not determined. Table adapted from ref. 45.

<table>
<thead>
<tr>
<th></th>
<th>HF/6-31G*</th>
<th>HF/6+31G*</th>
<th>MP2/6-31G*</th>
<th>BLYP/6-31G*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Gas Phase</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>R</td>
<td>0.00</td>
<td>0.00</td>
<td>0.00</td>
<td>0.00</td>
</tr>
<tr>
<td>RC</td>
<td>ND</td>
<td>-19.07</td>
<td>-36.57</td>
<td>-54.21</td>
</tr>
<tr>
<td>TS1B</td>
<td>ND</td>
<td>-15.46</td>
<td>-32.83</td>
<td>-45.66</td>
</tr>
<tr>
<td>I1B</td>
<td>-49.50</td>
<td>-31.28</td>
<td>-56.14</td>
<td>-64.35</td>
</tr>
<tr>
<td>TS2B</td>
<td>-42.17</td>
<td>-22.83</td>
<td>-50.05</td>
<td>-60.79</td>
</tr>
<tr>
<td>I2B</td>
<td>-47.68</td>
<td>-29.79</td>
<td>-45.49</td>
<td>-49.89</td>
</tr>
<tr>
<td>TS3B</td>
<td>-6.91</td>
<td>12.80</td>
<td>-9.18</td>
<td>-24.14</td>
</tr>
<tr>
<td>P</td>
<td>-115.15</td>
<td>-94.03</td>
<td>-107.07</td>
<td>-109.58</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Aqueous Phase</th>
</tr>
</thead>
<tbody>
<tr>
<td>R</td>
<td>ND</td>
</tr>
<tr>
<td>RC</td>
<td>-20.74</td>
</tr>
<tr>
<td>TS1B</td>
<td>-19.36</td>
</tr>
<tr>
<td>I1B</td>
<td>-23.14</td>
</tr>
<tr>
<td>TS2B</td>
<td>-20.04</td>
</tr>
<tr>
<td>I2B</td>
<td>-41.79</td>
</tr>
<tr>
<td>TS3B</td>
<td>-11.63</td>
</tr>
<tr>
<td>P</td>
<td>-100.55</td>
</tr>
</tbody>
</table>
Akin to the semibenzilic mechanism, the RC was determined as the initial point of contact for HO\(^{-}\) and \(\alpha\)-chlorobutanone in the gas phase. The activation barrier for proton abstraction (TS1B) ranged from 3.61 (HF/6+31G\(^*\)) to 8.55 kcal/mol (BLYP/6-31G\(^*\)), while that for the formation of cyclopropanone (TS2B) spanned from 3.56 (BLYP/6-31G\(^*\)) to 8.45 (HF/6+31G\(^*\)) kcal/mol. The rearrangement step through TS3B was established as the rate determining step in all methods of calculation; the energy barriers ranged from 42.59 (HF/6+31G\(^*\)) to 25.77 (BLYP/6-31G\(^*\)) kcal/mol. The cyclopropanation pathway for rearrangement in water manifested stationary point energies that were appreciably higher than in the gas phase, matching earlier observations for semibenzilic acid in water. A potential energy surface diagram denoting the cyclopropanone rearrangement route is provided in Figure 3.

**Figure 3.** A potential energy surface diagram of the cyclopropanone rearrangement mechanism, revealing relative energies (in kcal/mol) of stationary points on the reactive surface that were calculated at the HF/6-31G\(^*\) level. The broken line indicates the aqueous phase; the bold line, gas phase. Reproduced with permission by the ACS, ref. 45.
Similar to the semibenzilic mechanism above, a change in curvature of the incipient reaction steps inferred pronounced, solvation-incited encumbering effects on the reactivity of HO`. A very low energy barrier (TS1B, 1.4 kcal/mol) to forming the enolate intermediate I1B was manifested. The ensuing conversion to cyclopropanone through TS2B disclosed an energy barrier of 3.1 kcal/mol, a decline from the gas phase that was 7.2 kcal/mol. Lastly, the rearranging, rate determining step through TS3B, showed an energy barrier of 30.2 kcal/mol (I2B); this represented a steep drop compared to 40.8 kcal/mol in the gas phase. The exothermicity of the reaction was reduced by 15 kcal/mol compared to the gas phase. Akin to results for the semibenzilic mechanism, a general solvent influence on the cyclopropanone mechanism was to lower the activation barriers to intermediates and the corresponding product; this phenomenon was particularly noticeable on the PES (Figure 3). The semibenzilic mechanism appeared to be the most favored pathway to the Favorskii acid in either phase, by virtue of the drastically reduced energy barriers of rate determining steps to rearranged product. However, the inherently large ring strain of cyclobutanones, though not considered in the report, would be anticipated to play a substantial role in directing the preferred reactivity through the semibenzilic mechanism which requires fewer steps to the rearranged product.

IV. Variations to the classical Favorskii Reaction.

As alluded to earlier, several variations of the Favorskii reaction have been recognized and implemented in the realm of synthesis: 1) Non-enolizable ketones
can undergo rearrangements to acids/esters in what is termed the “quasi-Favorskii” reaction\(^\text{46}\) (Equation 4), this likened to a semibenzilic acid type of rearrangement. The primary distinction from the classical Favorskii is that no enolate or cyclopropanone intermediates are formed in the reaction course. 2) Ketones with \(\gamma\)-nucleofuges

\[ \text{R = H, alkyl} \]

Eq. 4

can undergo base-moderated rearrangements, the “homo-Favorskii” reaction\(^\text{47}\) (Scheme 23), to produce cyclobutanones, which, being more robust than cyclopropanone homologs, are isolable.\(^3\)

**Scheme 23.** Proposed mechanism of the homo-Favorskii reaction.
In general, reactions involving the formation of acids/esters from base facilitated rearrangements of α-nucleofuge substituted ketones can be considered the “Favorskii” sort. Up to this point, the focus has been on the ground state Favorskii reaction. The excited state complement is now presented.

V. The Photochemical Version of the Favorskii Rearrangement.

A more obscure, lesser known reaction involving rearrangements of α-substituted ketones to effect acids/esters in the excited state, the “Photo-Favorskii”, has, comparatively, received little attention.\(^\text{10}\) The earliest report of photolyses involving α-haloketones, practical antecedents for these types of rearrangements, was by Strachan et. al.,\(^\text{48}\) where chloroacetone was irradiated either neat or in non-hydroxylic solvents at 313 nm. Although a curious array of photoproducts was conveyed, including acetone, no rearrangements were demonstrated. However, the formation of acetone as a principal photoproduct intimated homolytic C-Cl bond scission and thus radical transients, which have been posited as viable precursors to Favorskii products. Though not broached in this report, it can be surmised that the excited state pathway for chloroacetone photolysis to photoproducts was through the triplet, as acetone is known to intersystem cross with an efficiency of near unity.\(^\text{49}\) Multiplicities will become a prominent interest later in this thesis. The seminal publication for the photo-Favorskii reaction was by Anderson and Reese\(^\text{50}\) almost 50 years ago and involved UV irradiation of 1% α-chlorooacetophenone derivatives in ethanol over a 1-2 hour time span. With certain substituents \((R = p-\text{OH}, p-\text{OCH}_3, \text{and})\)
two major products, acetophenone (72) and ethyl 2-phenylacetate (73) were obtained, the latter in abundance (Equation 5).

\[
\begin{align*}
\text{R} &= p\text{-OH}, p\text{-OMe}, o\text{-Me} \\
\text{Eq. 5}
\end{align*}
\]

The pathway to 72, as depicted in Equation 5, was hypothesized to occur by the homolytic cleavage of the C-Cl bond, yielding an \( \alpha \)-ketoradical (74) which would then abstract a hydrogen atom from ethanol, effecting 72. The provenience of this stemmed from the aforementioned Strachan\textsuperscript{48} studies and is presented in Equation 6 with the \( p\text{-OCH}_3 \) derivative. To obtain 73, the authors rationalized the intermediacy

\[
\begin{align*}
\text{Eq. 6}
\end{align*}
\]
of a spiro-cyclopropanone (75), with subsequent attack by EtOH at the carbonyl and, lastly, a ring-opening, 1,2-aryl shift, affording 73 that effectively amounted to a Favorskii rearrangement (Equation 7). Other intermediates were proposed, but no mechanistic explorations were related nor appeared subsequently in the literature. A potential ethyl ether substituent product, envisaged by the attack of EtOH at the methylene locus with concomitant chloride disjunction, was not reported, though precursor decay, presumably through ketene extrusion to chlorobenzene was formed as a minor product (Scheme 24).
Scheme 24. Posited mechanism to account for the formation of chlorobenzene from α-chloroacetophenone derivatives.

The electronic and positional influences of several substituents were discerned and are disclosed in Table 11. Strong electron donors, i.e., p-OH, o-OMe, p-OMe evinced the highest yields of Favorskii rearrangement product (73) while moderate donating (-Cl, -Me), generated none, except for, curiously o-Me (4%). In addition, electron withdrawing moieties (-CO₂Me) failed to reveal 73.
Table 11. Substituent effects on the photoprod
uct distributions from photolysis of substituted α-chloroacetophenones in ethanol, irradiated at 300 nm for 1-2 h. Table adapted from ref. 50.

<table>
<thead>
<tr>
<th>Substituent (X)</th>
<th>%Yield, acetophenone, 72</th>
<th>%Yield, ethyl phenylacetate, 73</th>
</tr>
</thead>
<tbody>
<tr>
<td>p-OH</td>
<td>26</td>
<td>32</td>
</tr>
<tr>
<td>p-OMe</td>
<td>30</td>
<td>32</td>
</tr>
<tr>
<td>o-OH</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>o-OMe</td>
<td>16</td>
<td>32</td>
</tr>
<tr>
<td>o-Me</td>
<td>58</td>
<td>4</td>
</tr>
<tr>
<td>H</td>
<td>53</td>
<td>0</td>
</tr>
<tr>
<td>p-CO₂Me</td>
<td>48</td>
<td>0</td>
</tr>
<tr>
<td>p-Cl</td>
<td>55</td>
<td>0</td>
</tr>
<tr>
<td>o-Cl</td>
<td>45</td>
<td>0</td>
</tr>
<tr>
<td>m-OMe</td>
<td>15</td>
<td>0</td>
</tr>
</tbody>
</table>

Since this seminal report, a paucity of research has been reported regarding the photo-Favorskii reaction. Sheehan\textsuperscript{51} investigated the use of the p-methoxyphenacyl chromophore as a phototrigger for carboxylic acids. This entity had previously been shown by Anderson and Reese to be a precursor for rearranged products and substrate release after irradiation (Equation 8). However, after irradiation at 254 nm in 1,4-dioxane or ethanol for 5-17 h, they reported that 76 was converted to free acid (77) and p-methoxyacetophenone (78).

\[ \text{R = PhCO}_2, \text{Boc-l-Ala, Boc-Gly, Z-l-Trp, Z-Gly,Gly} \]

Eq. 8
The theorized mechanism to explain the formation of photoproducts (Scheme 25) is in accord with those previously addressed,\textsuperscript{48,50} where upon irradiation, C-O bond homolysis occurs producing geminate radicals \textbf{79} that are posited to readily abstract H atoms from the solvent, yielding primary photoproducts \textbf{80} and \textbf{81} and products from solvent oxidation (e.g. acetaldehyde). Unlike Anderson and Reese’s earlier results, no Favorskii rearrangement esters were reported. The disparate results suggested a nucleofuge effect, but this premise was not further substantiated. Sonawane\textsuperscript{52} et. al., recognized the versatility of this method in the practical synthesis of useful phenylacetic acids. An early report concentrated on the nature and position of the substituent on the chromophore in determining photolysis products of $\alpha$-chloropropiophenone derivatives (Equation 9).\textsuperscript{53}
The principal photoproducts generated from 300 nm irradiation of a 3% solution of 82 in degassed methanol for 6 h were propiophenone (83) the o-chloropropiophenone, (84) the Favorskii ester (85) and the methylether (86). It was revealed that the substituent locus and electronic characteristics profoundly affected the product ratios (Table 12). Notably, both p and m-methyl derivatives of 82c (methyl as a mild electron donor) produced 85 as the primary product while the p-OCH₃ (stronger electron donor) 82a and m-OCH₃ (electron withdrawing) each afforded an exiguous amount of 85, the latter consistent with the previous findings of Anderson and Reese.

Table 12. Photoproduct % compositions from photolysis of 3% solutions of 82a-d in degassed CH₃OH. Table adapted from ref. 53.

<table>
<thead>
<tr>
<th>Compound</th>
<th>R</th>
<th>Product Compositions, %</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>83</td>
</tr>
<tr>
<td>82a</td>
<td>(m-OCH₃)</td>
<td>13</td>
</tr>
<tr>
<td>82b</td>
<td>(p-OCH₃)</td>
<td>12</td>
</tr>
<tr>
<td>82c</td>
<td>(m-OPh)</td>
<td>70</td>
</tr>
<tr>
<td>82d</td>
<td>(m-CH₃)</td>
<td>33</td>
</tr>
<tr>
<td></td>
<td>(p-CH₃)</td>
<td>8</td>
</tr>
<tr>
<td></td>
<td>(m-NO₂)</td>
<td>100</td>
</tr>
</tbody>
</table>
Since the late 1960’s, Givens and coworkers\textsuperscript{54} have gained considerable insight by studying photo-induced extrusion and rearrangement reactions, though those amenable to the photo-Favorskii rearrangement are the most recent among their studies. Early studies\textsuperscript{55} were aimed at comprehending stereoelectronic effects, i.e., the role of the nucleofuge (e.g., halogen, carboxylate, phosphate, and others) in determining the outcomes of photolyses of $\alpha$-substituted chromophores, such as isomers of 2-chloro bicyclo[2.2.2.]octanone (Scheme 26). These were not tailored specifically to investigate the photo-Favorskii rearrangement, that is, products derived from 1,2 migrations or cyclopropanone/oxyallyl intermediates. Nevertheless, a

**Scheme 26.** Product compositions from photolysis of corresponding $\alpha$-chloroketones.
detailed understanding of the rearrangement process that afforded esters, acids, or amides provided a stimulus for future endeavors with the ‘less-than-familiar’ Favorskii rearrangement. Photolysis of syn and anti 2-chlorobenzobicyclo[2.2.2]octadien-3-one isomers of 87 in methanol was shown to generate esters 89 and 90, though in dissimilar ratios. The compound syn-2-Chlorobicyclo[2.2.2]octenone, 91, produced the ester 92 as the major photoproduct, while from corresponding anti-91 isomer emerged a complex assortment of products. Rearrangements had occurred to produce esters, though a Favorskii-like, 1,2-aryl migration was not discerned in these systems. Additionally, a salient impact of the chloride orientation was demonstrably a determinant in the product compositions. It was assumed that the initial step in these reactions involved carbon-chloride bond scission, though whether through homolysis that would engender a geminate radical pair, or heterolysis to form chloride ion and an α-carbenium ion was speculative. A number of plausible radical and ionic intermediates were surmised and, ultimately, through sets of supplementary experiments, a conflation of radical/ion attributes for the intermediates was proposed to account for product distributions. The singlet was determined to be the reactive excited state in these reactions. Experiments involving triplet quenching with piperylene failed to divulge any vicissitudes in product yields and those involving triplet sensitization with acetophenone manifested marked declines in quantum yields, 0.03 for 87 (anti Cl) and 0.09 87 (syn Cl), than those from direct irradiation, 0.35 and 0.32, respectively.
The use of benzoin (desyl) as an \( \alpha \)-keto phototrigger, originally established by Sheehan and Wilson\(^5^8\) for caging phosphates,\(^5^9,^6^0\) was explored principally for its longer wavelength absorption (\( \lambda_{\text{max}} > 350 \text{ nm} \)), efficacious quantum yields (\( \Phi > 0.28 \)) for substrate release or “decaging”, and trenchant photochemistry; the primary photoproducts were the released substrate and benzofuran, derived from an internal benzoin photocyclization.

Despite its obvious utility as a phototrigger, the mechanism for substrate release and cyclization doesn’t abide by a “Favorskii-like” process, i.e., no cyclopropane intermediates nor 1,2-migrations were borne and would not warrant further discussion except for one salient feature. The triplet state preponderated and as such a plausible \( \alpha \)-keto carbocation intermediate\(^6^1\) appeared to be viable to product formation; a similar \( \alpha \)-keto carbocation intermediate is thought to be a possible precursor to the cyclopropanone intermediate in the Favorskii rearrangement. Independent studies\(^5^5,^6^2\) reported a Hammett correlation of \( \rho = -0.9 \) vs. \( \sigma \), imputing positive charge buildup (carbocation character) in the transition state. Indirect evidence for an \( \alpha \)-aryl carbocation intermediate was obtained by Givens and Matuszewski\(^5^9\) through \(^{18}\)O labeling experiments with corresponding arylmethyl phosphate esters. These studies disclosed that expeditive oxygen exchange between benzylic and phosphoryl units had occurred by, presumably, an ion-pair species. The same report\(^5^9\) revealed an observed photostereoisomerization of \((S)-(\cdot)-93\) in \( n \)-butanol, shown in Equation 10, that competed with the substitution reaction to
produce (S)-(−)-94, which itself was formed with a slight, net retention; the phenomenon strongly suggested the intermediacy of ion-pairs.

\[
\text{ produce (S)-(−)-94, which itself was formed with a slight, net retention; the phenomenon strongly suggested the intermediacy of ion-pairs.}
\]

\[
\begin{align*}
\text{Initial: } [\alpha]^2_{\text{D}} & \text{-39.7} \\
\text{After: } [\alpha]^2_{\text{D}} & \text{-28.5} \\
\text{Net racemization: } 28\% \\
\text{77\% conversion} \\
\text{[\alpha]^2_{\text{D}} & \text{-4.5} \\
\text{Net retention: } 5\%
\end{align*}
\]

\text{Eq. 10}

Givens’ seminal photo-Favorskii report\textsuperscript{63} concerned the \(p\)-hydroxyphenacyl chromophore as a cage for ATP (pHP ATP, 95) and for inorganic phosphate (pHP phosphate, 97) (Scheme 27). Photolysis of 95 and 97 at 300 nm in Tris buffer efficiently produced ATP and inorganic phosphate, \(\Phi_{\text{app1}} = 0.37\) and \(\Phi_{\text{app3}} = 0.12\), respectively, as well as the rearranged (Favorskii) \(p\)-hydroxyphenylacetic acid (97, \(\Phi_{\text{app2}} = 0.31\)). The process was shown to originate from the triplet surface by studies on the Stern-Volmer (S-V) quenching using 2-naphthalenesulfonate by the decline in \(\Phi_{\text{dis}}\) as the concentration of 2-naphthalenesulfonate was increased. The slope of the resulting linear dependence (\(K_{\text{SV}}\)) provided an indirect measure of the triplet lifetime and a rate constant for fragmentation, \(k_{\text{obs}} = 6.0 \times 10^8 \text{s}^{-1}\).
Scheme 27. Results from photolysis of pHP ATP and pHP diethyl phosphate in aqueous media. The quantum yields for these processes are included.

Unlike unsubstituted or \( m \)-substituted phenacyl esters studied by Anderson and Reese\(^\text{50} \) and Givens\(^\text{64} \) which primarily generated reduced acetophenones or radical-based products upon photolysis in aqueous media, \( p \)-hydroxyphenylacetate esters and acids (96) were the predominant products of photolysis of \( p \)-hydroxyphenacyl cages. The putative mechanism (Scheme 28) was derived from several consonant studies\(^\text{65,66} \) involving photo-induced solvolyses of benzhydryl chlorides, and involved a sequence beginning with prompt intersystem crossing to the triplet (98) followed by homolytic or heterolytic bond scission to generate geminate radical (99a) or ion pairs (99b). For the reduced pair, a neighboring group assisted,
electron transfer was a possible supervening step before or after the Favorinskii dienedione (100) and release of inorganic phosphate. These discrepancies were the subject of extensive investigations by Givens, Wirz, Wan and Corrie, Phillips and others and will be a major theme of the study reported here.

Scheme 28. Proposed mechanism for phosphate release from the pHp chromophore.

Lastly, solvent mediated cyclopropanone ring opening was proposed to occur to generate the Favorinskii rearrangement product, p-hydroxyphenylacetic acid (96). The relatively high efficacy for photo-stimulated substrate release from the pHp chromophore with accompanying Favorinskii rearrangement provided the thrust for Givens, et. al., not only to investigate the utility of the reaction as an effective
phototrigger for biologically relevant substrates, but also to experiment with the
design of better analogs and ultimately, to understand the mechanistic features of this
reaction. A follow up report\textsuperscript{67} concerned the photochemistry of pHp-caged
neurotransmitters, \(\gamma\)-aminobutyric acid (GABA) and \(l\)-glutamate (Glu), as depicted in
Equation 11. Photolysis of pHp GABA (101, \(\Phi = 0.35\)) or pHp Glu (102, \(\Phi = 0.12\))
in aqueous buffer at \(\lambda = 300\) nm cleanly educed the release of GABA and Glu with
the rearranged 96. Stability studies revealed that 101 and 102 resisted hydrolysis for

more than 24 h when placed in aqueous media, an important feature for application as
a phototrigger. The mechanism was posited to resemble that delineated in Scheme
27.

These findings served to expand the utility of pHp as a phototrigger for
carboxylates as well as phosphates and unequivocally demonstrated an operative
photo-Favorskii process, i.e., the discrete nucleofuge characteristics did not alter the
photorelease pathway. This phenomenon, as will be demonstrated later, is
inconsistent for nucleofuges whose conjugate acids are weaker than phenols. Other biologically significant carboxylic acids such as the oligopeptide bradykinin and the dipeptide (Al-Al) were caged by pH and demonstrated consistent release efficiencies through the photo-Favorskii route. In addition, pH was shown to be an effective and precise delivery agent for Glu and bradykinin in biochemical neuronal function assays. These excitatory neurotransmitters evoke \([Ca^{2+}]\) increases in sensory neurons. This effect was probed by \(Ca^{2+}\) sensitive dyes, FURA-2 and indo-1, that are loaded into the neurons prior to the assay. An enhancement in the fluorescence intensity of these dyes as a function of \([Ca^{2+}]\) indicates neurotransmitter action at the sensory neuron. Caged pH bradykinin and Glu are silent in this regards, i.e., they effectuate no significant changes in \([Ca^{2+}]\) as demonstrated by no significant variations in the fluorescence intensity of the dye. However, upon irradiation, abrupt fluorescence from the dyes was observed, indicating the neuronal release of augmented \([Ca^{2+}]\) in response to the newly liberated bradykinin or \(l\)-glutamate.

Falvey was among the first to introduce an alternative photo-Favorskii mechanism to account for photosolvolysis products he observed when conducting LFP studies with phenacyl phenylacetate 103 in CH\(_3\)CN (Equation 12).

\[
\text{Eq. 12}
\]
It was anticipated that if the presumed mechanism initiated with bond homolysis (Scheme 29) the phenylacetoxy radical (105) would rapidly decarboxylate to form the benzyl radical (106), then dimerize to 1,2-diphenylethane (107). However, 107 was detected only in minute amounts (<0.5%) by HPLC analysis after photolysis.

**Scheme 29.** Radical pair mechanism for the formation of dibenzyl (107) from phenacyl caged phenylacetic acid.

![Scheme 29](image)

Photolysis conditions for 103 were modulated with isopropanol, a H atom donor, or the H atom donor tributyltin hydride, each of which divulged markedly improved quantum yields for disappearance. A complex, hydrogen atom transfer 2-hydroxypropyl (or stannyl) radical addition mechanism was posited (Scheme 30).
Scheme 30. Hydrogen atom transfer mechanism for substrate release.

A transient absorption spectrum was obtained that indicated the presence of a triplet species \((^3103)\), which was further characterized by \(\text{O}_2\) quenching and the likeness of the LFP absorption spectrum \((\lambda_{\text{max}} = 340 \text{ nm} \text{ with a lifetime } \tau = 5.5 \text{ ns})\) to acetophenone’s known triplet \((\lambda = 330-350 \text{ nm, depending on solvent composition})\).²¹

It was hypothesized that the triplet state would abstract a hydrogen atom from isopropanol or tributyltin hydride to generate a ketyl radical transient \((108)\). Evidence for this came again from a transient absorption spectrum that exhibited a longer lived species with a \(\lambda_{\text{max}} = 310 \text{ nm}, \) corresponding nicely to known ketyl radicals obtained from other acetophenone derivatives.²¹ The triplet \(108\) was then
observed to decay to a transient with a lifetime, \( \tau \), of > 1 ms and a \( \lambda_{\text{max}} \sim 330 \) nm; this entity was tentatively assigned 109a or 109b, dependent on the H-atom source and termed a metastable, “light absorbing transient” (LAT).\(^{72}\) LAT promptly disintegrated to 104 and 105. Favley’s experiments explicitly demonstrated the importance of hydrogen atom abstraction to initiate events leading to the formation of fragmentation products from \( \alpha \)-substituted acetophenone derivatives.

Wan and Corrie\(^{73}\) conducted a series of photolyses with the analogous \( p \)-methoxyphenylacyl chromophore, specifically gauging the photorelease of a series carboxylic acids. A noteworthy discovery, however, was made in that no photolysis was seen to occur with \( p \)-methoxyphenacyl acetate in 1:1 \( \text{D}_2\text{O}/\text{CD}_3\text{CN} \) after protracted irradiation times. Earlier studies carried out by Givens\(^{62}\) had shown that photosolvolysis of \( p \)-methoxyphenacyl phosphate esters readily occurred after short irradiation times in mixed organic media. The nucleofuges (acetate and phosphate) were the only significant differences in these studies. It follows that the nucleofuge character seemed to have a significant impact on the photochemistry of \( p \)-methoxyphenacyl derivatives, akin to Bordwell’s earlier findings with ground state reactions. Dissimilar to “photo-inertness” of the \( p \)-methoxyphenacyl acetate, the \( p \)-hydroxy analog revealed a photosolvolysis and rearrangement to the \( p \)-hydroxyphenylacetic acid with a high efficiency in 1:1 \( \text{H}_2\text{O}/\text{CH}_3\text{CN} \), \( \Phi = 0.41 \). The discrepancy in photolysis behavior asserts the pertinence of the phenolic O-H in photosolvolysis of these precursors. It was reasoned, then, that initial homolytic scission was obviated by the assumption that the excited state electronic differences
between the -OH and -OCH₃ moieties were negligible and hence would not be expected to evoke such drastic changes in the product distribution.

Wan and Corrie went on to suggest a singlet excited-state proton transfer (ESPT) mechanism to describe the photo-Favorskii rearrangement of pH₆ acetate (Scheme 31).

**Scheme 31.** Singlet excited state proton transfer mechanism for acetate release.

Cogent evidence for this mechanism was empirically derived and included: (1) Insignificant declines in the quantum yields for decay of 110 in 1:1 H₂O/CH₃CN in the presence of known triplet quenchers piperylene and sodium sorbate, signaling the
singlet state as the reactive excited state. These findings were in direct contradiction to the previously mentioned report by Givens,\textsuperscript{63} where pHp ATP (95) imparted, unequivocally, a significant decline in $\Phi_{\text{dis}}$ in the presence of the triplet quencher, 2-napthylsulfonate. (2) A singlet excited state correlation with solution pH ($pK(S_1) \sim 1$) and quantum yield for acetate formation, $\Phi_p$, in 1:1 H$_2$O/CH$_3$CN was observed for pHp acetate. This phenomenon was ascribed to ESPT based on consistencies with other $pK(S_1)$ studies.\textsuperscript{74} (3) From laser flash photolysis studies, a strong absorption band appeared at $\lambda_{\text{max}} = 330$ nm that was not quenched by O$_2$. Absorption decay profiles indicated two transients, one shorter lived ($112$, $k_d = 6.3 \times 10^6$ s$^{-1}$)\textsuperscript{75} and one longer ($111$, $k_d = 1.8 \times 10^5$ s$^{-1}$)\textsuperscript{76,77} with similar $\lambda_{\text{max}}$ values (\textasciitilde 330 nm). When excited, 110 underwent rapid, adiabatic ESPT to generate 111*. This then abided two decay routes; one through solvent mediated deprotonation, restoring 110, while the other decayed by intramolecular proton transfer and acetate dissociation to generate 100, and subsequent solvolysis to 96. One additional, concerted mechanism was proposed, which entailed phenol deprotonation coupled with acetate departure through transition state 113 directly to 96 (Scheme 32).
Results from a comprehensive study into the identification of triplet excited state transients of pHP diethylphosphate were communicated by Givens and Wirz,78 employing both direct laser flash photolysis and indirect S-V quenching methods of analysis. Photolysis of *pHP diethyl phosphate (114)* in wet CH$_3$CN (≥ 5% water) disclosed only the production of 96, contrasting the product mixture reported in dry CH$_3$CN. The photoreaction was accelerated 5-fold in the presence of H$_2$O, while air saturation or 0.5 mM piperylene, a triplet quencher, educed a 4-fold deceleration. The quantum yield of diethyl phosphate release from 114, $\Phi_{\text{dis}}$, in wet CH$_3$CN, was conveyed as 0.94 from laser flash photolysis studies, which dwindled to 0.37 with 10 mM piperylene. To understand and interpret the laser flash photolysis results from 114, *p*-hydroxyacetophenone (115) was employed as a model system, lacking a leaving group. Triplet *p*-hydroxyacetophenone (3115) underwent singlet-triplet crossing with a rate constant, $k_{\text{ISC}} \sim 2.7 \times 10^{11}s^{-1}$, identified by signature $\lambda_{\text{max}} = 370$ nm triplet-triplet absorption band in dry CH$_3$CN that was red-shifted to $\lambda_{\text{max}} = 395$ nm.
in wet CH$_3$CN. The acidity of the triplet excited $p$-hydroxyacetophenone was registered by rapid protonation of bromophenol blue, suggesting a pKa $\sim$ -1.0 for the phenolic OH. The triplet energy of $^3$115 as determined to be 70.5 kcal/mol$^{79}$ by phosphorescence and accordingly was observed to transfer energy to added naphthalene, which has a triplet energy $E_T = 62$ kcal/mol.$^{78}$ The equilibrium constant for enolization of 115 with the hydroxyquinomethylene (115$'$), p$K_{E\alpha}$, was calculated to be 16.4 by DFT, and subsequent dissociation constant for 115$'$, $K_a = -$8.5, forming 115$'$. Ionization of $^3$115$'$ was shown to be suppressed in dry CH$_3$CN, but occurred adiabatically in 50% aq. CH$_3$CN to generate $^3$115, with $k_{ion} = 9.6 \times 10^6$ s$^{-1}$ (pKa 4.6); $^3$115$'$ showed absorption bands similar to $^3$115, $\lambda_{max} = 405$ nm and a weaker band at $\lambda = 500$ nm.

Excitation of ground state 115$^-$ at $\lambda = 308$ nm in 0.1 M NaOH manifested bands at $\lambda = 405$ nm and $\lambda = 500$ nm, signifying that these were due to $^3$115$^-$. Corresponding studies in acidic solutions demonstrated that $^3$115$^-$ underwent rapid protonation with a $k_{H^+} \sim 4 \times 10^{10}$ M$^{-1}$s$^{-1}$ from the decay of the band at $\lambda_{max} = 350$ nm. The enol protonated form was denoted as $^3$115$'$ based on the lack of naphthalene quenching and supporting DFT calculations that suggested a triplet energy of approximately 38.4 kcal/mol. These aforementioned results were assimilated into a thermodynamic cycle for the triplet manifold that parallels the ground state (Scheme 33).
Scheme 33. Proposed thermodynamic cycle of p-hydroxyacetophenone (115). Adapted from ref. 78.

The thermodynamic cycle of the triplet state proton tautomerization, determined from the laser flash photolysis studies of 115 and DFT modeling, was subsequently applied to pH diethyl phosphate 114. An intense absorption band at $\lambda = 395$ nm appeared in dry CH$_3$CN, which was ascribed to its corresponding triplet $^3$114 based on observed energy transfer to added naphthalene with a rate $k_{ET} \sim 7.8 \times 10^9$ M$^{-1}$s$^{-1}$ and phosphorescence spectra at 77 K in a solid matrix of ethyl ether, isopropanol, and isopentane (5:2:2). The self-quenching of $^3$114 accelerated with increasing concentrations of 114, by a second order decay, $k_{sq} = 8.5 \times 10^8$ M$^{-1}$s$^{-1}$. The decay rate of $^3$114 was accelerated by the addition of air or oxygen; $k_q = 3.0 \times 10^9$ M$^{-1}$s$^{-1}$, a 2.8 fold increase in oxygen or 0-1.25 x 10$^{-2}$ M piperylene that revealed $k_q \sim 4.5 \times 10^9$ M$^{-1}$s$^{-1}$, a 1.9 fold increase. The rate constant for phosphate release from 114
increased exponentially with water through 5% H2O, \( k_{\text{dis}} \sim 2.8 \times 10^7 \text{ s}^{-1} \). Above 5%, the rate was too fast to be measured by the LFP system utilized. A good correlation between the influence of piperylene on the steady state lifetime and the time resolved laser flash photolysis parameters adduced a reactive triplet state for release of diethyl phosphate from 114. The ratio of lifetimes of \( ^3\text{114} \) in 10 mM piperylene quencher (Q) in 5% aq. CH3CN were \( \tau_q/\tau_o = k_{\text{obs}}(k_{\text{obs}} + k Q) = 0.38 \). The quinomethylene (\(^3\text{114}'\)) was monitored by acid titration with HClO4 (0.1 - 10 mM) at half-protonation, 4.0 mM HClO4. In accord with Wan’s findings73, \( \Phi_{\text{dis}} \) of \( ^3\text{114} \) declined with increasing [H\(^+\)], due to protonation of \( ^3\text{114} \) to form \( ^3\text{114}' \). Collectively, these findings adduced the triplet as the reactive state of \( p\)-hydroxyacetophenone (\(^3\text{115} \)) for the photo-Favorskii rearrangement.

The lack of detail in the mechanistic interpretations, the easily synthesized pHp ester and the available TR\(^3\) spectrometer for detailed vibration spectra of triplet and other sufficiently long-lived intermediates, attracted Phillips and coworkers to enter the research arena of the pHp photoremovable protecting group. A series of empirical and theoretical investigations further probed the decay mechanism of pHp. Their first report80 examined the excited state structures of pHp diethyl phosphate (114) and \( p\)-hydroxyacetophenone (115) in N\(_2\) purged CH3CN, employing picosecond (ps) and nanosecond (ns) Time Resolved Resonance Raman spectroscopy (TR\(^3\)). No bands were observed in the 0-2 ps vibrational spectrum of 115 and 114. After 10 ps, however, a band at 1600 cm\(^{-1}\) was discerned for each, and its decay monitored on the nanosecond timescale. The inception of this band was established as 5.3 and 6.7 ps,
respectively, with a growth rate of 86 and $150 \times 10^9 \text{s}^{-1}$, respectively, and completely disappeared after $\sim 200$ ns (Figure 4). The decay rates for transients of 114 and 115 were hastened in air, implying excited triplets ($^3114$, $^3115$). This supposition was corroborated by Kerr-gated, time-resolved fluorescence (KGTRF) measurements of pHF-diethylphosphate (114) and $p$-hydroxyacetophenone (115) which decayed with rate constants $k_{\text{ISC}} \sim 5 \times 10^{11} \text{s}^{-1}$ that correlated with $\sim 3.8 \times 10^{11} \text{s}^{-1}$ for Givens.78 Perhaps more importantly, however, was that this value was congruent with the ps growth of the 1600 cm$^{-1}$ band seen by TR$^3$ above and attributed to the triplet state of 114 and 115. The pump $\lambda = 267$ nm was consistent with the high energy absorption band of 114 and 115, stemming from a $S_0$-$S_3$, $\pi\pi^*$ transition. Excitation wavelengths near 300 nm were characterized by $S_0$-$S_1$, $\pi\pi^*$ transitions. KGTRF data suggested

**Figure 4.** Left: Time dependent decay profile of $^3114$ band $\sim 1600 \text{ cm}^{-1}$ with ns $^3$TR. Right: Time dependent decay profile of $^3114$ band $\sim 1600 \text{ cm}^{-1}$ with ns $^3$TR. Reproduced with permission from ACS, ref. 80.
that intersystem crossing for both 114 and 115 adhered to the nπ* S1 to ππ* T1 pathway in both 1:1 CH3CN:H2O and neat CH3CN. The similar decay attributes of triplet pH3 diethyl phosphate (3114) and triplet pH3 p-hydroxyacetophenone (3115) gleaned from ns-TR3 and KGTRF analyses, and pH3 acetate (3110)81 in neat CH3CN, inferred that photophysical processes of 115 were thus independent of the nucleofuge. This disposition, however, was reversed in a later TR3 study82 with 114 and 115 in aqueous CH3CN that disclosed a strong correlation between the triplet lifetime and nature of the nucleofuge. Energies of 3115 were computed by DFT to be 69 kcal/mol (cp. experimental ~70.5 and Wirz78 ~69.6) and for 3114, 67 kcal/mol (cp. experimental ~ 70.674). TR3 spectra provided conformational analyses of 3115 and 3114; DFT applications were consistent with the TR3 results. TR3 spectra suggested that the carbonyl of 3115 was distorted ~16° out of plane (OOP) of the ring, in contrast to ground state, where the chromophore is planar. Similarly, a ~21° OOP orientation of the carbonyl relative to the aryl ring was found for 3114, compared to ~2° OOP for the ground state. From the planarity of the carbonyl and attached groups, the sp2 nature of the carbonyl for the triplet species was thought to be preserved.

A second report that year81 detailed a similar set of TR3, KGTRF, and DFT studies for pH3 acetate (110), akin to Corrie and Wan’s model.73 Coincident ps and ns TR3 spectra were obtained to those of 114 and 115 (Figure 5). No activity was observed after initial excitation (0-2 ps), after which a time-dependent band at 1601 cm−1 emerged at ~50 ps. The decay of this, analogous to 114 and 115, revealed a
good fit to a single exponential growth and a decay function by least squares analysis; time constant for growth was 12 ps and 137 ns for decay in N₂ purged CH₃CN, (Figure 5).

Figure 5. Time dependent decay profile of $^3110$ at ~1600 cm$^{-1}$ in ns-$^3$TR spectra. Circles indicate degassed, triangles open air, and squares oxygen. Reproduced with permission from the ACS, ref. 81.

In the presence of air, the decay constant decreased to 50 ns, and in oxygen to 16 ns, consistent with a likely triplet excited state. Once again, the decay summary of $^3110$ was consonant with triplet diethyl phosphate ($^3114$) and triplet $p$-hydroxyacetophenone ($^3115$) and was consistent with previous work with $^3110$. Additionally, the nanosecond rate of deactivation, as indicated by the disappearance of bands associated with $^3110$, was seen to accelerate as a function of pH concentration. This observation was consistent with earlier studies$^{78}$ and was ascribed to self-quenching from head-tail hydrogen abstraction by the carbonyl from a neighboring phenolic OH. To support the assignment of triplet state reactivity, KTRF
studies on 110 gave $k_{ISC} \sim 5.0 \times 10^{11}$ s$^{-1}$ that were congruous with the rapid growth of the $\sim 1600$ cm$^{-1}$ band seen in $^3$TR spectra and assigned to the carbonyl triplet as well as the results of previous studies.$^{74,75}$ This result was divergent from the 1.1 µs triplet lifetime of 110 that Corrie and Wan$^{73}$ reported. It was believed that Corrie and Wan’s value was obtained in thoroughly O$_2$ purged conditions and that residual O$_2$ was believed to be the cause of a shortened, 137 ns lifetime of $^3$110 in Phillips’ study.

A fluorescence decay of 2 ps was detected that conflicted with the 12 ps growth of the $^3$110 described earlier. Phillips, therefore, concluded that some other process early in the ps time regime must be active in developing the triplet TR$^3$ spectrum. This was designated as triplet relaxation of surplus energy brought forth by efficient intersystem crossing to an unrelaxed, upper triplet. Evidence for this assertion originated from frequency changes in the band centered at 1600 cm$^{-1}$ up to 200 ps; a band shift from 1570 cm$^{-1}$ at 4 ps to 1601 cm$^{-1}$ at 50 ps was observed. In addition, band widths were reported to narrow during this time, an indication of excess energy relaxation. The implications for this were as follows: The lowest triplet of 110 was considered to be much lower in energy than the corresponding singlet. The energy gap between $^1\pi\pi^*$ and $^3\pi\pi^*$ had been determined to be $\sim 1800$ cm$^{-1}$ for acetophenone;$^{83}$ this corresponds to $^3\pi\pi^*$ and $^3\pi\pi^*$ energy gap of analogous p-hydroxyacetophenone,$^{79}$ $\sim 2500$ cm$^{-1}$. Phillips reasoned that these values and the similarity of energies of $^1\pi\pi^*$ and $^1\pi\pi^*$ for 110, as determined by ground state UV absorption, would provide some insights into the $\pi\pi^*$ triplet energy. Several empirical studies reported a near degeneracy in lowest $n\pi^*$ and $\pi\pi^*$ triplet states of
acetophenone.\textsuperscript{84,79,83} Hence, the triplet state is thought to be easily perturbed by environmental changes, such as substituents, phase (gas, liquid, solid), solvent polarity, H-bonding, etc. For example, the lowest triplet energy of acetophenone is considered to be $n\pi^*$ in the gas phase\textsuperscript{85} and in non-polar solvents,\textsuperscript{79} and $\pi\pi^*$ in polar solvents and in the solid phase.\textsuperscript{79} Most $p$-substituted acetophenones including pHP diethyl phosphate (114), are believed to have a lowest $\pi\pi^*$ triplet both in polar and non-polar media.\textsuperscript{79} TR\textsuperscript{3} measurements for 110 revealed close similarities to 114, as did the corresponding phosphorescence profiles. These attributes, as well as the likeness of their triplet lifetimes, led Phillips to conclude that $3^1$110 is $\pi\pi^*$ in character. A triplet energy of 67 kcal/mol was calculated for pHP acetate (110) by DFT that was in agreement with previous studies.\textsuperscript{78} Conformational analysis imparted by DFT revealed a $\sim$20$^\circ$ OOP contortion of the carbonyl relative to the aromatic ring for $3^1$110 compared to a planar ground state, also consistent with other pHP analogs.

To discern the initial steps of substrate photorelease from pHP and concomitantly reconcile two disparate assertions regarding reactive state multiplicities, Phillips\textsuperscript{86} employed his psTR\textsuperscript{3} and nsTR\textsuperscript{3} methods for the analysis of $p$-hydroxyacetophenone (115) in aqueous media. The focus of this study was on the solvent-assisted deprotonation of 115, believed to be crucial in the rearrangement process. This investigation derived from results of an analogous study\textsuperscript{87} of the $p$-methoxy derivative of 115 in water, which substantiated a water-stimulated modification in properties of the triplet state vs. dry CH\textsubscript{3}CN. Dynamics of 115 from ps-TR\textsuperscript{3} analysis in water are depicted in Figure 6. Contrary to earlier results in dry
Figure 6. Left: 1) Bottom - psTR\textsuperscript{3} spectrum of 115 in water. 2) Top - psTR\textsuperscript{3} spectrum of 115 in water. Right: 1) Bottom - psTR\textsuperscript{3} spectrum of 115 in buffer, pH = 9.00. 2) Top - psTR\textsuperscript{3} spectrum of 115 in buffer, pH = 9.00. Reproduced with permission from the ACS, ref. 86.

CH\textsubscript{3}CN\textsuperscript{80} that evinced one transient species, two short-lived transients were conspicuous in the spectra: (1) one conveyed a band at 1600 cm\textsuperscript{-1} that correlated with previous studies\textsuperscript{80} in dry CH\textsubscript{3}CN, emerging at 2 ps and decaying within ~10 ns, (2) another showed bands at 1599, 1481, 1357, and 1167 cm\textsuperscript{-1} that decayed within 95 ns. When measured in N\textsubscript{2} purged water, the decay of bands associated with the second transient lengthened to 140 ns, a clear indication of O\textsubscript{2} quenching and evidence for a
triplet reactive state. The new transient was ascribed to H-bonding between the solvent and $^3{115}$, not evidenced in aprotic CH$_3$CN, and now distinguished as $^3{115}'$.

Results of ps-TR$^3$ analysis of 115 in buffer, pH = 9.00 (Figure 6) revealed that the formation of the new entity was faster than in neutral water, and was the sole entity that emerged in the ns spectrum. It was deduced that this species was the triplet $p$-hydroxyacetophenone anion ($^3{115}$). The direct excitation of a ground state anion of 115 under these conditions was precluded on the basis of its very low $\varepsilon$ at the pump $\lambda$ = 267 nm, and probe wavelengths of 400 and 415 nm. Control experiments in basic solutions with 115$^-$ as the conjugate base tautomer and using a pump wavelength of $\lambda$ = 341 nm, produced parallel spectra that had similar triplet decay dynamics, thus supporting the likelihood of $^3{115}^-$ as the intermediate. Full Raman spectral assignments of bands associated with $^3{115}^-$ and the corresponding protonated tautomer ($^3{115}$) manifested salient differences that were reproduced by DFT model calculations.

The kinetic profile of $^3{115}$ was shown to be a good fit for a two-exponential function comprised of a growth, 90 ns, and decay constant, 200 ns. Control TR$^3$ experiments with 115 in acidic media (pH = 1) demonstrated the presence of only this entity. A similar transient was observed from Wirz and Givens$^{78}$ at low pH, and was tentatively designated as a quinoid enol triplet (115') (Scheme 32). From this comprehensive study, it was inferred that an excited state deprotonation step was plausible for pH$_P$, and $^3{115}'$ a likely precursor to nucleofuge release. An optimized
conformation from DFT fit of TR$^3$ spectra for triplet $p$-hydroxyacetophenone $^{3115}$ in water is depicted in Figure 7.

**Figure 7.** DFT determined, optimized geometry and bond distances (Å) of $^{3115}$ in water. Reproduced with permission by the ACS, ref 86.

![Diagram showing optimized geometry and bond distances](image)

The results depicted extensive hydrogen bonding between water and $^{3115}$. Two water molecules associated with the phenolic O-H, affording 14.8 kcal/mol of stabilization energy; one water molecule interacted with the carbonyl moiety, affording 6.8 kcal/mol of stabilization energy. H-bonding with the phenolic O-H was thought to preface water-mediated deprotonation of triplet $p$-hydroxyacetophenone ($^{3115}$) to the corresponding the triplet anion, $^{3115}^-$. This event seemed plausible by the increased acidity of the phenolic proton of 115 in the triplet state, pKa = 3.6, compared to the ground state, pKa = 7.6. TR$^3$ analysis disclosed an unspecified transient (termed X) that emerged contemporaneously with $^{3115}^-+$; this entity exhibited
a decay constant of 88 ns, similar to 95 ns for $^{3}115^-$. UV measurements of the 115 after TR$^3$ analysis manifested no significant degradation, leading Phillips to conclude that X rapidly deactivates back to ground state 115. No further efforts were conducted to identify X. TR$^3$ and DFT results were assimilated to propose a decay and deprotonation mechanism for 115 (Scheme 34). This mechanism deviated from

**Scheme 34.** Phillips proposed mechanism for excited state processes of $p$-hydroxyacetophenone (115).
Wirz and Givens’ (Scheme 32), chiefly in regards to the identification of X. Givens and Wirz assigned this to the triplet enol (\(^{3}115\)) formed from a step involving solvent-induced protonation of \(^{3}115\). The aforementioned TR\(^{3}\) measurements of 115 in water manifested concomitant formation of \(^{3}115\) and X that represented the decay of \(^{3}115\). Experiments conducted in acidic media, where a reprotonation step is obviated, indicated a direct decay route of \(^{3}115\) through X. Having elucidated the sequence of photophysical and reaction events leading up to substrate release from pHp, Phillips turned to mechanism of substrate release, the heart of the photo-Favorskii reaction.\(^{88}\)

As before, ps and ns-TR\(^{3}\) spectroscopy were employed along with femtosecond transient absorption (fsTA) spectroscopy, for the investigations of pHp diethyl phosphate (114) and pHp diphenyl phosphate (116) in dry and 1:1 CH\(_{3}\)CN:H\(_{2}\)O (aq. CH\(_{3}\)CN) solutions. Results from fsTA analysis of 114 in both solvents revealed the formation of a strong absorption band at ~400 nm while one at ~320 nm decayed rapidly; one exponential function fit the simultaneous growth and decay components, with a time constant of 2.5 ps. This value matched the ISC constant derived previously\(^{80}\) and implied a rapid ISC to the triplet state. Pronounced changes in shape of the 400 nm band from 4 to 35 ps were observed in aq. CH\(_{3}\)CN but not in neat CH\(_{3}\)CN; the band was wide and structureless earlier than 4 ps, but better defined later. Phillips ascribed this occurrence to specific solvent-solute interactions, such as H-bonding. Investigations with fsTA of \(p\)-hydroxyacetophenone (115) in aq. CH\(_{3}\)CN exhibited no such shifts in the band shape at 400 nm. Concerning
116, fsTA measurements were congruent with 114, imputing structurally similar triplet reactive states. The triplet lifetimes of these, however, proved to be different, as determined by TR\(^3\) analysis; \(^3\)114 exhibited \(\sim 150\) ns lifetime in CH\(_3\)CN and \(\sim 350\) ps in aq. CH\(_3\)CN. Those for \(^3\)116 were \(\sim 27\) ns in CH\(_3\)CN, \(\sim 150\) ps in aq. CH\(_3\)CN. A synopsis for decay profiles of pHP diethyl phosphate (\(^3\)114) and pHP diphenylphosphate (\(^3\)116) as a function of water concentration are presented in Table 13, with pHP acetate (\(^3\)110) for comparison.\(^82\) Hydrogen ion dissociation constants, pKa, for the nucleofuges were included; these directly correlate with the respective anion stabilities in aqueous media as well as relative leaving group ability in the context of heterolysis in aqueous media. A higher pKa indicates a more stabilized, anionic leaving group. A noticeable trend was observed between the pKa and triplet lifetime; \(^3\)116, with the lowest nucleofuge pKa of 0.4, manifested the shortest triplet lifetime, 150 ps in aq. CH\(_3\)CN, while for \(^3\)110 with a pKa of 4.76, one order of magnitude longer at 2.130 ns.

**Table 13.** Time constants for triplet decays of 116, 114, and 110 in mixed CH\(_3\)CN/H\(_2\)O solvents and corresponding pKa’s of the nucleofuge. Table adapted from ref. 88.

<table>
<thead>
<tr>
<th></th>
<th>H(_2)O % (vol)</th>
<th>75%</th>
<th>50%</th>
<th>40%</th>
<th>30%</th>
<th>25%</th>
<th>15%</th>
<th>10%</th>
<th>pKa</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>triplet decay ((\tau), ps)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>pHP diphenyl phosphate, (^3)116</td>
<td>150</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>830</td>
</tr>
<tr>
<td>pHP diethyl phosphate, (^3)114</td>
<td>290</td>
<td>350</td>
<td>530</td>
<td>800</td>
<td>1280</td>
<td>2400</td>
<td>9000</td>
<td></td>
<td>1.39</td>
</tr>
<tr>
<td>pHP acetate, (^3)110</td>
<td>2130</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>4.76</td>
</tr>
</tbody>
</table>
A correlation between water concentration and triplet lifetime was also apparent; the lifetime of $^3$114 progressively shortened as the water content increased, with initial pronounced changes (10%-30% 11.2-fold, 30-50%, 2.2-fold, and 50-75%, 1.2-fold) and later leveling off as water was added. The same was observed for $^3$116, though not to the same caliber. Stern-Volmer analysis of 114 and 116 evinced a curved, near quadratic dependence on water concentration (Figure 8). Presupposing that water addition influences heterolysis in a non-linear fashion, the S-V results indicated a likely, triplet heterolytic dissociation route for phosphates from excited state 114 and 116. The salient influence of water concentration asserts that water is not only involved in early pHP deprotonation but also in the rearrangement (solvolysis) step. These studies adduce a 2nd order dependence of the triplet lifetime on water.

**Figure 8.** Stern Volmer analysis of $^3$114 lifetimes as a function of water concentration, in percentage. Reproduced with permission by the ACS, ref. 88.
Solvation of the nucleofuge, as well as the pH chromophore, was considered to be an important constituent in the correlation between triplet decay dynamics and leaving group ability, demonstrated in Table 9. The double bonded oxygen of phosphate, like the carbonyl, has strong basic character and hence would be expected to interact strongly with water as an H-bond acceptor. The effect of this interaction is an electrophilic pull towards ionization. Acidic and basic sites of the pH chromophore and nucleofuge are anticipated to be strongly solvated by water. To corroborate this “concerted” solvation premise, fsTA studies were performed with 114 in DMSO, a H-bond acceptor but poor donor, and trifluoroethanol (TFE), an H-bond donor but poor acceptor. In DMSO, the fsTA spectrum, denoting the triplet decay profile, was akin to that obtained in dry CH₃CN and demonstrated a triplet lifetime > 200 ns with an accompanying low quantum yield (<0.01). In TFE, a slightly shorter triplet lifetime was reported, as well as a low quantum yield (no data was provided for this). However, a 1:1 combination of DMSO/TFE revealed a significantly shortened lifetime for triplet pH diethyl phosphate (3114) and a quantum yield of 0.17 for phosphate release. These findings seem to support a general requirement for “concerted” solvation of the pH chromophore and nucleofuge for efficient nucleofuge release from the excited state pH chromophore. A comparison of spectra obtained from psTR³ analysis of 114 and an authentic sample of rearranged p-hydroxyphenylacetic acid photoproduct (95) (pump λ = 267 nm, probe λ = 200 nm) in 1:1 CH₃CN:H₂O was carried out to inspect the formation dynamics of 95. The formation and growth of bands following excitation of 114 were
undeniably associated with the formation of 95. A first order exponential curve fit resulted in a time constant of ~1100 ps. When comparing this to the triplet delay for 114 (~350 ps), a discrepancy was ostensible between the rearrangement reaction and triplet decay. The triplet decay was associated with nucleofuge (phosphate) disjunction; hence, to account for the delay time (~750 ps) between the triplet decay and formation of p-hydroxyphenylacetic acid (95), an intermediate (M) was proposed in Scheme 35 and thought to be produced immediately after nucleofuge disjunction but before the appearance of 95. An evasive, but accepted intermediate in the Favorskii reaction is cyclopropanone (100) which, as of yet, has not been detected. Thus, M, was perceived as a likely fit for 100, but necessitated corroborating data.

**Scheme 35.** Proposed intermediate precursor to rearranged product.

A formula was proposed to describe the formation of 95 from 3114 (Equation 13).

\[
[95] = \left\{ \frac{[3121]}{(k_2-k_1)} \right\} \ast [k_2(1-e^{-k_1t}) - (k_1(1-e^{-k_2t}))]
\]

**Eq. 13**
The rate for phosphate detachment and decay of $^3\text{114}$ to M, $k_1$, was deemed to be much faster than $k_2$. Efforts were concentrated on determining the rate of rearrangement, $k_2$, through psTR measurements in CH$_3$CN with various water at pump $\lambda = 267$ nm, probe $\lambda = 200$ nm. From exponential fits of the spectra, $k_2$ was determined to be $1.69 \times 10^{-9}$ s$^{-1}$ ($k_2 = 1/\tau_2$, $\tau_2 \sim 470$ ps) and was found to be insensitive to the water concentration. To quantify the formation of M, as a model system, equation 14, derived directly from equation 13, was applied. Though not empirically confirmed, a solvent assisted, “push-pull” neighboring group stimulus from the H-bonded solvent was postulated to afford M as a solvated complex containing features

$$[X] = [^3\text{114}]^*\left[\frac{k_1}{(k_2-k_1)}\right]^* (e^{-k_1t} - e^{-k_2t})$$

Eq. 14

**Figure 9.** Phillips’ proposed structure of unknown, M, as a solvation/contact ion pair complex.
of a contact ion pair of \( \alpha \)-protonated cyclopropanone (100) and liberated phosphate anion (Figure 9, with 114). Calculations with DFT at the random phase approximations (RPA) of 100 indicated an electronic transition with a large oscillator strength at \( \lambda \sim 274 \) nm, which was consistent with a spectral band at \( \lambda = 274 \) nm associated with M. Further empirical attempts to bolster this structural hypothesis, such as Kerr gated ps-TR\textsuperscript{3} were ineffectual, probably due to the relatively low concentration of M expected when \( k_2 \gg k_1 \) (steady state condition) as well as the lack of a strong absorption band between 300 to 700 nm when probed by fsTA. Incorporation of the mechanistic features acquired from these studies compelled Phillips\textsuperscript{88} to propose a new, detailed photo-Favorskii rearrangement to account for the formation of 95 (Scheme 36). The salient attributes of this mechanism are: (1) The rate of (a) is largely dependent on the water content and nature of the nucleofuge; (2) The rate of (b) is dependent on the nature of the nucleofuge but mainly independent of water concentration. Phillips conveyed two more comprehensive, though theoretical, investigations concerning the excited state photochemistry of pHP acetate (110)\textsuperscript{90} and the role of water in assisting the nucleofuge disjunction/solvolytic rearrangement steps of the photo-Favorskii rearrangement for \(^3\)114 and \(^3\)116.\textsuperscript{91} Information gleaned from these studies served to corroborate hypotheses derived from earlier, empirical data,\textsuperscript{80} and are beyond the scope of this introduction.
Scheme 36. Posited mechanism for phosphate release from the pHp chromophore.

In the latest installment of the ongoing exploration of the photo-Favorskii rearrangement, Givens and Wirz,\textsuperscript{92} reported the detection of an anionic pHp triplet biradical (117) in the reaction course of excited pHp diethyl phosphate (114), pHp tosylate (118) and pHp mesylate (119) from time resolved IR and UV studies. An amended, generalized mechanism was proposed and represented in Scheme 37.
It was revealed that the lifetime of $^3\text{114}$ reduced from ~0.4 ns in 50% aq. CH$_3$CN$^{88}$ to ~100 ps in 87% aq. CH$_3$CN and ~63 ps in 100% H$_2$O. Transient absorption spectra for 118 and 119 were consistent with 114. Decay kinetics of $^3\text{114}$ accorded with those reported by Phillips.$^{80}$ However, new transient absorption bands were detected at 445 and 420 nm, and were assigned to triplet biradical (117) based on the likeness of shape, location, and weak intensity of analogous phenoxy radicals.$^{93}$ MP2 calculations predicted the carbonyl of 117 to be contorted 27° out of plane from the aromatic ring. TDFT and open shell PPP SCF CI$^{94}$ calculations failed to detect the singlet analog of 117, implying an expeditious cyclization to 100 after ISC. Efforts to detect 100 by step scan FTIR, anticipated to be found at ~1880 cm$^{-1}$,$^{92}$ were ineffectual (Figure 10).
Figure 10. Step-scan FTIR of 114 in CD$_3$CN (2% D$_2$O) at 50 ns (red) and 1μs (blue) delay. Reproduced with permission from the ACS.

However, bands at 1450 and 1300 cm$^{-1}$ were immediately discerned after excitation and promptly decayed (first order) with a coincident emergence of another band at 1647 cm$^{-1}$, $k \sim 8.5 \times 10^6$ s$^{-1}$ (2% D$_2$O in CD$_3$CN). This rate of appearance was accelerated with additional water at the expense of band amplitude. This was tentatively designated as the $p$-quinone methide (120) and was congruent with results from recent work by Kresge$^{95}$ et. al., in the photolysis of $p$-hydroxybenzyl acetate.

Nanosecond LFP evaluation of pH diethyl phosphate (114) in wet CH$_3$CN supported the configuration of this transient; excitation of $^3$114, $\lambda = 395$ nm, produced an absorbing species, $\lambda = 276$ nm, which was relatively long-lived ($k \sim 0.01$ s$^{-1}$ in 5% H$_2$O, 0.05 s$^{-1}$ in 10% H$_2$O) and waned with increasing water content. Kresge$^{95}$ reported a rate constant of 3.3 s$^{-1}$ in 100% water. These observations provided strong
support for the triplet biradical (117). Low temperature $^1$H NMR, (-25°C, 2.5% D$_2$O in CD$_3$CN) of 114 conveyed the abrupt emergence of peaks associated with 120 upon irradiation. The likely route to the formation of 120 is through CO extrusion of spirodiendione (100). To further elucidate the role of water in the photoreaction, SKIE were probed with pH acetate (114) using ns-LFP (degassed, 10% aq. CH$_3$CN, 25°C), with a particular focus on the decay kinetics of $^3$114. In addition, a secondary photoproduct, p-hydroxybenzyl alcohol (121), was asserted from the putative p-quinone methide (120), derived from solvent kinetic isotope studies (SKIE) using laser flash photolysis (ns-LFP) in 87% aq. CH$_3$CN (Table 14).

Table 14. Rate constants, $k_{obs}$, for decay of $^3$114 as a function of the D/H mole fraction, n, using ns-LFP in 10% aq. CH$_3$CN at 25°C. Table adapted from ref. 92.

<table>
<thead>
<tr>
<th>n$^a$</th>
<th>average $k_{obs}$ (10$^6$ s$^{-1}$)</th>
<th>$k_H/k_n$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.00</td>
<td>11.4 ± 0.14</td>
<td>1.00</td>
</tr>
<tr>
<td>0.20</td>
<td>9.53 ± 0.07</td>
<td>1.20</td>
</tr>
<tr>
<td>0.40</td>
<td>8.36 ± 0.04</td>
<td>1.36</td>
</tr>
<tr>
<td>0.60</td>
<td>7.24 ± 0.03</td>
<td>1.57</td>
</tr>
<tr>
<td>0.80</td>
<td>6.29 ± 0.03</td>
<td>1.81</td>
</tr>
<tr>
<td>1.00</td>
<td>5.25 ± 0.06</td>
<td>2.17</td>
</tr>
</tbody>
</table>

$^a$n = mole fraction = n$_D$/(n$_D$ + n$_H$).

By employing the proton inventory method$^{96}$ of analysis the total SKIE was determined to be 2.17. Decay constants for $^3$114, $k_{obs}$, were then plotted against
**Figure 11.** Plot of $k_{\text{obs}}$ of $^{3114}$ vs. mole fraction of deuterium in degassed, 10% aq. CH$_3$CN at 25°C. The black line signifies the expected correlation for a single site isotope effect ($v = 1$), see equation 4. The lower line indicates a multisite, general solvation ($v = \infty$). The central two lines represent a two site effect (red, $v = 2$), and a three site effect (blue, $v = 3$). Reprinted with permission from the ACS.

![Figure 11](image)

deuterium mole fractions, $n_D$, depicted in Figure 11. An appropriate comparison (Equation 15) was exploited to model the solvent kinetic isotope results where $n$ is the mole fraction and $v$ represents the number of isotope effect sites on $^{3114}$.

$$k_n/k_H = (1 - n + n[k_H/k_D]^{-1})^v$$

Eq. 15

When $v = 1$, the entire isotope effect ($k_H/k_D = 2.17$, Table 10) is realized at one site. When $v = 2$, two sites are equally affected, as in conterminous, two-proton transfer ($k_H/k_D = 2.17^{1/2} = 1.47$). When $v = 3$, three sites are uniformly impacted, ($k_H/k_D =$
$2.17^{1/3} = 1.30$), and when $v = \infty$ a general solvation condition at multiple sites by water is effected.$^{96}$ The plot clearly exhibited a curved line, precluding a one site effect, while advancing a minimum of two site H-bonding between triplet pH acetate $^{3}114$ and the solvent. The phenolic proton of $^{3}114$ was calculated to be more than 4 orders of magnitude more acidic that $114$, whereas, the carbonyl moiety is more basic.$^{78}$ Two site SKIE results were in good agreement with these properties; a) one site, a robust H-bond donation of the phenolic proton to the enveloping water molecules and b) the other site, H-bond accepting of the carbonyl with the surrounding water molecules with concomitant phosphate departure. These results confirm that hydration is a sine qua non for the photo-Favorskii reaction, as previously asserted by Wan,$^{73}$ Givens,$^{78}$ and Phillips.$^{80,81,82,86,87,88,89}$

A prodigious quantity of information has been gleaned over the past decade with respect to the photo-Favorskii rearrangement. This has advanced a number of appropriate mechanisms to account for the steps leading to formation of the signature, rearranged phenylacetic acid (ester) photoproduct. Further studies are warranted, however, to further improve the overall comprehension of the photo-Favorskii rearrangement, much the same as Bordwell’s pervasive studies on the ground state Favorskii rearrangement, such as the impacts of substituents, added salt (ionic strength), and media among others, which have received little or no attention.
Statement of Problem

The $p$-hydroxyphenacyl (pHP) chromophore has received a considerable amount of attention over the past decade as a versatile photoremovable protecting group for a wide variety of substrates. The intriguing rearrangement of pHP that accompanies photoinduced substrate release has been equated with the well known, ground state Favorskii rearrangement. This excited state process, therefore, has been termed the “photo-Favorskii” rearrangement. Several working mechanisms for this process have been posited, primarily from theoretical and spectroscopic studies.

These mechanistic studies, though useful in bringing together preliminary details on the rearrangement of the pHP chromophore, are narrowly restricted in scope, which emphasize only a small pool of pHP derivatives and photolysis conditions. A number of programmed modifications of the photo-Favorskii substrates and conditions would add immeasurably to our understanding and would further elucidate the mechanistic picture of this rearrangement: 1) Would electron donating or withdrawing substituents on pHP impact the photorelease of substrates? 2) Is there a positional effect expressed by the substituents? If so, what are its origins? Fluorine, from its inherently high electronegativity and lipophilicity, is an attractive substituent for pHP for the investigation of electronic effects on the photochemistry, solvent and solubility property effects, and spectroscopic characteristics. 3) The $\lambda_{\text{max}}$ of pHP at 280 nm extends slightly past 300 nm at 1 mM, a concentration that is typical for photolysis reactions. One goal of this research has
been to extend the absorption profile of pHp beyond 350 nm, making it more amenable for biological studies. The $m$-methoxy moiety has been the sole auxochrome investigated at this time; other groups and additional approaches addressing this goal could provide favorable efficiencies and extended $\lambda_{\text{max}}$. 4) The range of leaving groups or nucleofuges examined in mechanistic studies of the photo-Favorskii reaction has also been confined. An expansion of substrates, especially to establish the limiting parameters of the nucleofuge and the determination of correlations between the nature of the nucleofuge and photorelease parameters, would be useful. 5) Lastly, the majority of studies tailored for the investigation of the mechanism of the photo-Favorskii rearrangement have employed water and/or aqueous acetonitrile as the media. The variations of standard physical organic parameters related to the media, such as ionic strength, pH, water content, and lipophilicity, that influence the photoreactivity and product outcomes of pHp caged compounds have not been established and would certainly improve our mechanistic understanding of the photo-Favorskii rearrangement.
Results

I. Synthetic Methodologies

A. pHGABA series (122-141) and 4-fluorophenacyl GABA (142). The novel pHGABA esters that were synthesized along with 4-fluorophenacyl GABA are depicted in Figure 12.

**Figure 12.** Novel pHGABA esters.

\[
\begin{align*}
\text{122} & : R_1 = F, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{123} & : R_1 = H, R_2 = F, R_3 = H, R_4 = H, R_5 = H \\
\text{124} & : R_1 = F, R_2 = F, R_3 = H, R_4 = H, R_5 = H \\
\text{125} & : R_1 = F, R_2 = H, R_3 = F, R_4 = H, R_5 = H \\
\text{126} & : R_1 = F, R_2 = H, R_3 = H, R_4 = F, R_5 = H \\
\text{127} & : R_1 = H, R_2 = F, R_3 = F, R_4 = H, R_5 = H \\
\text{128} & : R_1 = F, R_2 = F, R_3 = H, R_4 = F, R_5 = H \\
\text{129} & : R_1 = F, R_2 = H, R_3 = F, R_4 = F, R_5 = H \\
\text{130} & : R_1 = CF_3, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{131} & : R_1 = OC_3F_3, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{132} & : R_1 = NO_2, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{133} & : R_1 = OCH_3, R_2 = H, R_3 = H, R_4 = OCH_3, R_5 = H \\
\text{134} & : R_1 = OCH_3, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{135} & : R_1 = H, R_2 = OH, R_3 = H, R_4 = H, R_5 = H \\
\text{136} & : R_1 = OH, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{137} & : R_1 = CH_3, R_2 = H, R_3 = H, R_4 = H, R_5 = H \\
\text{138} & : R_1 = CH_3, R_2 = H, R_3 = H, R_4 = CH_3, R_5 = H \\
\text{139} & : R_1 = H, R_2 = CH_3, R_3 = H, R_4 = H, R_5 = H \\
\text{140} & : R_1 = H, R_2 = H, R_3 = H, R_4 = H, R_5 = CH_3 \\
\text{141} & : R_1 = H, R_2 = H, R_3 = H, R_4 = H, R_5 = CH_2CH_3
\end{align*}
\]

1) Synthesis of 4-(2-(3-fluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, \((m\text{-F-pH}GABA, 122)\). The synthesis of 122 (Scheme 38) was initiated with commercially available 4-bromo-2-fluorophenol, (122a), which was reacted with benzyl bromide to produce the benzyl protected 122b in high yield. Protection of the phenol was required prior to the supervening, acetylation step. The benzyl moiety is robust to severe conditions and easily removed.
when appropriate; thus it was a logical choice as a protecting group. Stille conditions, entailing the use of tributyl-1-ethoxy(vinyl)stannane and Pd(0) in an inert atmosphere were then applied to 122b, effectuating 122c in 89% yield. This approach to acetylation has received very little attention, in fact none for preparing \( p \)-hydroxyacetophenones from phenol precursors. We have refined this methodology to the point where we routinely obtain >80% yields of targeted compounds. The supervening step entailed \( \alpha \)-bromination of 122c through reaction with dioxane-dibromide at 0°C, furnishing 122d in ~95% yield as estimated by \(^1\)H NMR. Prudence was required to prevent \( \alpha,\alpha' \)-dibromination from occurring; the reaction was generally halted before complete conversion and the yield was estimated by GC/MS or \(^1\)H NMR. Separation of 122d from 122c was cumbersome and not essential to the subsequent, caging step; thus, in subsequent studies, no purification was attempted. The procedures through the \( \alpha \)-bromination step were routinely used for all pHp and 5-acetylsalicylic acid derivatives. The penultimate step, arguably the most critical, encompassed the “caging” or coupling of GABA with the pHp chromophore. Specifically, 122d underwent a facile S\( _{\text{N2}} \) reaction with the carboxylic acid terminus of N-Boc protected GABA (Boc, \( \text{tert-butoxycarbonyl} \) at room temperature in the presence of potassium carbonate, to afford the fully carbamate protected pHp-ester, 122e in 89% yield. This substrate coupling procedure was commonly used for all carboxylic acid containing substrates, such as GABA, Glu, and long chain alkanes. Complete deprotection, i.e., the removal of the Boc and benzyl moieties of 122e was effected with distilled trifluoroacetic acid (TFA), 0°C to room temperature, over a 24
Scheme 38. The synthesis of m-F-pHP GABA (122).

h period. After lyophilization, 122 was produced in 78% yield. The fabrication of 123-132 followed a congruent set of steps as 122.

The synthetic sequence for 133-139 was initiated with commercially available \( p \)-hydroxyacetophenones. The \( m,m' \)-OCH\(_3\)-pHP GABA (133) and \( m \)-OCH\(_3\)-pHP GABA (134) were assembled following the procedure in a prior report.\(^97\) A modified synthetic procedure for 135-139 was utilized compared to 133 in efforts to achieve a better overall yield.

2) Synthesis of 4-(2-(2,4-dihydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, (\( o \)-OH-pHP GABA, 135). The synthetic strategy for 135 is
depicted in Scheme 39. To ensure that complete benzylolation of commercially available 1-(2,4-dihydroxyphenyl)ethanone (135b) was achieved, 3 equivalents of benzyl bromide and 4 equivalents of potassium carbonate were used and reacted for 36 h under mild conditions. After purification, 135b was furnished in 97% yield. Bromination to 135c was accomplished under standard conditions in ~94% yield by 1H NMR. The coupling of N-Boc GABA with 135c was achieved under standard conditions, though exhibited lower yields than other pHP’s giving, 135d in 48% yield. The diminished yield for this step may originate from a steric effect of the o-benzyl group. One additional step involving the facile removal of the benzyl constituent of 135d was used. The method entailed dissolving 135d in ethyl acetate, adding activated Pd/C (30% w/w), and then plugging the reaction vessel with a septum. While lightly stirring, hydrogen gas (H₂) was suffused into the reaction vessel by balloon through a long needle positioned near the surface of the solution. A short needle was added as an outlet to permit gas to escape; thus the atmosphere becomes saturated with H₂ generally after one full balloon, at which time the short needle is removed to prevent air from entering the vessel. Prudence was required in the handling of the reaction not only due to the heightened flammability imparted by H₂ but also Pd/C, which itself is extremely flammable. After H₂ addition, the reaction was monitored closely by TLC to prevent undesired reduction of the carbonyl moiety. Complete conversion was achieved after 4 h. Following workup and drying, 135e was afforded in 97% yield. The last step differed slightly from that used for 122-131. 135e was immersed in 1:1 mixture of a TFA:CH₂Cl₂ solution at room temperature.

Significant effervescence was observed from the degradation of the Boc group to CO$_2$ and isobutylene; the reaction proceeded for ~15 min, and following lyophilization, effected 135 in 85% yield.

Compounds 136-139 were generated in a manner analogous to 135. When necessary, very slight variations and concentrations in the methodology were used. The yields in each step did not diverge significantly excepting the N-Boc coupling step. Here, yields of 136d-139d were much improved (>80%); presumably, no o-substituent of the magnitude of benzyl ether (135d) encumbered the substitution process.
Compounds 140 and 141 represented enantiomeric mixtures of α-methyl and ethyl-pHP GABA analogs (α-methyl and ethyl-pHP GABA, Figure 13).

Figure 13. α-methyl and ethyl-pHP GABA enantiomers, (R,S)-140 and (R,S)-141 respectively.

The synthetic procedure to (R,S)-140 is outlined in Scheme 40. The reaction series commenced with commercially available p-hydroxypropiophenone (140a) which was then benzylated to 140b, α-brominated to racemic 140c and coupled with N-Boc GABA to furnish (R,S)-140d as a racemate. Resolution of these using normal phase flash chromatography with various eluents and HPLC was unsuccessful. Subsequent benzyl deprotection to (R,S)-140e also defied resolution as did the products after Boc deprotection to generate (R,S)-140. Further efforts to isolate these enantiomers from one another were unsuccessful. The α-ethyl analogs, (R,S)-141, were synthesized by the same procedure as above and were not successfully resolved either.
Scheme 40. Synthesis of α-methyl-pHP GABA enantiomers, (R,S)-140.

3) Synthesis of 4-(2-(4-fluorophenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, (4-fluorophenacyl GABA, 142). The preparation of 4-(2-(4-fluoro-phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate (142) patterned closely after the sequence to generate 135 (Scheme 41); the starting
material consisted of 4-fluoroacetophenone (142a). The objective in preparing 142 was to examine the π electron donating ability of fluorine, a mimic of hydroxyl, in assisting the photorelease of GABA.

**Scheme 41.** Synthesis of 4-fluorophenacyl GABA (142).

![Synthesis Scheme 41.](image)

a. DDB, CH₂Cl₂, 0° to rt, 4 h, ~93%. b. N-Boc-GABA, K₂CO₃, CH₃CN, 82%. c. TFA:CH₂Cl₂ (1:1), 15 min, 93%.

**B. pH₃-caged l-Glu series.** A small suite of pH₃ l-glutamate (Glu) compounds were synthesized; these are listed in Figure 14.

**Figure 14.** pH₃ Glu derivatives, 143-145.

![Figure 14.](image)

The synthesis of pH₃ l-glutamate derivatives (pH₃ Glu, 143-145) involved a congruent set of steps with that of 122; the yields for corresponding intermediates
were equable with those for 122 as well. A minor alteration for Glu, however, was an additional tert-butyl moiety that served to protect the ancillary carboxylic acid (Scheme 42). The effective removal of both the N-Boc and tert-butyl ester from Glu required extra reaction time in TFA:CH₂Cl₂ (1:1), approximately 2 h at room temperature.

Scheme 42. Synthesis of m-F-pHP Glu (144).

C. Other pHP caged amino-acids. Four additional amino acids were caged to the pHP chromophore (146-149). These compounds are listed in Figure 15.
1) Synthesis of 2-(2-(4-hydroxyphenyl)-2-oxoethoxy)-2-oxoethanaminium 2,2,2-
trifluoroacetate, (pHP Gly, 146). The synthetic progression of steps towards pHP
Gly (146), are depicted in Scheme 43 and are similar to the sequence for pHP GABA.
The sequence commenced with the benzylation of p-hydroxyacetophenone using
standard conditions, affording 146b in 98%. Bromination with DDB efficaciously
generated 146c in ~95% yield as indicated by $^1$H NMR. The supervening coupling of
146c with a slight excess of N-Boc Gly in the presence of potassium carbonate,
afforded 146d in 95% yield, which, subsequently underwent Pd catalyzed
debenzylation to form pHP N-Boc Gly (146e) in 94%. The last short step was routine
Boc removal to furnish 146 in 79% yield. The assembly of 147-149 paralleled that for 146. The other amino acids all possessed Boc-protected amine functionalities, akin to Gly; accordingly they were caged to pHp through their carboxylate termini.

D. pHp Deoxycortic acid. The bile acid, deoxycortic acid (DOC), was caged to the pHp chromophore.

10,13-dimethyl-hexadeca-hydro-1H-cyclopenta[a]phenanthren-17-yl)pentanoate (150) is illustrated in Scheme 44. The caging step of DOC was similar to 122e.

Scheme 44. Synthesis of pHP DOC (150).

Scheme 44.

\[
\begin{align*}
\text{a. DOC, K}_2\text{CO}_3, \text{DMF, 22 h, 84\%} & \quad \text{b. Pd/C, H}_2, \text{EtOAc, 6 h, 93\%}.
\end{align*}
\]

but DMF was employed instead of CH$_3$CN for solubility reasons, producing 146d in 84\% yield. Reductive hydrogenation of 146d followed under standard conditions and afforded 150 in 93\% yield.
**E. pHp Diethyl phosphonate.** A diethyl phosphonate moiety was tethered to the pHp chromophore.

1) **Synthesis of diethyl 2-(4-hydroxyphenyl)-2-oxoethylphosphonate, (pHP diethyl phosphonate, 151).** The preparation of 2-(4-(hydroxyphenyl)-2-oxoethylphosphonate (151) to explore the nucleofuge ability of the phosphonate moiety, is displayed in Scheme 45. The initial reaction involved the Michaelis-Arbuzov protocol;\textsuperscript{98} 146c was heated to 100°C in dry triethylphosphite for 17 h, effectuating 151d in 86% yield. Catalytic debenzylation of 151d then afforded 151 in 96% yield.

**Scheme 45.** Synthesis of pHp diethyl phosphonate (151).

\[
\text{146c} \xrightarrow{\text{a}} \text{151d} \xrightarrow{\text{b}} \text{151}
\]

a. Triethylphosphite, 100°C, 17 h, 86%. b. Pd/C, H\textsubscript{2}, EtOAc, 2 h, 96%.

**F. pHp F.** Fluoride was caged by the pHp chromophore.

1) **Synthesis of 2-fluoro-1-(4-hydroxyphenyl)ethanone, pHp F 152.**
The short series of chemical transformations to make 2-fluoro-1-(4-hydroxy-phenyl)-ethanone (152) was a study on the efficacy of fluoride release and is represented in Scheme 46. A facile S\textsubscript{N}2 reaction between cesium fluoride and 146\textsubscript{c} in refluxing CH\textsubscript{3}CN for 3 h quantitatively produced 152\textsubscript{d} by GC/MS analysis, 90% isolated yield. This was followed by a standard Pd/C reduction protocol to debenzylation of 152\textsubscript{d}, engendering 152 in 92% isolated yield.

**Scheme 46.** Synthesis of pHF F (152).

G. 5-acetylsalicylate and salicylamide GABA’s. An array of novel 5-acetylsalicylate GABA and alkyl esters and one 5-acetylsalicylamide were assembled to distinguish possible antennae effects on GABA release. These, 153, 154, and 157-165 are represented in Figure 16. Compounds 155 and 156, which possess alkanoyl substrates in place of GABA, will be introduced later.
1) **Synthesis of 4-(2-(3-carboxy-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, (5-acetylsalicylic acid GABA, 153).** The synthetic approach to 4-(2-(3-carboxy-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate (153), for a substituent effect study on GABA release, is manifest in Scheme 47. The precursor, 5-acetylsalicylic acid (153a) was reacted with a surfeit of benzyl bromide and potassium carbonate, the same approach to generate 153b, to provide the dibenzylated analog; the ensuing string of steps to produce 153 were analogous to those previously described for GABA coupling and complete deprotection. The starting material utilized in the preparation of 154 was the methyl ester of 153a; an ordinary sequence of steps was followed for the coupling of GABA and subsequent deprotection.
Scheme 47. Synthesis of 5-acetylsalicylic acid GABA (153).

Scheme 47.

2) Synthesis of 4-(2-(3-(4-benzoylphenylcarbamoyl)-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, (5-acetyl salicylamide benzophenone GABA, 157). The procedure to compose 157 is divulged in Scheme 48. The initial step involved the benzyl protection of commercially available 157a under standard conditions that was then hydrolyzed in 6 M KOH for 2 h at room temperature generating benzyl protected 5-acetylsalicylic acid (157b) in 98% yield after workup with conc. HCl. The second step involved concerned a Steglich esterification,\(^9\) whereby 157b was reacted with DCC and 4-aminobenzophenone in
the presence of DMAP at 0°C for 12 h. After filtration to remove dicyclohexylurea (DCU), a byproduct of the reaction, and purification, 157c was generated in 74% yield. Subsequent conversion to the α-bromo analog (157d) was facile by standard procedure. N-Boc GABA coupling to 157d proved to be challenging. The reaction
was attempted several times; the number of equivalents of N-Boc GABA was altered from 1.3 to 2.5 and reaction times from 24 to 48 h. TLC did not provide sufficient evidence to determine the extent of conversion and hence was abandoned. The reaction yields hovered around 46% irrespective of changes in conditions. At this time no explanation can be provided to account for the relatively low yields. Standard deprotection procedures converted 157d to 157 in relatively high yield. The same protocol was utilized to assemble 158-165.

The limiting step in the succession proved to be Pd-mediated hydrogenation to restore the phenol functionality, i.e., step c. Due to the reduction prospect of the carbonyl moiety, the reaction was assiduously monitored by thin layer chromatography. Compounds 153-165 exhibited discrete retentions in a variety of eluent mixtures and hence permitted the reaction to be followed until completion. However, at least a half dozen other analogs were synthesized that contained a Boc-GABA tethered constituent but evinced unchanged retentions between the benzylated and free phenol forms in all eluent mixtures. These compounds are listed in Figure 17. In these instances, the reaction progress could not be followed; highly variable product mixtures resulted from “guesses” of reaction times. The segregation of these products was cumbersome as these reactions were conducted on very small scales. The relatively labile carbamate and ester linkages possessed by the precursors precluded other common approaches to debenzylation, such as TFA, TMSI, or AlCl₃. Several iterations with minor changes in conditions, such as solvent, Pd/C content,
**Figure 17.** 5-acetylsalicylate GABA compounds where the standard Pd/C protocol for debenzylolation wasn’t effective.

etc. were ineffectual. Hence, a new strategy is required to educate improved conversions for this step or simply the use of a different protecting group. These derivatives were not studied further.
H. 5-acetylsalicylic acid alkanoyl compounds. Two 5-acetylsalicylic acid alkanoyl compounds were synthesized, 155 and 156 (Figure 18).

Figure 18. Two 5-acetylsalicylic acid alkanoyl derivatives, 155 and 156.

1) Synthesis of 5-(2-(dodecanoyloxy)acetyl)-2-hydroxybenzoic acid, (5-acetylsalicylic acid dodecanoate, 155). The synthesis of 5-(2-(dodecanoyloxy)acetyl)-2-hydroxybenzoic acid (155) was completed to study its utility as a photocleavable surfactant \(^{100}\) (Scheme 49). The short series of steps commenced with 5-acetylsalicylic acid (153a). The same benzylation protocol was utilized as for the assembly of 153b but without the hydrolysis phase, to afford 155b in high yield. Standard bromination conditions were used to furnish 155c in high yield. The subsequent reaction of 155c with a 2 molar excess of dodecanoic acid in a standard coupling method generated 155d in 71% yield. Lastly, 155 was produced under common debenzylation conditions in high yield. The same procedure was adhered to in generating 2-hydroxy-5-(2-(tetradecanoyloxy)acetyl)benzoic acid (156).
Scheme 49. Synthesis of 5-acetylsalicylic acid dodecanoate (155).

153a  \[ \xrightarrow{\text{a. BnBr, K$_2$CO$_3$, CH$_3$CN, 24 h, 97\%}} \] 155b  \[ \xrightarrow{\text{b. DDB, CH$_2$Cl$_2$, 0\^\circ\text{C} \text{to rt.} 2 \text{h}, \sim 96\%}} \] 155c  \[ \xrightarrow{\text{c. Dodecanoic acid, K$_2$CO$_3$, CH$_3$CN, 30 h, 79\%}. \text{d. Pd/C, H$_2$, EtOAc, 92\%}.} \] 155d  \[ \xrightarrow{\text{d.} \] 155

a. BnBr, K$_2$CO$_3$, CH$_3$CN, 24 h, 97%. b. DDB, CH$_2$Cl$_2$, 0\(^{\circ}\text{C}\) to rt. 2 h, \sim 96%. c. Dodecanoic acid, K$_2$CO$_3$, CH$_3$CN, 30 h, 79%. d. Pd/C, H$_2$, EtOAc, 92%.

I. \(\alpha\)-alkyl-pHP diastereomers. Two \(\alpha\)-alkyl-pHP diastereomers were synthesized.

1) Synthesis of \(((R)-(R)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate, 166\) and \(((R)-(S)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate, 167\). The synthetic strategy for the construction of diastereomers \((R)-(R)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate (166) and \((R)-(S)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate (167), is outlined in Scheme 50. The starting material for the sequence consisted of racemic 1-(4-(benzyloxy)phenyl)-2-bromopropan-1-one (140c)
which underwent a S<sub>N</sub>2 reaction with (R)-2-acetoxy-2-phenylacetic acid in the presence of potassium carbonate to afford a mixture of 166a and 167a in 94% overall yield. Efforts to segregate these diastereomers were ineffectual. Standard debenzylation

**Scheme 50.** Synthesis of α-methyl-pHP diastereomers (166 and 167).

a. (R)-2-acetoxy-2-phenylacetic acid, K<sub>2</sub>CO₃, CH₃CN, 94%. b. Pd/C, H₂, EtOAc, 99%.
conditions permitted 166a and 167a to be converted to 166 and 167, respectively, in 99% yield. Attempts to separate these diastereomers by normal phase silica chromatography and reverse phase HPLC were unsuccessful. Further work on this series is expected upon completion of the thesis.

J. FITC-DOC. A molecule derived from fluorescein isothiocyanate and deoxycholic acid was constructed and given to collaborators for use in biological assays. Details from these assays will be provided in the Discussion section.

1) Synthesis of (R)-4-((3R,5R,8R,9S,10S,12S,13R,14S,17R)-3,12-dihydroxy-10,13-dimethylhexadecahydro-1H-cyclopenta[a]phenanthren-17-yl)-N-(6-(3-(3',6'-dihydroxy-3-oxo-3H-spiro[isobenzofuran-1,9'-xanthene]-5-yl)-thioureido)-hexyl)-pentanamide, (FITC-DOC, 168). The process for the fabrication of 168 is exhibited in Scheme 51. The series commenced with a modified Steglich esterification,\textsuperscript{101} where deoxycholic acid, 168a, was reacted with 1-ethyl-3-(3-dimethyl aminopropyl) carbodiimide hydrochloride (EDCI) modified N-Boc-1,6-hexaneamine with N-hydroxybenzotriazole as an initiator to furnish 168b. This compound was not isolated but further reacted with 4 M HCl in 1,4-dioxane for a short time to induce N-Boc degradation and form the ammonium trifluoroacetate species (168c) in good yield. The last step involved base (triethylamine) assisted, nucleophilic addition of 168c to fluorescein isothiocyanate (FITC) to afford 168 in moderate yield.
Scheme 51. Synthesis of FITC-DOC (168).

II. Exploratory Photochemistry

As discussed in the Introduction, photolysis of the pHP caged complexes in aqueous media generally results in facile substrate release and concurrent precursor
rearrangement to the major $p$-hydroxyphenylacetic acid chromophore and minor $p$-hydroxybenzyl alcohol (abbreviated Scheme 37, *Introduction*).

**Scheme 37.** (from the *Introduction*) Photolysis of $p$-hydroxyphenacyl diethyl phosphate (114) in water.

The formation of $p$-hydroxyphenylacetic acid (95) and $p$-hydroxybenzyl alcohol (121), a newly discovered minor product, are easily discerned by $^1$H NMR analysis when pHG caged compounds are dissolved in deuterium labeled hydroxylic solvents, as D$_2$O, then irradiated. The methylene unit of 95 has a characteristic chemical shift, $\delta = 3.6$ ppm and that corresponding to 121, $\delta = 4.5$ ppm. The peaks corresponding to the methylene constituents of pHG GABA precursors, 95 and 121, being conspicuous indicators of a photochemical reaction, are denoted as *signature* peaks. To investigate whether this phenomenon is consistent with caged pHG derivatives 122-167, preliminary photolyses monitored by $^1$H NMR were conducted in aqueous media under standard (non degassed, room temperature) conditions in NMR tubes.
A. pHGABA and 4-fluorophenacyl GABA ester investigations employing $^{1}$H NMR spectroscopy. Irradiation at 300 nm of 1-10 mM concentrations of pHGABA esters (122-142) in D$_{2}$O were followed up to 2 h. $^{1}$H NMR analysis was implemented to identify the constituents of the mixture. A number of pHGABA derivatives manifested an inertness to photolysis; these included, m-NO$_{2}$-pHP GABA (132), o-OH-pHP GABA (135), m-OH-pHP GABA (136) and 4-fluorophenacyl GABA (142). Protracted irradiation times up to 24 h at 254, 300, and 350 nm produced no detectible $p$-hydroxyphenylacetic acids, as determined from the lack of characteristic chemical shifts, and only small amounts of hydrolysis products, i.e., $\alpha$-hydroxyl (acyloin) acetophenones, presumably resulting from the thermal, ground state reaction. Compounds 122-131 and 137-141 evinced clean photorelease of GABA, with $p$-hydroxyphenylacetic acids as the major products estimated to be $\geq$95% and $p$-hydroxybenzyl alcohol as the minor product estimated to be $\leq$5%. No other photoproducts from these precursors were observed by $^{1}$H NMR. An example of this is illustrated with $^{1}$H NMR spectra associated with photolysis of 0.02 M m-OCF$_{3}$-pHP GABA (131) in D$_{2}$O (Figures 19-23). Prior to photolysis, 131 manifested two signature sets of signals; the methylene component, labeled a at $\delta$ = 5.47 ppm, and the tethered GABA region in the aliphatic region, assigned b. The large signal at $\delta$ $\sim$ 4.80 ppm represents HOD, which will be commonly seen in all spectra where D$_{2}$O was utilized as the primary solvent. After 2 h of irradiation, pronounced vicissitudes in
Figure 19. $^1$H NMR spectrum of $m$-OCF$_3$-pHP GABA (131) in D$_2$O.

The $^1$H NMR spectrum was evident (Figure 20). The signal associated with a is absent and two new peaks have materialized; the signal labeled e at $\delta = 3.66$ ppm represents the methylene unit of $p$-hydroxyphenylacetic acid (131$_1$) and the small peak labeled d at $\delta = 4.56$ ppm that corresponds to the methylene group of $p$-hydroxybenzyl alcohol (131$_2$). The bands that are assigned e correlate with free GABA, which divulge subtle differences in chemical shifts compared to caged GABA. The peak at c is incontrovertibly linked to the methylene of 131$_1$ based on
spiking experiments with an *authentic sample*; this entailed adding a pure sample of *p*-hydroxyphenylacetic acid to the photolysis mixture. Supervening $^1$H NMR analysis manifested increases in intensities of peaks linked to this entity. In addition, to corroborate the assignment of d to $131_2$, *p*-hydroxy-*m*-trifluoromethoxybenzyl alcohol was independently synthesized and the authentic sample was then added into the photolysis mixture for $^1$H NMR analysis, shown in Figure 21. The $^1$H NMR band
Figure 21. Spiking experiment with $p$-hydroxy-$m$-trifluoromethoxybenzyl alcohol ($131_2$).

The intensity of $d$ at $\delta = 4.55$ ppm is pronounced and is associated with the increase in the concentration of $131_2$, unequivocally confirming the presence of the benzyl alcohol. It should be noted that the peak near $\delta = 4.90$ ppm could constitute another variant, the acyloin $131_3$. Another, authentically synthesized sample of the $m$-OCF$_3$-acyloin was prepared and added to the sample to discern whether it was a photoproduct; the appearance of a new signal at $e$ with $\delta = 4.95$ ppm next to the 4.90 ppm signal clearly demonstrated the absence of $131_3$, precluding the acyloin as a photoproduct. Likewise, $^{19}$F NMR’s were examined where spiking experiments bolstered the $^1$H
NMR results. The results of photolyses of $m$-OCH$_3$-pHP GABA (134) and $m,m'$-diOCH$_3$-pHP GABA (133) were also investigated by $^1$H NMR analysis. These had been reported to exhibit preponderant GABA release pathways that were not related to pHP rearrangement at 300 nm.$^{97}$ A 0.03 M solution of 134 in D$_2$O was prepared (Figure 22 the signature band is designated a, the methylene protons at $\delta = 5.53$ ppm)

**Figure 22.** Pre-photolysis $^1$H NMR spectrum of $m$-OCH$_3$-pHP GABA (134) in D$_2$O.
and then irradiated for 90 min at 300 nm (Figure 23). The salient chemical entity in the photolysis mixture was unmistakably unreacted starting material; it held true to the reported very low quantum efficiency for conversion. The major photoproduct, however, was the rearranged p-hydroxy-m-methoxyphenylacetic acid, its signature resonance band for methylene protons labeled b δ = 3.69 ppm. In addition, a band labeled c at δ = 4.5 ppm, likely to be associated with the methylene protons of p-hydroxy-m-methoxybenzyl alcohol, was apparent. There were a few minor signals due to the presence of other photoproducts which were not identified. The photolysis of nearly the same concentration of m,m’-diOCH3-pHP GABA (133) in D2O with 300 nm lamps for 90 minutes produced new bands at δ = 3.67 and δ = 4.56 ppm; the former probably indicated the p-hydroxyphenylacetic acid analog of 133 and the latter the p-hydroxybenzyl alcohol. The primary chemical species in the mixture was unreacted 133. It is noteworthy to point out the significant disparities in photochemical behavior of m-OCF3-pHP GABA (131) and m-OCH3-pHP GABA (133); the former manifests a much higher efficacy for chromophore rearrangement and GABA release. The primary structural deviations are fluorine moieties in 131 that commute with hydrogens in 133; the provenience will be discussed later in the Discussion section.
Figure 23. $^1$H NMR spectrum of the photolysis mixture of 134 at 300 nm in D$_2$O.

B. $^1$H NMR examination of $\alpha$-alkyl-pHP GABA esters. Compounds (R,S)-140, $\alpha$-methyl-pHP GABA enantiomers, and (R,S)-141, $\alpha$-ethyl-pHP GABA enantiomers, exhibited uncomplicated and complete photoconversions under similar conditions as described above. For example, a 0.05 M solution of a racemic mixture of (R,S)-140 in D$_2$O (Figure 24) was irradiated for 1 h with two 300 nm lamps (Figure 25) and showed nearly complete conversion to photoproducts by diminutive a with $\delta = 6.08$
Figure 24. $^1$H NMR spectrum of α-methyl-pHP GABA enantiomers (R,S)-140 in D$_2$O prior to irradiation.

ppm and new bands for free GABA and photoproducts with resonance bands at b $\delta = 5.22$ ppm, c $\delta = 3.63$ ppm and d $\delta = 3.23$ ppm. A spiking experiment with 2-(4-hydroxyphenyl)propanoic acid (Figure 26) revealed an enhanced band intensity at d with $\delta = 3.63$, the identity of the p-hydroxyphenylpropionic acid photoproduct. The signals at b are tentatively assigned to the methine protons of α-hydroxy (acyloin) ketone of pHP based on the downfield chemical shift of aromatic hydrogen atoms.
Figure 25. $^1$H NMR spectrum of the product mixture after 1 h irradiation of (R,S-140) at 300 nm, in D$_2$O.

i.e., $\delta = 7.89$ ppm, that are similar to 140 and impute a ketone moiety. The resonance band at d could not be identified.
Figure 26. $^1$H NMR spectrum of photolysis mixture with added 2-(4-hydroxyphenyl)propanoic acid. The enhanced peak intensity of c confirms the structure of the photoproduct.

C. Examinations of pHP-l-glutamate derivatives (143-145) by $^1$H NMR analysis. The photochemistry of pHP Glu derivatives, o-F-pHP Glu (143), m-F-pHP Glu (144), and m-CF$_3$-pHP Glu (145) was congruent with the corresponding GABA analogs 122, 123 and 130, respectively, employing identical photolysis conditions. Free l-glutamate along with rearranged p-hydroxyphenylacetic acids were released as major
Figure 27. Pre photolysis $^1$H NMR spectrum of $m$-CF$_3$-pHP Glu (145) in 1:1 D$_2$O: CD$_3$OD.

products; the sole minor products were $p$-hydroxybenzyl alcohols. For example, $^1$H NMR analysis of 0.01 mM of 145 in 50% aq. CD$_3$OD (Figure 27) that was photolyzed for 1 h at 300 nm (Figure 28) revealed the complete disappearance of the signature methylene bands labeled a at $\delta = 5.45$ ppm and Glu labeled b, with appearance of bands that correspond to methylene protons of $p$-hydroxy-$m$-trifluoromethylphenylacetic acid, labeled c at $\delta = 3.66$ ppm and $p$-hydroxy-$m$-
trifluoromethylbenzyl alcohol, labeled d at δ = 4.55 ppm, as well as free Glu (bands labeled e).

**Figure 2.** $^1$H NMR analysis of products from photolysis of $m$-CF$_3$-pHP Glu (145) at 300 nm in 1:1 CD$_3$OD:D$_2$O.

D. **Investigations of pHP-caged amino acids (146-148) by $^1$H NMR analysis.** The photochemistry of pHP caged glycine (146), valine (147), tryptophan (148) and phenylalanine (149) paralleled that of GABA under the same photolysis conditions.
Clean transformations to $p$-hydroxyphenylacetic acid and $p$-hydroxybenzyl alcohol, along with substrate release, were observed.

**E. $^1$H NMR analysis for the photolysis of pHp deoxycholic acid, (pHP DOC, 150).** A sample of 0.009 M pHp DOC (150) in 1:9 D$_2$O:CD$_3$OD was prepared (Figure 29, the signature band is denoted a for methylene protons at $\delta = 5.37$ ppm and

**Figure 29.** $^1$H NMR spectrum of 150 in 1:9 D$_2$O:CD$_3$OD prior to photolysis.
b for methylene protons at $\delta = 2.49$ ppm) and was irradiated at 300 nm over 4 h (Figure 30); complete disappearance of precursor was divulged by the absence of a. New bands appeared at b with $\delta = 4.37$ ppm and c at $\delta = 3.57$ ppm which are posited to represent the methylene protons of rearranged $p$-hydroxyphenylacetic acid and $p$-hydroxybenzyl alcohol entities respectively. Additionally, free DOC was observed from signature resonance bands b at $\delta = 2.38$ ppm.

Figure 30. $^1$H NMR spectrum of the product mixture after irradiation of 150 for 4 h at 300 nm in 1:9 D$_2$O:CD$_3$OD.
F. $^1$H NMR exploration of pH diethyl phosphonate (151). Irradiation of 0.02 M of 151 in D$_2$O for 2, 4, 16, and 36 h at 300 nm lamps failed to disclose any photolysis products. Additional photolysis studies with 0.015 M 151 in D$_2$O at 254 nm over 12 h evinced no photoproducts.

G. $^1$H NMR analysis of pH F (152). A 0.01 M sample of 152 in 1:3 D$_2$O:CD$_3$OD

Figure 31. $^1$H NMR analysis of 152 in 1:3 D$_2$O:CD$_3$OD prior to irradiation.
(Figure 31, with signature, discrete signals for each the methylene group alpha to the ketone a at δ = 5.72 and 5.61 ppm respectively) was irradiated at 300 nm for 1 h (Figure 32). Noticeably, the bands at a were significantly diminished in intensity. Prominent new signals were also observed at δ = 3.56 and 3.52 ppm. These are located at a similar position in the spectrum as the methylene protons of p-hydroxyphenylacetic acid; the exhibition of two signals in this spectrum, however, differs with the acid, which typically reveals only one resonance band. A second

**Figure 32.** $^1$H NMR assessment of the product mixture after 1 h photolysis of 152 at 300 nm in 1:3 D$_2$O:CD$_3$OD.
prominent signal is conveyed at $\delta = 4.36$ ppm and could not be identified. Further studies are required to better understand the photochemistry of pH F (152).

H. $^1$H NMR evaluation of 5-acetylsalicylic acid (153) and salicylate GABA derivatives (154, 157-165). A 0.18 M solution of 5-acetylsalicylic acid GABA (153) in D$_2$O (Figure 33, with signature methylene bands a at $\delta = 5.46$ ppm and GABA

Figure 33. $^1$H NMR spectrum of 153 in D$_2$O prior to photolysis.
alkyl bands \( b \) at \( \delta = 3.10, 2.69 \) and 2.04 ppm) was irradiated for 1 h at 300 nm (Figure 34) manifesting the clean photorelease of GABA, \( d \). This was evidenced by the appearance of resonance bands associated with free GABA at \( \delta = 3.03, 2.54, \) and 1.96 ppm, which are slightly offset in the spectrum from caged GABA. A considerable amount of starting material, 153, remained due to the relatively short irradiation time.

**Figure 34.** \(^1\)H NMR analysis of the product mixture from 1 h photolysis of 153 at 300 nm in D\(_2\)O.
A new, prominent signal at $\delta = 3.66$ ppm suggests the presence of rearranged phenylacetic acid analog labeled $c$, analogous to the previously assigned derivatives. Salicylic esters and amide derivatives of 153, compounds 154 and 157-165, exhibited congruent photochemical behaviors with 153. Several of these exhibited limited water solubilities and required, first, dissolution in a CD$_3$OD or CD$_3$CN, followed by dilution with D$_2$O.

1. $^1$H NMR studies of (5-(2-(dodecanoyloxy)acetyl)-2-hydroxybenzoic acid, 155) and (2-hydroxy-5-(2-(tetradecanoyloxy)acetyl)benzoic acid, 156). The photochemical behavior of model photocleavable surfactants 155 and 156 were investigated in 1:9 D$_2$O:CD$_3$OD. Similar to antennae compounds 157-165, water solubility was a restriction. However, it was found that this was obviated by dissolution in basic buffer, which will be elaborated on later. An $^1$H NMR study of
Figure 35. $^1$H NMR analysis of 155 in d$^6$-DMSO prior to irradiation.

155 consisted a 0.03 M solution in 1:9 D$_2$O:CD$_3$OD (Figure 35), with methylene protons labeled a at $\delta = 5.40$ ppm and b at $\delta = 2.40$ ppm, was irradiated for 1 h at 300 nm (Figure 36) to give three new resonance signals centered at $\delta = 4.42$ (d), 3.64 (c), and 3.59 ppm (c). A significant amount of 155 was retained, evidenced by the band at a. It was deduced that dodecanoic acid was released from the 5-acetylsalicylic acid chromophore from the appearance of new bands at e at $\delta = 2.31$ ppm,
Figure 36. $^1$H NMR spectrum of the product mixture after irradiation of 155 in 1:9 D$_2$O:CD$_3$OD.

... corresponding to free dodecanoic acid, which were slightly offset from the caged analog. The chemical shifts of other products indicate that b might represent the methylene protons of the $p$-hydroxyphenylacetic acid derivative, and c or d correspond to methylene protons of $p$-hydroxybenzyl alcohol derivative.
J. \textsuperscript{1}H NMR investigations of ((R)-((R)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate, 166) and ((R)-((S)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl) 2-acetoxy-2-phenylacetate, 167). A 0.05 M solution of a mixture of pHp

**Figure 37.** \textsuperscript{1}H NMR spectrum of a mixture of 166 and 167 in 1:1 D\textsubscript{2}O:CD\textsubscript{3}OD.

phenylacetate diastereomers (166 and 167) in 1:1 D\textsubscript{2}O:CD\textsubscript{3}OD (Figure 37) with signature methine protons for each at a and b that overlap at $\delta = 6.08$ and 6.04 ppm was irradiated with at 300 nm lamps for 2 h. Complete conversion of precursors (Figure 38) were apparent from the absence of signals corresponding to a and b. New
resonance signals were manifested at $c$ with $\delta = 5.92$ ppm that corresponded to the released $(R)$-2-acetoxy-2-phenylacetic acid with resonance signals at 5.27, 4.34, 3.71 and 3.67 ppm. Comparing these to those obtained from the product mixture of $(R-S)$

above, the bands at $\delta = 5.27$ ppm are posited as the methine protons of the $\alpha$-hydroxy (acyloin) derivative $d$; those at $\delta = 3.71$ and 3.67 ppm likely represent the methylene protons of rearranged $p$-hydroxyphenylacetic acid derivative $e$, which had been confirmed previously from spiking experiments. In addition, the small resonance bands labeled $f$ at 4.34 ppm could represent the methylene protons of $p$-hydroxybenzyl alcohol analogs.
Figure 38. $^1$H NMR spectrum of the product mixture from irradiation of 166 and 167 at 300 nm in 1:1 CD$_3$OD:D$_2$O.

K. $^{19}$F NMR analysis of fluoro pHp and 5-acetylsalicylic derivatives. $^{19}$F NMR spectroscopy provided a useful technique to assess the progress of photoreactions of certain pHp derivatives. The compounds that were anticipated to manifest the most conspicuous changes in $^{19}$F resonance band positions
were those that possessed an o-fluoro moiety; the powerful inductive effect imparted by the carbonyl of pHp ketone on the o-fluoro atom is quite disparate from the corresponding p-hydroxyphenylacetic acid, where this effect is obtunded by a more remote and modified carboxylate functionality. For example, a 0.03 M D₂O solution of 2,3,5,6-tetrafluoro-pHP GABA (129) (Figure 39) with signature ¹⁹F NMR bands for the m-fluoro moieties labeled a (δ = -162 ppm), the o-fluoro moieties labeled b (δ
-142 ppm), and the trifluoro constituent of the trifluoroacetate anion labeled c (δ = -76 ppm) was photolyzed for 15 min with at 300 nm (Figure 40). Incomplete

**Figure 40.** $^{19}$F NMR spectrum of the product mixture from photolysis of 129 at 300 nm in D$_2$O.

conversion to the rearranged $p$-hydroxy-2,3,5,6-tetrafluorophenylacetic acid, with resonance signals assigned d for o-fluorines (δ = -147 ppm) and e for m-fluorines (δ = -164 ppm), and $p$-hydroxy-2,3,5,6-tetrafluorobenzyl alcohol, with resonance signals
assigned f for o-fluorines (δ = -147 ppm) and g for m-fluorines (δ = -164 ppm) are clearly discernible with respect to those from 129. The intensity of the band e is greater than f, which is expected, as the former is the major product. The corresponding 1H NMR studies divulged that the major photoproduct of 129 was rearranged p-hydroxy-2,3,5,6-tetrafluorophenylacetic acid and thus the 19F intensities would be major at δ = -147 (d) and at δ = -164 ppm (e). The p-hydroxy-2,3,5,6-tetrafluorobenzyl alcohol product, exiguously formed, would be identified with signals at lower intensities, i.e., f and g. A sizeable upfield shift in the o-fluoro resonance band, a to d of -5 ppm for the phenylacetic acid and a to g -6 ppm for the benzyl alcohol, is due to the loss of deshielding from the carbonyl. Additionally, minor vicissitudes in chemical shifts of m-fluoro moieties were perceived; 1 ppm for b to e and f, respectively, were due to the loss of the resonance and inductive withdrawing contribution of the carbonyl. The resonance and inductive bearing of the carbonyl ketone is underscored by these chemical shift changes.

III. UV-Vis Spectroscopic Studies.

A. pH and p-fluorophenacyl GABA esters 122-142 in water and buffers, pH = 7.3 and 8.2. UV-Vis spectroscopic analysis of compounds 122-142 in water and pH = 8.2 buffer were completed and results presented in Table 15.
Table 15. UV-Vis properties of 122-142 with corresponding pKa values.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>λ_{max}/nm (log ε[M^-1cm^-1])^a</th>
<th>λ_{max}/nm (log ε[M^-1cm^-1])b</th>
<th>λ_{max}/nm (log ε[M^-1cm^-1])c</th>
<th>pKa^d</th>
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</thead>
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<td>122</td>
<td>274 (4.00), 335 (2.97)</td>
<td>330 (4.06)</td>
<td>351 (3.67), 328 (3.96)</td>
<td>6.5</td>
</tr>
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<td>123</td>
<td>271 (3.97)</td>
<td>322 (3.87), 284 (3.91)</td>
<td>328 (3.90), 319 (3.91)</td>
<td>7.2</td>
</tr>
<tr>
<td>124</td>
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<td>323 (4.10)</td>
<td>323 (4.07)</td>
<td>5.9</td>
</tr>
<tr>
<td>125</td>
<td>278 (3.26), 326 (3.92)</td>
<td>328 (4.06)</td>
<td>326 (4.12)</td>
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</tr>
<tr>
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<td>331 (4.12)</td>
<td>330 (4.14)</td>
<td>5.3</td>
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<tr>
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<td>313 (4.03)</td>
<td>313 (4.06)</td>
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<tr>
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<td>324 (4.11)</td>
<td>323 (4.01)</td>
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<td>317 (4.08)</td>
<td>316 (4.10)</td>
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<td>329 (4.18), 322 (4.18)</td>
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<td></td>
<td></td>
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<td></td>
<td>254 (3.92)</td>
<td>397 (3.20), 334 (4.14)</td>
<td>399 (3.22), 335 (4.20)</td>
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</tr>
<tr>
<td>133^e</td>
<td>303 (3.90), 355 (3.55)</td>
<td>ND</td>
<td>ND</td>
<td>7.8</td>
</tr>
<tr>
<td>134^e</td>
<td>279 (3.97), 307 (3.90)</td>
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<td>ND</td>
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<tr>
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<td>334 (3.77), 318 (3.59),</td>
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<td></td>
</tr>
<tr>
<td>135</td>
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<td>ND</td>
<td>7.8</td>
</tr>
<tr>
<td>136</td>
<td>309 (3.97), 279 (3.86)</td>
<td>ND</td>
<td>ND</td>
<td>8.0</td>
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<tr>
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<td>284 (3.84)</td>
<td>283 (3.77), 307 (3.65)</td>
<td>356 (3.75), 334 (3.97)</td>
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<td>304 (3.79), 287 (3.96)</td>
<td>316 (3.40), 305 (3.81),</td>
<td>362 (3.91), 348 (4.05)</td>
<td>8.2</td>
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<tr>
<td></td>
<td></td>
<td>287 (3.88)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>139</td>
<td>287 (4.06)</td>
<td>303 (3.45), 287 (3.98)</td>
<td>343 (3.86), 328 (4.10)</td>
<td>8.0</td>
</tr>
<tr>
<td>140</td>
<td>281 (4.10)</td>
<td>329 (3.98), 284 (3.92)</td>
<td>333 (4.02)</td>
<td>7.9</td>
</tr>
<tr>
<td>141</td>
<td>282 (4.07)</td>
<td>328 (3.96), 284 (3.94)</td>
<td>334 (4.12)</td>
<td>7.9</td>
</tr>
<tr>
<td>142</td>
<td>273 (3.57), 267 (3.89), 248(4.12)</td>
<td>ND</td>
<td>ND</td>
<td>-</td>
</tr>
</tbody>
</table>

^a In water. ^b In 0.01 M HEPES, 0.1 M LiClO4, pH = 7.3. ^c In 0.01 M HEPES, 0.1 M LiClO4, pH = 8.2. ^d Determined for the corresponding p-hydroxyacetophenones, see Experimental for details. ^e See reference 1

The impetus for spectral investigations in buffers relates to the protonated versus conjugate base forms of the pH chromophore, i.e., to discern whether the conjugate bases manifest disparate photophysical properties to their protonated counterparts.

Several other buffers of identical concentration and adjusted ionic strengths were utilized; these included TRIS pH 7.5, tetrasodium pyrophosphate pH 7.4, sodium phosphate pH 7.0, ammonium acetate, pH = 7.0 and HEPES pH = 7.2 with 0.1 M LiCl. The UV-Vis parameters for these were consistent with those obtained
with HEPES, pH = 7.3, shown in Table 15. The pKa of the phenol influenced the UV-Vis spectra for all pHp esters; many derivatives manifested two or more absorption bands in unbuffered, aqueous media, resulting from the equilibrium between the protonated and conjugate base concentrations of the phenol. In HEPES buffered solution at pH = 8.2 the conjugate bases of all pHp esters dominated. The long wavelength maxima were bathochromically shifted by ~50 nm compared with the protonated form. For example, o-F-pHP GABA (123) revealed a $\lambda_{\text{max}} = 271$ nm in water, signifying the protonated form; in HEPES buffer, pH = 8.2, the $\lambda_{\text{max}}$ shifts to 328 and 319 nm (Figure 41), respectively, a clear indication of the conjugate base. The photophysical parameters that influence substrate release depend on the nature of the protonated and conjugate base of the pHp chromophore and will be discussed later.

By manipulating the pH of the photoreaction media and the lamps for the Rayonet reactors (300 and 350 nm), the photoreactivity of the two forms of 122-141 allow the accurate quantization of quantum yields and rate constants for each form individually. In water, 122-141 exist dually as a mixture which makes the analysis complex (two competing photoreactions). Furthermore, as the reaction progresses, two additional changes are occurring. The pH of the aqueous solution will change, since the Favorskii rearrangement both releases an acid as the leaving group and generates a phenylacetic acid, each have different pKa’s than the phenol. The second factor is a consequence of this change in that the absorption spectrum will shift to represent a higher proportion of the original protonated form of the caging
Figure 41. Comparison of UV-Vis spectra of 123 in water (red) and pH = 8.2 buffer (pink).

For example, o-F-pHP GABA (123) revealed a $\lambda_{\text{max}}$ of 271 nm (Figure 41) at low concentrations ($10^{-5}$ M) in water, with a tapering off at 320 nm. At 1 mM concentrations, an extended absorbance profile used for photolysis was manifested (Figure 42) and reached the baseline at 350 nm. The cutoff was at 350 nm for excitation of the protonated forms of 122-132 and 135-141 in water (the $m$-methoxy compounds 133 and 134 extended past 370 nm at 5 mM concentrations). Such was not the case for the corresponding conjugate bases of pHP chromophores 122-141 (Table 15). Congruent with their protonated counterparts, saturation UV-Vis profiles of 5 mM concentrations of 122-141 conjugate bases, all or nearly all of the light was
being absorbed at wavelengths up to 400 nm and therefore the 350 nm lamps would seem to be amenable for photorelease. For example, when probed at 5 mM in 0.01 M HEPES, 0.1 M LiClO₄, pH = 8.2, 123 showed absorbances that extended past 350 nm (Figure 43), with a cut off nearer to 370 nm.

**Figure 42.** UV-Vis spectroscopic analysis of 1 mM solution of 123 in unbuffered aqueous solution. The spectrum indicates a saturation condition.
The photo-conversion process of pHG GABA esters 122-141 was probed by UV-Vis spectroscopy. Recall the p-fluorophenacyl GABA 142 demonstrated photoinertness at $\lambda = 254$ and 300 nm as determined by $^1$H NMR and thus was not examined. All of the caged GABA derivatives were converted to products with a high efficiency when irradiated as 2 mM aqueous solutions at 300 nm (light output, 0.01 mE/mol) where complete transformation to photoproducts occurred within 20 min (Table 16). This was verified by HPLC (UV-detection) through analysis of
Table 16. Times for 100% conversion to photoproducts of 1 mM concentrations of 122-141 in water.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Time&lt;sup&gt;a&lt;/sup&gt; (min)</th>
<th>Time&lt;sup&gt;b&lt;/sup&gt; (min)</th>
<th>Time&lt;sup&gt;c&lt;/sup&gt; (min)</th>
<th>Time&lt;sup&gt;d&lt;/sup&gt; (min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>122</td>
<td>7</td>
<td>9</td>
<td>70</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>123</td>
<td>5</td>
<td>6</td>
<td>45</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>124</td>
<td>6</td>
<td>12</td>
<td>45</td>
<td>&gt;8h</td>
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<td>125</td>
<td>10</td>
<td>10</td>
<td>80</td>
<td>&gt;8h</td>
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<td>&gt;8h</td>
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<td>127</td>
<td>13</td>
<td>15</td>
<td>70</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>128</td>
<td>9</td>
<td>9</td>
<td>20</td>
<td>&gt;8h</td>
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<tr>
<td>129</td>
<td>6</td>
<td>9</td>
<td>40</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>130</td>
<td>12</td>
<td>13</td>
<td>60</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>132&lt;sup&gt;e&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>133&lt;sup&gt;f&lt;/sup&gt;</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>134&lt;sup&gt;f&lt;/sup&gt;</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>135&lt;sup&gt;g&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>136&lt;sup&gt;g&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>137</td>
<td>8</td>
<td>7</td>
<td>50</td>
<td>&gt;8h</td>
</tr>
<tr>
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</tr>
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<td>140</td>
<td>20</td>
<td>24</td>
<td>90</td>
<td>&gt;8h</td>
</tr>
<tr>
<td>142&lt;sup&gt;e&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

<sup>a</sup>Two 300 nm lamps, in water, estimated by HPLC/UV analysis in water. <sup>b</sup>Two 300 nm lamps, in 0.01 M HEPES, 0.1 M LiClO<sub>4</sub>, pH = 7.3, estimated by HPLC/UV analysis. <sup>c</sup>Two 300 nm lamps, in 0.01 M HEPES, 0.1 M LiClO<sub>4</sub>, pH = 8.2, estimated by HPLC/UV analysis. <sup>d</sup>16 350 nm lamps, in 0.01 M HEPES, 0.1 M LiClO<sub>4</sub>, pH = 8.2, estimated by HPLC/UV analysis, overnight irradiation. <sup>e</sup>No photolysis was observed. <sup>f</sup>Not determined.

aliquots every 30 seconds. For 122-141 in 0.01 M HEPES buffer with 0.1 M LiClO<sub>4</sub> at pH = 8.2 (Table 15), the conjugate bases were irradiated with two 300 nm lamps over longer time periods also achieved complete conversion to photoproducts. Exposure of the conjugate bases to sixteen 350 nm lamps (0.09 mE/min) led to
complete conversion to photoproducts but at much longer times, i.e., >8 hours. An illustration of a typical photolysis, as probed by UV-Vis spectroscopy, is shown in Figure 44. Here, a 10^{-4} M sample of 2,3,5,6-tetrafluoro-pHP GABA (129) in water was irradiated with at 300 nm and analyzed at 3 minute intervals to complete conversion at 9 min. For 129, the absorption decrease at \( \lambda \sim 316 \) nm was

**Figure 44.** UV-Vis probe of the photolysis of 129, predominantly the conjugate base in water, monitored at 3 minute intervals.
accompanied by an increase at $\lambda \sim 220$ nm due to the principal photoproduct, $p$-hydroxy-2,3,5,6-tetrafluorophenylacetic acid (pHP Acid). An isosbestic point at 225 nm further validates that $p$-hydroxy-2,3,5,6-tetrafluorophenylacetic acid is the primary photoproduct of 129.

**B. pHP l-glutamate derivatives (143-145) in water and buffer pH = 8.20.** As anticipated, the UV-Vis analysis of pHP Glu derivatives (143-145) paralleled the UV-Vis profiles in water and buffer of their GABA counterparts 123, 122, 130 respectively. Corresponding $\lambda_{\text{max}}$ values, times for complete conversions (at 300 nm) in water, and pKa’s of 143-145 are shown in Table 17.

**Table 17.** UV-Vis parameters and estimated times for 100% conversion to photoproducts from irradiation at 300 nm and associated pKa’s for 143-145.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$\lambda_{\text{max}}$/nm (log $\varepsilon$[M$^{-1}$cm$^{-1}$])$^a$</th>
<th>$\lambda_{\text{max}}$/nm (log $\varepsilon$[M$^{-1}$cm$^{-1}$])$^b$</th>
<th>time (min)$^c$</th>
<th>time (min)$^d$</th>
<th>pKa</th>
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<td>143</td>
<td>272 (4.01)</td>
<td>323 (4.02)</td>
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<td>70</td>
<td>6.5</td>
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<tr>
<td>145</td>
<td>328 (4.09)</td>
<td>326 (4.05)</td>
<td>6</td>
<td>30</td>
<td>5.5</td>
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</tbody>
</table>

$^a$Measured in water. $^b$Measured in 0.01 M HEPES, 0.1 M LiClO$_4$, pH = 8.20. $^c$In water, estimated by HPLC/UV-Vis analysis. $^d$In 0.01 M HEPES, 0.1 M LiClO$_4$, pH = 8.20, estimated by HPLC/UV-Vis analysis.
C. Other pHp amino acids (glycine (146), valine (147), tryptophan (148), phenylalanine (149)), deoxycholic acid (150), diethyl phosphonate (151) and fluoride (152). The UV-Vis profiles for 146-152 in water and buffered media paralleled those

Table 18. UV-Vis parameters and 100% conversion times for 146-152 from irradiation at 300 nm.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$\lambda_{\text{max}}$/nm (log $\varepsilon$[M$^{-1}$cm$^{-1}$])$^{a}$</th>
<th>$\lambda_{\text{max}}$/nm (log $\varepsilon$[M$^{-1}$cm$^{-1}$])$^{b}$</th>
<th>time (min)$^{c}$</th>
<th>time (min)$^{d}$</th>
<th>pKa</th>
</tr>
</thead>
<tbody>
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<td>146</td>
<td>280 (4.09)</td>
<td>330 (3.94), 282 (3.89)</td>
<td>7</td>
<td>5</td>
<td>7.9</td>
</tr>
<tr>
<td>147</td>
<td>279 (4.07)</td>
<td>327 (3.89), 281 (3.76)</td>
<td>7</td>
<td>6</td>
<td>7.9</td>
</tr>
<tr>
<td>148</td>
<td>272 (4.12), 300 (3.89), 310 (3.67)</td>
<td>ND</td>
<td>15</td>
<td>13</td>
<td>7.9</td>
</tr>
<tr>
<td>149</td>
<td>282 (4.09)</td>
<td>328 (3.98), 282 (3.91)</td>
<td>7</td>
<td>6</td>
<td>7.9</td>
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<td>279 (4.00)$^{e}$</td>
<td>ND</td>
<td>20$^{c}$</td>
<td>ND</td>
<td>7.9</td>
</tr>
<tr>
<td>151$^{f}$</td>
<td>280 (4.03)</td>
<td>ND</td>
<td>-</td>
<td>-</td>
<td>7.9</td>
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<tr>
<td>152</td>
<td>282 (4.12)$^{g}$</td>
<td>331 (3.65), 282 (4.05)$^{h}$</td>
<td>12</td>
<td>10</td>
<td>7.9</td>
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</tbody>
</table>

$^{a}$Measured in water. $^{b}$Measured in 0.01 M HEPES, 0.1 M LiClO$_4$, pH = 7.3. $^{c}$In water, estimated by HPLC/UV-Vis analysis. $^{d}$In 0.01 M HEPES, 0.1 M LiClO$_4$ pH = 7.3, estimated by HPLC/UV-Vis analysis. $^{e}$Measured in 5% aq. CH$_3$OH. $^{f}$No photoreaction occurred. $^{g}$50% aq. CH$_3$OH. $^{h}$20% aq. CH$_3$OH.

for GABA analogs 140 and 141 above except for pHp Trp (148). The tryptophan moiety contributes to the absorptivity of the caged derivative. The nucleofuges for each, however, were quite distinct, resulting in variable conversion times. UV-Vis parameters and times for complete conversion are depicted in Table 18.

D. 5-acetylsalicylic acid and salicylate GABA and alkanoyl derivatives, 153-165.

UV-Vis spectra were obtained for compounds 5-acetylsalicylic acid GABA and alkanoyl analogs 153, 155-156, and 5-acetylsalicylate/amide GABA derivatives 154,
The parameters associated with these, along with times for complete conversion of 2 mM concentrations of precursor in water to photoproducts at 300 and 350 nm are illustrated in Table 19. The pKa’s for these were not determined. The principal intentions for synthesizing 157-165 were: a) To lengthen the $\lambda_{\text{max}}$ of the chromophore towards 350 nm and b) to construct and evaluate a potential for an energy relay (antennae) system from the added auxochrome to enhance the photochemical efficiency. Most of the entities manifested higher molar absorptivities at 300 nm rather than at 350 nm. The full transformations to photoproducts at 300 nm, however, were markedly longer than the pH analogs lacking the auxochrome.

### Table 19. UV-Vis parameters, 100% conversion times from irradiation at 300 nm, and % conversion after 24 h from irradiation at 350 nm for 153-165.

<table>
<thead>
<tr>
<th>Compound</th>
<th>$\lambda_{\text{max}}$/nm (log $\varepsilon$ [M$^{-1}$ cm$^{-1}$])$^a$</th>
<th>time (min)$^b$</th>
<th>% conversion$^c$</th>
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<td>153</td>
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<td>25</td>
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<tr>
<td>154$^d$</td>
<td>272 (4.09), 310 (3.22), 330 (3.94)</td>
<td>12</td>
<td>ND</td>
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<tr>
<td>155$^e$</td>
<td>282 (3.96)</td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>156$^e$</td>
<td>281 (4.00)</td>
<td>20</td>
<td>20</td>
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<tr>
<td>157$^f$</td>
<td>340 (3.76), 310 (3.92), 274 (4.15)</td>
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<tr>
<td>158$^g$</td>
<td>343 (2.61), 320 (3.16), 300 (3.36), 271 (3.80)</td>
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<tr>
<td>159</td>
<td>305 (3.52), 269 (3.92)</td>
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<td>160</td>
<td>306 (3.46), 269 (3.96), 229 (4.17)</td>
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<td>307 (3.38), 271 (3.82), 234 (4.11)</td>
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<td>25</td>
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<tr>
<td>164$^g$</td>
<td>312 (3.52), 273 (3.89)</td>
<td>17</td>
<td>25</td>
</tr>
<tr>
<td>165$^f$</td>
<td>313 (3.64), 273 (3.95), 227 (4.20)</td>
<td>11</td>
<td>50</td>
</tr>
</tbody>
</table>

$^a$ Measured in water. $^b$ Two 300 nm lamps, estimated by HPLC with UV-Vis detection in a gradient of methanol/water. $^c$ Sixteen 350 nm lamps, estimated by $^d$ $^1$H NMR analysis after 24 h, in 1:1 CD$_3$OD:D$_2$O. $^e$ Previously determined. $^f$ 90% aq. CH$_3$OH. $^g$ 75% aq. CH$_3$OH. $^i$ 95% aq. CH$_3$OH.
Excitation at 350 nm effectuated only marginal substrate release and necessitated more than 24 h for sufficient substrate release.

E. α-Alkyl pHp diastereomers (R,R)-166, and (R,S)-167. UV-Vis assessment of (R,R) 166 and (R,S) 167 produced an absorbance summary akin to 101 above. Exposure to two 300 nm lamps in 50% aq. CH$_3$OH conveyed complete conversion to photoproducts at 10 minutes for (R,R)-166 and 11 minutes (R,S)-167, as determined by HPLC/UV-Vis analysis.

IV. Fluorescence Spectroscopic Studies

The pHp chromophore does not fluoresce sufficiently for detection on a conventional, benchtop Varian Eclipse® spectrofluorimeter. The corresponding conjugate bases were weakly fluorescent. An example of fluorescence from m-CF$_3$-pHP GABA (130) in 0.01 M HEPES, 0.1 M LiClO$_4$ at pH = 8.2 is shown in Figure 45. The excitation wavelength was 325 nm, which produced a single, broad emission signal $\lambda_{em}$ at 380 nm that tailed out past 500 nm. Examinations of the fluorescence properties of pHp GABA derivatives, 122-131 and the unsubstituted parent in the
Figure 45. Fluorescence spectrum of 130 in 0.01 M HEPES, 0.1 M LiClO$_4$ pH = 8.2, $\lambda_{ex} = 325$ nm.

same media as 130 gave the same results as 130 and are reported in Table 20.

Table 20. Fluorescence of 122-131 in 0.01 M HEPES, 0.1 M LiClO$_4$ at pH 8.2.

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<thead>
<tr>
<th>Compound</th>
<th>$\lambda_{em}$ (nm)</th>
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</thead>
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<tr>
<td>122</td>
<td>395</td>
</tr>
<tr>
<td>123</td>
<td>382</td>
</tr>
<tr>
<td>124</td>
<td>390</td>
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<tr>
<td>129</td>
<td>384</td>
</tr>
<tr>
<td>130</td>
<td>380</td>
</tr>
<tr>
<td>131</td>
<td>395</td>
</tr>
<tr>
<td>Parent (101)</td>
<td>382</td>
</tr>
</tbody>
</table>
V. *Time correlated single photon counting, TCSPC*. TCSPC (see experimental), studies were carried out to obtain the fluorescence lifetimes for conjugate bases of 122-131. Exponential decay profiles and the fluorescence lifetimes were obtained (Table 21). The transients were determined by the Maximum Entropy Method.

**Table 21.** TCSPC derived fluorescence decay profiles of 101, 122-131 in aq. 0.1 M LiOH.

<table>
<thead>
<tr>
<th>Compound</th>
<th># transients</th>
<th>t₁ (ns)</th>
<th>t₂ (ns)</th>
</tr>
</thead>
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<td>0.129</td>
<td></td>
</tr>
<tr>
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<td>2</td>
<td>0.115</td>
<td></td>
</tr>
<tr>
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<td>2</td>
<td>0.156</td>
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<td>0.197</td>
<td></td>
</tr>
<tr>
<td>126</td>
<td>2</td>
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<tr>
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<td>2</td>
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<tr>
<td>130</td>
<td>3</td>
<td>0.039</td>
<td>1.803</td>
</tr>
<tr>
<td>131</td>
<td>2</td>
<td>0.013</td>
<td></td>
</tr>
</tbody>
</table>

An illustration of a TCSPC decay profile is provided in Figure 46 for a 0.1 mM solution of 130 in 0.1 M LiOH. In this example, a curve fitting application (red) correlated the fluorescence decay (black) to a single exponential best fit with a lifetime of 1.80 ns respectively.
**Figure 46.** TCSPC decay profile for \( m \)-CF\(_3\)-pHP GABA (130) in 0.1 M LiOH.

![Fluorescence Decay Profile, Conjugate Base of 130](image)

**VI. Quantitative Photochemical Analysis**

The quantum yields, \( \Phi \), equal the ratio \( n_x/n_p \), where \( n_x \) depicts the number of photochemical or photophysical occurrences \( x \), effectuated by light absorbed, \( p \), by the precursor at a particular wavelength \( \lambda \).\(^{102}\) The value can be considered as a percentage; for example, a \( \Phi \) of 0.20 for pHP GABA disappearance signifies that 20% of the excited state pHP GABA molecules decayed by substrate release. It follows that the higher the quantum yield, the greater efficacy for the overall photochemical process. Quantum yields were determined for those compounds 122-167 that conveyed substrate photorelease from preliminary \(^1\)H NMR studies; these included reactant disappearance (\( \Phi_{\text{dis}} \)), substrate appearance (\( \Phi_{\text{app}} \)), and appearance of
$p$-hydroxyphenylacetic acids ($\Phi_{\text{acid}}$). The quantum yields of the minor $p$-
hydroxybenzyl alcohols were not quantified. Analyses were accomplished by
HPLC/UV and LC/MS/MS techniques and were done in triplicate.

A. Quantum yields of pHGABA derivatives 122-141 from photolysis at 300
nm. The determination of quantum yields, $\Phi_{\text{dis}}$, $\Phi_{\text{app}}$, $\Phi_{\text{acid}}$ for 1-10 mM of 122-141
and the unsubstituted parent (101) at 300 nm were determined in an assortment of
media. The principal objectives for these studies were to distinguish any pronounced
variations in the quantum yields as a function of the media, i.e., effects due to buffer,
pH, added salts, and changes from organic to aqueous solvent ratios. Initial
photolysis studies involved the use of water buffered by 0.01 M ammonium acetate
that spanned from pH 5.0 to pH 9.0. Not all precursors were probed in these media.
The results are presented in Table 22. Blank spaces indicate “not determined”. In all
cases, the quantum yields for precursor disappearance, $\Phi_{\text{dis}}$, were commensurate with
those for the emergence of GABA, $\Phi_{\text{app}}$. The quantum yields for appearance of $p$-
hydroxyphenylacetic acids, $\Phi_{\text{acid}}$, were always slightly reduced; this phenomenon was
expected from a competing pathway to the minor product, $p$-hydroxybenzyl alcohol.
As such, $\Phi_{\text{dis}}$ will be emphasized from this point on with the assumption that $\Phi_{\text{app}}$ and
$\Phi_{\text{acid}}$ do not diverge significantly from $\Phi_{\text{dis}}$.

In general, values for $\Phi_{\text{dis}}$ were optimum in water or 0.01 M ammonium
acetate solutions at pH 5.0. At pH 6.0, the values for some are lower; for example, $m$-
F-pHGABA (123) exhibited a $\Phi_{\text{dis}} = 0.28$ (±0.02) at pH 5.0 and 0.22 (±0.01).
Table 22: Quantum yields for 122-141 and the unsubstituted parent (101) from photolyses in water and 0.01 M ammonium acetate buffer, pH 5.0 to pH 9.0.

<table>
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<tr>
<th>Compd</th>
<th>pKa</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
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<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
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</thead>
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</table>

*18 MΩ, 0.01 M Ammonium Acetate
This downward trend continued at pH 7.0; only the unsubstituted parent displayed unchanged yields from pH 5.0 to pH 7.0 (\(\Phi_{\text{dis}} = 0.20 \, (\pm 0.01)\)) to 7.0 (\(\Phi_{\text{dis}} = 0.20 \, (\pm 0.01)\)). At pH 9.0, \(\Phi_{\text{dis}}\) for 122-141 were minimal; the conjugate base forms of 122-141 prevail in this medium. This important trend correlating the pKa’s of 122-141 with solution pH will be analyzed further in the Discussion section.

A series of studies with 122-141 were performed in several other, commonly used buffers including TRIS, HEPES, and sodium phosphate, ranging in pH from 7.0 to 9.0. In some cases, salts were added to control the ionic strength. The findings from these are depicted in Table 23. Blank spaces indicate “not determined”. Collectively, \(\Phi_{\text{dis}}\) were relatively uniform in neutral pH ~ 7 media, regardless of the buffer type or modified ionic strength. A slightly enhanced \(\Phi_{\text{dis}}\) was observed when added LiClO_4 was used instead of LiCl in HEPES buffer. When photolyses were conducted in HEPES buffer pH = 8.2, the efficiencies universally declined; for example, the unsubstituted parent (101) manifested a \(\Phi_{\text{dis}}\) of 0.21 \((\pm 0.01)\) in 0.01 M HEPES and 0.1 M LiClO_4 at pH 7.3, which plummeted to 0.09 \((\pm 0.01)\) under identical conditions at pH 8.2. Additionally, when photolyses were executed in 0.01 M TRIS buffer with 0.1 M LiClO_4 at pH 9.0, where the conjugate bases are the sole species, most precursors evinced low quantum yields (>0.05). The declines in \(\Phi_{\text{dis}}\) of the conjugate bases were congruent with the results of the ammonium acetate buffers.

Another assemblage of photolysis experiments involved the use of organic media and mixed aqueous organic solvents. Degassing before photolysis with argon was also examined for O_2 effects on the quantum yields. The former studies
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<th>Cmpd</th>
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<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
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<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
<th>$\Phi_{\text{acid}}$</th>
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<td>0.20</td>
<td>0.20</td>
<td>0.18</td>
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*0.01 M HEPES, 0.1 M LiClO$_4$, pH = 7.3. *0.01 M HEPES, 0.1 M LiCl, pH = 7.2. *0.05 M Sodium Phosphate, pH = 7.0.

*0.01 M TRIS, 0.1 M LiClO$_4$, pH = 7.5. *0.01 M HEPES, 0.1 M LiClO$_4$, pH = 8.2. *0.01 M TRIS, 0.1 M LiClO$_4$, pH = 9.0.
were performed to determine whether hydroxylic solvents, excepting water, were sufficient to induce substrate photorelease from pHp caged compounds. Of particular note, all pHp GABA esters displayed full solubility in 1-pentanol and perhaps more remarkably, in 1-octanol. The results are provided in Table 24. Blank spaces indicate “not determined”.

Reasonable Φ_dis values were observed for nearly all precursors in 1-pentanol and 1-octanol, though in general these were lower than in water. Exceptions to this were 2,3,6-triF-pHP GABA (128) where Φ_dis were essentially the same as in water (0.09 (±0.01) and 0.08 (±0.008) respectively) and m,m′-diOCH3-pHP GABA (133) which displayed a pronounced improvement compared to water from (Φ_dis = 0.07 (±0.003) to 0.03 (±0.005) respectively). In 25% aq. CH3CN, marginal differences were observed for Φ_dis compared to water. However, 3,5-diF-pHP GABA (126) and 2,3,5,6-tetraF-pHP GABA (129) conveyed marked improvements (0.16 (±0.01) vs. 0.11 (±0.01) for the former, 0.14 (±0.01) vs. 0.10 (±0.01) for the latter).

Nearly all precursors manifested low quantum yields in DMSO; the fact that substrate photorelease occurred at all in this aprotic media is noteworthy. Interestingly, 128 exhibited an enhanced quantum yield in DMSO compared to water (0.12 (±0.01) and 0.08 (±0.008), respectively); this was further ameliorated in 10% aq. DMSO to Φ_dis = 0.15 (±0.02). Several other pHp GABA derivatives manifested slightly augmented yields in 10% aq. DMSO compared to water, including 2,5-diF-pHP GABA (125, 0.27 (±0.01) and 0.24 (±0.01), respectively), 126 (0.14 (±0.01) and
Table 24 Quantum yields for 122-131, 133-134 and the unsubstituted parent from photolysis in organic media and degassed aq. solution.

<table>
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<tr>
<th>Cmpd</th>
<th>1-pentanol</th>
<th>1-octanol</th>
<th>CH$_3$CN$^a$</th>
<th>DMSO</th>
<th>DMSO$^b$</th>
<th>Degassing$^c$</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>$\Phi_{\text{dis}}$</td>
<td>$\Phi_{\text{app}}$</td>
<td>$\Phi_{\text{dis}}$</td>
<td>$\Phi_{\text{app}}$</td>
<td>$\Phi_{\text{dis}}$</td>
<td>$\Phi_{\text{app}}$</td>
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</table>

$^a$25% aqueous. $^b$10% aqueous. $^c$Argon purge for 45 minutes.
0.11 (±0.01), respectively), 129 (0.18 (±0.01) and 0.11 (±0.01) respectively), and \textit{m}-OCH$_3$-pHP GABA (131 (0.12 (±0.01) and 0.09 (±0.01) respectively)).

Degassing, that entailed purging the aqueous solution of the pHP GABA derivative for 45 minutes with argon, failed to change the quantum yields.

Photolyses of 122-141 were conducted in water with modified ionic strengths to distinguish whether a \textit{salt effect} would be reflected in $\Phi_{\text{dis}}$. The first set of experiments focused on LiClO$_4$ as the ancillary salt, ranging from 0.008 to 1.0 M (Table 25). Remarkably, nearly all pH derivatives manifested enhancements in quantum yield at 1.0 M LiClO$_4$ in water. For example, a $\Delta \Phi_{\text{dis}}$ of 0.14, from 0.16 (±0.01) to 0.30 (±0.02) or a 94% increase for 122 was observed. Several pH derivatives evinced improved quantum yields in 0.01 M LiClO$_4$ relative to water as well, such as 126 with a $\Delta \Phi_{\text{dis}} = 0.07$, from 0.11 (±0.008) in water to 0.18 (±0.01) in 0.01 M LiClO$_4$.

Other ancillary salts (NaClO$_4$ and Mg(ClO$_4$)$_2$) were explored for similar influences on a few pH GABA derivatives, including the parent, 122, 123, and 124. In addition, experiments involving the addition of free GABA (1 and 10 equivalents) to aqueous solutions of these four pH-GABA precursors were performed to discern any changes in $\Phi_{\text{dis}}$ due to a \textit{common ion effect}. The findings from these studies are depicted in Table 26. Each of the precursors gave increased quantum yields in 1.0 M NaClO$_4$ and Mg(ClO$_4$)$_2$ relative to water, consistent with the results in LiClO$_4$. A curious observation, however, was the marked decline in $\Phi_{\text{dis}}$ at low concentrations of Mg(ClO$_4$)$_2$. For example, a decline in $\Phi_{\text{dis}}$ was observed for the 2,3-diF-pHP GABA
Table 25. Quantum yields for 122-141 and the unsubstituted parent from photolyses in water with adjusted ionic strengths from 0.008 to 1.0 M LiClO₄.

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<tr>
<td>135</td>
<td>0.19</td>
<td>0.16</td>
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<tr>
<td>136</td>
<td>0.15</td>
<td>0.14</td>
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<tr>
<td>137</td>
<td>0.08</td>
<td>0.07</td>
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<tr>
<td>138</td>
<td>0.10</td>
<td>0.09</td>
<td></td>
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</tbody>
</table>

*LiClO₄
Table 26. Quantum yields for 122-124 and the unsubstituted parent (101) from photolysis in water with adjusted ionic strengths and added GABA at 300 nm.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{acid}}$</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{acid}}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parent</td>
<td>0.30</td>
<td>0.28</td>
<td>0.10</td>
<td>0.08</td>
<td>0.32</td>
<td>0.30</td>
<td>0.20</td>
<td>0.18</td>
</tr>
<tr>
<td>122</td>
<td>0.25</td>
<td>0.24</td>
<td>0.08</td>
<td>0.06</td>
<td>0.25</td>
<td>0.23</td>
<td>0.17</td>
<td>0.16</td>
</tr>
<tr>
<td>123</td>
<td>0.35</td>
<td>0.32</td>
<td>0.14</td>
<td>0.13</td>
<td>0.34</td>
<td>0.31</td>
<td>0.29</td>
<td>0.27</td>
</tr>
<tr>
<td>124</td>
<td>0.31</td>
<td>0.28</td>
<td>0.12</td>
<td>0.10</td>
<td>0.32</td>
<td>0.29</td>
<td>0.25</td>
<td>0.23</td>
</tr>
</tbody>
</table>

*NaClO$_4$, $^b$Mg(ClO$_4$)$_2$.

(124), from 0.24 ($\pm$0.02) in water to 0.12 ($\pm$0.01) in 0.01 M Mg(ClO$_4$)$_2$. At 1.0 M Mg(ClO$_4$)$_2$, however, $\Phi_{\text{dis}}$ increased to 0.31 ($\pm$0.02). Added GABA imparted no change in $\Phi_{\text{dis}}$ relative to solutions without added GABA.

**B. Quantum yields of pH-GABA derivatives, 122-131, from photolysis at 350 nm.** The quantum yields for 1-10 mM concentrations of 122-131 in 0.01 M TRIS, 0.1 M LiClO$_4$ at pH 9.0 were determined by photolyses at 350 nm (Table 27), where the conjugate bases preponderate.
Table 27. Quantum yields for 122-131 in 0.01 M TRIS, 0.1 M LiClO₄ at pH 9.0 at 350 nm.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>Φ&lt;sub&gt;dis&lt;/sub&gt;</th>
<th>Φ&lt;sub&gt;app&lt;/sub&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parent (101)</td>
<td>0.0041</td>
<td>0.0038</td>
</tr>
<tr>
<td>122</td>
<td>0.0086</td>
<td>0.0083</td>
</tr>
<tr>
<td>123</td>
<td>0.0046</td>
<td>0.0045</td>
</tr>
<tr>
<td>124</td>
<td>0.0017</td>
<td>0.0013</td>
</tr>
<tr>
<td>125</td>
<td>0.0029</td>
<td>0.0024</td>
</tr>
<tr>
<td>126</td>
<td>0.0019</td>
<td>0.0018</td>
</tr>
<tr>
<td>127</td>
<td>0.0037</td>
<td>0.0035</td>
</tr>
<tr>
<td>128</td>
<td>0.0042</td>
<td>0.0035</td>
</tr>
<tr>
<td>129</td>
<td>0.0069</td>
<td>0.0067</td>
</tr>
<tr>
<td>130</td>
<td>0.0081</td>
<td>0.0074</td>
</tr>
<tr>
<td>131</td>
<td>0.0058</td>
<td>0.0054</td>
</tr>
</tbody>
</table>

Conspicuously, the quantum yields were substantially diminished in contrast to those obtained with 300 nm lamps.

C. Quantum yields for pHP-/-glutamate derivatives, 143-145, determined by photolysis at 300 nm. The quantum yields of 143-145 and unsubstituted parent (102) were ascertained from photolysis of 1-10 mM in water, 0.01 M ammonium acetate, 0.01 M HEPES buffers with 0.1 M LiClO₄ at pH 7.3 and pH 8.2 (Table 28) at 300 nm. The determination of Φ<sub>dis</sub>, Φ<sub>app</sub> (Glu) and Φ<sub>acid</sub> were consistent with those from pHP GABA’s. Only Φ<sub>dis</sub> will be discussed. No significant variations were observed in Φ<sub>dis</sub> in water and with neutral pH ~ 7 buffers, apart from 145, which gave lower Φ<sub>dis</sub> in pH 7.3 buffer (0.11 (±0.01) versus 0.18 (±0.01) in water). All showed declined Φ<sub>dis</sub> in pH 8.2 buffer, associated with conjugate bases, and coherent to findings from those pHP GABA derivatives.
D. Quantum yields of pH caged amino acids glycine, valine, phenylalanine, and tryptophan, 146-149, at 300 nm. The quantum yields (Φ_dis and Φ_acid) associated with 146-149 at 300 nm at 1-10 mM concentrations in water and 0.01 M HEPES with 0.1 M LiClO₄ at pH 7.3 were determined. In addition, salt effects were explored for 149 (0.1 and 1.0 M LiClO₄ aqueous solutions). The findings from these studies are summarized in Table 29. No noteworthy variations were discerned for Φ_acid and Φ_dis under any conditions. In HEPES buffer pH 7.3, an enhancement in the quantum yields was observed for 146-148. Added salts did not significantly improve Φ_dis.

Table 28. Quantum yields of 143-145 and unsubstituted parent (102) from photolysis in water, pH 7.0, pH 7.3 and pH 8.2 buffers at 300 nm.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>water⁴</th>
<th>pH 7.0⁵</th>
<th>pH 7.3⁶</th>
<th>pH 8.2⁶</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Φ_dis</td>
<td>Φ_app</td>
<td>Φ_acid</td>
<td>Φ_dis</td>
</tr>
<tr>
<td>Parent</td>
<td>0.16</td>
<td>0.16</td>
<td>0.12</td>
<td>0.16</td>
</tr>
<tr>
<td>143</td>
<td>0.15</td>
<td>0.15</td>
<td>0.12</td>
<td>0.18</td>
</tr>
<tr>
<td>144</td>
<td>0.16</td>
<td>0.15</td>
<td>0.13</td>
<td>0.14</td>
</tr>
<tr>
<td>145</td>
<td>0.18</td>
<td>0.18</td>
<td>0.15</td>
<td>0.17</td>
</tr>
</tbody>
</table>

⁴18 MΩ. ⁵0.01 M Ammonium Acetate. ⁶0.01 M HEPES, 0.1 M LiClO₄
Table 29. Quantum yields of 146-149 in water, pH 7.3 buffer, and water with adjusted ionic strengths.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>water</th>
<th>pH 7.3</th>
<th>0.1 M</th>
<th>1.0 M</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$\Phi_{\text{dis}}$</td>
<td>$\Phi_{\text{acid}}$</td>
<td>$\Phi_{\text{dis}}$</td>
<td>$\Phi_{\text{acid}}$</td>
</tr>
<tr>
<td>146</td>
<td>0.17</td>
<td>0.14</td>
<td>0.22</td>
<td>0.18</td>
</tr>
<tr>
<td>147</td>
<td>0.17</td>
<td>0.13</td>
<td>0.22</td>
<td>0.17</td>
</tr>
<tr>
<td>148</td>
<td>0.16</td>
<td>0.15</td>
<td>0.20</td>
<td>0.16</td>
</tr>
<tr>
<td>149</td>
<td>0.10</td>
<td>0.08</td>
<td>0.09</td>
<td>0.13</td>
</tr>
</tbody>
</table>

*18 MΩ. b0.01 M HEPES, 0.1 M LiClO₄. cLiClO₄

E. Quantum yields of pHp DOC (150) and pHp F (152) at 300 nm. Photolysis of 1-5 mM of 150 in 5% aq. CH₃OH produced $\Phi_{\text{dis}}$ of 0.06 (±0.007) and $\Phi_{\text{acid}}$ of 0.04 (±0.006) at 300 nm. For 152, $\Phi_{\text{dis}}$ was determined to be 0.10 (±0.01) at 1 mM in 50% aq. CH₃OH at 300 nm.

F. Quantum yields of 5-acetylsalicylic acid derivatives 153-165 at 300 nm. The quantum yields for disappearance, $\Phi_{\text{dis}}$, of 1-10 mM of 5-acetylsalicylic acid GABA (153) in an assortment of media at 300 nm are listed in Table 30. The quantum yields for appearance of GABA were found as the same $\Phi_{\text{dis}}$. The quantum yields were relatively uniform, regardless of the nature photolysis medium, though slightly enhanced in aqueous 1.0 M LiClO₄ and 1.0 NaClO₄ solutions. In a previous report, the $\Phi_{\text{dis}}$ of the $m$-carbomethoxy derivative of 5-acetylsalicylic acid GABA (154) was given as 0.31 from photolysis in water at 300 nm; this diverges significantly from $\Phi_{\text{dis}}$.
Table 30. $\Phi_{\text{dis}}$ of 153 in various media from photolysis at 300 nm.

<table>
<thead>
<tr>
<th>Compound</th>
<th>water$^a$</th>
<th>pH 7.2$^b$</th>
<th>pH 7.3$^c$</th>
<th>pH 7.0$^d$</th>
<th>pH 9.0$^e$</th>
<th>0.01M$^f$</th>
<th>0.05M$^g$</th>
<th>0.1M$^h$</th>
<th>0.5M$^i$</th>
<th>1.0M$^j$</th>
<th>0.1M$^k$</th>
<th>1.0M$^l$</th>
</tr>
</thead>
<tbody>
<tr>
<td>153</td>
<td>0.09</td>
<td>0.08</td>
<td>0.09</td>
<td>0.11</td>
<td>0.09</td>
<td>0.10</td>
<td>0.11</td>
<td>0.12</td>
<td>0.10</td>
<td>0.12</td>
<td>0.12</td>
<td></td>
</tr>
</tbody>
</table>

$^a$18 MΩ. $^b$0.01 M HEPES, 0.1 M LiCl. $^c$0.01 M HEPES, 0.1 M LiClO₄. $^d$0.05 M Sodium Phosphate. $^e$LiClO₄. $^f$NaClO₄.

$= 0.09 \pm 0.01$ for 153. The quantum yields for 154 were re-evaluated at 300 nm in an assortment of media. The results are depicted in Table 31. Evidently, the quantum yields for 154 are consistent with those determined for the parent and substantially lower than the $\Phi_{\text{dis}} = 0.31$ previously reported. Additionally, insignificant variations in these values were observed in all solvents tested.

Table 31. The values for $\Phi_{\text{dis}}$ and $\Phi_{\text{app}}$ of 154 from photolyses at 300 nm in a variety of media.

<table>
<thead>
<tr>
<th>Compound</th>
<th>water$^a$</th>
<th>pH 7.3$^b$</th>
<th>pH 9.0 M$^b$</th>
<th>1.0 M$^c$</th>
</tr>
</thead>
<tbody>
<tr>
<td>154</td>
<td>0.08</td>
<td>0.07</td>
<td>0.09</td>
<td>0.08</td>
</tr>
</tbody>
</table>

$^a$18 MΩ. $^b$0.01 M HEPES, 0.1 M LiClO₄. $^c$LiClO₄.

The quantum yields for the disappearance ($\Phi_{\text{dis}}$) of 5-acetylsalicylic acid dodecanooyl (155) and tetradecanoyl (156) esters were determined to be 0.04 ($\pm 0.005$) and 0.06 ($\pm 0.006$) respectively, from photolyses at 300 nm in 5% aqueous CH₃OH.
Additionally, $\Phi_{\text{dis}}$ in 0.01 M HEPES, 0.1 M LiClO$_4$, at pH 8.2 were 0.10 (±0.01) and 0.09 (±0.01) respectively, consistent with the GABA caged 153.

The quantum yields for the disappearance ($\Phi_{\text{dis}}$) of 157-165 “antennae” derivatives of 5-acetylsalicylic acid GABA, and appearance ($\Phi_{\text{app}}$) of GABA were determined in 3:1 H$_2$O:CH$_3$OH at 300 nm (Table 32). The values were slightly irregular and except for 157 ($\Phi_{\text{dis}} = 0.01$ (±0.002)), 159 ($\Phi_{\text{dis}} = 0.02$ (±0.002)), and 162 ($\Phi_{\text{dis}} = 0.04$ (±0.006)), did not vary appreciably from the parent 153 ($\Phi_{\text{dis}} = 0.09$ (±0.01)). When irradiated at 350 nm in water and 0.01 M HEPES, 0.1 M LiClO$_4$ at pH 8.2, 157-165 gave very low $\Phi_{\text{dis}}$ and $\Phi_{\text{app}}$ values (<0.01).

Table 32. $\Phi_{\text{dis}}$ and $\Phi_{\text{app}}$ for 157-165 in 3:1 H$_2$O:CH$_3$OH at 300 nm.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$\Phi_{\text{dis}}$</th>
<th>$\Phi_{\text{app}}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>157</td>
<td>0.01</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>158</td>
<td>0.08</td>
<td>0.07</td>
</tr>
<tr>
<td>159</td>
<td>0.02</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>160</td>
<td>0.06</td>
<td>0.05</td>
</tr>
<tr>
<td>161</td>
<td>0.08</td>
<td>0.07</td>
</tr>
<tr>
<td>162</td>
<td>0.04</td>
<td>0.04</td>
</tr>
<tr>
<td>163</td>
<td>0.11</td>
<td>0.11</td>
</tr>
<tr>
<td>164</td>
<td>0.07</td>
<td>0.06</td>
</tr>
<tr>
<td>165</td>
<td>0.12</td>
<td>0.11</td>
</tr>
</tbody>
</table>

G. Quantum yields of $\alpha$-alkyl-pHP caged diastereomers (R,R) 166 and (R,S) 167 at 300 nm. The quantum yields for a mixture of (R,R) 166 and (R,S) 167 diastereomers was determined from photolysis at 300 nm of 5 mM concentrations of
precursor in 50% aq. CH₃OH; these were found to be $\Phi_{\text{dis}} = 0.18 \ (\pm 0.01)$ and $\Phi_{\text{acid}} = 0.16 \ (\pm 0.01)$, consistent with unsubstituted pHP-GABA, $0.20 \ (\pm 0.01)$ and $0.16 \ (\pm 0.01)$ respectively.

**VII. Determination of Rate Constants for Substrate Release by Stern-Volmer Quenching**

The Stern-Volmer (S-V)⁷⁸ quenching method to indirectly determine the rate constants for substrate release was employed for GABA esters of 122-131, 137-139, and 153. One to five mM of each was prepared in water or buffer, photolyzed at 300 nm and $\Phi_{\text{dis}}$ determined. Potassium sorbate, a triplet quencher $Q$, was added in increments and $\Phi_{\text{dis}}$ determined. Using linear regression analysis, plots of the ratios $\Phi_o$ (quantum yield without $Q)/\Phi$ (quantum yield with $Q)$ vs. $[Q]$ gave linear S-V plots with $K_{SV}$ as the slope. For example, a 5 mM aqueous solution of 2,3-diF-pHP GABA (124) was photolyzed at 300 nm lamps along with three other solutions containing 20 mM, 40 mM, and 60 mM of potassium sorbate. A plot of corresponding $\Phi_o/\Phi$ vs. $[Q]$ values is illustrated in Figure 47. The slope, $K_{SV}$, is shown as the product of the diffusion rate, $k_{q}$, of the quencher in water (estimated to be $7.4 \times 10^9 \text{ M}^{-1}\text{s}^{-1}$)¹ and the triplet lifetime of chromophore, $\tau^3$. The rate of sorbate quenching is assumed to be diffusion controlled, i.e., $k_{\text{diff}} = k_{q}$. From the plot, $\tau^3$ was calculated to be $4.14 \times 10^9$ s or 4.14 ns, and the rate of GABA release, $k_r$, = $5.75 \times 10^7$ s⁻¹. Congruent analyses were completed for the other pHP GABA derivatives and
in water and 0.01 M HEPES, 0.1 M LiClO$_4$ at pH = 7.3. The results are listed in Table 33.

Figure 47. Stern-Volmer relationship from the photolysis of 2,3-diF-pHP GABA (124) in water with potassium sorbate as the quencher, Q.

$\Phi_o/\Phi = 1 + K_{SV}[Q]$  
Slope = $K_{SV} = 30.61$  
$k_{diff} = k_q = 7.4 \times 10^9 \text{ s}^{-1}$  
$K_{SV} = k_q \times \tau^3$  
$\tau^3 = 4.14 \times 10^{-9} \text{ s}$  
$k_r = \Phi_o/\tau^3 = 5.75 \times 10^7 \text{ s}^{-1}$

<table>
<thead>
<tr>
<th></th>
<th>$\Phi$</th>
<th>$\Phi_o/\Phi$</th>
<th>[Sorbate], M</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial ($\Phi_o$)</td>
<td>0.24</td>
<td>1.00</td>
<td>0.00</td>
</tr>
<tr>
<td>Series #1</td>
<td>0.14</td>
<td>1.76</td>
<td>0.02</td>
</tr>
<tr>
<td>Series #2</td>
<td>0.11</td>
<td>2.30</td>
<td>0.04</td>
</tr>
<tr>
<td>Series #3</td>
<td>0.08</td>
<td>2.86</td>
<td>0.06</td>
</tr>
</tbody>
</table>

The rate constants for GABA release spanned about 1 order of magnitude, from $k_r = 2 \times 10^7 \text{ s}^{-1}$ for $m$-OCF$_3$-pHP GABA (131), to $k_r = 1.58 \times 10^8 \text{ s}^{-1}$ for $o$-F-pHP GABA (123), in water. An overall decline in the rate constants was observed when the
solvent was changed to 0.01 M HEPES pH 7.3, except for 2,6-diF-pHP GABA (126) and 2,3,5,6-tetraF-pHP GABA (129), which evinced a near doubling of rate constants (4.7 to 8.9 and 2.7 to 5.5 $\times 10^7$ s$^{-1}$, respectively).

Table 33. Parameters derived from Stern-Volmer quenching analysis for 122-131, 137-139, and 153 in water and 0.01 M HEPES, 0.1 M LiClO$_4$ at pH = 7.3.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$K_{SV}$ (M$^{-1}$)</th>
<th>$\tau^3$ ($10^9$ s)</th>
<th>$k_q (10^7$ s$^{-1}$)</th>
<th>$K_{SV}$ (M$^{-1}$)</th>
<th>$\tau^3$ ($10^9$ s)</th>
<th>$k_q (10^7$ s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>101 (parent)</td>
<td>24 ± 1$^a$</td>
<td>3.2 ± 0.4</td>
<td>6.8 ± 0.4</td>
<td>26 ± 1$^a$</td>
<td>3.6 ± 0.4</td>
<td>5.5 ± 0.1</td>
</tr>
<tr>
<td>122</td>
<td>28 ± 2</td>
<td>3.8 ± 0.2</td>
<td>4.2 ± 0.2</td>
<td>23 ± 1</td>
<td>3.1 ± 0.3</td>
<td>3.6 ± 0.2</td>
</tr>
<tr>
<td>123</td>
<td>13 ± 0.2</td>
<td>1.9 ± 0.1</td>
<td>15.8 ± 0.1</td>
<td>19 ± 1</td>
<td>2.5 ± 0.1</td>
<td>8.4 ± 0.2</td>
</tr>
<tr>
<td>124</td>
<td>44 ± 2</td>
<td>6.0 ± 0.6</td>
<td>4.0 ± 0.4</td>
<td>39 ± 1</td>
<td>5.3 ± 0.4</td>
<td>2.1 ± 0.3</td>
</tr>
<tr>
<td>125</td>
<td>15 ± 0.8</td>
<td>2.1 ± 0.1</td>
<td>10.7 ± 0.1</td>
<td>51 ± 2</td>
<td>6.9 ± 0.2</td>
<td>1.6 ± 0.1</td>
</tr>
<tr>
<td>126</td>
<td>29 ± 0.8</td>
<td>4.0 ± 0.1</td>
<td>2.5 ± 0.1</td>
<td>32 ± 1</td>
<td>4.3 ± 0.1</td>
<td>1.1 ± 0.2</td>
</tr>
<tr>
<td>127</td>
<td>25 ± 0.5</td>
<td>3.3 ± 0.1</td>
<td>4.7 ± 0.1</td>
<td>8 ± 1</td>
<td>1.1 ± 0.1</td>
<td>8.9 ± 0.1</td>
</tr>
<tr>
<td>128</td>
<td>34 ± 2</td>
<td>4.6 ± 0.4</td>
<td>1.8 ± 0.1</td>
<td>36 ± 1</td>
<td>4.8 ± 0.2</td>
<td>1.3 ± 0.2</td>
</tr>
<tr>
<td>129</td>
<td>30 ± 0.5</td>
<td>4.1 ± 0.1</td>
<td>2.7 ± 0.1</td>
<td>14 ± 1</td>
<td>1.9 ± 0.3</td>
<td>5.5 ± 0.3</td>
</tr>
<tr>
<td>130</td>
<td>20 ± 2</td>
<td>2.7 ± 0.3</td>
<td>7.4 ± 0.7</td>
<td>30 ± 2</td>
<td>4.1 ± 0.5</td>
<td>3.0 ± 0.4</td>
</tr>
<tr>
<td>131</td>
<td>33 ± 0.7</td>
<td>4.4 ± 0.2</td>
<td>2.0 ± 0.1</td>
<td>29 ± 2</td>
<td>4.0 ± 0.3</td>
<td>1.4 ± 0.3</td>
</tr>
<tr>
<td>137</td>
<td>17 ± 1</td>
<td>2.2 ± 0.2</td>
<td>6.7 ± 0.3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>138</td>
<td>23 ± 0.8</td>
<td>3.1 ± 0.5</td>
<td>4.9 ± 0.7</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>139</td>
<td>15 ± 1</td>
<td>2.1 ± 0.3</td>
<td>5.2 ± 0.4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>153</td>
<td>11 ± 0.5</td>
<td>1.4 ± 0.4</td>
<td>6.6 ± 0.3</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$^a$18 MΩ, $^b$0.01 M HEPES, 0.1 M LiClO$_4$. $^c$Standard deviation.

VIII. Computational Studies with analogs of pHGABA (122-131) and pHGlu (143-145).

Several computational studies were conducted with derivatives of pHGABA (122-131, 133-134) and pHGlu (143-145) in order to provide insights into relevant
parameters of solubility (ClogP), correlations between structure and relative energies, and electronic parameters such as spin densities and charge densities.

A. Determination of calculated logP (ClogP)\textsuperscript{108} values for 122-131, 133-134 and 143-145. The \textit{log} of the partition coefficient for 1-octanol versus water, \( P \), which represents the ratio of the concentration of a particular substance \( S \) in 1-octanol (o) and water (w) (\( P = [S]_o/[S]_w \)) is a commonly employed indicator of compound hydrophobicity.\textsuperscript{105} This criterion is particularly important in predicting the biological compatibility of the compound, e.g., solubility in lipids, metabolism, receptor interaction, and bioavailability, among others.\textsuperscript{105} 1-Octanol has been traditionally the preferred organic constituent because of its likeness to lipids and its easy access. A positive ClogP value indicates a greater lipophilic bearing, i.e., a higher concentration of the substance in the 1-octanol layer versus the water layer. Using the CLogP\textsuperscript{®} software package, version 3.4\textsuperscript{106} implemented in SYBYL 7.3,\textsuperscript{107} which models these values for ClogP, were calculated for \textbf{122-131, 133-134} and \textbf{143-145}. In addition, ClogP values were determined for the conjugate bases of \textbf{122-131}, since these play a prominent role in many of the substituted pHp derivatives. These are displayed in Table 34. Within this series of compounds, \textit{m-CF}_3-pHP GABA (\textbf{130}) evinced the
Table 34. ClogP values for GABA derivatives (101, 122-131, 133-134), Glu derivatives (102, 143-145) and the conjugate bases of pHP GABA’s 122-131.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>ClogP</th>
<th>ClogP (conjugate base)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parent (101)</td>
<td>1.19</td>
<td></td>
</tr>
<tr>
<td>122</td>
<td>1.06</td>
<td>-2.71</td>
</tr>
<tr>
<td>123</td>
<td>0.81</td>
<td>-3.15</td>
</tr>
<tr>
<td>124</td>
<td>0.73</td>
<td>-2.82</td>
</tr>
<tr>
<td>125</td>
<td>0.8</td>
<td>-2.75</td>
</tr>
<tr>
<td>126</td>
<td>1.05</td>
<td>-2.3</td>
</tr>
<tr>
<td>127</td>
<td>0.55</td>
<td>-3.2</td>
</tr>
<tr>
<td>128</td>
<td>0.76</td>
<td>-2.44</td>
</tr>
<tr>
<td>129</td>
<td>0.66</td>
<td>-2.52</td>
</tr>
<tr>
<td>130</td>
<td>2.02</td>
<td>-1.9</td>
</tr>
<tr>
<td>131</td>
<td>1.93</td>
<td>-1.59</td>
</tr>
<tr>
<td>133</td>
<td>-3.81</td>
<td></td>
</tr>
<tr>
<td>134</td>
<td>-3.38</td>
<td></td>
</tr>
<tr>
<td>Parent (102)</td>
<td>-0.03</td>
<td></td>
</tr>
<tr>
<td>143</td>
<td>-0.21</td>
<td></td>
</tr>
<tr>
<td>144</td>
<td>0.04</td>
<td></td>
</tr>
<tr>
<td>145</td>
<td>0.99</td>
<td></td>
</tr>
</tbody>
</table>

highest ClogP value at 2.02 while the $m,m'$-di-OCH$_3$-pHP GABA (133) the lowest at -3.81; the implications from these data are that 130 is the most lipophilic and 133 the least lipophilic in the series. The extra carboxylic acid moiety that distinguishes $l$-Glu
from GABA was shown to impart drastic hydrophilic changes in the ClogP values; 1.06 to 0.04 for \( m\)-F-pHP, 0.81 to -0.21 for \( o\)-F-pHP, and 2.02 to 0.99 for \( m\)-CF\(_3\)-pHP. The conjugate bases, as anticipated from their charges, demonstrated sharp declines in ClogP values in reference to the protonated 122-131.

**B. Energies and Geometries of Oxyallyl-Phenoxy Triplet Biradicals.** Previous studies by Givens and Wirz\(^7^8\) and Phillips\(^8^2\) revealed a significant role for hydroxylic component of water in the photo-Favorskii reaction. Specifically, water is thought to accept the phenolic proton from the triplet excited state of pHP, which helps drive the detachment of the hydrated nucleofuge, tantamount to a photosolvolysis process for the leaving group release and an extrusion process for the chromophore. In order for these processes to obey the Wigner spin conservation rule, if it were an overall single step for a concerted process, these coupled steps generate at least one product that possesses unpaired spins, here the a phenoxy-oxyallyl triplet biradical (117) is the likely candidate. A posited requirement for nucleofuge departure in this process is an out of plane (OOP) contortion of the acetate moiety relative to the planar, phenoxy ring. This generates an oxyallyl-phenoxy biradical as the extended triplet fragment. To investigate the energetics of the OOP contortions thus determine the associated relative stabilization energies of the variety of substituted triplet oxyallyl biradicals (117), Gaussian 03\(^®\) MP2 calculations with 6-31G* basis set were performed for the biradical component 117 generated from 122-131.\(^1^0^8\) Additionally, the relative energies were determined for spirodienediones (100) that emerged after intersystem
crossing of the biradical triplet to a fleeting singlet followed by collapse to 100. The geometries of both the spirodienone 117 and the 100 are composed of nearly orthogonal oxyallyl and phenoxy constituents. The results from these calculations are listed in Table 35. In all cases, the triplet biradical (117) was calculated to be substantially more stable than the corresponding spirodienone by at least 78 kcal/mol. In addition, all precursors displayed an OOP contortion of at least 22°. The

Table 35. The out of plane (OOP) contortion angles and relative energies of the oxyallyl-phenoxy triplet biradicals (117) of 122-131.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>Singlet(^a) (Hartrees)</th>
<th>Triplet(^b) (Hartrees)</th>
<th>(\Delta(T-S)^c) (Hartrees)</th>
<th>(\Delta(T-S)^c) (kcal/mol)</th>
<th>Twist Angle</th>
</tr>
</thead>
<tbody>
<tr>
<td>122</td>
<td>-554.847</td>
<td>-554.970</td>
<td>-0.124</td>
<td>-78.068</td>
<td>22.0</td>
</tr>
<tr>
<td>123</td>
<td>-554.848</td>
<td>-554.973</td>
<td>-0.125</td>
<td>-78.392</td>
<td>26.4</td>
</tr>
<tr>
<td>124</td>
<td>-653.685</td>
<td>-653.812</td>
<td>-0.126</td>
<td>-79.252</td>
<td>25.3</td>
</tr>
<tr>
<td>125</td>
<td>-653.677</td>
<td>-653.811</td>
<td>-0.134</td>
<td>-83.997</td>
<td>47.5</td>
</tr>
<tr>
<td>126</td>
<td>-653.690</td>
<td>-653.816</td>
<td>-0.125</td>
<td>-78.578</td>
<td>22.3</td>
</tr>
<tr>
<td>127</td>
<td>-653.678</td>
<td>-653.813</td>
<td>-0.137</td>
<td>-86.091</td>
<td>57.2</td>
</tr>
<tr>
<td>128</td>
<td>-752.528</td>
<td>-752.654</td>
<td>-0.126</td>
<td>-79.197</td>
<td>25.0</td>
</tr>
<tr>
<td>129</td>
<td>-851.346</td>
<td>-851.485</td>
<td>-0.139</td>
<td>-86.984</td>
<td>57.8</td>
</tr>
<tr>
<td>130</td>
<td>-791.614</td>
<td>-791.741</td>
<td>-0.127</td>
<td>-79.895</td>
<td>22.8</td>
</tr>
<tr>
<td>131</td>
<td>-866.474</td>
<td>-866.602</td>
<td>-0.128</td>
<td>-80.280</td>
<td>23.1</td>
</tr>
</tbody>
</table>

\(^a\)Spirodienodione. \(^b\)Oxyallyl-phenoxy biradical. \(^c\)Energy difference between the triplet and singlet species.
oxyallyl-phenoxy biradical associated with 2,3,5,6-tetrafluoro-pHP, \textbf{tetraF 117}, exhibited the greatest OOP contortion at 57.8° (Figure 48).

\textbf{Figure 48.} Out of plane orientation of the biradicals of \textbf{tetraF 117}, fluorine (yellow), oxygen (red), carbon (gray), and white (hydrogen).

\[
\Delta G (S-T) = -86.98 \text{ kcal/mol}
\]

\begin{figure}
\centering
\includegraphics[width=0.7\textwidth]{Figure48.png}
\end{figure}

\textbf{C. Spin and Charge Density Calculations.} The spin and charge densities of the oxyallyl-phenoxy transients were also obtained. The spin density is the amount of unpaired electron density\textsuperscript{109} at specific atoms in the radical.\textsuperscript{9} The spin and charge densities were determined by from Gaussian 03\textsuperscript{®} at the B3LYP level using the 6-31G\textsuperscript{*} basis set of the oxyallyl-phenoxy biradicals transients of the \textit{p}-hydroxyphenacyl analogs of \textbf{122}, \textbf{129-131}, \textbf{134}, and \textbf{154} and are represented by \textbf{tetraF 117}, the
oxyallyl-phenoxy triplet biradical from 129, in Table 36. The carbonyl carbon, C9, evinced the lowest spin density (-0.171987) i.e., the atom with the lowest probability on which one of the unpaired electrons would dwell. In contrast, the alpha carbon to

Table 36. Calculated spin and charge densities for tetraF 117.

<table>
<thead>
<tr>
<th>Atom</th>
<th>Type</th>
<th>Spin D</th>
<th>Charge D</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>C</td>
<td>0.25365</td>
<td>0.288447</td>
</tr>
<tr>
<td>2</td>
<td>C</td>
<td>-0.149423</td>
<td>0.305392</td>
</tr>
<tr>
<td>3</td>
<td>C</td>
<td>0.379063</td>
<td>-0.100035</td>
</tr>
<tr>
<td>4</td>
<td>C</td>
<td>-0.14822</td>
<td>0.287707</td>
</tr>
<tr>
<td>5</td>
<td>C</td>
<td>0.265947</td>
<td>0.290756</td>
</tr>
<tr>
<td>6</td>
<td>C</td>
<td>-0.087923</td>
<td>0.305876</td>
</tr>
<tr>
<td>7</td>
<td>O</td>
<td>0.409811</td>
<td>-0.440011</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>0.03294</td>
<td>-0.239243</td>
</tr>
<tr>
<td>9</td>
<td>C</td>
<td>-0.171987</td>
<td>0.408392</td>
</tr>
<tr>
<td>10</td>
<td>O</td>
<td>0.419314</td>
<td>-0.409834</td>
</tr>
<tr>
<td>11</td>
<td>F</td>
<td>0.031952</td>
<td>-0.23914</td>
</tr>
<tr>
<td>12</td>
<td>F</td>
<td>-0.006142</td>
<td>-0.271304</td>
</tr>
<tr>
<td>13</td>
<td>F</td>
<td>-0.009262</td>
<td>-0.251713</td>
</tr>
<tr>
<td>14</td>
<td>C</td>
<td>0.85803</td>
<td>-0.312481</td>
</tr>
<tr>
<td>15</td>
<td>H</td>
<td>-0.039206</td>
<td>0.194389</td>
</tr>
<tr>
<td>16</td>
<td>H</td>
<td>-0.038543</td>
<td>0.184801</td>
</tr>
</tbody>
</table>

the carbonyl, C14, manifested the highest spin density (0.85803). The oxygen atoms O10 (0.419314) and O7 (0.409811) conveyed nearly equivalent, significant probabilities for the location of an unpaired electron. The most favored (lowest energy) biradical distribution would place one electron at C14 and the other electron at O10 or O7 or at C3 and O10 (Figure 49).
**Figure 49.** Predicted radical positions for the tetraF 117, from MP2 calculations at the B3LYP level using the 6-31G* basis set.

**IX. Laser Flash Photolysis Studies of pHG GABA derivatives (101, 122, 125, 128-131), 5-acetylsalicylic acid GABA (153), and 5-acetyl-methylosaliclylate GABA (154).**

To obtain excited state parameters for 122, 125, 128-131, the unsubstituted parent 101, and 153, such as transient absorptions and rates of decay, laser flash photolysis (LFP) studies were conducted through a long standing collaboration with the Wirz group at the University of Basel. By pump-probe optical spectroscopy with a pump wavelength of 266 nm (see Experimental), bands at 320, 420, and 440 nm were manifested for some derivatives and assigned to the appearance of the triplet oxyallyl phenoxy biradical (Table 37). The rate constants for intersystem crossing for all derivatives were consistently ~10^{-11} s^{-1}, and the triplet decay rate constants clustered around, ~10^8 s^{-1}. The time to reach maximum absorption for the triplet state, however, was variable (5 ps for 2,3,5,6-tetraF-pH GABA (129) to 174 ps for m-OCF_3-pH GABA (131) in water). Three dimensional, time dependent full spectral profiles for
excitation and decay from LFP $m$-F-pHP GABA (122) in aqueous CH$_3$CN are illustrated in Figures 50 and 51.

Table 37. Pump probe spectroscopy results from 101, 122, 125, 128-131, and 153 in water and mixed media.

<table>
<thead>
<tr>
<th>Cmpd</th>
<th>$k_1$ (ISC, s$^{-1}$)$^a$</th>
<th>$k_2$ ($\tau^3$ decay, s$^{-1}$)$^b$</th>
<th>$k_3$ ($\tau^3$ decay, s$^{-1}$)$^b$</th>
<th>$\Phi_{diss}$, s$^{-1}$)$^c$</th>
<th>340 nm decay (s$^{-1}$)$^d$</th>
<th>$\tau^3$ max. (ps)$^e$</th>
<th>$\tau^3$</th>
<th>Biradical decay (s$^{-1}$)$^e$</th>
</tr>
</thead>
<tbody>
<tr>
<td>101$^h$</td>
<td>4.4E+11</td>
<td>2.90E+09</td>
<td>5.8E+08</td>
<td>1.3E+08</td>
<td>70</td>
<td>400, 510</td>
<td>6.4E+08</td>
<td></td>
</tr>
<tr>
<td>122$^i$</td>
<td>3.6E+11</td>
<td>1.60E+09</td>
<td>2.6E+08</td>
<td>1.6E+08</td>
<td>100</td>
<td>394</td>
<td></td>
<td></td>
</tr>
<tr>
<td>125$^h$</td>
<td>2.60E+09</td>
<td>2.60E+09</td>
<td>5.7E+08</td>
<td>73</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>125$^i$</td>
<td>1.60E+09</td>
<td>3.5E+08</td>
<td>4.0E+07</td>
<td>89</td>
<td>400</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>128$^h$</td>
<td>8.30E+08</td>
<td>6.6E+07</td>
<td>73</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>128$^i$</td>
<td>3.9E+11</td>
<td>9.00E+08</td>
<td>7.2E+07</td>
<td>80</td>
<td>398, 391</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>129$^i$</td>
<td>3.40E+09</td>
<td>3.7E+08</td>
<td>390, 400</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>130$^h$</td>
<td>2.60E+09</td>
<td>4.4E+08</td>
<td>5</td>
<td>410, 400</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>130$^i$</td>
<td>3.0E+11</td>
<td>2.60E+09</td>
<td>4.4E+08</td>
<td>60</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>131$^h$</td>
<td>2.20E+09</td>
<td>2.0E+08</td>
<td>2.1E+08</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>131$^i$</td>
<td>1.40E+09</td>
<td>1.3E+08</td>
<td>6.5E+06</td>
<td>174</td>
<td>397</td>
<td>6.5E+06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>153$^h$</td>
<td>9.90E+08</td>
<td>9.9E+07</td>
<td>397</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>153$^i$</td>
<td>2.1E+11</td>
<td>7.00E+08</td>
<td>7.0E+07</td>
<td>1.0E+06</td>
<td>422</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>154$^h$</td>
<td>1.30E+09</td>
<td>1.30E+08</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>154$^i$</td>
<td>3.2E+11</td>
<td>1.30E+09</td>
<td>1.0E+06</td>
<td>140</td>
<td>407</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$^a$Intersystem crossing. $^b$Triplet decay. $^c$Rate constant for GABA release
$^d$Rate of unknown transient decay. $^e$Time to reach triplet $\lambda_{\text{max}}$. $^f$Triplet
$^g$Decay at 420 and 445 nm. $^h$Water. $^i$Water with unknown amount of CH$_3$CN

Following excitation, the initially formed singlet at 319 nm rapidly undergoes intersystem crossing to the triplet as observed by the decay of the singlet band at 319 nm within 3 ps (the singlet maximum was observed at ~ 1 ps). The rise of the triplet band at 394 nm, reaches its maximum at ~10 ps. The triplet decay occurs with a rate constant of $1.6 \times 10^9$ s$^{-1}$. In some instances a transient is detected at $\lambda = 340$ nm that
Figure 50. Three dimensional, time resolved absorption contour of 122 in aqueous CH$_3$CN.

Reaches a maximum at 20 ns and subsequently decays with a rate constant, 1.6 x 10$^8$ s$^{-1}$. A part of this transient has tentatively also been assigned to the triplet oxyallyl-phenoxy biradical, m-F 117. Other absorptions of transients at 420 and 440 nm were revealed for several other precursors and were also assigned to the transient 117. Further elaboration of these findings can be found in the Discussion section.
Figure 51. Three dimensional triplet decay contour of 122 in aqueous CH$_3$CN.

A cross section of this diagram demonstrates the triplet of 122 at 394 nm (green), an unassigned band at 420 nm (red), and unknown transient at 340 nm (violet) (Figure 52).
**Figure 52.** Cross section of the three dimensional, time resolved absorption contour of 122 in aqueous CH$_3$CN.

Triplet decay parameters for 101 were also obtained from LFP in aqueous CH$_3$CN. The three dimensional contour is presented in Figure 53. The triplet displayed two $\lambda_{\text{max}}$ at 400 nm and 500 nm, the former much stronger than the latter, which reached their maximum absorption maxima at 8 ps and decayed with rate constants of 2.6 x $10^9$ s$^{-1}$. 
**Figure 53.** Three dimensional, time resolved triplet decay contour of 101 in aqueous CH$_3$CN.

Consistent with 122, the emergence of a band at 340 nm was observed, which decayed with a rate constant of $1.3 \times 10^8$ s$^{-1}$. Corresponding one dimensional plots represent triplet decays at both wavelengths, 400 nm in Figure 54 and 500 nm in Figure 55. The curve fit is represented by the green line, derived from data points that are in red.
Figure 54. Triplet decay profile of 101 at 400 nm in aqueous CH₃CN.

Figure 55. Triplet decay profile of 101 at 500 nm in aqueous CH₃CN.
For comparison, the triplet decay profiles of 2,3,6-trifluoro-pHP GABA (128) from LFP in aqueous CH$_3$CN are included. The three dimensional contour of 128 is shown in Figure 56. Evidently, the band at 340 nm is significantly diminished compared to other pHP GABA derivatives.

**Figure 56.** Three dimensional, time resolved triplet decay contour of 128 in aqueous CH$_3$CN.

Two triplet wavelengths were determined at 391 and 398 nm, the former much stronger in intensity than the latter, which reached their maximum absorbance at 80 ps and decayed with a rate constant of $8 \times 10^8$ s$^{-1}$. A cross section of this triplet decay plot is exhibited in Figure 57. In this representation, the unknown transient at with
\( \lambda_{\text{max}} \) at 340 nm is readily apparent. Both triplet bands at 391 and 398 nm are clearly depicted as well. The corresponding one dimensional triplet decay of the band at 391 nm is presented in Figure 58. The green line represents the curve fit of the data points, which are in red.
Figure 58. Triplet decay plot of 128 in aqueous CH$_3$CN.

The triplet decay profiles for 5-acetysalicylic acid GABA (153) in water were obtained by LFP. A cross section of the three dimensional plot is represented in Figure 59.
The triplet at 422 nm reaches its maximum absorption at 100 ps and decays with a rate constant of $6.0 \times 10^8$ s$^{-1}$. Similar to the pHGABA esters, a band emerges at 340 nm that decays, however, with a markedly decelerated rate constant, $1.0 \times 10^6$ s$^{-1}$, versus unsubstituted pHGABA, $1.3 \times 10^8$ s$^{-1}$, a 2 order of magnitude difference. The corresponding one-dimensional triplet decay profile for 153 in water is depicted in Figure 60.
LFP studies were carried out with the 5-acetyl-methylsalicylate GABA (154) in aqueous CH$_3$CN. The absorption profile (Figure 61) revealed the singlet (red) at 326 nm which was seen to decay to two triplet bands at 330 and 396 nm.
A short time later, a cross section (Figure 62) revealed the emergence of the commonly detected 340 nm band (violet), an unassigned band (red), and the triplet of 154 at 416 nm, which reaches its maximum at 140 ps.
**Figure 62.** Cross section of the three dimensional absorption profile of 154 in aqueous CH$_3$CN.

Lastly, a one dimensional decay plot of the triplet at 416 nm (Figure 63) displayed the rate of decay as $1.3 \times 10^9$ s$^{-1}$. 
Figure 63. Triplet decay profile of 154 at 416 nm in aqueous CH$_3$CN.
A. Synthetic Strategies

The methodology used to synthesize pHGABA esters 122-131 was outlined in Scheme 38 (Introduction) The synthesis of m-fluoro pHGABA (122).

Scheme 38. (Introduction) The synthesis of m-fluoro pHGABA (122).

in Scheme 38 (Introduction) with m-F-pHP GABA 122. Briefly, the commercially available 4-bromo-2-fluorophenol (122a) was benzylated with benzyl bromide in an excess of potassium carbonate to afford 122b that then underwent a transmetallation
reaction with a slight excess of tributyl(1-ethoxyvinyl)tin (1.1 equivalents), catalyzed by Pd(0), to afford the acetylated derivative (122c).

It should be noted that heretofore, this Stille protocol for acetylation had not received any use with phenols. In our study, the Stille coupling is an essential step in the synthesis of ketones 122-131. Several other acylation procedures were initially attempted, including Friedel Crafts acylation with acetyl chloride and aluminum trichloride, the Fries rearrangement with an acetyl-substituted phenol in the presence of aluminum trichloride, and lastly, transmetallation with n-butyl lithium followed by addition of acetyl chloride. None of these methods provided yields of acetylated targets in excess of 20%.

Several modifications of the Stille protocol were made to improve the acetylation. Palladium (0) as the tetrakistriphenylphosphine complex, replaced the Pd(II) bischlorotriphenylphosphine complex (we thank Professor Helena Malinakova for this suggestion as well as advice on Stille protocols). The former is air sensitive and oxidizes rapidly, whereas the latter is stable. Presumably, when the latter is dissolved, prompt reduction to Pd (0) occurs from chloride departure, activating palladium for oxidative addition with the aryl bromide. When employed for this purpose with 122, for example, the color turned from yellow to black within 15 minutes, even after 30 minutes of solution degassing, and the yields of 122c never surpassed 10%. Palladium black, as it is commonly known, is an inactive form of palladium.
Another modification involved the addition of saturated aqueous KF to the solution mixture after acetylation. The byproduct of transmetallation, tributyltin bromide, is a liquid that proved difficult to separate from the target compound. Column chromatography with a broad range of eluents failed to completely remove this material. However, in the presence of KF, fluoride exchange readily occurred with bromide to generate tributyltin fluoride, a white, insoluble precipitate that was easily removed by filtration and did not react with the target. The use of Pd(0) as the catalyst for acetylation and fluoride for purification permitted the yields of 122-131c to routinely surpass 80%.

Once 122c (Scheme 38) was synthesized, supervening α-bromination was accomplished with dioxane-dibromide, DDB, in 1 h to afford 122d in high yield that subsequently underwent facile S_N2 substitution with N-Boc protected GABA to generate 122e. At this point, two divergent approaches were used. Early synthetic strategies, including the preparation of the p-hydroxyphenacyl GABA’s, utilized distilled TFA over 24 h to eliminate both the benzyl and Boc protecting groups. Though this approach was effective in generating the fluorinated targets 122-131, oftentimes, wet, adhesive precipitates resulted that necessitated drying under reduced pressure for more than a week. A later strategy was to first debenzylate with Pd/C and H₂ in ethyl acetate, which was quick (2 to 4 h) and efficient (>95% without the need for further purification), followed by Boc deprotection with TFA:CH₂Cl₂ (1:1) for 10-15 minutes. After lyophilization the products were generally flaky, white precipitates.
The synthetic approach for pHG GABA esters (132-141), 4-fluorophenacyl GABA (142), pHG DOC (150), and α-alkyl-pHG diastereomers (166 and 167) were patterned after that for 122 but did not require the use of the Stille protocol, as the corresponding p-hydroxyacetophenones and 4-fluoroacetophenone (142) were commercially available. In the cases where there were no protecting groups on the substrate, such as DOC for 150 and enantiomers for 166 and 167, the last step in the sequence involved Pd/C mediated hydrogenation.

The highlight for the synthesis of pHG diethyl phosphonate (151) entailed the Arbuzov reaction. Here, the benzyl protected, α-brominated 146c was refluxed neat in triethylphosphite over 17 hours to afford the phosphonate that was easily reduced to the corresponding phenol 151 via Pd/C mediated hydrogenation.

The key step in the synthesis of pHG F (152) consisted of a SN2 reaction between 146c and CsF in refluxing acetonitrile for 3 h to afford 152 after deprotection.

B. General photochemistry of pHG and 5-acetylsalicylic acid caged derivatives

The general photochemistry of pHG caged compounds was summarized in Scheme 37 in the Introduction.

The results from photolysis of pHG GABA and Glu esters, the pHG caged amino acids, deoxycholic acid, and diastereomers at 300 nm in aqueous media were consistent with past studies; the only photo products observed were liberated
substrate and rearranged $p$-hydroxyphenylacetic acids as major products, along with minor $p$-hydroxybenzyl alcohols.

The $p$-hydroxybenzyl alcohols had been overlooked previously. In current studies, however, a small, signature band at $\delta \sim 4.5$ ppm was universally discerned in the $^1$H NMR spectra following photolysis and was verified in several cases by spiking the product mixture with an authentic sample of the corresponding $p$-hydroxybenzyl alcohol. Certain derivatives were found to be unreactive, however. These will be discussed later.

The photochemistry of $m$-methoxy derivatives 133 and 134 were re-examined. The photoproducts from these, in addition to GABA, had been reported to be predominantly $m$-methoxy-$p$HP-acyloin and $m$-methoxy-$p$-hydroxyacetophenones. The rearranged $p$-hydroxyphenylacetic acid was said to be minor or non-existent. When re-evaluated, the primary photoproduct obtained for each was, in fact, the rearranged $p$-hydroxyphenylacetic acid; this was evidenced by the intense $^1$H NMR signal at $\delta \sim 3.7$ ppm (see Figure 22 in Results). The acyloin and acetophenone byproducts were also observed at $\delta = 4.99$ and $\delta = 2.60$ ppm, respectively, however only as minor photoproducts, along with the recently discovered $p$-hydroxybenzyl alcohol at $\delta = 4.60$ ppm. Nevertheless, the discrepancies in photoproduct distributions for 133, 134, and a few others were the first indicators of possible substituent-directed mechanistic changes in substrate photorelease from pH.

The photochemistry of 5-acetylsalicylic acid GABA (153), 5-acetyl-methylsalicylate GABA (154), 5-acetylsalicylic acid alkanoyl acids (155 and 156),
and antennae compounds (157-165) produced primary photoproducts, besides the liberated substrate, that were consistent in chemical shift with the corresponding p-hydroxyphenylacetic acid (δ = 3.64 ppm, Figure 35 in Results). In addition, minor resonance signals were generally observed at δ ~ 4.50 ppm that approximated those for p-hydroxybenzyl alcohol.

C. Preliminary Mechanistic Studies

1) Excited state assignments for pHGABA esters. Stern-Volmer quenching studies were conducted with 1-10 mM of pHGABA esters (101, 122-131, and 137-139) using 0-100 mM of potassium sorbate as a triplet quencher in water, in pH 7.3 buffer and the reactions were analyzed by HPLC (see Table 33 and Figure 46 in Results). Significant drops in the quantum efficiencies were observed with increasing sorbate concentration regardless of the medium, providing good empirical evidence for a reactive triplet excited state for all of these pH derivatives. The m-methoxy and m,m'-dimethoxy pHGABA derivatives (134 and 133, respectively) had been analyzed previously97,111 and manifested a prevalent reactive triplet state.

In addition to S-V quenching, LFP studies were conducted on the fluorinated pHGABA esters in both aqueous CH₃CN and water. Absorbance profiles revealed rapid intersystem crossing from the singlet to triplet states (~10¹¹ s⁻¹) with efficiencies near unity. Degassing experiments failed to manifest any changes in the quantum efficiencies, due to the rapid decay rate (10⁷-10⁹ s⁻¹) of the pH triplet and the inherently low concentration of O₂ in water (1.28 mM).112
In accord with previous findings by Givens and Wirz,\textsuperscript{78,92} and Phillips,\textsuperscript{80-82,86-88,99,91} the reactive excited state for these pHP GABA esters is unequivocally the triplet. In 2008, Givens and Wirz\textsuperscript{92} suggested the possibility of an adiabatic nucleofuge disjunction (Scheme 37) from the observation of a series of new bands at 330, 420 and 445 nm derived from LFP of pHP diethyl phosphate, mesylate, and tosylate, in aqueous CH\textsubscript{3}CN. An oxyallyl-phenoxy triplet biradical was assigned to the transient intermediate based on these absorptions. The same bands were consistently revealed for the other LFP probed pHP GABA esters, suggesting a common triplet oxyallyl-phenoxy biradical intermediate to the entire series.

2) \textbf{Excited state assignments for 5-acetylsalicylic acid GABA and its esters.} The sole compound probed by S-V quenching analysis and LFP studies was 5-acetylsalicylic acid GABA (153). The carbomethoxy analog (154) was analyzed by LFP studies and was reported previously as exhibiting an operative triplet state.\textsuperscript{113} As shown in the Results section, 153 was readily quenched by potassium sorbate in water, manifesting a linear S-V correlation constant, $K_{SV}$. LFP studies in both water and aqueous CH\textsubscript{3}CN confirmed this and clearly showed the singlet rapidly crossing to the triplet, $k = 2.1 \times 10^{11}$ s\textsuperscript{-1}. The triplet maximum occurred at $\lambda = 422$ nm within 100 ps and decayed with a rate constant of $6.0 \times 10^{8}$ s\textsuperscript{-1}. These findings confirm the triplet as the reactive state. Additionally, short-lived absorption bands at 330, 420, and 445 nm further substantiated an triplet oxyallyl-phenoxy biradical intermediate. The remaining 5-acetylsalicylic acid derivatives (155-165) were not probed by S-V quenching or LFP, but, nonetheless, are anticipated to react from the triplet state.
Scheme 52 Preliminary mechanism of the photo-Favorskii rearrangement, with 153

[Diagram showing the mechanism with reactions and intermediates labeled]
3) **A preliminary photo-Favorskii mechanism.** Integrating the empirical data obtained from $^1$H NMR photochemical analyses in D$_2$O, which revealed the rearranged $p$-hydroxyphenylacetic acid and released substrates as major products in near quantitative yields and a trace amount of $p$-hydroxybenzyl alcohol, with the S-V quenching and LFP established functional triplet state and a probable oxyallyl phenoxy triplet biradical that leads to the rearranged and decarbonylation product, a preliminary photo-Favorskii mechanism is posited (Scheme 52, with 153).

The process is as follows: Initial excitation of 153 to the singlet excited state is followed by rapid ($2.1 \times 10^{11}$ s$^{-1}$) intersystem crossing to the triplet, (153$^3$). Significant H-bonding interactions with water assist the complex in skewing the carbonyl OOP with respect to the ring, a necessary stereoelectronic arrangement for facile substrate release. In this manner, the triplet undergoes solvent assisted deprotonation of the phenol with concomitant substrate departure ($6.0 \times 10^8$ s$^{-1}$), forming a triplet oxyallyl-phenoxy biradical that has an orthogonal OOP orientation.

DFT calculations of 101, 122-131 (Table 35, Results) confirmed the OOP geometry of biradical and also predicted this to be much lower in energy than the corresponding ground state spirodieneone. Once formed, the biradical then rapidly relaxes with intersystem crossing and collapses to the spirodiendione. The completion of the route to the major, $p$-hydroxyphenylacetic acid product is through water-mediated fragmentation of the spirodiendione; the minor $p$-hydroxybenzyl alcohol is formed by an initial CO extrusion from the spirodiendione to form the $p$-quinone methide, which then undergoes conjugate addition of H$_2$O.
4) **Alternative mechanisms and photoproducts** The \( m \)-methoxy and \( m,m' \)-dimethoxy pHP GABA derivatives (134 and 133, respectively) were the only derivatives of pHP that manifested other exiguous photoproducts, such as \( p \)-hydroxyacetophenone (115) and \( \alpha \)-hydroxy acetophenone, an acyloin derivative (\( \alpha \)-OH-115) (Figure 64). It is obvious that divergent mechanisms are operative in these.

**Figure 64.** Photoproducts of 134 a) \( m,m' \)-diOCH\(_3\)-115, b) \( m,m' \)-diOCH\(_3\)-\( \alpha \)-OH-115 and 133 c) \( m \)-OCH\(_3\)-115, d) \( m \)-OCH\(_3\)-\( \alpha \)-OH-115.

Yousef\(^{111}\) previously explored the photochemistry of 134 in aqueous media and had noted the appearance of these photoproducts. His mechanistic interpretations were congruent with results from disparate studies of Sheehan,\(^{51}\) and Pincock\(^{114}\) (Scheme 53 with 133). The photoconversion course is proposed as follows: Initial excitation of 133 to the singlet state is followed by rapid ISC, \( k = 2.7 \times 10^{11}\text{ s}^{-1} \) to the triplet (133\(^3\)), which undergoes homolytic scission to produce the geminate radical pair (133\(^g\)). The geminate radical pair can recombine to reproduce 133, or can diverge into two disparate mechanisms. 1) Electron transfer occurs between the radicals to generate a geminate ion pair (133\(^1\)), which can regenerate 133 by recombination (not
**Scheme 53** Modified version of Yousef's proposed discrete mechanistic pathways to \(m,m'-\text{diOCH}_3\)-115 and \(m,m'-\text{diOCH}_3\)-\(\alpha\)-OH-115

\[
\begin{align*}
\text{H}_3\text{CO} & \quad \text{OH} & \quad \text{OCH}_3 \\
\text{O} & \quad \text{O} & \quad \text{CO} & \quad \text{NH}_3^+ \\
\text{H}_3\text{CO} & \quad \text{OH} & \quad \text{OCH}_3 \\
\text{O} & \quad \text{O} & \quad \text{CO} & \quad \text{NH}_3^+ \\
\text{H}_3\text{CO} & \quad \text{OH} & \quad \text{OCH}_3 \\
\text{O} & \quad \text{O} & \quad \text{CO} & \quad \text{NH}_3^+ \\
\text{H}_3\text{CO} & \quad \text{OH} & \quad \text{OCH}_3 \\
\text{O} & \quad \text{O} & \quad \text{CO} & \quad \text{NH}_3^+ \\
\text{H}_3\text{CO} & \quad \text{OH} & \quad \text{OCH}_3 \\
\text{O} & \quad \text{O} & \quad \text{CO} & \quad \text{NH}_3^+ \\
\end{align*}
\]

\(133^1 \xrightarrow{\text{ISC}} 133^3 \)

\(k = 2.7 \times 10^{11} \text{s}^{-1}\)

solvent separated radical pair

geminate radical pair

geminate ion pair

electron transfer

solvent mediated hydrogen atom abstraction

\(m,m'-\text{dimethoxy-115}\)

GABA

\(m,m'-\text{dimethoxy-\(\alpha\)-OH-115}\)

solvolyis

\(\text{\(\alpha\)-OH-115}\)}
shown) or solvent separate and hydrolyze to afford GABA and \( m,m'-\text{diOCH}_3-\alpha-\text{OH} \) (115). Pincock\(^{114}\) demonstrated this route was feasible from photolyses of a series of arylmethylacetates in methanol. 2) A second possible mechanism manifests solvent incited separation of the geminate radical pair to engender a solvent separated radical pair, which H atom abstracts from the mixture, analogous to Sheehan’s\(^{51}\) suggested mechanism, from photolysis with \( p \)-methoxyphenacyl benzoates in EtOH, to furnish \( m,m'-\text{diOCH}_3 \) 115 and GABA.

D. Substituent effects on the photo-Favorskii rearrangement

1) Quantum Efficiencies of Fluoro-pHP GABA’s: The role of the conjugate base. The position and extent of fluoro substitution on the pHP chromophore was found to manifest stark changes in the quantum yields for GABA release, \( \Phi_{\text{dis}} \), in aqueous solutions. The quantum yields, determined at 300 nm, for GABA and \( p \)-hydroxyphenylacetic acid appearance were nearly identical to \( \Phi_{\text{dis}} \) in all cases, thus \( \Phi_{\text{dis}} \) will be principally considered. In water (Table 22, Results), \( m \)-F-pHP GABA (123 (\( \Phi_{\text{dis}} = 0.28 \))), 2,3-diF-pHP GABA (124 (\( \Phi_{\text{dis}} = 0.24 \))), and 2,5-diF-pHP GABA (125 (\( \Phi_{\text{dis}} = 0.22 \))) were the only derivatives that demonstrated improved yields relative to the unsubstituted parent (101 (\( \Phi_{\text{dis}} = 0.20 \))), and \( o \)-F-pHP GABA (122 (\( \Phi_{\text{dis}} = 0.16 \))) and 2,6-diF-GABA (127 (\( \Phi_{\text{dis}} = 0.16 \))) were nearly the same as 101. The common factor for these five was the \( \lambda_{\text{max}} \sim 280 \) nm which indicated the protonated pHP dominated. In contrast, 3,5-diF-GABA (126 (\( \Phi_{\text{dis}} = 0.11 \))), 2,3,6-triF-GABA (128 (\( \Phi_{\text{dis}} = 0.08 \))), and 2,3,5,6-tetraF-GABA (129 (\( \Phi_{\text{dis}} = 0.11 \))), evinced
the predominant conjugate base of the chromophore, which absorbs at $\lambda_{\text{max}} > 300$ nm. In buffer pH 7.3 (Table 23, Results) all fluoro pHp derivatives manifested predominant conjugate bases, except 123, which exhibited both the conjugate base and protonated forms with absorptivities nearly the same. Accordingly, $\Phi_{\text{dis}}$ at this pH were lower for all fluoro pHp GABA esters, e.g., 124 ($\Phi_{\text{dis}} = 0.11$) and 125 ($\Phi_{\text{dis}} = 0.10$), while that for 123 dipped only slightly, ($\Phi_{\text{dis}} = 0.21$).

The unsubstituted parent, 101, which was predominantly protonated, evinced a constant $\Phi_{\text{dis}} = 0.21$. At pH 8.2, all fluoro pHp GABA’s were conjugate bases and all manifested low $\Phi_{\text{dis}}$. Conspicuously, the pH of the media impacts the efficiency of photoconversion 122-131 by controlling the equilibrium between the conjugate bases and protonated forms; thus, there exists a clear correlation between the pKa of the pHp and $\Phi_{\text{dis}}$. In water, pH = 6.5, those derivatives with pKa’s 5.5 and lower (Table 15 Results) 126, 128, 129, 130, conveyed the preponderant conjugate base, with accompanying lower $\Phi_{\text{dis}}$. Those with pKa’s between 5.5 and 6.7 showed both forms, i.e., 122, 124, 125, 131, while those with pKa’s 6.8 and higher, 123, and 127, predominantly the protonated form.

Two interesting trends are noticeable from comparisons of fluoro pHp GABA’s to the unsubstituted pHp GABA (101). First, those with a single ortho fluorine, i.e., 123, 124, and 125 with pKa’s 7.2, 5.9, and 5.7 respectively, exhibited higher $\Phi_{\text{dis}}$ than unsubstituted 101 in water, with no control on the pH. A sufficient explanation for this ortho-effect is not available. Second, the $\Phi_{\text{dis}}$ for $m$-OCF₃-pHP GABA (131 (0.09)) was double that of its proton counterpart $m$-OCH₃-pHP GABA
(134 (0.04)). A possible rationalization for this might be due to the reduced electron donating capacity of the -OCF₃ vs. -OCH₃, an influence that Givens, et. al., had previously reported.¹⁰⁴

Attempts to discern linear correlations between pKa and Φ_{dis} in water (blue) and Φ_{dis} at pH 7.3 (pink) (Figure 65) showed little difference between the two sets of data. However, the notion of discrete differences in the branching between decay of the excited state and progress towards release for the protonated versus conjugate bases is consistent with our earlier conclusion. The efficiency for photorelease is apparently independent of the pKa of the chromophore within the narrow range exhibited in this fluorinated pHp GABA series.

**Figure 65.** Plot of pKa of fluoro pHp GABA’s vs. Φ_{dis} in water (blue diamonds) and buffer pH 7.3 (pink squares). Unsubstituted pHp GABA (101) is included for comparison.
The singlet excited state of the conjugate base also rapidly intersystem crosses to the triplet state, \( k_{\text{ISC}} = 3.0 \times 10^{11} \text{ s}^{-1} \). A small fraction of the singlet also deactivates by fluorescence (\textit{Results}, Table 20). The triplet deprotonation step, deemed to be crucial in initiating the photo-Favorskii rearrangement, is obviated from the singlet. These two distinctions separate the conjugate base pathways from the triplet reaction of the protonated pHp. However, a rapid deprotonation of the singlet excited protonated pHp is precluded as there is no fluorescence observed from this species. The diminished \( \Phi_{\text{dis}} \) for the conjugate bases must result from a more efficient decay pathway. Since the products are the same, an operative photo-Favorskii rearrangement mechanism is the likely pathway.

2) Rate Constants of Fluoro pHp GABA esters. The rates constants, \( k_r \), obtained from S-V quenching for GABA release, disclosed marginal differences in aqueous media. From Table 33, \( k_r \) spanned from \( 1.8 \times 10^7 \text{ s}^{-1} \) for 2,3,6-trifluoro-pHP GABA (128) to \( 1.6 \times 10^8 \text{ s}^{-1} \) for \( \alpha \)-F-pHP GABA (123) in water and \( 1.1 \times 10^7 \text{ s}^{-1} \) for 3,5-diF-pHP GABA (127) to \( 8.9 \times 10^7 \text{ s}^{-1} \) for 123 in pH 7.3 buffer. These essentially identical rapid rates parallel the nearly constant quantum efficiencies. The lack of change in the rate constants as might be anticipated in other linear free energy relationships such as Hammett \( \sigma-\rho \) correlation militates against an ionic charge buildup in the transition state for the release of the substrate. At first pass, the result might appear implausible since the fragmentation step for covalent bond breaking results in the release of the carboxylate ion rather than a carbonyl radical. The latter has been ruled
Figure 66. A. Plot of rate constants for release, $k_r$, of pHP GABA’s vs. pKa’s in water. B. Plot of rate constants for release, $k_r$, of pHP GABA’s vs. pKa’s in pH 7.3 buffer.

out due to the absence of radical decarboxylation processes in the photo-Favorskii reaction (see Introduction). Attempts to correlate $k_r$ with corresponding pKa’s in water and pH 7.3 buffer demonstrated no association (Figure 66).
3) **Comparisons of the Stern-Volmer derived rate constants with the LFP values.** Rate constants obtained independently from LFP (Table 37) were, in general, higher by up to one order or magnitude relative to those obtained by S-V quenching studies. The triplet lifetimes obtained by LFP are accordingly shorter than those obtained by S-V quenching with sorbate. The origin of this effect may be due to the method, i.e., the nature of the quencher and estimated rate of diffusion used in calculation of the rate constants. For example, Givens and Wirz,\(^92\) and Phillips\(^81\) showed that added organic solvents greatly increase the lifetime of the pHp triplet. The diffusion rate constant may be an overestimate. As sorbate is added in a S-V run, the “organic nature” of the solvent will increase, lengthening the lifetime of pHp triplets. This renders them more susceptible to bimolecular quenching and gives the appearance of a slower competing rate for the reaction out of the triplet manifold. A second factor may be the relative protonation state of the pHp chromophore. A slower quenching diffusion rate for the conjugate base will have a corresponding effect on the S-V constant, \(K_{SV}\), which will consequently result in a lower estimated lifetime of its triplet.

4) **The role of H\(_2\)O.** As shown previously by Givens\(^78\) and Phillips\(^45\), the triplet pHp ester is influenced by water, primarily assisting in the removal of the phenolic proton but also in assisting the concomitant loss of the leaving group, GABA, through
Scheme 54 Proposed mechanism of GABA photorelease from the conjugate base of 101
solvation. Added organic quenchers will negatively influence the solvation and deprotonation processes causing a decrease in the triplet reactivity.

5) Other sources of variance. An additional difference lies in the excitation source. For Stern-Volmer quenching, analyses were conducted with 300 nm, RPR phosphor coated lamps that span a range from 250 to 350 nm. Both the phenol and its conjugate base, if present, will be excited at longer wavelengths near 300 nm. In contrast, the pump probe spectroscopy studies were conducted with a monochromatic laser excitation of 266 nm having a dynamic range of ± 5 nm. This wavelength would predominantly excite the protonated forms (the absorptivities are very low at 266 nm for the conjugate bases).

6) Photochemistry of the conjugate bases. Photolysis of 122-131 at 350 nm in 0.01 M TRIS pH 9, where the conjugate bases predominate, evinced $\Phi_{\text{dis}}$ that were well below 0.01, ranging from 0.001 (124) to 0.008 (130). These low values indicate that other deactivation pathways are more efficient than GABA photorelease.

7) Proposed mechanism for conjugate base. An alternative mechanism is proposed for the conjugate bases of pH (Scheme 54, vide supra with the conjugate base of pH GABA (101)). Red values highlighted in this mechanism are the result of this work and may differ from the mechanism for protonated pH GABA.

E. Miscellaneous substituent effects on pH GABA ester photochemistry

Several other substituents were explored for their effects on the photochemistry.\textsuperscript{115} The nitro group is notorious for its deleterious effects on ketone
photochemistry, frequently shutting down all photoluminescent and photochemical activity. On the other hand, in an o-nitrobenzyl motif, the hydrogen abstraction photochemistry is the key step for this most popular photo-removable protecting group.\textsuperscript{104} For nitro substituted pHp, m-NO$_2$-pHP GABA (132), the -NO$_2$ moiety extended the $\pi$-$\pi^*$ absorption past 400 nm, but did not lead to GABA release when photolyzed between 254-350 nm in water. Here, the nitro group served as a deactivator.

Ortho and m-OH-pHP GABA’s (135 and 136, respectively) effectively quenched the photorelease of GABA as well. Hydrogen bonding with the excited ketone carbonyl for the ortho OH, a stabilizing interaction, could hinder the stereoelectronic alignment to the necessary OOP rotation angle of the carbonyl for substrate release (Figure 67).

**Figure 67.** Proposed H-bonding between the $\omega$-OH and ketone carbonyl of 135.

The m-OH isomer also failed to effect GABA photorelease in water. H-bonding with the $p$-OH moiety could interfere with the crucial proton transfer step of the substrate release (Figure 68).
The meta-methyl substituted pHp GABA’s manifested $\Phi_{\text{dis}}$ in water of 0.15. A small absorbance band shift to $\lambda_{\text{max}} = 304$ nm for 138 (24 nm) vs. pHp GABA did not change the $\Phi_{\text{dis}}$, 0.15 and 0.20 respectively. At pH 8.2, $\Phi_{\text{dis}}$ drop to 0.08 and 0.11, respectively, congruent with declines previously seen with conjugate bases of all the derivatives.

Interestingly, the $o$-CH3-pHP GABA (139) evinced a 45% decline in $\Phi_{\text{dis}}$ (0.11) compared to the unsubstituted analog (101) (0.20) and a 26% decline compared to meta isomer (0.15). Although the origins of this were not examined, two effects, one steric effect and the other a reversible Norrish Type II hydrogen atom abstraction may interfere with GABA release. The steric effect of CH3 may force the acetyl moiety to adopt a conformation that is less amenable to GABA release. Low level MM2 calculations116 predicted the carbonyl of decaged 139 to adopt an OOP orientation with the ring (angle not specified), whereas the unsubstituted analog 115 was planar (Figure 69). The reversible nature of $o$-arylmethyl ketone photochemistry has been well documented117 and has been pictured.
Figure 69. Lowest energy conformations of decaged 139 and 115 by low level MM2 calculations. The OOP twist angle was not determined.

as an “energy wasting” mechanism, much the same as the alkene cis-trans isomerization process (Equation 16).

Eq. 16
The α-methyl and ethyl substituted pHGABA’s 140 and 141, respectively, showed low $\Phi_{\text{dis}}$, 0.05 and 0.06, respectively. These are secondary α-centers to the ketone, which may alter the stability of the initially formed radical intermediates. The photoproducts included released GABA and the analogous, rearranged p-hydroxyphenylacetic acid.

The impetus for the synthesis and subsequent probing of the photochemistry of 4-fluorophenacyl GABA (142) was based on the well documented $p$ electron donation of fluorine to the $\pi$ aromatic system of aromatic fluorine compounds.\textsuperscript{118} It was reasoned, then, that the $p$-fluoro moiety may be a surrogate for the $p$-OH group in inducing photorelease of GABA and rearranging to the corresponding 4-fluorophenylacetic acid. However, this compound proved to be inert at irradiation wavelengths of 254 and 300 nm.
F. Substituent effects on 5-acetylsalicylic acid GABA derivatives (154, 157-165).

The quantum yield of methyl 5-acetyl-salicylate GABA (154) at 300 nm spanned from $\Phi_{\text{dis}} = 0.08$ to 0.11 between pH 7 to 9, respectively, and the rate constants for GABA release, $k_r = 1.3 \times 10^8$ s$^{-1}$ were equable. These were consistent with 5-acetylsalicylic acid GABA (153). Replacing the methyl with auxochromes (157-165) gave quantum yields between 0.01 and 0.12. Though all of these compounds demonstrated strong absorbances at $\lambda = 350$ nm at 1 mM in water, when irradiated at 350 nm in water, they gave $\Phi_{\text{dis}}$ well below 0.01. An energy threshold for efficient GABA release from the pHp and 5-acetylsalicylic acid phototriggers may have been reached when the triplet energy of the system drops from 70 kcal/mol (pHp) to that effectuated by the auxochrome.

G. Nucleofuge effects on the photo-Favorskii rearrangement. The primary research objectives were on comprehending substituent, pH and media effects on the photo-Favorskii rearrangement. Another crucial aspect, however, concerns the nature of the nucleofuge. Carboxylic acids were emphasized in studies presented in this thesis, i.e., $\gamma$-aminobutyric acid, $l$-glutamic acid, glycine, tyrosine, phenylalanine, valine, deoxycholic acid, and alkanoyl acids.

1) $l$-Glutamate. The $l$-glutamate nucleofuge exhibited the same behavior as GABA.

2) Other amino acids. The amino-acids glycine, valine, phenylalanine, and tryptophan manifested the same behavior as GABA.
3) **Deoxycholic acid.** The quantum yield for the disappearance of pHp DOC (150) was determined to be 0.06 in 5% aq. CH₃OH. DOC is highly lipophilic, contrary to the hydrophilic GABA salt. Solvent assistance, primarily through H-bonding with the nucleofuge, is considered to be central to nucleofuge departure. Though CH₃OH can hydrogen bond, it is less much less efficient than water; less effective solvation could complicate nucleofuge departure, which would direct deactivation to other pathways. However, the rearranged \( p \)-hydroxyphenylacetic acid was observed, suggesting that the operative mechanism is the “photo”-Favorskii pathway.

4) **Alkanoyl acids.** The nucleofuge capacities of dodecanoic and tetradecanoic acids were probed as substrates caged to the 5-acetylsalicylic acid chromophore (155 and 156, respectively). The motivation here involved the application of these as photocleavable surfactants. Caprioli and coworkers had synthesized structurally similar compounds to 155 and reported successful cell membrane protein extraction, improved ionization efficiencies and resolution of these proteins by MALDI analysis and facile surfactant removal by photolysis. Only a handful of photocleavable surfactants have been reported in the literature. The water solubility of 155 and 156 was low. The photolysis of 1 mM solutions of each in 5% aqueous CH₃OH at 300 nm evinced \( \Phi_{\text{dis}} \) of 0.04 and 0.06, respectively. However, 155 and 156 promptly dissolved in 0.01 M HEPES with 0.1 M LiClO₄ at pH = 8.2, suggesting the carboxylic acid was deprotonated. Irradiation of 1 mM of each in this buffer divulged \( \Phi_{\text{dis}} \) that were 0.10 and 0.09, respectively, *commensurate with the GABA derivative*. The improved quantum yields in buffer, substantiate water-assisted substrate
disjunction, i.e., higher water content promotes substrate disjunction because of a higher degree of solvation. This observation corroborates the hydrated triplet complex depicted in Scheme 52. Though the photocleavable surfactant properties of 155 and 156 seem favorable, i.e., each are soluble in pH 8.2 buffered media and exhibit efficient photodegradation, their biological assessments have not been determined.

H. Nucleofuge effects - linear free energy relationships. Most nucleofuges studied for this thesis were carboxylic acids. Dr. Cope, a recent graduate of the Givens group, explored the how the nature of nucleofuge impacts the photo-Favorskii rearrangement by synthesizing and probing the photochemistry of pHp mesylate (168), tosylate (169), benzoate (170a), p-OCH₃-benzoate (170b), p-CF₃-benzoate (170c) and formate (171). Through a combination of Stern-Volmer quenching and

![Chemical structures](image-url)
LFP studies, a *Brønsted* $\beta_{LG}$ (*nucleofuge*), *linear free energy relationship* (*LFER*) was determined.

The nucleofuge capacity has also been determined to correlate with the log of the reaction rate constant (Equation 17), where $\beta_{LG}$ signifies the response of the rate constant to the nucleofuge acidity.

\[
\log(k) = \beta_{LG}(pK_a) + \log(C) \quad \text{Eq. 17}
\]

The nucleofuge capacity is enhanced by a decrease in pKa; thus a negative $\beta_{LG}$ would signify favorable nucleofuge properties. Applied to a linear free energy setting, $\beta_{LG}$ indicates the degree of bond breaking (nucleofuge departure) in the rate determining step.\textsuperscript{120} The rate determining step in the photo-Favorskii is believed to be nucleofuge disjunction, so the acquisition of a $\beta_{LG}$ value is key to an improved comprehension of this step. Parameters of other esters and sulphonates with those of pHGABA, pHG $p$-CN-phenolate\textsuperscript{121} and pHG-phenolate\textsuperscript{121} (for comparison), derived in aqueous CH$_3$CN, are presented in Table 38.
Table 38. Parameters of 168-171 and other pHp derivatives. Adapted and modified from ref.119.

<table>
<thead>
<tr>
<th>pHp derivative</th>
<th>LG pKa</th>
<th>Φ_{dis}</th>
<th>k_{c3} decay (s^{-1})</th>
<th>τ^3 (ns)</th>
<th>log (Φ_{dis}/τ^3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mesylate 168</td>
<td>-1.54</td>
<td>0.932</td>
<td>8.5 x 10^9</td>
<td>0.118</td>
<td>9.70</td>
</tr>
<tr>
<td>Tosylate 169</td>
<td>-0.43</td>
<td>1.04</td>
<td>9.6 x 10^9</td>
<td>0.104</td>
<td>10.0</td>
</tr>
<tr>
<td>Diethyl Phosphate 114</td>
<td>0.71</td>
<td>0.40</td>
<td>2.9 x 10^9</td>
<td>0.345</td>
<td>9.06</td>
</tr>
<tr>
<td>pCF₃ Benzoate 170c</td>
<td>3.69</td>
<td>0.201</td>
<td>1.6 x 10^9</td>
<td>0.625</td>
<td>8.51</td>
</tr>
<tr>
<td>Formate 171</td>
<td>3.75</td>
<td>0.94</td>
<td>1.5 x 10^9</td>
<td>0.667</td>
<td>9.15</td>
</tr>
<tr>
<td>pOCH₃ Benzoate 170b</td>
<td>4.09</td>
<td>0.288</td>
<td>1.2 x 10^9</td>
<td>0.833</td>
<td>8.54</td>
</tr>
<tr>
<td>Benzoate 170a</td>
<td>4.21</td>
<td>0.316</td>
<td>8.6 x 10^8</td>
<td>1.16</td>
<td>8.44</td>
</tr>
<tr>
<td>GABA 101</td>
<td>4.00</td>
<td>0.21</td>
<td>2.9 x 10^9</td>
<td>0.345</td>
<td>8.78</td>
</tr>
<tr>
<td>pCN Phenolate</td>
<td>7.17</td>
<td>0.11</td>
<td>6.9 x 10^8</td>
<td>1.45</td>
<td>7.88</td>
</tr>
<tr>
<td>Phenolate</td>
<td>9.8</td>
<td>0.04</td>
<td>2.6 x 10^8</td>
<td>3.85</td>
<td>7.02</td>
</tr>
</tbody>
</table>

^a log k_r = log(Φ_{dis}/τ^3)

The logs of the rates of disappearance, k_r, were plotted against the pKa of the conjugate acid of the nucleofuge to obtain a slope (β_{LG}) that was -0.2419^{121} (Figure 70). A good correlation between log of k_r and nucleofuge pKa’s was observed, R^2 = 0.9061. The moderate, negative value for β_{LG}, indicates that some bond scission between pHp and the nucleofuge occurs in the transition state of the rate determining step. The pKa of the nucleofuge, one direct measure of leaving group ability, is shown to be a significant factor on determining the rate of nucleofuge detachment. The data points in pink originate from the fluoro pHp GABA’s; these demonstrate that substituent effects are operative in modifying k_r.
Figure 70. A plot of rates of disappearance, \((\Phi_{\text{dis}}/\tau^3)\) vs. the pKa of pH esters, establishing a Bronsted linear free energy relationship. The values in pink represent the fluoro pH GABA series.

\[
\log \left( \frac{\Phi_{\text{dis}}}{\tau^3} \right) \text{ vs. pKa}
\]

\[
y = -0.2419x + 9.5657
\]

\(R^2 = 0.9061\)

I. Media effects on the photo-Favorskii rearrangement.

1) **Background.** The media in which pH cages are photolyzed is crucial to the efficiency of the photo-Favorskii rearrangement. Wan and Corrie,\(^{73}\) Givens and Wirz,\(^{78,92}\) and Phillips\(^{86,88}\) conveyed the pertinence of water to induce the release of phosphate and acetate from the pH chromophore. Phillips,\(^{88}\) accentuated the importance of both the hydrogen bond donor and acceptor nature of water in his modeling of the photo-Favorskii reaction. For example, in 1:1 H\(_2\)O:CH\(_3\)CN, \(\Phi_{\text{dis}}\) for pH diethyl phosphate\(^{114}\) was 0.40. When measured in 1:1 DMSO:CF\(_3\)CH\(_2\)OH, a solvent mixture with a hydrogen bond acceptor (DMSO) and a hydrogen bond donor (CF\(_3\)CH\(_2\)OH), \(\Phi_{\text{dis}}\) reduced to 0.17 but was still efficient. However, when measured
in neat DMSO or CF<sub>3</sub>CH<sub>2</sub>OH, the reaction did not occur. Phillips<sup>88</sup> demonstrated a correlation between triplet lifetime, 3τ, of pHP diethyl phosphate (114) and the water content in CH<sub>3</sub>CN (Table 13); from neat CH<sub>3</sub>CN, where 3τ was 150 ns,<sup>80</sup> the triplet lifetime progressively attenuated to less than 290 ps in 75% aq. CH<sub>3</sub>CN, a factor of 517 decreases in 3τ. Likewise, the 3τ of pHP acetate 110 dropped to a more modest extent, from 137 ns in neat CH<sub>3</sub>CN to 2.1 ns in 50% aq. CH<sub>3</sub>CN, a factor of 65.

The triplet lifetimes are directly related to the rates of substrate release by the equation k<sub>r</sub> = (Φ<sub>dis</sub>3τ). High Φ<sub>dis</sub> have been communicated even at very low concentrations of H<sub>2</sub>O.<sup>78,119</sup> Wan and Corrie<sup>73</sup> ascertained Φ<sub>dis</sub> for pHP acetate (110) as 0.41 in 1:1 H<sub>2</sub>O:CH<sub>3</sub>CN pH 7 and revealed a correlation between media pH and Φ<sub>dis</sub> (Figure 71). At low pH (0-1), Φ<sub>dis</sub> for 110 spanned from 0.12 to 0.32, respectively. At pH 10.5 to 12.5, Φ<sub>dis</sub> spanned from 0.30 to 0 (no acetate appearance), respectively. Earlier, a media pH dependence on Φ<sub>dis</sub> was underscored for fluoro pHGABA’s 122-131, where the ratios of protonated vs. conjugate bases altered the Φ<sub>dis</sub>. Despite solvent modulated triplet lifetimes of pHP, Φ<sub>dis</sub> appeared to remain steady. For example, Givens and Wirz<sup>78</sup> reported a Φ<sub>dis</sub> for pHP diethyl phosphate (114) of 0.40 in 5% aq. CH<sub>3</sub>CN with a triplet lifetime of ~ 9 ns.<sup>88</sup> Phillips<sup>88</sup> reported a Φ<sub>dis</sub> of 0.40 for the same compound 1:1 H<sub>2</sub>O:CH<sub>3</sub>CN, though the triplet lifetime shortened to 0.350 ns. From the equation, k<sub>r</sub> = (Φ<sub>dis</sub>3τ), the rate constants for nucleofuge disjunction change as a function of triplet lifetime. One plausible explanation is that though k<sub>r</sub> vary according to water content, they remain
fast enough to predominate as a major triplet deactivation pathway for pHP versus other deactivation pathways.

**Figure 71.** Titration curves obtained from $^1$H NMR analysis depicting the dependence of quantum yields for acetate appearance, $\Phi_{\text{app}}$ ($\Phi_p$ in plot) on pH for photolysis of pHP acetate (114) in 1:1 CH$_3$CN (circles). The circles represent $p$-methoxyphenacyl acetate. Reproduced with permission from the ACS, ref. 73.

2) **Organic and mixed organic media.** The quantum yields for disappearance, $\Phi_{\text{dis}}$, of pHP GABA’s were assessed in 25% aq. CH$_3$CN, DMSO, 10% aq. DMSO, 1-pentanol, and 1-octanol at 300 nm (Table 24).
a) In 25% aq. CH$_3$CN, the $\Phi$$_{\text{dis}}$ for 101, 122-125 and 127-131 were unchanged from those for 100% water, except the marked increase for 2,6-diF-pHP (126) (0.16 vs. 0.11 in H$_2$O). UV-Vis analysis of 126 in this media revealed two absorbance bands, one at 277 nm and the other at 325 nm, reflecting the presence of both the protonated phenol and conjugate base. Recall that 126 in H$_2$O, only the conjugate base was manifested, at $\lambda_{\text{max}}$ = 331 nm. The implication from these findings is that 25% H$_2$O is sufficient to effectively solvate and consequently promote GABA release and pHP rearrangement as if in H$_2$O. Further, the CH$_3$CN-H$_2$O mixture forces the protonation equilibrium away from the conjugate base and toward the more reactive protonated pHP form.

b) DMSO-H$_2$O. In accord with Phillips findings$^{88}$ with pHP diethyl phosphate (114), the quantum yields, $\Phi$$_{\text{dis}}$, for 101, 122-128, 130-131 were considerably lower than with those in 100% H$_2$O. However, 2,3,6-triF-GABA (128) expressed a $\Phi$$_{\text{dis}}$ of 0.12, an increase vs. H$_2$O (0.08). Congruent with 25% aq. CH$_3$CN above, the $\Phi$$_{\text{dis}}$ for most pHP GABA derivatives in 10% aq. DMSO were similar to those in H$_2$O. Marked increases, however, were observed for 2,3,6-trifluoro-pHP GABA (128 (0.15 vs. 0.08 in H$_2$O)) and 2,3,5,6-tetraF-GABA (129 (0.18 vs. 0.11 in H$_2$O)). All pHp GABA’s revealed absorbances in this medium at ~280 nm, the expected absorption for the predominantly protonated phenol; 128 and 129 have the lowest pKa’s (4.5 and 3.9, respectively) and therefore consist predominantly of the deprotonated form at higher H$_2$O concentrations. The results here represent $\Phi$$_{\text{dis}}$ for protonated 128 and 129.
c) **1-pentanol.** 1-pentanol is a model solvent for pHGABA esters, employed here to
test the relevance of the calculated ClogP’s to their photorelease efficacy in lipophilic
solvents. The calculated ClogP values for 101, 122-131, 133-134 revealed only m-
CF₃-pHP GABA (130 (2.02)) and m-OCF₃-pHP GABA (131 (1.93)) (Table 34
Results) to be larger (more lipophilic) than pHGABA (101 (1.19)). The ClogP’s of
the remaining fluoro pHGABA’s 122-129, though less than for 101, were,
nonetheless, more positive. In contrast, the m-methoxy pH derivatives had ClogP
values of -3.81 and -3.38, respectively. Nevertheless, 1 mM solutions of 133 and 134
in 1-pentanol were prepared without difficulty. The Φᵦᵩ values were reduced by 25-
50% compared to H₂O. The rest were unchanged within experimental error.

d) **1-octanol.** The quantum yields in 1-octanol were consistent with 1-pentanol.
Photolysis experiments of a 1mM sample of m-OCF₃-pHP GABA (131) in 1-octanol
with ¹⁹F NMR analysis revealed complete disappearance after 30 minutes at 300 nm,
evidenced from the lack of signal at -59.68 ppm. New bands were seen at -59.49 and
-59.24 ppm, the former much larger than the latter. Spiking with an authentic sample
of p-hydroxy-m-OCF₃-benzyl alcohol produced a new signal at -59.09 ppm, ruling this
out as a photoprodut. Spiking with an authentic sample of p-hydroxy-m-
trifluoromethoxy-acetophenone effected a substantial increase in the signal at -59.49
ppm, identifying this as the major, reduced photoprodut and comparable to the
results from photolysis of m-OCH₃-pHP GABA (134) in H₂O. A radical mechanism
is imputed for the photorelease of GABA in 1-octanol, the same as shown in Scheme
53.
J. Salt and ionic strength effects.

1) Background. Phillips first broached the notion of contact ion pair recombination (Figure 9, Introduction) as a possible way to reestablish the pH. In this example, the intimate or contact spiroketone-diethyl phosphate ion pair, formed prior to deprotonation of the phenol, was posited to either recombine to pH diethyl phosphate as one triplet decay mechanism or completely separate to allow intersystem crossing and solvolysis in forming the rearranged \( p \)-hydroxyphenylacetic acid. A direct result of ion-pair recombination is a lower \( \Phi_{\text{dis}} \), and hence this process represents one pathway for triplet deactivation.

A conceivable method to attenuate this ion recombination is to adjust the ionic strength of the solution by adding a photostable, non-nucleophilic salts, e.g., Li\(^+\), Na\(^+\),
or K$^+$ ClO$_4^-$, which could intercede between the contact ion pair and hinder substrate return.

Winstein\textsuperscript{122} suggested that “salt effects” could enhance observed rates of solvolysis.\textsuperscript{123,124,125} The Winstein ion-pair mechanism (Scheme 55) is often applied to $S_N1$ solvolysis reactions to demonstrate ion pair return. Initial ionization of RX

**Scheme 55.** Winstein’s ion-pair, solvolysis mechanism with added LiClO$_4$.

![Scheme 55 diagram](image)

produces an intimate ion pair (IP) that can recombine (internal return) or further solvate through solvent insertion to form the solvent separated ion pairs (SSIP). The SSIP can also recombine to IP or progress on to the solvolysis products. Further stages of solvent separation to unpaired discrete ions (DI) may intervene and these can recombine to SSIP. In the presence of added salts, the increased ionic strength
encourages the separation of the ions. Normally, these added salts do not affect the intimate ion pair dissociation. However, LiClO\textsubscript{4} is an exception. It is particularly effective at dissociating SSIP by Li\textsuperscript{+} coordination with the nucleofuge through a salt scavenged pair (ScIP) that prevents external ion pair return from occurring, generally resulting in ameliorated rates of solvolysis. Furthermore, in certain instances, it has been shown to intercept IP and is the origin of the “special salt effect”.

Tidwell\textsuperscript{126} and coworkers recently reported results from examinations of added LiClO\textsubscript{4} on the rates of trifluoroacetolysis of 1,1,1-trifluoro-2-phenylpropan-2-yl 4-methylbenzenesulfonate (172) (Equation 18).

![Equation 18](image)

A positive linear correlation between the unimolecular rates of trifluoroacetolysis of 172 (1) with NaO\textsubscript{2}CCF\textsubscript{3} in CF\textsubscript{3}CO\textsubscript{2}H and [LiClO\textsubscript{4}] was observed (Figure 72), suggesting that LiClO\textsubscript{4} effectively scavenges or further separates SSIP and thus hinders external ion pair return, resulting in augmented rates of trifluoroacetolysis.
Figure 72. Observed enhancements of $k_{\text{obsd}}$ of trifluoroacetolysis of 172 ($R = \text{CH}_3$) as a function of [LiClO$_4$]. Reproduced with permission from the ACS, ref. 126.

2) Effects of added LiClO$_4$ on the quantum yields. Quantum yields at 300 nm were determined in the presence of [LiClO$_4$] from 0 to 1.0 M. It was reasoned that added salt would influence any pHp GABA ionization processes including any recombination events. An increased $\Phi_{\text{dis}}$ could be evidence for a contact ion-radical pair of the triplet biradical and the nucleofuge. Conversely, if GABA departure occurred prior to phenol deprotonation of pHp, a triplet ion pair intermediate would be obtained. UV-Vis analysis of all pHp esters indicated the prevalence of protonated pHp in the ground states. Nearly all pHp GABA’s, excepting the $m$-methoxy pHp derivatives (133 and 134) evinced enhancements in $\Phi_{\text{dis}}$ with added LiClO$_4$ (Table
One example, pHGABA (101), revealed a linear increase $\Phi_{\text{dis}}$ with an increase in $[\text{LiClO}_4]$ between 0 to 1.0 M (Figure 73) and a slope of 0.0153 M$^{-1}$ or 15.3 mM$^{-1}$.

**Figure 73.** Plot of $\Phi_{\text{dis}}$ vs. $[\text{LiClO}_4]$ for pHGABA (101).

This was also obtained with 2,3-difluoro pHGABA (123), which disclosed a linear relationship with a slope 0.0187 M$^{-1}$ (18.7 mM$^{-1}$) (Figure 74). Note that units are not bimolecular rate constants because quantum yields are in mM/Ein and are not rates.
In contrast to these findings, other derivatives showed an early rise in the increased efficiency dependence in LiClO$_4$ concentration followed by a change in slope, such as 2,5-diF-pHP GABA (125) (Figure 75). Initially, a special salt effect was considered. This is characterized by rapid changes induced at low concentrations of exclusively LiClO$_4$ as the added salt, followed by a leveling off at higher concentrations. However, a similar result with observed with NaClO$_4$, which precluded the “special salt” effect associated with Li$^+$. The form of the quantum yield
dependence on the concentration of LiClO\textsubscript{4} for 125 is similar to a saturation curve which would be expected for the ionic strength effect on quantum yields.

**Figure 75.** Plot of $\Phi_{\text{dis}}$ vs. [LiClO\textsubscript{4}] for 2,5-diF-pHP GABA (125).

No salt effects were observed for pHP Trp (149), which exhibited a $\Phi_{\text{dis}}$ of 0.12 in 1.0 M LiClO\textsubscript{4} vs. 0.10 in water.

Other perchlorate salts, as NaClO\textsubscript{4} and Mg(ClO\textsubscript{4})\textsubscript{2}, were probed for influences on $\Phi_{\text{dis}}$ for 2,3-diF-pHP GABA (124) and 2,5-diF-pHP GABA (125). For 125, concentrations up to 1.0 M NaClO\textsubscript{4} produced a correlation that differed with LiClO\textsubscript{4}. A logarithmic fit ($R^2 = 0.99$) was revealed that possibly signifies saturation effect. Similarly, 124 manifested a logarithmic correlation ($R^2 = 0.95$) between $\Phi_{\text{dis}}$ and
[NaClO₄]. When assessed in KClO₄, 125 manifested a logarithmic correlation ($R^2 = 0.99$) that paralleled that with LiClO₄. When assessed in Mg(ClO₄)₂, significant drops in $\Phi_{\text{dis}}$ were observed at low concentrations, but increased at 1.0 M Mg(ClO₄)₂. For example, 2,3-diF-pHP GABA (124) divulged a drop in $\Phi_{\text{dis}}$ from 0.24 in H₂O to

![Figure 76. Plot of $\Phi_{\text{dis}}$ vs. [Mg(ClO₄)₂] for 2,3-diF-pHP GABA (124).](image)

0.12 at 0.01 M Mg(ClO₄)₂ that augmented to 0.31 at 0.1 M and remained constant up to 1.0 M (Figure 76). The distinction that the doubly charged Mg$^{2+}$ vs. singly charged Li$^+$ or Na$^+$ attenuates the $\Phi_{\text{dis}}$ could be through stabilizing interactions with GABA and the ketone compared to the ClO₄$^-$ influence at low concentrations (Figure 77).
The rate constants and lifetimes measured by LFP for the triplet excited state prior to rearrangement were also examined to discern an effect of LiClO₄ on pHp mesylate (168) in water and 0.66 M aqueous LiClO₄. This was accomplished by LFP and found to cause no change in the rate for intersystem crossing, $3.4 \times 10^{11} \text{s}^{-1}$ vs. $3.9 \times 10^{11} \text{s}^{-1}$, the rate for triplet decay, $1.6 \times 10^{11} \text{s}^{-1}$ and $1.1 \times 10^{11} \text{s}^{-1}$ or the rate for the decay of the triplet biradical (117), $1.4 \times 10^{9} \text{s}^{-1}$ and $2.0 \times 10^{9} \text{s}^{-1}$. The results suggest that (1) either the ion radical recombination is occurring in those instances that show an increased $\Phi_{\text{dis}}$ with added LiClO₄ or (2) a delicate balance or compensation between the increase in the rate of rearrangement relative to the rate of triplet pHp GABA decay that does not alter the sum of these two processes is occurring. The yields of triplet (ISC vs. singlet decay) are not likely to be a factor here since $\Phi_{\text{ISC}}$ and $k_{\text{ISC}}$ values are already maximal.

3) Common ion effect. As noted above, the recombination of the released GABA and either the protonated or unprotonated triplet biradical may serve as pathways for the decreased efficiencies in the absence of LiClO₄. This suggests that a “common
ion-like effect$^{125}$ might also operate here. In reference to Scheme 58, added $X^-$, the common ion serves to enhance the recombination steps for SSIP and DI to regenerate RX, the consequence of which is a reduced apparent rate of solvolysis. It was rationalized that added GABA would reduce $\Phi_{\text{dis}}$ by recombination of GABA with either the protonated or the conjugate base triplet pHP. However, addition of 1 to 10 equivalents of GABA (Table 26) made no significant changes on $\Phi_{\text{dis}}$. This observation could be due to the lack of intervention at the SSIP or IP stages. It might also result from rather poor nucleophilicity of GABA. A larger quantity of GABA or better nucleophilic trapping agent, such as azide $N_3^-$, may be needed to educe recombinations reflected in $\Phi_{\text{dis}}$.

K. Rationalized mechanism of the photo-Favorskii rearrangement. A summary of the findings from studies reported here; a) the singular formation of $p$-hydroxyphenylacetic acid and $p$-hydroxybenzyl alcohol photoproducts from the pHP chromophore determined by $^1\text{H}$ NMR and HPLC analysis; b) a discrete deactivation pathway for the protonated pHP ($\lambda_{\text{max}} \sim 280$ nm, marginal fluorescence); c) a different, less efficient pathway for the pHP conjugate base ($\lambda_{\text{max}} \sim 330$ nm, $\Phi_{\text{dis}} \leq 0.10$, fluorescence $\lambda_{\text{em}} > 382$ nm, $k_F 10^{10}$ s$^{-1}$); d) a narrow range for all GABA release efficiencies ($\Phi_{\text{dis}}$) from 0.08 to 0.28, as well as for 5-acetylsalicycic acid caged carboxylic acids (GABA, l-Glu, Gly, etc.) at 300 nm; e) a narrow range for the Stern-Volmer and LFP derived rate constants for intersystem crossing, ($k_{\text{ISC}} \sim 10^{11}$ s$^{-1}$) and GABA release ($k_r \sim 10^7$-$10^8$ s$^{-1}$); f) the confirmatory and diagnostic absorbance bands
for the transient intermediate, a triplet biradical (330, 420 and 445 nm); g) the DFT derived, out of plane (OOP) motion of the carbonyl during formation of the oxyallyl phenoxy triplet biradical to the ring; h) a reinforcement of the recently discovered Brönsted $K_p$, linear free energy relationship for the rate of nucleofuge release versus the pKa of the nucleofuge; i) a measure of the change in the reactivity of the pH photorelease process when solvents such as organic alcohols (1-pentanol and 1-octanol) elicit GABA release at reasonable efficiencies and j) a dramatic salt effect from added LiClO₄ and related salts that improve $\Phi_{\text{dis}}$, possibly signifying intimate ion pairs. These results are summarized by representative mechanisms for the photo-Favorskii rearrangement in Scheme 56 for protonated pHGABA and Scheme 57 for corresponding conjugate base.
Scheme 56. Proposed photo-Favorskii mechanism for protonated pHp GABA esters. Red indicates findings from current studies. Italicized processes indicate that further confirmation is needed.

R-pHP GABA
\( \lambda_{\text{max}} = 271-287 \text{ nm} \)

\[ \text{hv} \longrightarrow \text{Aq. Solvents, ClO}_4^- \]

\( \lambda_{\text{max}} = 330, 420, 445 \text{ nm} \)

spiroidienone

\( \text{H}_2\text{O} \)

p-hydroxyphenylacetic acid
\(^1\text{H} \text{NMR: } \delta = 3.6 \text{ ppm} \)

\( \text{CO} \)

p-quinone methide
\(^1\text{H} \text{NMR: } \delta = 4.6 \text{ ppm} \)

oxyaryl-phenoxo

OOP twisted biradical

\( \Phi_{\text{dis}} = 0.08 - 0.28 \)

\( k_c = 1.8 - 15.8 \times 10^7 \text{ s}^{-1} \)

Internal Return

p-hydroxybenzyl alcohol

\( \text{H}_2\text{O} \)

pHP GABA
\( \text{ClO}_4^- \)

\( \text{H}_2\text{O} \)

Intimate Ion Pair

\( k = 10^{11} \text{s}^{-1} \)

pKa = 4.00
**Scheme 57.** Proposed photo-Favorskii mechanism for the conjugate base of pHp GABA esters. Red indicates findings from current studies. Italicized processes indicate that further confirmation is needed.

- *R*-pHP GABA
  - $\lambda_{max}$ 316-332 nm

- Aq. Solvents, $pH \geq 8$
  - $\text{ClO}_4^-$
  - hv
  - $\lambda_{em}$ 380-399 nm
  - $k_f \sim 10^{10} - 10^{11} \text{ s}^{-1}$

- ISC

- OOP twisted biradical

- Internal Return

- Intimate Ion Pair

- p-hydroxyphenylacetic acid
  - $^1H$ NMR: $\delta \sim 3.6$ ppm

- p-quinone methide
  - $p$-hydroxybenzyl alcohol
  - $^1H$ NMR: $\delta \sim 4.6$ ppm
L. Biological studies.

1) Neuronal assays. Neuronal studies were conducted by the Kandler Groups at the University of Pittsburgh. To test the biological activity of pHGABA esters, \( m\text{-CF}_3\text{-pHP GABA (130)} \) \( m\text{-OCF}_3\text{-pHP GABA (131)} \) and were assessed in neurons of cortical sections of mice using patch clamp recordings. Confined photolysis with 10-50 ms UV pulses through a narrow optical fiber generated whole cell inward currents. Increasing concentrations of photolyzed 131 effectuated membrane currents of sizeable amplitudes and long durations (Figure 78). To compare the sensitivity of

Figure 78. Amplified current as a function of photoreleased GABA from \( m\text{-OCF}_3\text{-pHP GABA (131)} \).

photolyzed 130 and 131 with previously studied \( m\text{-OCH}_3\text{-pHP GABA (134)} \), concentration response curves for neurons subjected to 130, 131, and 134 were generated by fixing the responses to Hill’s equation with subsequent
determinations of half maximal effective concentration, EC\textsubscript{50}, values of 49 µM, 120 µM and 93 µM, respectively, for 131, 134, and 130, respectively (Figure 79). The

**Figure 79.** Population data of peak currents normalized to the maximum peak response and fitted to Hill’s equation to derive EC\textsubscript{50} values for 130, 131, and 134.

The specific activation of the GABA\textsubscript{A} receptor antagonist SR 95531 (Garbazine; Tocris, Ellisville, MI) was probed with m-OCF\textsubscript{3}-pHP GABA (131). Photolysis of 131 produced membrane currents that were completely blocked by Garbazine (Figure 80), signifying that the response is governed by activation of the GABA\textsubscript{A} receptor. Additionally, the current-voltage relationships of the responses divulged a reverse potential of -14.2 ± 2.9 mV (Figure 81), which is near the

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**Figure 80.** Membrane currents elicited from photolysis of 131 are completely obstructed by Garbazine, indicating the activation origin as the GABA<sub>A</sub> receptor.

**Figure 81.** Reverse potential educed from the photolysis of 131. Theoretical potential of -20 mV, as determined from the Nernst equation for chloride concentration between the internal (60 mM) and external (133 mM) solution. Collectively, membrane currents evoked from photolysis of 130 and 131 are moderated by activating the GABA<sub>A</sub> chloride receptor channel.
To determine whether caged GABA could function as an agonist for the GABA\textsubscript{A} receptor, 130 and 131 were examined for their effects on membrane input resistance by monitoring the current responses with short command steps from a holding potential of -70 mV to step potential of -65 mV during 100 ms in the presence and absence of GABA (100 µM) or 130 and 131 (100 µM). Neither 130 nor 131 altered the holding current (Figure 82) or reduced the membrane input resistance.

**Figure 82.** Probing the effect of 131 on the holding currents or membrane input resistance. No effect was noticed.

![Graph showing current responses](image)

However, the employ of GABA on the same neurons educed inward currents and decreased membrane resistance due to activation of GABA\textsubscript{A} chloride channels. Thus, only photoreleased GABA and not caged GABA can activate the GABA\textsubscript{A} chloride channels.

2) **FITC-DOC.** FITC-DOC was synthesized and distributed to the Picking group\textsuperscript{129} for use. The crux of their research involves the *Shigella flexneri* organism,
specifically the induction mechanism to bacillary dysentery in humans. Bacillary dysentery is a debilitating and potentially fatal affliction educed by this organism and is common in the third world. The type III secretion system (TTS) is a crucial apparatus for infection in this organism. Among other constituents of TTS, a surface needle is protuberant. The far terminus of the needle manifests a ‘tip complex’ that consists of an invasion plasmid antigen D (IpaD), a regulatory body for TTS and necessary for the engagement and organization of the translocator protein IpaB at the needle tip in the presence of deoxycholate (DOC) or other bile salts. Until recently, the role of DOC in IpaB recruitment to the TTSA needle tip was unknown. However, current studies revealed that IpaD specifically binds to FITC-DOC, specifically at the N-terminal-domain of IpaD at its central coiled-coil, a locus that also may be involved in needle-tip interactions.  

M. Future Studies. Substituent and nucleofuge influences on the photo-Favorskii rearrangement have been probed. The pKa’s of substituted pH’s and variation of the nucleofuge play key roles in the efficiencies and rates of nucleofuge release. Time resolved resonance Raman, TR, and laser flash photolysis, LFP, studies have provided a wealth of information concerning the structure and reactivity of excited state of pH and the corresponding transient intermediates. Assimilating these data into detailed mechanisms for the photo-Favorskii rearrangement of the pH chromophore and its substituted derivatives are condensed into Schemes 56 and 57.
Nevertheless, there are several studies that are essential for further elucidation of these processes.

1) **Organic and mixed organic media.** The preliminary studies reported here conveyed nucleofuge release in 10% aq. DMSO, 1-pentanol and 1-octanol, but the minor the photoproducts in the latter two studies have not been identified. The overall effects of organic or mixed organic media on the photo-Favorskii reaction would be useful information, especially with regard to future studies related to highly lipophilic, biologically relevant nucleofuges and media.

2) **Salt effects.** Added perchlorate salts clearly manifested enhanced $\Phi_{\text{dis}}$ for nearly all pHp GABA’s. The preliminary interpretation of these effects involve an influence of salts possibly on intimate ion-radical pairs, paralleling Winstein's $S_N1$ solvolyses. However, further investigations are required to probe the origin of the effects, particularly focusing on rates of release. Although LFP studies on pHp mesylate showed no changes in either the rates of ISC or triplet decay with 0.66 M LiClO$_4$, this experiment was the only one tested and involved a superb nucleofuge, where return is expected to be minimal. Expanded LFP studies on other pHp derivatives are necessary.

3) **Chiral nucleofuge.** A photochemical investigation of (R,R)-166 or (R,S)-167 would have been useful for following the configurational changes at a stereocenter that is intimately involved in the cyclopropanone formation and collapse during the photo-Favorskii rearrangement. Measurements at partial conversion to photoproducts could test for ion-radical return as well. This could be strong evidence for the
intermediacy of an intimate ion pair prior to the triplet biradical formation. A suggested stepwise nucleofuge departure and phenol deprotonation instead of the currently posited concomitant process may emerge.

4) **Structurally similar entities to pHr.** Yousef\textsuperscript{111} reported low $\Phi_{\text{dis}}$ for a series of substituted 1,4 and 2,6 naphthols despite favorably lengthening the $\lambda_{\text{max}}$. Nitrogen analogs of pHr, such as 4-aminoacetophenone were briefly studied but gave radical products.\textsuperscript{131} Some structurally analogous entities to pHr should be investigated.
Conclusion

The $p$-hydroxyphenacyl (pHP) chromophore is an efficacious phototrigger for the temporal, spatial and concentration control over substrate release. The release mechanism is underscored by an intriguing rearrangement of pHP that resembles the well known Favorskii reaction. Previous quenching and spectroscopic studies with various pHP cages were effective in providing an early mechanistic picture of the photo-Favorskii rearrangement, including a) the clean photoconversion, b) favorable quantum efficiencies for release (~0.2 and higher) at 300 nm, c) the triplet reactive excited state, d) the dependence of the triplet lifetime on the water content of the solvent, e) the pertinence of water in inducing substrate release and proton abstraction from pHP, f) the rapid rates ($10^7$-$10^9$ s$^{-1}$) of substrate release and g) very limited structure-reactivity explorations.

However, the expanded studies reported here are essential for a more detailed, mechanistic comprehension. These include 1) distinguishing nucleofuge influences, 2) media effects (pH, buffer, mixed solvents and added salts), 3) structure-reactivity relationships (electron withdrawing and electron donating groups) and 4) modulating the pKa of the phenol to discern pH induced changes in, for example, the quantum yields for substrate release. The pHP chromophore suffers from a low molar absorptivity at $\lambda > 300$ nm, which limit it’s usefulness in the biological arena because of possible secondary photodegradations and the paucity of inexpensive lasers that
utilize wavelengths between 300-320 nm. Extending the $\lambda_{\text{max}}$ of pHP by substitution with auxochromes is a plausible motive to overcome these deficiencies.

Several pHP GABA derivatives were specifically designed and synthesized to probe structure-reactivity relationships, primarily between the nature of the substituent and quantum yield for GABA release, $\Phi_{\text{dis}}$. Electron donating groups, such as methoxy and hydroxy, appeared to attenuate $\Phi_{\text{dis}}$ (relative to the unsubstituted parent) or completely quench GABA photorelease. Electron withdrawing, $m$-substituted carbonyl moieties also reduced $\Phi_{\text{dis}}$ by half. The rate constants for GABA release, $k_r$, did not vary significantly from the unsubstituted parent.

Several fluoro pHP GABA esters were designed and synthesized to explore how modulating the pKa of pHP would result in pH induced changes by the media on $\Phi_{\text{dis}}$ and $k_r$. The pH of the media was found to directly control the equilibrium between the protonated and conjugate bases of these fluoro pHP derivatives as evidenced by a) $\lambda_{\text{max}}$ values (<300 nm for protonated and >300 nm for conjugate bases) and b) the quantum yields for GABA release at 300 nm ($\Phi_{\text{dis}} > 0.15$ for protonated, $\Phi_{\text{dis}} < 0.10$ for conjugate bases). Stern-Volmer and LFT studies determined rate constants for release that narrowly spanned from $10^7$ to $10^8$ s$^{-1}$. At 350 nm, the conjugate bases revealed $\Phi_{\text{dis}}$ that were less than 0.01. The conjugate bases are fluorescent and the fluorescent lifetimes of 1-10 ns for these were obtained. A mechanism for GABA release from the conjugate bases of pHP was posited.

A series of 5-acetylsalicylate GABA’s with UV absorbing appendages (antennae compounds) were generated in efforts to lengthen the $\lambda_{\text{max}}$ of pHP. Though
this objective was accomplished, the $\Phi_{\text{dis}}$ were low at 300 nm (0.01 to 0.11) and very low (<0.01) at 350 nm.

The quantum efficiencies at 300 nm of several pHGABA’s in aqueous DMSO and CH$_3$CN were determined to be commensurate with those in water. However, GABA photorelease was quenched in neat DMSO. Additionally, photolyses of these compounds were conducted in 1-pentanol and 1-octanol, which disclosed reasonable efficiencies (~0.1), though the primary photoproducts were the reduced $p$-hydroxyacetophenone, signifying an alternate mechanistic pathway than the photo-Favorskii rearrangement.

When perchlorate salts were added, significant increases in $\Phi_{\text{dis}}$ were observed for nearly all pHGABA’s. A corollary from this was a proposed transient, solvent separated, triplet pH and GABA radical or ion pair. This phenomenon suggests a possible nucleofuge ionization step prior to proton abstraction from the pH phenol, a pathway contrary to the currently accepted concerted process.

Laser flash photolysis on several pHGABA’s revealed time resolved absorption bands at 330, 420, and 445 nm, which were ascribed to a triplet oxyallyl-phenoxy biradical, a consequence of adiabatic nucleofuge disjunction. DFT calculations of the triplet oxyallyl-phenoxy biradical determined the radicals to be contorted out of plane from 22° to 58° depending on the substituents and their position on the aryl ring. Additionally, the triplet biradicals were calculated to be at least 78 kcal/mol lower in energy than their singlet spirodienedione counterparts. Time resolved FTIR spectroscopy was conducted with pH diethyl phosphate and
disclosed a band at 1647 cm\(^{-1}\) that was consistent with a transient \(p\)-quinone-methide, a suitable precursor to the minor \(p\)-hydroxybenzyl alcohol.

Finally, studies on biological processes that are initiated by release of an active substrate such as the neurotransmitter GABA have shown the power of spatial and concentration control in neuronal preparations. These studies were performed with three pH\(p\) GABA derivatives, which demonstrated the pH\(p\) chromophore as being an effective phototrigger for facile release of GABA in neuronal stimulation.

In summary, previous work and the studies reported here have provided a clearer understanding of the photochemical and photophysical processes of the \(p\)-hydroxyphenacyl chromophore. The photochemical details of the Favorskii rearrangement have been shown to include a transient oxyallyl-phenoxy triplet biradical, formed during the release of the conjugate base as substrate. The role of \(H_2O\), solvents, pH and added salts on the photochemistry and photophysics as well as substituent effects on the chromophore have supported the general mechanistic schemes presented here.
Experimental

Methods

Melting points were conducted with open ended capillary tubes using a non-calibrated Thomas-Hoover melting point apparatus. Lyophilization was performed using a Labconco FreezeZone device set at -52 °C and 0.04 mbar. Solution pH values were determined using a Fisher Science pH 510 meter calibrated with certified Fisher buffer solutions of pH 4, 7, and 10. Products of all reactions were assessed for purity using the following instrumental techniques: For $^{1}H$, $^{13}C$, and $^{19}F$ NMR, Bruker DRX 400 and DRX 500 MHz spectrometers were utilized with trifluoroacetic acid as an internal standard ($\delta = -76.55$ ppm). GC/MS analysis was done on an Agilent 6890N Network GC System equipped with a Quattro Micromass Triple Quadrupole Electron Impact mass spectrometer. Exact masses were performed on a Quattro Micromass Triple Quadrupole Electro Spray Ionization mass spectrometer. UV/Vis data was obtained on a Cary Bio100 instrument using 1.5 ml quartz cuvettes. Fluorescence data were acquired on a Cary Eclipse Fluorescence Spectrophotometer using a 10 mm quartz cell. IR data were obtained using a Shimadzu FT-IR 8400 S instrument with pressed potassium bromide (KBr) pellets or a prefabricated sodium chloride (NaCl) chloroform cell for solid phase analysis, and NaCl prefabricated plates for oil analysis. Ground state $pK_a$’s were determined through sample titration with increasing amounts of 0.0461 M NaOH (standardized with potassium hydrogen phthalate), correlating pH vs. [OH$^-$], and ascertaining the half equivalence point as the $pK_a$.  

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Product separation was achieved by flash chromatography using EM Science silica gel and gradient hexanes/ethyl acetate as eluting solvents. All reactions were run under ambient conditions unless otherwise stated. $^1$H NMR spectroscopy was utilized to monitor the progress of photolysis of all new pHp cages. In general, a 1-10 millimolar sample of pHp-caged GABA was prepared in 2 ml D$_2$O, approximately 2 ml was placed in an NMR tube. This was positioned in a photoreactor equipped with two 15 W, 3000 Å Rayonet lamps and a merry-go-round. Irradiation for 30 min generally led to 100% conversion as judged by the absence of the ester proton signals. Photoproduction identification was achieved through spiking the samples of the irradiated mixture with authentic samples of the resolved GABA. Quantitative photolysis conditions for determination of quantum yields and Stern-Volmer quenching constants ($K_{SV}$) were as follows: The lamp light output (in mEinstein/min) was established using the potassium ferrioxalate method. 7 Milligram quantities of caged compounds and caffeine or acetamidophenol were weighed out on a Fisher brand Microbalance and dissolved in 4 ml of 18 MΩ ultrapure water, salt solutions of various concentrations, buffers with or without adjusted ionic strengths, or purified organic solvents were then added to a quartz tube and vortexed, resulting in a homogenous solution of the caged compound and internal standard. Concentrations of the caged compounds ranged from 1-9 mM. These tubes were then placed in a carousel within a Rayonet Photochemical Reactor equipped with two 3000 Å, 15 W Rayonet Photochemical Reactor RPR3000 Mercury Lamps as light sources.
100 µl samples were removed at 30 s intervals up to 5 minutes using a 250 µl Hamilton micro-syringe and diluted to 1 ml with water using 1 ml volumetric flasks.

**Picosecond Pump-probe Spectroscopy**

Picosecond transient absorption was measured with the pump–supercontinuum probe technique using a Ti/Sa laser system (Clark MXR CPA-2001; 775 nm, pulse energy 0.9 mJ, full width at half maximum 150 fs, operating frequency 426 Hz). Part of the beam was fed into a Clark MXR NOPA. The output at 532 nm was frequency-doubled by a β-barium borate (BBO) crystal to 266 nm and, after compression, provided pump pulses with an energy of 1 µJ and <150 fs pulse width. A probe beam continuum was produced by focusing the 775-nm beam in front of a CaF$_2$ of 4 mm path length. The pump and probe beams were focused to a 0.04 mm$^2$ spot on the sample that was flowing in an optical cell of 0.3 mm thickness. The absorbance was in the range of 0.3–1, depending on the solubility of the compound. The probe beam and a reference signal (passing the solution besides the pump beam) were spectrally dispersed and registered with two photodiode arrays (512 pixels). Transient absorption spectra were calculated from ratio of the two beams. Spectra recorded with time delays ranging from 0 to 1.8 ns (to observe the triplet decay) were typically recorded with 20 ps steps, those ranging from 0-8 ps (to observe ISC) with 80 fs steps. The shorter measurements were corrected for chirp using a program (SPAN) kindly provided by Prof. N. Ernsting, Berlin. To improve the signal-to-noise ratio, the
data were averaged over multiple pump-probe scans (3-6 scans with 400 shots per temporal point).

**Quantum Efficiency Determinations**

Quantitative analysis was achieved by HPLC/UV or HPLC/MS/MS. The LC/MS/MS instrument was a Waters 2695 Liquid Chromatographer equipped with a Quattro Micromass Triple Quadrupole Electro Spray Ionization Mass Spectrometer, outfitted with an autosampler. UV-Vis detection consisted of a Waters 2497 type with a dual wavelength detector set at 220 and 240 nm. The reservoirs used were as follows: A) 99% water, 1% methanol, 10 mM ammonium formate and 0.06% formic acid. B) 99% methanol, 1% water, 10 mM ammonium formate, and 0.06% formic acid. The column was a reverse-phase (C18), 4 µm mesh Altech Altima, 50 mm in length. Injections of 100 µl were made with an automated sampler for each run for a total of 3 injections per vial. A mobile phase gradient was utilized to optimize compound separation. The flow rate was set at 300 µl/min. Data analysis was performed by Mass Lynx Ultima software and Microsoft Excel. Smoothing functions were used for peak analysis of the chromatographic peaks. Calibration curves to obtain $R$ values from linear least-squares regression were determined at concentrations of the reactants and products in photolyses by systematic increases of pHP-caged GABA, free GABA, and $p$-hydroxyphenylacetic acid concentrations to determine correlations with internal standards caffeine or 4-acetamidophenol. The quantum efficiencies
were then calculated from the ratio of the reactant or product concentrations to the photons absorbed using the actinometer values obtained as indicated above.

**Lifetime and Rate Measurements**

The Stern–Volmer quenching method was employed to determine the triplet lifetimes of pH derivatives. Potassium sorbate served as the quenching agent. Solutions of pH-GABA, 0.001–0.01 M, were diluted with sorbate solutions of increasing concentration (0–0.1 M), and photolyzed under the aforementioned conditions to ascertain the change in quantum efficiencies of GABA release. $\Phi_0/\Phi$ vs. [Q], and the Stern–Volmer constant, $K_{SV}$, were determined. To determine the triplet lifetime, $\tau^3$, the rate of quenching, $k_q$, was assumed to be the rate of bimolecular diffusion, $k_{diff} \sim 7.2 \times 10^9$ s$^{-1}$ (water). The rates of photorelease, $k_r$, were then computed as $k_r = \Phi_{rel} \tau^3$.

**Calculations**

The atom coordinates for each molecule were introduced into SYBYL for molecular mechanics optimization using the Tripos Force Field. The ensuing structures withstood full geometric optimizations in Gaussian 03 at the B3LYP level using the 6-31G* basis set, soliciting the lowest energy triplet state in each case. The triplet stabilization energy was determined by comparing the final energy of optimized structure of each compound to that of the lowest energy singlet state at the triplet-optimized geometry. In order to examine differences in the carbonyl out-of-
plane angles generated by the B3LYP calculation (vs. the MP2-level models reported elsewhere), a subset of these triplet-state species were geometrically optimized at the MP3 level\textsuperscript{138} in Gaussian using the 6-31G* basis set.

**Time Correlated Single Photon Counting Fluorescence**

The experiments were conducted on a custom built instrument using a mode-locked, cavity-dumped Mira Optima 900f Ti:Sapphire system pumped by a 10W Verdi Laser from Coherent, Inc. (Santa Clara, CA) as the excitation source. The pulses were frequency tripled using an Inrad harmonic separator. The excitation wavelength was 281 nm and the fluorescence emission was measured at the emission maximum determined with a fluorescence spectrophotometer. The detection system was fitted with Oriel 27320 polarizers set to the magic angle (54.7) for fluorescent lifetime measurements, single-pass Sciencetech 9030 monochrometers, and Hamamatsu microchannel plate PMTs. Sample temperature was regulated using a Peltier thermoelectric temperature controller from Quantum Northwest (Spokane, WA). Data collection was performed using an SPC-630 PCI card from Becker and Hickl GmbH (Berlin, Germany). Data was collected to at least 60,000 peak counts for all of the decays. The fluorescent lifetimes were fit using the maximum entropy method implemented in Pulse5T (Maximum Entropy Data Consultants, Inc.). The analysis used 200 logarithmically spaced exponentials with a maximum of 15 ns. Discrete fitting was performed using standard iterative deconvolution techniques written in
Microsoft Excel. Samples were prepared by dissolving requisite quantities of precursors in 25 ml of 0.1 M LiOH to furnish ~ 1 mM concentrations.

**Materials**

All starting materials were obtained from Aldrich or Matrix Scientific, unless otherwise indicated. Solvents were distilled prior to use, employing phosphorus pentoxide (P₂O₅) and stored in flame-dried, argon purged round-bottom flasks containing activated molecular sieves. Ultrapure (18 MΩ) water was used in all instances.

**Syntheses**

1-(benzyloxy)-4-bromo-2-fluorobenzene, 122b. The general method of Frechét, et al.¹³⁹ was followed. A solution of 4-bromo-2-fluorophenol (122a, 3.83 g, 20.0 mmol), benzyl bromide (2.38 ml, 20.0 mmol) and potassium carbonate (6.91 g, 50.0 mmol) in CH₃CN (30 ml) was stirred at room temperature for 15 h. The solution was diluted with 50 ml CH₂Cl₂, washed with water (3 × 30 ml), dried (magnesium sulfate), and concentrated to give 5.24 g (93%) of 1-(benzyloxy)-4-bromo-2-fluorobenzene, 122b, as a white precipitate: Mp: 65–67 °C; IR (CHCl₃): 3020, 2987, 2684, 2304, 1498, 1421, 1265, 1217, 1051, 896, 738, 695 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.41 (m, 5H), 7.27 (dd, J = 6.36, J = 2.28, 1H), 7.26 (dt, J = 4.37, J = 2.00, 1H), 6.89 (t, J = 8.76, 1H), 5.13 (s, 2H); ¹³C NMR (125 MHz, CDCl₃) δ 154.23, 151.73, 146.06, 136.02, 128.88, 127.73, 120.14, 117.23, 115.86, 113.02, 71.90; ¹⁹F NMR (376 MHz,
CDCl$_3$ + drop CF$_3$CO$_2$H) $\delta$ -130.78; HRMS (EI): Calc’d for C$_{13}$H$_{10}$FBrO: 279.9899. Found: 279.9905.

1-(benzyloxy)-3-fluorophenyl)ethanone, 122c. The general method of Kosugi, et al.$^{140}$ was used. The experimental procedure is described with 129 (vide infra).

White ppt., 89%, Mp: 79–81 °C; IR (CHCl$_3$): 3053, 2987, 1679, 1610, 1514, 1498, 1421, 1265, 1217, 1052, 896, 738, 696 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 7.74 (dd, $J = 11.33, J = 2.38$, 1H), 7.69 (dd, $J = 9.04, J = 1.59$, 1H) 7.44, (m, 5H), 7.04 (t, $J = 8.44$, 1H), 5.23 (s, 2H), 2.55, (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) $\delta$ 196.05, 153.66, 151.16, 135.82, 130.92, 128.95, 127.57, 125.76, 116.36, 114.22, 71.25, 26.52; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) -133.02; HRMS (M + H): Calc’d for C$_{15}$H$_{13}$FO$_2$: 245.0978. Found: 245.0951.

1-(4-(benzyloxy)-3-fluorophenyl)-2-bromoethanone, 122d. The general method of Paul, et al.$^{142}$ was followed. A solution of 122c (700 mg, 2.86 mmol) in 20 ml CH$_2$Cl$_2$ was cooled to 0 °C in an ice bath. Dioxane dibromide (773 mg, 3.15 mmol) was added and the resulting mixture stirred for 30 min. The ice bath was removed with continued stirring for another 60 min. GC/MS indicated the reaction to be complete.

The solution was washed with water and CH$_2$Cl$_2$, the aqueous layer removed, and the residual layer dried with anhydrous MgSO$_4$, filtered, and evaporated under reduced pressure to yield an orange precipitate, 900 mg (2.58 mmol). Purification was cumbersome and unnecessary for completion of the supervening step, and hence was not pursued. ~95%. IR (CHCl$_3$): 3055, 2988, 1679, 1615, 1511, 1498, 1417, 1267, 1219, 1050, 891, 748, 696 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 7.77 (dd, $J =$
10.00, J = 2.38, 1H), 7.72 (dd, J = 7.05, J = 2.78, 1H), 7.42 (m, 5H), 7.07 (t, J = 8.43, 1H), 5.24 (s, 2H), 4.37 (s, 2H); $^{19}$F NMR (376 MHz, CDCl$_3$ + a drop CF$_3$CO$_2$H) δ (ppm), -133.12.

2-(4-(benzyloxy)-3-fluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 122e. The general method of Hiyama, et al.$^{143}$ was utilized. A solution of 1-(4-(benzyloxy)-3-fluorophenyl)-2-bromoethanone, (122d, 900 mg, 2.78 mmol), potassium carbonate (1.15 mg, 8.35 mmol), 4-(tert-butoxycarbonylamino)butanoic acid (N-Boc-GABA, 678 mg, 3.33 mmol) in 50 ml of CH$_3$CN was stirred for 24 h at room temperature. The solution was washed with EtOAc, water, the aqueous phase discarded, and the resulting phase evaporated under reduced pressure. Flash column chromatography (5:1 Hexanes/EtOAc) afforded a white precipitate, 1.10 g (89%).

Mp: 97-99 °C; IR (KBr): 3300, 3100-2800, 1742, 1701, 1600, 1517, 1437, 1400 1309, 1200, 1165, 953, 750, 700, 668 cm$^{-1}$; $^1$H NMR (400 MHz, CD$_3$CN) δ (ppm), 7.80 (dd, J = 7.05, J = 2.47, 1H), 7.78 (dd, J = 7.05, J = 1.99, 1H), 7.47 (dd, J = 4.45, J = 1.78, 1H), 7.42 (m, 5H), 5.31 (s, 2H), 5.25 (s, 2H), 2.97 (t, J = 6.79), 2.45 (t, J = 7.59 Hz, 2H), 1.76 (m, 2H), 1.40 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 190.67, 172.51, 155.60, 152.30, 150.80, 135.82, 128.39, 128.14, 127.68, 126.93, 125.60, 115.28, 114.82, 77.45, 70.36, 66.12, 39.64, 30.63, 28.22, 24.96; $^{19}$F NMR (376 MHz, CD$_3$CN + drop CF$_3$CO$_2$H) δ (ppm), -133.44; HRMS (M + Na): Calc’d for C$_{24}$H$_{28}$FNO$_6$Na: 468.1798. Found 468.1796.

4-(2-(3-fluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 122. The general method of Marsh, et al.$^{144}$ was followed. In a
flame-dried, 50 ml round bottomed flask containing 2-(4-(benzyloxy)-3-fluorophenyl)-2-oxoethyl-4-(tert-butoxy-carbonylamino)-butanoate, (122e, 850 mg, 1.91 mmol) was added 20 ml of freshly distilled TFA. Stirring continued for 24 h at room under ambient conditions. The solution was evaporated under reduced pressure and washed with EtOAc/water. The water layer was then extracted, frozen, then lyophilized, affording an adhesive precipitate, 550 mg (78%). IR (KBr): 3434, 3269, 3100-2800, 1741, 1693, 1610, 1553, 1420, 1201, 1050, 823, 759, 719 cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm), 7.70 (d, J = 9.03, 2H), 7.12 (t, J = 9.94, 1H), 5.50 (s, 2H), 3.09 (t, J = 7.59, 2H), 2.66 (t, J = 7.08, 2H), 2.05 (m, 2H); ¹³C NMR (125 MHz, MeOD) δ (ppm), 190.05, 172.06, 161.60, 152.27, 151.07, 150.33, 126.03, 125.33, 117.30, 115.33, 65.95, 38.54, 30.02, 22.48; ¹⁹F NMR (376 MHz, D₂O) δ (ppm), -133.58; HRMS (M⁺): Calc’d for C₁₂H₁₅FNO₄: 256.0985. Found 256.0977.

The methodology for generating compounds 123–132 was analogous to that for 122 unless otherwise indicated; the yields were consistent, as well, unless stated otherwise. Only spectroscopic data are included for each intermediate and the final caged derivatives.

1-(4-benzyloxy)-2-fluorophenylethanone, 123c. 123b is commercially available. White ppt., 90%, Mp: 97-99 °C; IR (CHCl₃): 3056, 2992, 1701, 1616, 1510, 1487, 1415, 1275, 1207, 1055, 881, 728, 683 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), δ 7.93 (t, J = 7.76, 1H), 7.40 (m, 5H), 6.85 (dd, J = 8.96, J = 2.49, 1H), 6.72 (dd, J = 13.25, J = 2.39, 1H), 5.15 (d, J = 4.98, 2H), 2.70 (s, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 198.76, 165.25, 159.13, 135.54, 132.70, 128.78, 127.75, 118.13,
115.45, 114.03, 103.11, 70.96, 31.08; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ (ppm), -104.64; HRMS (M + H): Calc’d for C$_{13}$H$_{14}$FO$_2$: 245.0978. Found: 245.0962.

1-(4-(benzyloxy)-2-fluorophenyl)-2-bromoethanone, 123d. Oil, ~92%; IR (NaCl): 3058, 2990, 1711, 1621, 1510, 1486, 1413, 1265, 1063, 890, 725, 680 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), δ 7.95 (t, $J = 8.84$, 1H), 7.40 (m, 5H), 6.89 (dd, $J = 8.84$, $J = 2.31$, 1H), 6.73 (dd, $J = 13.36$, $J = 2.50$, 1H), 5.15 (s, 2H), 4.49 (d, $J = 4.49$, $J = 2.50$, 2H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm), 189.31, 165.37, 159.17, 135.51, 132.68, 128.80, 127.75, 115.66, 115.45, 114.03, 102.99, 71.01, 35.73; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ (ppm), -104.68; HRMS (M + H): Calc’d for C$_{13}$H$_{13}$FO$_2$Br: 323.0083. Found: 323.0101.

2-(4-(benzyloxy)-2-fluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 123e. White ppt., 86%, Mp: 105–107 °C; IR (KBr): 3303, 3100–2800, 1742, 1710, 1659, 1437, 1408, 1309, 1203, 1167, 955, 747, 700, 668 cm$^{-1}$; $^1$H NMR (400 MHz, d$_6$-DMSO ) δ (ppm), δ 7.85 (t, $J = 8.94$, 1H), 7.40 (m, 5H), 7.10 (dd, $J = 13.59$, $J = 2.30$, 1H), 7.01 (dd, $J = 8.86$, $J = 2.42$, 1H), 6.88 (t, $J = 5.70$, 1H), 5.24 (s, 2H), 5.22 (d, $J = 3.25$, 2H), 2.98 (t, $J = 6.81$, 2H), 2.42 (t, $J = 7.60$, 2H), 1.70 (m, 2H), 1.38 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 188.97, 172.17, 164.26, 162.13, 155.60, 135.85, 131.75, 128.53, 127.96, 127.50, 115.05, 112.37, 102.82, 77.45, 70.04, 68.32, 39.30, 30.78, 28.22, 24.93; $^{19}$F NMR (376 MHz, CD$_3$CN + drop CF$_3$CO$_2$H) δ (ppm), -106.01; HRMS (M + Na): Calc’d for C$_{24}$H$_{28}$FNO$_6$Na: 468.1798. Found: 468.1787.
4-(2-(2-fluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoro-acetate, 123. Adhesive solid. IR (KBr): 3434, 3244, 3100–2800, 1741, 1693, 1610, 1523, 1421, 1400, 1309, 1269, 1198, 1166, 1050, 824, 800, 760, 720. cm⁻¹, ¹H NMR (400 MHz, D₂O) δ (ppm), δ 7.85 (t, J = 7.58, 2H), 6.73 (d, J = 9.09, 1H), 5.40 (d, J = 3.03, 2H), 3.10 (t, J = 7.64, 2H), 2.64 (t, J = 7.12, 2H), 2.09 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm), 189.35, 172.10, 165.34, 165.02, 163.13, 133.68, 113.65, 113.39, 102.58, 68.71, 38.60, 30.06, 22.48; ¹⁹F NMR (376 MHz, D₂O) δ (ppm), -106.45; HRMS (M+): Calc’d for C₁₂H₁₅FNO₄: 256.0985. Found 256.0976.

1-(benzyloxy)-4-bromo-2,3-difluorobenzene, 124b.¹⁴⁵ Oil, 94%; IR (CHCl₃): 3058, 2982, 1600, 1510, 1465, 1250, 1202, 1050, 835, 740, 695 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.42 (m, 5H), 7.19 (dt, J = 8.05, J = 2.50, 1H), 6.90 (dt, J = 8.00, J = 1.82, 1H), 5.16 (s, 2H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm) 161.58, 149.74, 147.53, 135.60, 128.75, 127.50, 126.39, 117.38, 113.62, 101.35, 72.09; ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm) -129.51, -154.65; HRMS (EI): Calc’d for C₁₃H₉F₂OBr: 297.9805. Found: 297.9822.

1-(4-(benzyloxy)-2,3-difluorophenyl)ethanone, 124c. White ppt., 88%, Mp: 96–98 °C; IR (KBr): 3055, 2990, 1695, 1614, 1464, 1448, 1380, 1275, 1202, 1051, 880, 725, 680 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.69 (dt, J = 9.39, J = 2.24, 1H), 7.39 (m, 6H), 6.90 (dt, J = 8.00, J = 1.82, 1H), 5.26 (s, 2H), 2.71 (d, J = 5.38, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm) 198.21, 160.44, 141.88, 140.59, 134.96, 128.75, 127.46, 118.93, 116.46, 113.62, 109.67, 71.67, 30.86; ¹⁹F NMR (376 MHz,
CDCl$_3$ + drop CF$_3$CO$_2$H) δ (ppm) -132.74, -158.39; HRMS (M + H): Calc’d for C$_{13}$H$_{13}$F$_2$O$_2$: 263.0884. Found: 263.0887.

1-(4-(benzyloxy)-2,3-difluorophenyl)-2-bromoethane, **124d**. Oil, ~96%; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm) 7.71 (dt, $J$ = 8.44, $J$ = 2.38, 1H), 7.41 (s, 5H), 6.92 (dt, $J$ = 7.80, $J$ = 1.90, 1H), 5.27 (s, 2H), 4.46 (d, $J$ = 2.48, 2H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm) 198.61, 160.37, 153.65, 140.38, 135.11, 128.93, 125.60, 117.97, 113.44, 109.93, 71.96, 31.02; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ (ppm), -132.93, -158.06.

2-(4-(benzyloxy)-2,3-difluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, **124e**. White ppt., 84%, Mp: 115–117 °C; IR (KBr): 3300, 3100–2800, 1742, 1701, 1664, 1619, 1436, 1406, 1200, 1169, 1136, 955, 749, 700, 670, cm$^{-1}$; $^1$H NMR (400 MHz, d$_6$-DMSO) δ (ppm) 7.71 (dt, $J$ = 8.38, $J$ = 2.22, 1H), 7.41 (m, 5H), 7.29 (dt, $J$ = 8.22, $J$ = 1.57, 1H), 6.88, (t, $J$ = 5.76, 1H), 5.34 (s, 2H), 5.25 (d, $J$ = 2.82, 2H), 2.96 (t, $J$ = 6.85, 2H), 2.44 (t, $J$ = 7.55, 2H), 1.67 (m, 2H), 1.37 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm) 188.76, 172.13, 155.60, 152.08, 149.66, 142.12, 141.00, 139.04, 128.60, 128.02, 124.86, 116.21, 110.59, 72.91, 69.81, 69.72, 40.79, 32.09, 29.01, 26.49; $^{19}$F NMR (376 MHz, CD$_3$CN + drop CF$_3$CO$_2$H) δ (ppm) -158.62, -133.05; HRMS (M + Na): Calc’d for C$_{24}$H$_{27}$F$_2$NO$_6$Na: 486.1704. Found: 486.1689.

4-(2-(2,3-difluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, **124**. Adhesive ppt., 92%; IR (KBr): 3435, 3246, 3100–2800, 1739, 1694, 1610, 1525, 1420, 1400, 1310, 1269, 1200, 1165, 1049, 825, 800, 759, 720.
cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm) 7.43 (d, J = 6.57, 1H), 6.75 (d, J = 9.34, 1H), 5.18 (d, J = 2.64, 2H), 3.04 (t, J = 6.81, 2H), 2.60 (t, J = 7.53, 2H), 1.97 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm) 191.36, 173.96, 163.08, 151.57, 140.64, 138.70, 128.71, 119.79, 117.47, 113.20, 68.83, 38.34, 30.09, 21.61; ¹⁹F NMR (376 MHz, D₂O) δ (ppm) -133.71, -162.58; HRMS (M⁺): Calc’d for C₁₂H₁₄F₂NO₄: 274.0891. Found: 274.0884.

1-(benzyloxy)-4-bromo-2,5-difluorobenzene, 125b. White ppt., 91%, Mp: 58–60 °C; IR (CHCl₃): 3002, 2943, 1618, 1508, 1419, 1406, 1375, 1334, 1188, 1170, 1039, 840, 748, 702; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.37 (m, 5H), 7.29 (dd, J = 6.64, J = 3.82, 1H), 6.82 (dd, J = 7.19, J = 2.68, 1H), 5.12 (s, 2H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 154.82, 150.45, 148.49, 147.28, 135.96, 129.35, 128.03, 120.47, 104.54, 98.93, 72.22; ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm), -111.53, -137.66; HRMS (EI): Calc’d for C₁₃H₀F₂BrO: 297.9805. Found: 297.9833.

1-(4-(benzyloxy)-2,5-difluorophenyl)ethanone, 125c. White ppt., 89%, Mp: 89–91 °C; IR (CHCl₃): 3060, 3002, 2943, 1681, 1625, 1508, 1411, 1398, 1375, 1334, 1186, 1159, 1039, 744, 702; ¹H NMR (400 MHz, CD₃CN) δ (ppm), 7.60 (dd, J = 6.88, J = 4.78, 1H), 7.43 (m, 5H), 7.00 (dd, J = 6.73, J = 5.93, 1H), 5.20 (s, 2H), 2.54 (d, J = 5.18, 3H); ¹³C NMR (125 MHz, CD₃CN) δ (ppm), 194.65, 159.45, 158.50, 148.92, 136.38, 130.69, 129.43, 118.88, 117.28, 115.01, 104.46, 72.74, 31.59; ¹⁹F NMR (376 MHz, CD₃CN + drop CF₃CO₂H) δ (ppm), -110.97, -140.38; HRMS (EI): Calc’d for C₁₅H₁₂F₂O₂: 262.0805. Found: 262.0791.
1-(4-(benzyloxy)-2,5-difluorophenyl)-2-bromoethanone, 125d. Oil ~94%; IR (NaCl): 3055, 3008, 2942, 1691, 1624, 1395, 1357, 1332, 1203, 1181, 1150, 1039, 790, 744, 702; H NMR (400 MHz, CDCl₃) δ (ppm), 7.70 (dd, J = 6.69, J = 4.59, 1H), 7.43 (m, 5H), 6.78 (dd, J = 6.50, J = 5.73, 1H), 5.22 (s, 2H), 4.46 (d, J = 2.89, 3H).

2-(4-(benzyloxy)-2,5-difluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino) butanoate, 125e. White ppt., 92%, Mp: 118–120 °C; IR (KBr): 3303, 3100–2800, 1741, 1701, 1657, 1623, 1512, 1437, 1407, 1203, 1165, 1144, 953, 897, 750, 699, 667, cm⁻¹; H NMR (400 MHz, d₆-DMSO) δ (ppm), 7.69 (dd, J = 6.66, J = 4.91, 1H), 7.43 (m, 6H), 6.89 (t, J = 5.62, 1H), 5.32 (d, 2H), 5.23 (d, J = 3.33, 2H), 2.97 (t, J = 6.79, 2H), 2.44 (t, 2H), 1.68 (m, 2H), 1.38 (s, 9H); C NMR (125 MHz, d₆-DMSO) δ (ppm), 188.50, 172.14, 159.73, 155.59, 151.91, 149.05, 147.12, 135.26, 128.48, 128.10, 115.36, 114.11, 103.96, 77.46, 70.96, 68.25, 38.97, 30.61, 28.22, 24.92; HRMS (M + Na): Calc’d for C₂₄H₂₇F₂NO₆Na: 486.1704. Found: 486.1702.

4-(2-(2,5-difluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 125. Adhesive ppt., 75%; IR (KBr): 3440, 3246, 3100-2800, 1736, 1695, 1610, 1525, 1422, 1398, 1310, 1269, 1200, 1167, 1049, 825, 800, 760, 718. cm⁻¹; H NMR (400 MHz, MeOD) δ (ppm), 7.62 (dd, J = 6.40, J = 4.73, 1H), 6.81 (d, J = 6.75, 1H), 5.26 (d, J = 3.54, 2H), 3.10 (t, J = 7.59, 2H), 2.67 (t, J = 7.08, 2H), 2.05 (m, 2H); C NMR (125 MHz, MeOD) δ (ppm), 188.62, 172.02, 160.69, 158.70, 152.48, 149.40, 147.48, 115.53, 113.04, 104.71, 68.77, 38.55, 29.99, 22.48; F NMR
(376 MHz, MeOD) δ (ppm), -110.81, -141.90; HRMS (M+): Calc’d for C_{12}H_{14}NF_2O_4: 274.0891. Found: 274.0878.

2-(benzyloxy)-5-bromo-1,3-difluorobenzene, 126b. Oil, 98%; IR (NaCl): 3091, 2948, 1595, 1581, 1494, 1454, 1382, 1220, 1190, 1037, 840, 750, 696; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.41, (m, 5H), 7.08 (d, J = 7.70, 2H), 5.17 (s, 2H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 155.74, 136.37, 134.91, 129.60, 118.28, 116.52, 113.03, 109.48, 66.85; ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm), -125.93; HRMS (EI): Calc’d for C_{13}H_{9}F₂BrO: 297.9805. Found: 297.9815.

1-(4-(benzyloxy)-3,5-difluorophenyl)ethanone, 126c. Oil, 91%; IR (NaCl): 3033, 2927, 1689, 1616, 1577, 1508, 1454, 1433, 1360, 1197, 1041, 975, 750, 696; ¹H NMR (400 MHz, CD₃CN) δ (ppm), 7.49 (dd, J = 9.24, 2H), 7.41 (m, 5H), 7.36 (d, 1H), 5.30 (s, 2H) 2.57 (s, 3H); ¹³C NMR (125 MHz, CD₃CN) δ (ppm), 194.88, 156.50, 154.11, 135.99, 131.77, 128.39, 128.18, 113.32, 111.00, 75.48, 25.49; ¹⁹F NMR (376 MHz, CD₃CN + drop CF₃CO₂H) δ (ppm), -127.25; HRMS (EI): Calc’d for C_{15}H_{12}F₂O₂: 262.0805. Found: 262.0816.

1-(4-(benzyloxy)-3,5-difluorophenyl)-2-bromoethanone, 126d. Oil, ~95%; IR (NaCl): 3035, 2922, 1692, 1621, 1577, 1516, 1446, 1425, 1369, 1205, 1041, 965, 749, 696; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.55 (d, J = 6.54, 2H), 7.43 (m, 5H), 5.34 (s, 2H) 4.41 (s, 3H); ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm), -127.88.

2-(4-(benzyloxy)-3,5-difluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino) butanoate, 126e. Adhesive ppt., 93%; IR (KBr) 3302, 3230, 3100-2800, 1742, 1700,
1660, 1621, 1518, 1426, 1407, 1202, 1152, 951, 748, 698, 672 cm⁻¹; ¹H NMR (400 MHz, CD₃CN) δ (ppm), 7.56 (d, J = 6.32, 2H), 7.43 (m, 5H), 5.32 (s, 2H), 5.08 (s, 2H), 3.09 (t, J = 6.83, 2H), 2.45 (t, J = 7.58, 2H), 1.76 (m, 2H), 1.40 (s, 9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 193.45, 172.21, 160.02, 156.10, 151.72, 149.22, 146.32, 135.30, 128.76, 128.18, 115.54, 113.99, 105.01, 76.91, 69.95, 68.01, 39.30, 30.61, 28.20, 24.85; ¹⁹F NMR (376 MHz, CD₃CN) δ (ppm), -127.82; HRMS (M + Na): Calc’d for C₂₄H₂₇F₂(NO₆)Na: 486.1704. Found: 486.1700.

4-(2-(3,5-difluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium2,2,2-trifluoroacetate, 126. White ppt., 82%, Mp: 85–87 °C; IR (KBr): 3435, 3238, 3100–2800, 1740, 1695, 1608, 1524, 1420, 1399, 1310, 1270, 1200, 1165, 1050, 823, 800, 759, 720. cm⁻¹; ¹H NMR (400 MHz, MeOD) δ (ppm), 7.63 (d, J = 7.41, 2H), 5.39 (d, 2H), 3.07 (t, J = 7.33, 2H), 2.64 (t, J = 7.09, 2H), 2.05 (m, 2H); ¹³C NMR (125 MHz, MeOD) δ (ppm), 194.16, 175.94, 157.12, 155.17, 155.12, 144.00, 132.59, 128.13, 115.39, 69.88, 42.47, 33.93, 26.42; ¹⁹F NMR (376 MHz, MeOD) δ (ppm), -133.62; HRMS (M⁺): Calc’d for C₁₂H₁₄F₂NO₄: 274.0891. Found: 274.0884.

1-(4-benzyloxy)-2,6-difluorophenylethanone, 127c. 5-(benzyloxy)-2-bromo-1,3-difluorobenzene, 127b, was generated by the same method as for 122b, determined to be pure by ¹H NMR and GC/MS and utilized in the synthesis of 127c. Oil, 97%; IR (NaCl): 3035, 2930, 1696, 1610, 1571, 1500, 1451, 1433, 1360, 1199, 1043, 978, 748, 695; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.39 (m, 5H), 6.62 (dt, J = 8.07, J = 5.08, 2H), 5.08 (s, 2H) 2.56 (dd, J = 1.85, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 193.78, 163.12, 162.20, 161.02, 134.26, 128.74, 127.74, 111.05, 99.54; ¹⁹F NMR
(376 MHz, CD$_3$CN + drop CF$_3$CO$_2$H) δ (ppm), -108.33; HRMS (EI): Calc’d for C$_{13}$H$_{12}$F$_2$O$_2$: 262.0805. Found: 262.0812.

1-(4-benzyloxy)-2,6-difluorophenyl)-2-bromoethanone, 127d. Oil, ~95%; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 7.45 (m, 5H), 6.61 (dt, $J = 10.47$, $J = 5.32$, 2H), 5.10 (s, 2H), 4.35 (s, 2H).

2-(4-(benzyloxy)-2,6-difluorophenyl)-2-oxoethyl-4-((tert-butoxycarbonyl-amino)-butanoate, 127e. White ppt., 90%, Mp: 69–71°C; IR (KBr): 3304, 3100–2800, 1742, 1703, 1652, 1630, 1437, 1407, 1201, 1153, 1133, 1049, 953, 750, 699, 667, cm$^{-1}$; $^1$H NMR (400 MHz, d$_6$-DMSO) δ (ppm), 7.43 (m, 5H), 6.97 (dd, $J = 11.14$, $J = 4.46$, 1H), 6.89, (t, $J = 5.95$, 1H), 5.22 (s, 2H), 5.08 (s, 2H), 2.95 (t, $J = 6.80$, 2H), 2.41 (t, $J = 7.59$, 2H), 1.65 (m, 2H), 1.37 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 188.75, 172.12, 163.10, 162.68, 160.75, 155.59, 135.51, 128.88, 128.33, 128.06, 106.77, 106.49, 100.06, 76.33, 69.63, 67.69, 38.01, 29.40, 24.70; $^{19}$F NMR (376 MHz, d$_6$-DMSO) δ (ppm), -108.41; HRMS (M + Na): Calc’d for C$_{24}$H$_{27}$F$_2$NO$_6$Na: 486.1704. Found: 486.1682.

4-2-(2,6-difluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 127. White ppt., 86%, Mp: 85–87°C; IR (KBr): 3438, 3245, 3100-2800, 1741, 1694, 1610, 1525, 1420, 1400, 1310, 1268, 1200, 1165, 1050, 825, 800, 758, 720. cm$^{-1}$; $^1$H NMR (400 MHz, MeOD) δ (ppm) 7.41 (d, 1H), 6.54 (d, $J = 13.06$, 2H), 5.10 (s, 2H), 3.09 (t, $J = 7.59$, 2H), 2.59 (t, $J = 7.05$, 2H), 2.00 (m, 2H); $^{13}$C NMR (125 MHz, D$_2$O) δ (ppm) 188.59, 171.98, 163.70, 162.48, 161.56, 128.75,
117.38, 99.68, 68.98, 38.52, 30.07, 22.45; $^{19}$F NMR (376 MHz, D$_2$O) δ (ppm) - 109.55; HRMS (M+): Calc’d for C$_{12}$H$_{14}$F$_2$NO$_4$: 274.0891. Found: 274.0887.

2-(benzyloxy)-5-bromo-1,3,4-trifluorobenzene, 128b. Oil, 94%; IR (NaCl): 3033, 2952, 1629, 1598, 1487, 1475, 1456, 1190, 1089, 1018, 744, 696 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ 7.38 (m, 5H), 7.11-7.09 (qd, J = 2.54, 1H), 5.22 (s, 2H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ 152.73, 150.75, 135.53, 129.06, 128.65, 128.45, 115.38, 114.56, 113.11, 102.41, 98.93, 66.31; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ -131.82, -133.66, -147.45; HRMS (EI): Calc’d for C$_{13}$H$_8$F$_3$BrO: 315.9711. Found: 315.9712.

1-(4-(benzyloxy)-2,3,5-trifluorophenyl)ethanone, 128c. White ppt., 84%, Mp: 45–47 °C; IR (CHCl$_3$): 3002, 2943, 1691, 1627, 1500, 1375, 1346, 1081, 1039, 918, 750, 702 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ 7.45 (m, 6H), 5.37 (s, 2H), 2.68 (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ 194.74, 152.20, 150.23, 147.94, 143.62, 135.58, 128.77, 119.68, 118.10, 115.83, 113.56, 66.45, 31.39; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ -131.10, -136.69, -149.56. HRMS (EI): Calc’d for C$_{15}$H$_{11}$F$_3$O$_2$: 280.0711. Found: 280.0717.

1-(4-(benzyloxy)-2,3,5-trifluorophenyl)-2-bromoethanone, 128d. Oil, ~94%; IR (CHCl$_3$): 3000, 2945, 1698, 1624, 1502, 1369, 1352, 1081, 1045, 921, 750, 699. $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm) 7.53–7.51 (qd, J = 2.30, 1H), 7.40 (m, 5H), 5.25 (d, 2H), 4.45 (s, 2H).

2-(4-(benzyloxy)-2,3,5-trifluorophenyl)-2-oxoethyl-4-(tert-butoxy-carbonyl-amino)butanoate, 128e. White ppt., 93%, Mp: 110–112 °C; IR (KBr) 3302, 3100-
2800, 1743, 1703, 1651, 1629, 1437, 1407, 1202, 1170, 1147, 950, 748, 698, 667 cm−1; 1H NMR (400 MHz, d6-DMSO) δ (ppm), 7.62-7.60 (qd, J = 2.30, 1H), 7.41 (m, 5H), 6.88 (t, J = 6.00, 1H), 5.40 (s, 2H), 5.26 (s, 2H), 2.96 (t, J = 6.82, 2H), 2.44 (t, J = 7.60, 2H), 1.67 (m, 2H), 1.38 (s, 9H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 188.33, 172.09, 155.60, 151.48, 149.53, 148.47, 146.44, 144.63, 142.83, 140.05, 128.55, 116.87, 110.58, 77.46, 75.71, 68.22, 30.53, 28.21, 24.89; HRMS (M + Na): Calc’d for C24H26F3NO6Na: 504.1610. Found: 504.1593.

4-oxo-4-(2-oxo-2-(2,3,5-trifluoro-4-hydroxyphenyl) ethoxy) butan-1-aminium 2,2,2-trifluoroacetate, 128. White ppt., 90%, Mp: 131–133 °C; IR (KBr): 3434, 3244, 3100-2800, 1738, 1695, 1610, 1523, 1420, 1400, 1309, 1267, 1199, 1166, 1049, 824, 800, 759, 720 cm−1; 1H NMR (400 MHz, D2O) δ (ppm), 7.52-7.50 (qd, J = 2.35, 1H), 5.37 (d, 2H), 3.14 (t, J = 7.06, 2H), 2.72 (t, J = 6.71, 2H), 2.07 (m, 2H); 13C NMR (125 MHz, D2O) δ (ppm), 190.89, 174.05, 163.12, 149.21, 147.28, 141.03, 139.81, 117.19, 115.16, 109.89, 69.88, 42.47, 33.93, 26.42; 19F NMR (376 MHz, D2O) δ (ppm), -136.48, -137.81, -156.79; HRMS (M+): Calc’d for C12H13F3NO4: 292.0797. Found: 292.0794.

1-(benzyloxy)-4-bromo-2,3,5,6-tetrafluorobenzene, 129b. White ppt., 96%, Mp: 43–45 °C; IR (CHCl3): 3002, 2943, 1560, 1458, 1442, 1407, 1380, 1039, 918, 750; 1H NMR (400 MHz, CDCl3) δ (ppm) 7.39 (m, 5H), 5.28 (s, 2H); 13C NMR (125 MHz, CDCl3) δ (ppm) 146.18, 144.22, 144.11, 140.70, 135.25, 129.04, 128.72, 93.26, 76.51; 19F NMR (376 MHz, CDCl3 + drop CF3CO2H) δ (ppm) -131.37, -155.12. HRMS (El): Calc’d for C13H7F4BrO: 333.9616. Found: 333.9606.
1-(4-(benzyloxy)-2,3,5,6-tetrafluorophenyl)ethanone, 129c. A solution of 129b (1.00 g, 2.98 mmol), tributyl(1-ethoxyvinyl)stannane (1.44 ml, 4.27 mmol) and 25 ml toluene was degassed for 30 min while stirring. Tetrakistriphenylphosphine palladium (0) (205 mg, 0.178 mmol) was then added and the solution heated to 100 °C overnight with stirring. Fifty ml of 50% HCl (aq) were added and the solution stirred for 6 h, followed by filtration through Celite that effectively removed the palladium black. The resulting toluene filtrate was charged with 50 ml of saturated aqueous potassium fluoride and continuously stirred for 12 h. The insoluble tributylfluorostannane was then filtered, the aqueous layer removed, and the organic layer concentrated. Flash column chromatography (6:1 Hex/Et₂O) afforded 1-(4-(benzyloxy)-2,3,5,6-tetrafluorophenyl)ethanone, 129c, in 85% yield (755 mg) as a faint yellow solid. Mp: 47–49 °C; IR (CDCl₃): 2921, 2852, 1712, 1643, 1461, 1377, 1190, 1100, 742, 723, 696 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.38 (m, 5H), 5.27 (s, 2H), 2.59 (t, J = 1.90, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm) 192.07, 146.39, 143.95, 142.52, 139.89, 135.28, 129.27, 128.46, 66.07, 32.70; ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm) -144.22, -161.88; HRMS (M + H): Calc’d for C₁₅H₁₁F₄O₂: 299.0695. Found: 299.0702.

1-(4-(benzyloxy)-2,3,5,6-tetrafluorophenyl)-2-bromoethanone, 129d. Oil, ~99%; IR (CHCl₃): 2919, 2856, 1721, 1645, 1453, 1379, 1190, 1107, 745, 715, 696; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.40 (m, 5H), 5.35 (s, 2H), 4.32 (t, J = 1.20, 2H).

2-(4-(benzyloxy)-2,3,5,6-tetrafluorophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)butanoate, 129e. White ppt., 87%, Mp: 106–108 °C; IR (KBr) 3301, 3100-
2800, 1744, 1703, 1655, 1615, 1487, 1437, 1407, 1210, 1153, 1050, 749, 700, 667 cm$^{-1}$; $^1$H NMR (400 MHz, d$^6$-DMSO) δ (ppm) 7.42 (m, 5H), 6.88, (t, $J = 5.92$, 1H), 5.43 (s, 2H), 5.02 (t, $J = 1.43$, 2H), 2.93 (t, $J = 6.74$, 2H), 2.41 (t, $J = 7.57$, 2H), 1.65 (m, 2H), 1.37 (s, 9H); $^{13}$C NMR (125 MHz, d$^6$-DMSO) δ (ppm), 188.04, 172.14, 155.59, 145.97, 143.96, 141.32, 139.47, 135.39, 128.64, 128.39, 109.10, 77.46, 76.14, 68.81, 30.39, 28.82, 24.82; HRMS (M+ Na): Calc’d for C$_{24}$H$_{25}$F$_4$NO$_6$Na: 522.1516. Found: 522.1493.

4-oxo-4-(2-oxo-2-(2,3,5,6-tetrafluoro-4-hydroxyphenyl)ethoxy)butan-1-aminium-2,2,2-trifluoroacetate, 129. White ppt., 83%, Mp: 107–109 °C; IR (KBr): 3440, 3241, 3100–2800, 1735, 1690, 1610, 1525, 1420, 1400, 1310, 1265, 1200, 1166, 1050, 824, 800, 759, 720. cm$^{-1}$; $^1$H NMR (400 MHz, D$_2$O) δ (ppm), 5.25 (t, $J = 1.67$, 2H), 3.07 (t, $J = 7.61$, 2H), 2.66 (t, $J = 7.06$, 2H), 2.01 (m, 2H); $^{13}$C NMR (125 MHz, D$_2$O) δ (ppm), 189.10, 173.25, 162.36, 147.43, 139.18, 129.31, 117.78, 113.12, 104.46, 69.40, 38.59, 30.13, 22.22; $^{19}$F NMR (376 MHz, D$_2$O) δ (ppm), -142.76, -163.05; HRMS (M+): Calc’d for C$_{12}$H$_{12}$F$_4$NO$_4$: 310.0702. Found: 310.0701.

1-benzyloxy-4-bromo-2-trifluoromethylbenzene, 130b. Oil, 93%; IR (NaCl): 3033, 2935, 1602, 1490, 1454, 1139, 1053, 908, 811, 732 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 7.71 (d, $J = 2.56$, 1H), 7.56 (d, $J = 2.56$, 1H), 7.38 (m, 5H), 6.90 (d, $J = 8.91$, 1H), 5.19 (s, 2H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm), 155.66, 135.89, 130.29, 129.19, 128.58, 127.02, 124.31, 121.59, 120.81, 115.22, 112.45; $^{19}$F NMR (376 MHz, CDCl$_3$ + drop CF$_3$CO$_2$H) δ (ppm), -63.49; HRMS (EI): Calc’d for C$_{14}$H$_{10}$F$_3$BrO: 329.9867. Found: 329.9850.
1-(4-benzyloxy-3-trifluoromethylphenyl)ethanone, **130c**. White ppt., 84%, Mp: 119-121 °C; IR (CHCl₃): 3055, 2987, 2829, 1685, 1610, 1421, 1265, 1141, 896, 738, 696 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.23 (d, J = 2.10, 1H), 8.11 (dd, J = 8.78, J = 2.29, 1H), 7.44 (m, 5H), 7.09 (d, J = 8.80, 1H), 5.29 (s, 2H), 2.59 (s, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 201.19, 159.89, 135.05, 129.10, 128.5, 127.04, 124.49, 121.78, 119.35, 116.09, 113.26, 110.42, 71.02, 26.11; ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm), -63.77; HRMS (EI): Calc’d for C₁₆H₁₄F₃O₂: 295.0946. Found: 295.0927.

1-(4-benzyloxy-3-trifluoromethylphenyl)-2-bromoethanone, **130d**. Oil, ~93%; IR (CHCl₃): 3058, 2989, 2821, 1687, 1605, 1421, 1255, 1139, 892, 740, 690 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.90 (dd, J = 8.96, J = 2.15, 2H), 7.43 (m, 5H), 7.06 (d, J = 8.90, 1H), 5.31 (s, 2H), 4.39 (s, 2H); ¹⁹F NMR (376 MHz, CDCl₃ + drop CF₃CO₂H) δ (ppm), -63.81.

2-(4-benzyloxy-3-trifluoromethylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)butanoate, **130e**. White ppt., 87%, Mp: 112-114°C; IR (KBr) 3304, 3100-2800, 1741, 1701, 1659, 1612, 1508, 1437, 1407, 1218, 1167, 1137, 1053, 1008, 741, 699, 667; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.26 (dd, J = 8.64, J = 2.20, 1H), 8.15 (d, J = 1.50, 1H), 7.43 (m, 5H), 7.34 (t, J = 7.15, 1H), 6.88 (t, J = 7.25, 1H), 5.48 (s, 2H), 5.41 (s, 2H), 2.98 (t, J = 6.74, 2H), 2.44 (t, J = 7.63, 2H), 1.70 (m, 2H), 1.38 (s, 9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ(ppm), 190.79, 172.18, 159.87, 155.60, 135.73, 133.10, 132.00, 131.42, 128.69, 127.94, 126.44, 124.27, 122.10, 119.93, 117.63, 114.27, 77.45, 70.21, 66.16, 38.97, 28.22, 26.47, 24.96; ¹⁹F NMR (376 MHz,
CD$_3$CN + drop CF$_3$CO$_2$H) $\delta$ (ppm), -61.41; HRMS (M + Na): Calc’d for C$_{25}$H$_{28}$F$_3$NO$_6$Na: 518.1766. Found 518.1756.

2-(4-hydroxy-3-(trifluoromethyl)phenyl)-2-oxoethyl4-(tert-butoxy-carbonyl)-aminobutanoate, 130e’. The general method of Theodorakis$^{146}$ was employed. To a solution of 130e (1.10g, 2.22 mmol) in 25 ml EtOAc was added 220 mg of 10% Pd/C, and the heterogenous mixture stoppered with a septum. While stirring, hydrogen (H$_2$) was infused through a needle into the solution. The reaction was completed ~ 2h as indicated by TLC. The residual Pd/C was filtered off through Celite and the organic phase concentrated under reduced pressure to yield 875 mg of 130e’ as an adhesive precipitate (97%); $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 11.60 (s, 1H), 8.10 (dd, $J = 8.80, J = 2.10$, 1H), 7.25 (t, $J = 8.00$, 1H), 7.15 (d, $J = 7.70$, 1H), 6.90 (t, $J = 7.20$, 1H), 5.48 (s, 2H) 2.96 (t, $J = 6.90$, 2H), 2.42 (t, $J = 7.50$, 2H), 1.66 (m, 2H), 1.38 (s, 9H).

4-(2-(4-hydroxy-3-trifluoromethylphenyl)-2-oxoethoxy)-4-oxobutan-1-amonium 2,2,2-trifluoroacetate, 130. A modified preparation of Marsh’s$^{13}$ technique was utilized. To a round bottomed flask containing 130e’ (800 mg, 1.97 mmol) was added 20 ml of 1:1 TFA/CH$_2$Cl$_2$. The reaction persisted for 15 min while stirring and at room temperature. At this time, excess solvent was removed under reduced pressure, washed with 50 ml of 1:1 CH$_2$Cl$_2$/H$_2$O and the water layer retained and lyophilized to afford 130 as a white solid, (72%); Mp: 129-131 °C; IR (KBr): 3439, 3271, 3100-2800, 1742, 1692, 1605, 1516, 1431, 1380, 1311, 1253, 1222, 1200, 1166, 1057, 822, 760 cm$^{-1}$; $^1$H NMR (400 MHz, D$_2$O) $\delta$ (ppm), 8.25 (d, $J = 2.10$, 1H), 5.48 (s, 2H) 2.96 (t, $J = 6.90$, 2H), 2.42 (t, $J = 7.50$, 2H), 1.66 (m, 2H), 1.38 (s, 9H).
8.12 (d, J = 9.29, 1H), 7.20 (d, J = 8.98, 1H), 5.55 (s, 2H), 3.15 (t, J = 8.05, 2H), 2.73 (t, J = 9.29, 2H), 2.06, (m, 2H); \(^{13}\)C NMR (125 MHz, MeOD) \(\delta\) (ppm), 191.05, 172.35, 161.92, 161.05, 133.82, 128.25, 125.01, 122.32, 120.16, 117.42, 115.71, 65.97, 38.51, 30.09, 22.38; HRMS (M+): Calc’d for C\(_{13}\)H\(_{15}\)F\(_3\)NO\(_4\): 306.0953. Found 306.0953.

The same steps were utilized to make 131 and 132.

1-(benzyloxy)-4-bromo-2-(trifluoromethoxy)benzene, 131b. Oil, 93%; IR (NaCl): 3033, 2916, 1595, 1500, 1454, 1382, 1190, 1135, 1024, 804, 736, 696. \(^1\)H NMR (400 MHz, CDCl\(_3\)) \(\delta\) 7.36 (m, 8H), 6.88 (d, J = 8.9, 1H), 5.15 (s, 2H); \(^{13}\)C NMR (125 MHz, CDCl\(_3\)) \(\delta\) 150.33, 138.80, 135.79, 129.06, 127.13, 121.61, 119.55, 117.49, 115.45, 112.36, 110.92, 71.13; \(^{19}\)F NMR (376 MHz, CDCl\(_3\) + drop CF\(_3\)CO\(_2\)H) \(\delta\) -59.08; HRMS (EI): Calc’d for C\(_{14}\)H\(_{10}\)F\(_3\)BrO\(_2\): 345.9816. Found: 345.9812.

1-(4-benzyloxy)-3-(trifluoromethoxy)phenyl)ethanone, 131c. Oil, 92%; IR (NaCl): 3035, 2929, 1681, 1606, 1581, 1514, 1454, 1427, 1380, 1205, 1170, 1135, 813, 738, 696; \(^1\)H NMR (400 MHz, CDCl\(_3\)) \(\delta\) 7.87 (dd, J = 8.88, J = 2.01, 2H), 7.43 (m, 5H), 7.26 (d, J = 8.57, 1H), 5.25 (s, 2H) 2.56 (s, 3H); \(^{13}\)C NMR (125 MHz, CDCl\(_3\)) \(\delta\) 195.65, 154.73, 137.40, 135.84, 129.34, 127.44, 122.68, 119.53, 117.49, 115.74, 113.85, 111.20, 70.64, 25.56; \(^{19}\)F NMR (376 MHz, CDCl\(_3\) + drop CF\(_3\)CO\(_2\)H) \(\delta\) -58.69; HRMS (EI): Calc’d for C\(_{16}\)H\(_{13}\)F\(_3\)O\(_3\): 310.0817. Found: 310.0828.

1-(4-benzyloxy-3-trifluoromethoxyphenyl)-2-bromoethanone, 131d. Oil, ~93%; IR (CDCl\(_3\)): 3035, 2928, 1680, 1606, 1580, 1516, 1444, 1423, 1371, 1215, 1150, 1125, 827, 730, 692; \(^1\)H NMR (400 MHz, CDCl\(_3\)) \(\delta\) (ppm), 8.26 (d, J = 2.00, 1H),
8.14 (dd, \( J = 8.74, J = 2.20 \), 1H), 7.41 (m, 5H), 7.12 (d, \( J = 8.58 \), 1H), 5.26 (s, 2H) 4.37 (s, 2H); \(^{19}\)F NMR (376 MHz, CDCl\(_3\) + drop CF\(_3\)CO\(_2\)H) \( \delta \) (ppm), -58.73.

2-(4-benzyloxy-3-trifluoromethoxy)phenyl-2-oxoethyl-4-(\(t\)-butyoxycarbonyl-amino)butanoate, 131e. White ppt., 85\%, Mp: 112-114\(^\circ\)C; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \( \delta \) (ppm), 8.03 (dd, \( J = 8.98, J = 2.20 \), 1H), 7.89 (t, \( J = 2.63 \), 1H), 7.42 (m, 5H), 6.88 (t, \( J = 6.30 \), 1H), 5.44 (s, 2H) 5.35 (s, 2H) 2.98 (t, \( J = 6.79 \), 2H), 2.44 (t, \( J = 7.59 \), 2H), 1.68 (m, 2H), 1.38 (s, 9H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) \( \delta \) (ppm), 190.79, 172.18, 159.87, 155.60, 134.40, 133.10, 132.00, 128.91, 127.23, 126.44, 124.27, 122.10, 119.93, 117.38, 114.27, 77.45, 70.32, 66.16, 39.47, 30.62, 28.22, 24.96; \(^{19}\)F NMR (376 MHz, CD\(_3\)CN + drop CF\(_3\)CO\(_2\)H) \( \delta \) (ppm), -61.14; HRMS (M + Na): Calc’d for C\(_{25}\)H\(_{28}\)F\(_{3}\)NO\(_7\)Na: 534.1716. Found 534.1712.

2-(4-hydroxy-3-(trifluoromethoxy)phenyl)-2-oxoethyl-4-(\(t\)-butyoxycarbonyl-amino)butanoate, 131e. White ppt., 97\%; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \( \delta \) (ppm), 7.87 (dd, \( J = 8.60, J = 2.20 \), 1H), 7.81 (s, 1H), 7.11 (d, \( J = 8.62 \), 1H), 6.89 (t, \( J = 5.50 \), 1H), 5.38 (s, 2H), 2.97 (t, \( J = 6.80 \), 2H), 2.42 (t, \( J = 7.65 \), 2H), 1.70 (m, 2H), 1.38 (s, 9H).

4-(2-(4-hydroxy-3-trifluoromethoxyphenyl)-2-oxoethoxy)-4-oxobutan-1-ammonium-2,2,2-trifluoroacetate, 131. White ppt., 96\%, Mp: 121-123 \(^\circ\)C. IR (KBr): 3439, 3271, 3100-2800, 1742, 1692, 1605, 1516, 1431, 1380, 1311, 1253, 1222, 1200, 1166, 1057, 822, 760 cm\(^{-1}\); \(^1\)H NMR (400 MHz, D\(_2\)O) \( \delta \) (ppm), 7.90 (t, \( J = 1.50 \), 1H), 7.85 (dd, \( J = 8.82, J = 2.13 \), 1H), 7.14 (d, \( J = 8.80 \), 1H), 5.47 (s, 2H),
3.08 (t, \( J = 7.65, 2H \)), 2.68 (t, \( J = 7.08, 2H \)), 2.02 (m, 2H); \(^{13}\)C NMR (125 MHz, D\(_2\)O) \( \delta \) (ppm), 193.76, 174.15, 163.12, 154.43, 136.51, 132.19, 129.16, 125.96, 123.47, 121.99, 119.37, 117.73, 66.67, 38.53, 30.23, 21.96; HRMS (M+): Calc’d for C\(_{13}\)H\(_{15}\)F\(_3\)NO\(_5\): 322.0902. Found: 322.0905.

**1-(4-(benzyloxy)-3-nitrophenyl)ethanone, 132c.** 3-nitro-4-hydroxyacetophenone is commercially available. 96\%. *This compound is known.* \(^1\)H NMR (400 MHz, CDCl\(_3\)) \( \delta \) (ppm), 8.45 (dd, \( J = 9.22, J = 2.38, 1H \)), 8.14 (dd, \( J = 9.15, J = 1.69, 1H \)), 7.42 (m, 6H), 5.34 (s, 2H), 2.60 (s, 2H).

**1-(4-(benzyloxy)-3-nitrophenyl)-2-bromoethanone, 132d.** Oil, \( \sim97\% \); \(^1\)H NMR (400 MHz, CDCl\(_3\)) \( \delta \) (ppm), 8.34 (s, 1H), 8.00 (d, 1H), 7.28 (m, 6H), 5.20 (s, 2H), 4.23 (s, 2H).

**2-(4-(benzyloxy)-3-nitrophenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)butanoate, 132e.** White ppt., 67\%, Mp: 144-146\(^\circ\)C. IR (KBr): 3310, 3100-2800, 1740, 1700, 1655, 1603, 1433, 1400, 1204, 1170, 955, 745, 697, 665; \(^1\)H NMR (400 MHz, CD\(_3\)CN) \( \delta \) (ppm), 8.37 (dd, \( J = 7.12, J = 2.57, 1H \)), 8.12 (d, \( J = 1.50, 1H \)), 7.43 (m, 6H), 6.85 (t, \( J = 7.21, 1H \)), 5.34 (s, 2H), 3.09 (t, \( J = 6.79, 2H \)), 2.46 (t, \( J = 7.60, 2H \)), 1.77 (m, 2H), 1.40 (s, 9H); \(^{13}\)C NMR (125 MHz, CD\(_3\)CN) \( \delta \) (ppm), 195.22, 173.42, 160.65, 158.77, 139.23, 136.54, 135.40, 133.98, 131.78, 128.36, 125.89, 119.24, 116.82, 66.72, 38.44, 30.51, 29.11, 21.88; HRMS (M + Na): Calc’d for C\(_{24}\)H\(_{28}\)N\(_2\)O\(_8\)Na: 495.1743. Found: 495.1755

**4-(2-(4-hydroxy-3-nitrophenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 132.** White ppt., 78\%, Mp: 76-78\(^\circ\)C; IR (KBr): 3441, 3265, 3100-
2800, 1740, 1690, 1605, 1520, 1430, 1377, 1309, 1252, 1225, 1199, 1166, 1055, 820, 760. cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm), 8.72 (d, J = 2.16, 1H), 8.20 (d, J = 9.29, 1H), 7.32 (d, J = 8.85, 1H), 5.56 (s, 2H), 3.14 (t, J = 7.80, 2H), 2.72 (t, J = 9.18, 2H), 2.08 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm), 193.16, 174.10, 163.13, 157.81, 135.75, 134.33, 126.61, 120.50, 117.44, 115.12, 66.69, 38.54, 30.23, 21.96; HRMS (M +): Calc’d for C₁₂H₁₅N₂O₆: 283.0925. Found: 283.0914.

4-(2-(4-hydroxy-3,5-dimethoxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 133. The synthesis of 133 was accomplished from use of the procedure by Conrad, et. al.⁹⁷

4-(2-(4-hydroxy-3-methoxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 134. The synthesis of 134 was accomplished from use of the procedure by Conrad, et. al.⁹⁷

1-(2,4-bis(benzyloxy)phenyl)ethanone, 135b. 2,4-dihydroxyacetophenone is commercially available. Benzylolation of this utilized the same procedure as in the synthesis of 122b. White ppt., 97%. This compound is known. ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.86 (t, J = 7.63, 1H), 7.39 (m, 11H), 6.64 (dd, J = 8.71, J = 2.33, 1H), 5.12 (s, 2H), 5.10 (s, 2H), 2.57 (s, 3H).

1-(2,4-bis(benzyloxy)phenyl)-2-bromoethanone, 135c. An identical procedure was used as to construct 122d. ~ 94%. ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.78 (s,1H), 7.35 (m, 10H), 6.60 (d, 2H), 5.12 (s, 2H), 5.09 (s, 2H), 4.51 (s, 1H), 4.46 (1H).

2-(2,4-bis(benzyloxy)phenyl)-2-oxoethyl 4-(tert-butoxycarbonylamino)butanoate, 135d. The same procedure was utilized as to synthesize 122e. Adhesive ppt, 48%; IR
(KBr):  3310, 3100-2800, 1743, 1700, 1658, 1601, 1436, 1400, 1200, 1165, 955, 745, 742, 700, 697, 665, 651;  $^1$H NMR (400 MHz, d$^6$-DMSO) $\delta$ (ppm),  7.93 (t, $J = 8.79$, 1H), 7.43 (m, 10H), 7.14 (dd, $J = 8.74$, $J = 2.15$, 1H), 6.88 (t, $J = 5.70$, 1H), 6.62 (d, $J = 4.66$, 1H), 5.37 (s, 2H), 5.29 (s, 1H), 5.21 (s, 1H), 5.10 (s, 2H), 2.94 (t, $J = 6.63$, 2H), 2.37 (t, $J = 7.51$, 2H), 1.76 (m, 2H), 1.37 (s, 9H); $^{13}$C NMR (125 MHz, d$^6$-DMSO) $\delta$ (ppm),  190.50, 172.15, 164.65, 159.78, 155.58, 135.99, 132.04, 128.35, 127.51, 118.10, 107.59, 103.90, 100.33, 77.43, 70.72, 69.19, 39.95, 30.68, 28.22, 24.88; HRMS (M + H):  Calc’d for C$_{31}$H$_{36}$NO$_7$:  534.2492.  Found: 534.2504.

2-(2,4-dihydroxyphenyl)-2-oxoethyl-4-(tert-butoxycarbonylamino)butanoate, 135e.  An identical procedure was used as to engender 130’e. Adhesive ppt., 97%; $^1$H NMR (400 MHz, d$^6$-DMSO) $\delta$ (ppm), 11.43, (s, 1H), 11.34 (s, 1H), 7.88 (s,1H), 7.70 (d, 1H), 6.87 (t, 1H), 6.38 (d, 1H), 5.28 (s, 1H), 2.96 (t, 2H), 2.40 (t, 2H), 1.67 (m, 2H), 1.38 (s, 9H).

4-(2-(2,4-dihydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 135.  An identical procedure was used as to produce 130. White ppt., 85%, Mp: 136-138 °C; IR (KBr):  3432, 3274, 3100-2800, 1739, 1691, 1602, 1520, 1421, 1377, 1309, 1298, 1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613 cm$^{-1}$; $^1$H NMR (400 MHz, D$_2$O) $\delta$ (ppm), 7.97 (d, $J = 6.81$, 1H), 7.72 (d, $J = 8.84$, 1H), 6.53 (d, $J = 7.63$, 1H), 5.48 (s, 1H), 5.43 (s, 1H), 3.14 (t, $J = 7.64$, 2H), 2.72 (t, $J = 7.05$, 2H), 2.08 (m, 2H); $^{13}$C NMR (125 MHz, D$_2$O) $\delta$ (ppm), 193.77, 174.15, 162.75, 151.02, 144.22, 125.92, 122.70, 119.65, 117.37, 115.58, 66.60, 38.51, 30.28, 22.00; HRMS (M +):  Calc’d for C$_{12}$H$_{16}$NO$_5$:  254.1028.  Found: 254.1038.
The same methodology was utilized to assemble 136-141.

1-(3,4-bis(benzyloxy)phenyl)ethanone, 136b. 3,4-dihydroxyacetophenone is commercially available. Benzylaion of this utilized the same procedure as in the synthesis of 127b. 98%. This compound is known. $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 7.63 (dd, $J = 8.77$, $J = 2.32$, 1H), 7.40 (m, 11H), 6.94 (dd, $J = 9.10$, $J = 1.90$, 1H), 5.25 (s, 2H), 5.22 (s, 2H), 2.52 (s, 3H).

1-(3,4-bis(benzyloxy)phenyl)-2-bromoethanone, 136c. Oil ~96%; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 7.68 (s, 1H), 7.41 (m, 11H), 6.96 (d, 1H), 5.24 (s, 2H), 5.22 (s, 2H), 4.46 (s, 2H).

2-(3,4-bis(benzyloxy)phenyl)-2-oxoethyl-4-(tert-butoxy-carbonylamino)-butanoate, 136d. White ppt., 90%, Mp: 112-114°C; IR (KBr): 3310, 3100-2800, 1743, 1700, 1658, 1601, 1436, 1400, 1200, 1165, 955, 745, 742, 700, 697, 665, 651; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.62 (dd, $J = 7.25$, $J = 2.38$, 1H), 7.56 (d, $J = 1.72$, 1H), 7.38 (m, 10H), 7.19 (d, $J = 8.13$, 1H), 6.92 (t, $J = 7.21$, 1H), 5.40 (s, 2H), 5.27 (s, 2H), 5.21 (s, 2H), 2.97 (t, $J = 6.78$, 2H), 2.40 (t, $J = 7.70$, 2H), 1.70 (m, 2H), 1.38 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 197.09, 171.80, 155.60, 152.85, 147.98, 136.85, 128.41, 127.94, 127.58, 126.84, 122.64, 113.04, 112.51, 77.47, 70.00, 66.07, 39.61, 30.67, 28.22, 24.54; HRMS (M + H): Calc’d for C$_{31}$H$_{36}$NO$_7$: 534.2492. Found: 534.2484.

2-(2,4-dihydroxyphenyl)-2-oxoethyl-4-(tert-butoxycarbonylamino)butanoate, 136e. Adhesive ppt., 98%; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.32 (d, 2H),
6.93 (s, 1H), 6.89 (t, 1H), 5.39 (s, 2H), 2.98 (t, 2H), 2.38 (t, 2H), 1.68 (m, 2H), 1.40 (s, 9H).

4-(2-(3,4-dihydroxyphenyl)-2-oxoethoxy)-4-oxobut-1-aminium-2,2,2-trifluoroacetate, 136. White ppt., 92%, Mp: 56-58°C; IR (KBr): 3432, 3274, 3100-2800, 1739, 1691, 1602, 1421, 1377, 1309, 1298, 1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613 cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm), 7.35 (s, 1H), 7.33 (s, 1H), 6.91 (d, J = 7.10, 1H), 5.36 (s, 2H), 3.07 (t, J = 6.80, 2H), 2.65 (t, J = 7.76, 2H), 2.00 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm), 194.29, 174.13, 163.06, 150.88, 144.11, 125.88, 122.69, 119.74, 117.42, 115.10, 66.56, 38.30, 30.23, 21.95; HRMS (M +): Calc’d for C₁₂H₁₆NO₅: 254.1028. Found: 254.1032.

1-(4-(benzyloxy)-3-methyl-phenyl)ethanone, 137b. 3-methyl-4-dihydroxyacetophenone is commercially available. Benzylation of this comprised the same procedure as in the synthesis of 122b. white ppt., 95%. This compound is known. ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.82 (s 1H), 7.80 (s, 1H), 7.40 (m, 5H), 6.90 (d, J = 8.75, 1H), 5.17 (s, 2H), 2.56 (s, 3H), 2.33 (s, 3H).

1-(4-(benzyloxy)-3-methylphenyl)-2-bromoethanone, 137c. Oil, ~97%; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.82 (d, 2H), 7.40 (m, 5H), 6.88 (d, J = 8.80,1H), 5.19 (s, 2H), 4.33 (s, 2H), 2.33 (s, 3H).

2-(4-(benzyloxy)-3-methylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)butanoate, 137d. White ppt., 94%, Mp: 105-106 °C; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 7.82 (d, J = 9.14, 2H), 7.40 (m, 5H), 7.14 (d, J = 8.10,
1H), 6.87 (t, J = 7.40, 1H), 5.39 (s, 2H), 5.25 (s, 2H), 2.96 (t, J = 6.81, 2H), 2.42 (t, J = 7.62, 2H), 2.24 (s, 3H), 1.70 (m, 2H), 1.38 (s, 9H); \(^{13}\text{C NMR}\) (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 191.14, 172.20, 161.40, 155.60, 136.67, 131.00, 130.10, 129.26, 128.50, 127.34, 126.45, 125.07, 111.62, 77.45, 69.55, 66.03, 39.33, 30.68, 28.18, 24.98, 16.04; HRMS (M + Na): Calc’d for C\(_{25}\)H\(_{31}\)NO\(_6\)Na: 464.2049. Found: 464.2017.

2-(4-hydroxy-3-methylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 137e. Adhesive ppt., 98%; \(^1\text{H NMR}\) (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 10.76 (s, 1H), 7.86 (d, J = 9.26, 2H), 7.66 (d, J = 8.04, 1H), 6.89 (t, 1H), 5.38 (s, 2H) 2.96 (t, J = 6.94, 2H), 2.42 (t, J = 7.88, 2H), 2.24 (s, 3H), 1.70 (m, 2H), 1.38 (s, 9H).

4-(2-(4-hydroxy-3-methylphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 137. White ppt., 89%, Mp: 118-120\(^\circ\)C. IR (KBr): 3432, 3274, 3100-2800, 1739, 1691, 1602, 1520, 1421, 1377, 1309, 1298, 1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613. cm\(^{-1}\); \(^1\text{H NMR}\) (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 7.86 (s, 3H), 7.72 (d, J = 6.22, 1H), 7.67 (dd, J = 9.03, J = 1.79, 1H), 5.39 (s, 1H), 2.88 (t, J = 6.97, 2H), 2.56 (t, J = 7.55, 2H), 2.16 (s, 3H), 1.87 (m, 2H); \(^{13}\text{C NMR}\) (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.67, 171.72, 160.98, 158.27, 130.76, 127.76, 125.10, 120.30, 117.94, 114.76, 66.11, 38.09, 30.15, 22.10, 15.80; HRMS (M+): Calc’d for C\(_{13}\)H\(_{18}\)NO\(_4\): 252.1230. Found: 252.1212.

1-(4-(benzyloxy)-3,5-dimethylphenyl)ethanone, 138b. 4-hydroxy-3,3-dimethyl-acetophenone is commercially available. Benzylation of this involve the same procedure as in the synthesis of 135b. Oil, 95%. This compound is known. GC/MS
(EI): One peak, retention time, 8.38 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 254.0 m/z.

1-(4-(benzyloxy)-3,5-dimethylphenyl)-2-bromoethanone, 138c. White ppt., ~95%.
This compound is known. GC/MS (EI): One peak, retention time, 9.87 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 331.9, 333.9 m/z.

2-(4-(benzyloxy)-3,5-dimethylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 138d. Adhesive ppt., 96%; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; 1H NMR (400 MHz, d6-DMSO) δ (ppm), 7.70 (d, J = 8.60, 1H), 7.50-7.40 (m, 5H), 6.89 (t, J = 7.20, 1H), 5.42 (s, 2H), 4.88 (s, 2H), 2.97 (t, J = 6.90, 2H), 2.43 (t, J = 7.80, 2H), 2.29 (s, 6H), 1.68 (m, 2H), 1.36 (s, 9H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 191.84, 172.19, 160.04, 155.60, 137.06, 131.49, 129.59, 128.68, 128.36, 128.13, 127.85, 77.45, 73.52, 66.20, 38.97, 30.50, 28.18, 24.97, 16.20; HRMS (M + Na): Calc’d for C26H33NO6Na: 478.2206; Found: 478.2191. (M + H): Calc’d for C26H34NO6: 456.2386. Found: 456.2388.

2-(4-hydroxy-3,5-dimethylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 138e. Adhesive ppt., 99%; IR (KBr): 3548, 3301, 3100-2800, 1739, 1702, 1661, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; 1H NMR (400 MHz, d6-DMSO) δ (ppm), 9.40 (s, 1H), 7.59 (d, J = 8.60, 2H), 6.89 (t, J = 7.15, 1H), 5.36 (s, 2H), 2.98 (t, J = 7.00, 2H), 2.55 (t, J = 7.50, 2H), 2.20 (s, 6H), 1.92 (m, 2H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 190.91, 172.20, 159.67, 155.59, 128.60,
125.25, 123.74, 77.44, 65.91, 38.96, 30.90, 28.18, 24.98, 16.52; HRMS (M + Na): Calc’d for C_{19}H_{27}NO_{6}Na: 388.1736. Found: 388.1735.

4-(2-(4-hydroxy-3,5-dimethylphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 138. White ppt., 92%, Mp: 125-127°C; IR (KBr): 3550, 3301, 3100-2800, 1739, 1702, 1661, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 9.36 (s, 1H), 7.86 (s, 3H), 7.62 (d, \(J = 8.60\), 2H), 5.36 (s, 2H), 2.98 (t, \(J = 7.00\), 2H), 2.43 (m, 5H), 2.22 (s, 6H), 1.69 (m, 2H), 1.37 (s, 9H); \(^13\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 189.74, 170.61, 157.71, 156.71, 127.72, 124.97, 123.15, 116.95, 65.02, 37.10, 28.88, 21.48, 15.55; HRMS (M+): Calc’d for C\(_{14}\)H\(_{20}\)NO\(_4\): 266.1392. Found: 266.1389.

1-(4-(benzyloxy)2-methylphenyl)ethanone, 139b. 4-hydroxy-2-methylacetophenone is commercially available. Benzylaion of this entailed the same procedure as in the synthesis of 135b. White ppt., 93%, This compound is known. GC/MS (EI): One peak, retention time, 8.24 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 240.0 m/z.

1-(4-(benzyloxy)-2-methylphenyl)-2-bromoethanone, 139c. White ppt., ~92%. This compound is known. GC/MS (EI): One peak, retention time, 9.77 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 317.9, 319.9 m/z.

2-(4-(benzyloxy)-2-methylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 139d. Adhesive ppt., 91%; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 7.86 (d, \(J = 9.10\), 1H), 7.47-7.40 (m, 5H), 6.97 (t, \(J = 6.50\), 2H),
6.87 (t, J = 7.35, 1H), 5.28 (s, 2H), 5.19 (s, 2H), 2.97 (t, J = 6.70, 2H), 2.42 (m, 5H), 1.68 (m, 2H), 1.37 (s, 9H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) δ (ppm), 193.63, 172.21, 161.05, 155.60, 141.48, 136.40, 132.04, 128.47, 127.42, 125.06, 118.30, 111.85, 107.74, 77.45, 69.30, 67.05, 38.83, 30.68, 28.22, 24.54, 21.11; HRMS (M + Na): Calc’d for C\(_{25}\)H\(_{31}\)NO\(_6\)Na: 464.2049. Found: 464.2040.

2-(4-hydroxy-2-methylphenyl)-2-oxoethyl-4-(tert-butoxycarbonyl-amino)-butanoate, 139e. Adhesive ppt., 98%; IR (KBr): 3460, 3300, 3100-2800, 1740, 1699, 1662, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) δ (ppm), 10.30 (d, 1H), 7.78 (d, J = 9.10, 1H), 6.89 (t, J = 7.50, 1H), 6.71 (t, J = 6.30, 1H), 5.26 (s, 2H), 2.98 (t, J = 6.70, 2H), 2.41 (m, 5H), 1.69 (m, 2H), 1.39 (s, 9H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) δ (ppm), 192.91, 172.21, 160.96, 155.60, 141.83, 131.87, 125.29, 118.74, 112.51, 77.45, 66.86, 38.97, 30.46, 28.22, 24.95, 21.68; HRMS (M + Na): Calc’d for C\(_{18}\)H\(_{25}\)NO\(_6\)Na: 374.1580. Found: 374.1577.

4-(2-(4-hydroxy-2-methylphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 139. White ppt., 95%, Mp: 110-112°C; IR (KBr): 3460, 3300, 3100-2800, 1740, 1699, 1662, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) δ (ppm), 7.86 (s, 3H), 7.80 (d, J = 9.00, 1H), 6.74 (t, J = 6.20, 1H), 5.28 (s, 2H), 2.96 (t, J = 6.70, 2H), 2.54 (t, J = 7.60, 2H), 2.40 (s, 3H), 1.87 (m, 2H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) δ (ppm), 192.71, 171.72, 161.19, 158.64, 141.88, 131.90, 124.55, 118.81, 117.56, 112.58, 67.02, 37.84, 30.14, 22.54, 20.34; HRMS (M +): Calc’d for C\(_{13}\)H\(_{18}\)NO\(_4\): 252.1236. Found: 252.1227.
2-(4-(benzyloxy)phenyl)-2-oxoethyl propionate, 140b. 2-(4-(benzyloxy)phenyl)-2-oxoethyl propionate is commercially available. Benzylation of this involved the same procedure as in the synthesis of 135b. This compound is known. 97%, GC/MS (EI): One peak, retention time, 7.63 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 262.0 m/z.

2-(4-(benzyloxy)phenyl)-2-oxoethyl 2-bromopropanoate, 140c (racemic). ~94%. GC/MS (EI): One peak, retention time, 8.10 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 317.0, 319.0, m/z.

(S)-tert-butyl 5-(4-(benzyloxy)phenyl)-4-methyl-5-oxopentylcarbamate and (R)-tert-butyl-5-(4-(benzyloxy)phenyl)-4-methyl-5-oxopentylcarbamate, (R,S)-140d.
Adhesive ppt., 95%; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; 1H NMR (400 MHz, d6-DMSO) δ (ppm), 7.98 (d, J = 8.89, 2H), 7.40 (m, 5H), 7.14 (d, J = 9.03, 2H), 6.87 (t, J = 7.35, 1H), 5.95 (m, 1H), 5.21 (s, 2H), 2.93 (t, J = 7.00, 2H), 2.37 (t, J = 7.51, 2H), 1.63 (m, 2H), 1.39 (s, 9H), 1.17 (d, J = 7.20, 3H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 194.93, 172.10, 162.61, 155.61, 136.31, 130.67, 128.48, 128.03, 127.53, 126.38, 114.93, 77.47, 69.89, 63.47, 36.16, 30.12, 28.18, 20.70, 17.08; HRMS (M + Na): Calc’d for C25H31NO6Na: 464.2049. Found: 464.2049.

(S)-tert-butyl-5-(4-hydroxyphenyl)-4-methyl-5-oxopentylcarbamate and (R)-tert-butyl-5-(4-hydroxyphenyl)-4-methyl-5-oxopentylcarbamate, (R,S)-140e.
Adhesive ppt., 92%; 1H NMR (400 MHz, d6-DMSO) δ (ppm), 7.87 (d, J = 9.05, 2H), 6.88 (m, 3H), 5.90 (m, 1H), 2.94 (t, J = 6.86, 2H), 2.34 (t, J = 7.72, 2H), 2.24 (s, 3H),
1.63 (m, 2H), 1.39 (s, 9H), 1.18 (d, J = 6.88, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 195.55, 173.02, 163.50, 157.24, 132.90, 128.93, 115.54, 78.43, 71.95, 71.95, 40.00, 31.57, 29.14, 25.82, 16.81; HRMS (M + Na): Calc’d for C$_{18}$H$_{25}$NO$_6$Na: 374.1567. Found: 374.1567.

(R)-4-(1-(4-hydroxyphenyl)-1-oxopropan-2-ylxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate and (S)-4-(1-(4-hydroxyphenyl)-1-oxopropan-2-ylxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, (R,S)-140. Adhesive ppt., 83%; IR (KBr): 3448, 3274, 3100-2800, 1739, 1691, 1602, 1520, 1421, 1377, 1309, 1298, 1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613 cm$^{-1}$; $^1$H NMR (400 MHz, D$_2$O) δ (ppm), 7.99 ($J = 8.69$, 2H), 7.00 (d, $J = 8.10$, 2H), 6.06 (m, 1H), 3.06 ($J = 7.10$, 2H), 2.64 (t, $J = 7.70$, 2H), 2.00 (m, 2H), 1.54 (d, $J = 7.05$, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 190.67, 171.72, 160.98, 158.27, 130.76, 127.76, 125.10, 120.30, 117.94, 114.76, 66.11, 38.09, 30.15, 22.10, 15.80; HRMS (M +): Calc’d for C$_{13}$H$_{18}$NO$_4$: 252.1230. Found: 252.1236.

1-(4-(benzyloxy)phenyl)butan-1-one, 141b. 1-(4-(benzyloxy)phenyl)butan-1-one is commercially available. Benzylation of this involved the same procedure as in the synthesis of 122b. This compound is known. White ppt., 97%. $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 7.96 (d, $J = 8.64$, 2H), 7.40 (m, 5H), 7.02 (d, $J = 8.50$, 2H), 5.14 (s, 2H), 2.90 (t, $J = 7.43$, 2H), 1.76 (m, 2H), 1.01 (t, $J = 6.73$, 3H).

1-(4-(benzyloxy)phenyl)-3-bromobutan-1-one, 141c. Racemic, 88%; GC/MS (EI): One peak, retention time, 9.58 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 331.8, 333.9 m/z.
(R)-tert-butyl 4-(4-(benzyloxy)benzoyl)hexylcarbamate and (S)-tert-butyl 4-(4-(benzyloxy)benzoyl)hexylcarbamate (R,S)-141d. Adhesive ppt., 89%; IR (KBr): 3307, 3100-2800, 1742, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, d$_6$-DMSO) δ (ppm), 7.99 (d, $J = 8.62$, 2H), 7.39 (m, 5H), 7.13 (d, $J = 8.88$, 2H), 6.87 (t, $J = 7.21$, 1H), 5.88 (m, 1H), 5.22 (s, 2H), 2.94 (t, $J = 6.88$, 2H), 2.39 (t, $J = 7.51$, 2H), 1.70-1.60 (m, 4H), 1.37 (s, 9H), 0.94 (t, $J = 7.05$, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 194.42, 172.51, 162.60, 155.58, 136.23, 130.63, 128.49, 128.03, 127.85, 127.10, 115.45, 77.77, 75.64, 70.45, 39.14, 30.77, 28.21, 24.91, 20.73, 9.49; HRMS (M + Na): Calc’d for C$_{26}$H$_{33}$NO$_6$Na: 478.2206. Found: 464.2189.

(R)-tert-butyl 4-(4-(hydroxybenzoyl)hexylcarbamate and (S)-tert-butyl 4-(4-(hydroxybenzoyl)hexylcarbamate, (R,S)-141e. Adhesive ppt., 98%; $^1$H NMR (400 MHz, d$_6$-DMSO) δ (ppm), 10.66 (s, 1H), 7.89 (d, $J = 8.75$, 2H), 6.88 (m, 3H), 5.80 (m, 1H), 2.94 (t, $J = 6.95$, 2H), 2.37 (t, $J = 7.33$, 2H), 1.70-1.60 (m, 4H), 1.38 (s, 9H), 0.94 (t, $J = 6.85$, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 194.04, 172.26, 162.52, 155.58, 130.86, 125.29, 115.45, 77.59, 75.59, 39.13, 30.95, 28.35, 24.92, 21.02, 9.54; HRMS (M + Na): Calc’d for C$_{17}$H$_{27}$NO$_6$Na: 388.1736. Found: 388.1721.

(R)-4-(1-(4-hydroxyphenyl)-1-oxopropan-2-yloxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate and (S)-4-(1-(4-hydroxyphenyl)-1-oxopropan-2-yloxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, (R,S)-141. Adhesive ppt., 80%; IR (KBr): 3450, 3274, 3100-2800, 1739, 1691, 1602, 1520, 1421, 1377, 1309, 1298,
1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613. cm$^{-1}$; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.88 ($J = 8.23$, 2H), 6.88 (d, $J = 7.75$, 2H), 5.83 (m, 1H), 2.85 ($J = 7.01$, 2H), 2.40 (t, $J = 7.62$, 2H), 1.82-1.68 (m, 4H), 0.91 (d, $J = 6.82$, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 193.38, 171.76, 162.69, 158.41, 130.87, 125.42, 117.47, 115.50, 75.82, 38.08, 30.17, 24.57, 22.55, 9.48; HRMS (M+): Calc’d for C$_{14}$H$_{20}$NO$_4$: 266.1392. Found: 266.1389.

2-bromo-1-(4-fluorophenyl)ethanone, 142b. 4-fluoroacetophenone is commercially available. The same conditions were used as in the synthesis of 141c. This compound is known. Oil, ~95% yield by GC/MS (EI): One peak, retention time, 4.79 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 215.9, 217.9 m/z.

2-(4-fluorophenyl)-2-oxoethyl 4-(tert-butoxycarbonylamino)butanoate, 142c. The same general procedure to make 122e was followed. White ppt., 82%, Mp: 85-87°C; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 8.04 (m, 2H), 7.39 (t, $J = 8.80$, 2H), 6.87 (t, $J = 7.40$, 1H), 5.46 (s, 2H), 2.96 (dd, $J = 6.82$, $J = 6.65$, 2H), 2.42 (t, $J = 7.94$, 2H), 1.68 (m, 2H), 1.38 (s, 9H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 191.51, 172.19, 166.35, 164.34, 155.60, 130.63, 116.09, 77.46, 66.26, 39.14, 30.62, 28.22, 24.95; HRMS (M + Na): Calc’d for C$_{17}$H$_{22}$FNO$_5$Na: 362.1380. Found: 362.1377

4-(2-(4-fluorophenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate 142. The same general method to make 130 was followed. Adhesive ppt., 93%; IR (KBr): 3460, 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167,
955, 743, 700, 667. $^1$H NMR (400 MHz, D$_2$O) $\delta$ (ppm), 8.06 (m, 2H), 7.32 (t, $J = 8.80$, 2H), 5.56 (s, 2H), 3.12 (dd, $J = 6.82$, $J = 6.65$, 2H), 2.72 (t, $J = 7.94$, 2H), 2.06 (m, 2H); $^{19}$F NMR (376 MHz, D$_2$O) $\delta$ (ppm), -104.12. $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 191.46, 171.72, 166.40, 164.39, 158.67, 130.91, 116.12; HRMS (M+): Calc’d for C$_{12}$H$_{15}$FNO$_3$: 240.1030. Found: 240.0998.

5-(2-(4-(benzyloxy)-2-fluorophenyl)-2-oxoethyl)-1-tert-butyl-2-(tert-butoxy-carbonylamino)pentanedioate, 143e. The series to construct 143e paralleled those for 122e. Adhesive ppt., 91%; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.85 (t, $J = 8.88$, 1H), 7.40 (m, 5H), 7.22 (d, $J = 7.64$, 1H), 7.10 (d, $J = 10.50$, 1H), 7.01 (d, $J = 6.61$, 1H), 5.24 (s, 2H), 5.22 (s, 2H), 3.89 (m, 1H), 2.45 (t, $J = 6.69$, 2H), 1.99 (m, 1H), 1.89 (m, 1H), 1.39 (s, 18H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 198.83, 171.32, 170.32, 164.28, 162.14, 154.64, 135.99, 131.42, 128.63, 127.95, 115.84, 112.37, 111.80, 102.83, 80.42, 78.11, 70.18, 68.41, 59.74, 30.73, 29.76, 28.15, 25.91; HRMS (M+Na): Calc’d for C$_{29}$H$_{36}$FNO$_8$Na: 568.2323. Found: 568.2296.

1-tert-butyl-5-(2-(2-fluoro-4-hydroxyphenyl)-2-oxoethyl)-2-(tert-butoxycarbonylamino)pentanedioate, 143e'. An identical procedure was followed to compose 130e'. Adhesive ppt., 94%; IR (KBr): 3460, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, MeOD) $\delta$ (ppm), $\delta$ 7.86 (t, $J = 8.24$, 1H), 6.90 (t, $J = 6.55$, 1H), 6.73 (d, $J = 7.70$, 1H), 6.64 (d, $J = 9.10$, 1H), 5.24 (s, 2H), 3.99 (m, 1H), 2.56 (t, $J = 7.05$, 2H), 2.10 (m, 1H), 1.97 (m, 1H),
1.51 (s, 18H); $^{13}$C NMR (125 MHz, MeOD) δ (ppm), 189.27, 172.44, 171.59, 165.31, 163.29, 158.70, 131.71, 115.88, 113.74, 102.51, 81.28, 79.14, 68.50, 54.80, 47.10, 29.73, 27.33, 26.47; HRMS (M + Na): Calc’d for C$_{22}$H$_{31}$FNO$_8$: 456.2034. Found: 456.2034. Calc’d for C$_{22}$H$_{30}$FNO$_8$Na: 478.1853. Found: 478.1866.

1-carboxy-4-(2-(2-fluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 143. The same procedure was used as in the synthesis of 130. Adhesive ppt., 68%; IR (KBr): 3460, 3274, 3100-2800, 1739, 1691, 1602, 1520, 1421, 1377, 1309, 1298, 1201, 1178, 1128, 1055, 1027, 1008, 823, 762, 717, 613 cm$^{-1}$, $^1$H NMR (400 MHz, MeOD) δ (ppm), δ 7.84 (t, $J = 8.20$, 1H), 6.75 (d, $J = 7.52$, 1H), 6.62 (d, $J = 9.22$, 1H), 5.22 (s, 2H), 3.96 (m, 1H), 2.53 (t, $J = 6.92$, 2H), 2.09 (m, 1H), 1.97 (m, 1H); $^{13}$C NMR (125 MHz, MeOD) δ (ppm), 189.55, 172.46, 169.99, 163.47, 161.33, 130.12, 117.24, 115.37, 114.10, 112.03, 103.37, 68.81, 54.27, 29.33, 26.48; HRMS (M+): Calc’d for C$_{13}$H$_{15}$FNO$_6$: 300.0878. Found 300.0859.

The same sequence of steps was followed to generate 144 and 145.

5-(2-(4-(benzyloxy)-3-fluorophenyl)-2-oxoethyl)-1-tert-butyl-2-(tert-butoxy-carbonylamino)pentanedioate, 144e. Adhesive ppt., 82%; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, CD$_3$OD) δ (ppm), δ 7.82 (t, $J = 8.85$, 1H), 7.44 (m, 6H), 7.21 (d, $J = 12.02$, 1H), 5.43 (s, 2H), 5.31 (d, 2H), 3.89 (m, 1H), 2.49 (t, $J = 6.88$, 2H), 1.99 (m, 1H), 1.89 (m, 1H), 1.39 (s, 18H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) δ (ppm), 190.58, 171.74, 170.32, 155.53, 152.29, 150.89, 150.34, 135.82, 128.56, 127.93, 126.91,
125.61, 115.29, 114.83, 80.42, 78.11, 70.36, 66.23, 53.45, 29.75, 27.90, 27.60, 25.92; HRMS (M + Na): Calc’d for C_{29}H_{36}FNO_8Na: 568.2323. Found: 568.2275.

1-tert-butyl-5-(2-(3-fluoro-4-hydroxyphenyl)-2-oxoethyl)-2-(tert-butoxy carbonylamino)pentanedioate, 144e'. Adhesive ppt., 94%; IR (KBr): 3460, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; \textsuperscript{1}H NMR (400 MHz, d\textsuperscript{6}-DMSO) \delta (ppm), \delta 7.72 (t, J = 8.52, 1H), 7.67 (d, J = 8.63, 1H), 7.22 (d, J = 7.64, 1H), 7.05 (t, J = 8.14, 1H), 5.39 (s, 2H), 3.82 (m, 1H), 2.48 (t, J = 6.88, 2H), 1.95 (m, 1H), 1.82 (m, 1H), 1.40 (s, 18H); \textsuperscript{13}C NMR (125 MHz, d\textsuperscript{6}-DMSO) \delta (ppm), 190.30, 171.74, 171.33, 155.53, 151.56, 150.50, 148.63, 125.62, 117.23, 115.70, 80.41, 78.10, 66.11, 54.37, 29.70, 28.15, 27.90, 25.79; HRMS (M + Na): Calc’d for C_{22}H_{31}FNO_8: 456.2034. Found: 456.2034. Calc’d for C_{22}H_{30}FNO_8 Na: 478.1853. Found: 478.1815.

1-carboxy-4-(2-(3-fluoro-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 144. Adhesive ppt., 84%; IR (KBr): 3455, 3269, 3100-2800, 1741, 1693, 1610, 1553, 1420, 1201, 1050, 823, 759, 719 cm\textsuperscript{-1}; \textsuperscript{1}H NMR (400 MHz, D\textsubscript{2}O) \delta (ppm), 7.72 (d, J = 8.98, 2H), 7.08 (t, J = 9.52, 1H), 5.41 (s, 2H), 4.57 (t, J = 7.42, 1H), 2.65 (t, J = 7.14, 2H), 2.34 (m, 1H), 2.12 (m, 1H); \textsuperscript{13}C NMR (125 MHz, D\textsubscript{2}O) \delta (ppm), 190.12, 172.06, 170.06, 161.44, 152.57, 150.54, 131.17, 126.14, 119.58, 117.22, 115.40, 67.74, 53.99, 30.02, 23.51; HRMS (M\textsuperscript{+}): Calc’d for C_{13}H_{15}FNO\textsubscript{6}: 300.0878. Found 300.0892.

5-(2-(4-(benzyl oxy)-3-(trifluoromethyl)phenyl)-2-oxoethyl)-1-tert-butyl-2-(tert-butoxycarbonylamino)pentanedioate, 145e. This was synthesized in the same
manner as 143e by a laboratory associate. However, no characterization was done preceding the next synthetic step.

1-tert-butyl-5-(2-(4-hydroxy-3-(trifluoromethyl)phenyl)-2-oxoethyl)-2-(tert-butoxycarbonylamino)pentanedioate, 145e. Adhesive ppt., 91%; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 11.73 (s, 1H), 8.08 (s, 1H), 8.06 (s, 1H), 7.21 (d, J = 7.52, 1H), 7.13 (d, J = 9.50), 5.55 (s, 2H), 3.91 (dd, J = 8.40, J = 5.80, 1H), 2.49 (t, J = 9.05, 2H), 1.94, (m, 2H), 1.82 (m, 1H), 1.40 (s, 9H), 1.36 (s, 9H).

1-carboxy-4-(2-(4-hydroxy-3-(trifluoromethyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 145. White ppt., 55%, Mp: 59-61°C; IR (KBr): 3405, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm$^{-1}$; $^1$H NMR (400 MHz, MeOD) $\delta$ (ppm), 8.13 (d, J = 1.50, 1H), 8.06 (dd, J = 8.70, J = 2.10, 1H), 7.06 (d, J = 8.50, 1H), 5.41 (s, 2H), 4.55 (d, J = 9.10, 1H), 2.61 (t, J = 7.70, 2H), 2.36, (m, 1H), 2.12 (m, 1H); $^{13}$C NMR (125 MHz, MeOD) $\delta$ (ppm), 192.31, 173.69, 162.42, 159.48, 159.19, 134.81, 128.68, 126.79, 123.89, 118.70, 118.07, 116.42, 67.78, 53.46, 31.21, 27.19; HRMS (M+): Calc’d for C$_{14}$H$_{15}$F$_3$NO$_6$: 350.0851. Found 350.0863.

1-(4-(benzyloxy)phenyl)ethanone, 146b. The same procedure to construct 135b was utilized but only one equivalent of benzyl bromide was required. White ppt, 96%. This compound is known. GC/MS (EI): One peak, retention time, 7.43 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 226.09 m/z.

1-(4-(benzyloxy)phenyl)-2-bromoethanone, 146c. The same method to generate 135c was used. Oil, ~94%. This compound is known. GC/MS (EI): One peak,
retention time, 8.22 min (initial temp 50°C, ramp 25°C/min until 300°C), MS; 304.01, 306.01 m/z.

2-(4-(benzyloxy)phenyl)-2-oxoethyl-2-(tert-butoxycarbonylamino)acetate, 146d. 
The same general method to produce 135d was used but with N-Boc-Gly. White ppt., 
95%, Mp: 100-102°C; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; 1H NMR (400 MHz, CDCl3) δ (ppm), 7.90 (d, J = 8.49, 2H), 7.41 (m, 5H), 7.05 (d, J = 8.82, 2H), 5.37 (s, 2H), 5.15 (s, 2H), 5.04 (s, 1H), 4.12 (d, J = 7.02, 2H), 1.45 (s, 9H); 13C NMR (125 MHz, CDCl3) δ (ppm), 189.94, 170.09, 163.32, 155.72, 135.92, 132.28, 130.12, 128.77, 128.04, 127.15, 114.99, 80.10, 70.25, 66.30, 42.31, 28.33; HRMS (M + Na): Calc’d for C22H25NO6 Na: 422.1580. Found: 422.1559.

2-(4-hydroxyphenyl)-2-oxoethyl 2-(tert-butoxycarbonylamino)acetate, 146e. The 
same general method to produce 135e was used. White ppt., 94%, Mp: 125-127 °C; 
1H NMR (400 MHz, d6-DMSO) δ (ppm), 10.48 (s, 1H), 7.85 (d, J = 8.66, 2H), 7.27 (t, J = 5.63, 1H), 6.88 (d, J = 8.23, 2H), 5.42 (s, 2H), 3.82 (d, J = 6.95, 2H), 1.38 (s, 9H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 190.36, 170.15, 162.87, 155.75, 130.55, 125.32, 115.41, 78.48, 66.25, 41.58, 28.04; HRMS (M + Na): Calc’d for C15H19NO6 Na: 332.1110. Found: 332.1101.

2-(2-(4-hydroxyphenyl)-2-oxoethoxy)-2-oxoethanaminium-2,2,2-trifluoroacetate, 
146. The same general method to produce 135 was followed. Adhesive ppt, 79%; IR (KBr): 3405, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm⁻¹; 1H NMR (400 MHz, MeOD) δ (ppm), 7.88 (d,
$J = 8.42, 2H), 6.88 (d, $J = 8.06, 2H), 5.57 (s, 1H), 5.47 (s, 1H), 4.24 (s, 1H), 4.04 (s, 1H); ^{13}$C NMR (125 MHz, MeOD) $\delta$ (ppm), 191.21, 168.34, 167.11, 163.54, 130.40, 125.88, 117.38, 115.41, 66.63, 40.52; HRMS (M+): Calc’d for $C_{10}H_{12}NO_4$: 210.0766. Found: 210.0764.

The same succession of steps was followed to produce 147-149. The amino acids all were N-Boc protected.

2-(4-(benzyloxy)phenyl)-2-oxoethyl-2-(tert-butoxycarbonylamino)-3-methylbutanoate, 147d. White ppt., 95%, Mp: 119-120°C; IR (KBr): 3307, 3100-2800, 1742, 1701, 1659, 1606, 1437, 1407, 1209, 1167, 955, 743, 700, 667; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.96 (d, $J = 8.60, 2H), 7.40 (m, 5H), 7.13 (m, 3H), 5.49 (d, $J = 8.24, 1H), 5.42 (d, J = 8.20, 1H), 5.15 (s, 2H), 4.05 (t, $J = 6.75, 1H), 2.14 (m, 1H), 1.39 (s, 9H), 0.97 (d, $J = 7.05, 6H); ^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 190.88, 171.51, 162.60, 155.73, 136.37, 130.15, 128.89, 128.03, 127.79, 126.89, 114.97, 78.19, 69.53, 66.34, 59.07, 29.78, 27.91, 19.03; HRMS (M + Na): Calc’d for $C_{25}H_{31}NO_6Na$: 464.2049. Found: 464.2061.

2-(4-hydroxyphenyl)-2-oxoethyl-2-(tert-butoxycarbonylamino)-3-methylbutanoate, 147e. Adhesive ppt., 75%; $^1$H NMR (400 MHz, MeOD) $\delta$ (ppm), 7.90 (d, $J = 8.75, 2H), 6.88 (m, 3H), 5.50 (d, $J = 8.22, 1H), 5.46 (d, J = 8.15, 1H), 4.22 (t, $J = 6.85, 1H), 2.25 (m, 1H), 1.48 (s, 9H), 1.05 (d, $J = 7.10, 6H); ^{13}$C NMR (125 MHz, MeOD) $\delta$ (ppm), 191.03, 171.84, 163.00, 156.88, 130.73, 125.81, 114.80, 79.19, 66.02, 59.08, 30.58, 27.17, 18.26; HRMS (M + Na): Calc’d for $C_{18}H_{25}NO_6Na$: 374.1580. Found: 374.1592.
1-(2-(4-hydroxyphenyl)-2-oxoethoxy)-3-methyl-1-oxobutan-2-aminium-2,2,2-trifluoroacetate, 147. Adhesive ppt, 72%; IR (KBr): 3405, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm\(^{-1}\); \(^1\)H NMR (400 MHz, MeOD) \(\delta\) (ppm), 7.86 (d, \(J = 8.29\), 2H), 6.88 (d, \(J = 8.02\), 2H), 5.59 (d, \(J = 8.22\), 1H), 5.42 (d, \(J = 8.15\), 1H), 4.09 (t, \(J = 6.72\), 1H), 2.49 (m, 1H), 1.06 (d, \(J = 7.15\), 6H); \(^13\)C NMR (125 MHz, MeOD) \(\delta\) (ppm), 188.64, 167.78, 161.50, 158.73, 129.15, 123.81, 115.85, 113.58, 65.78, 61.36, 27.84, 16.29; HRMS (M+): Calc’d for C\(_{13}\)H\(_{18}\)NO\(_4\): 252.1230. Found: 252.1222.

1-(2-(4-hydroxyphenyl)-2-oxoethoxy)-3-(1H-indol-3-yl)-1-oxopropan-2-aminium 2,2,2-trifluoroacetate, 148. The precursor, 148e, was obtained from an associate, without characterization. White ppt., 92%, Mp: 75-77\(^\circ\)C; IR (KBr): 3405, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 11.11 (s, 1H), 10.70, (s, 1H), 8.46, (s, 3H), 7.87 (d, \(J = 12.1\), 2H), 7.60 (d, \(J = 8.10\), 1H), 7.41, (d, \(J = 8.04\), 1H), 7.33 (d, \(J = 4.23\), 1H), 7.10 (t, \(J = 7.95\), \(J = 1H\)), 7.05 (t, \(J = 7.85\), 1H), 6.92 (d, \(J = 8.20\), 2H), 5.61 (dd, \(J = 16.30\), \(J = 12.20\), 2H), 4.45 (t, \(J = 7.05\), 1H), 3.32 (m, 2H); \(^13\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 189.81, 169.06, 162.96, 158.82, 136.25, 130.50, 128.10, 126.90, 125.12, 125.01, 121.20, 118.89, 117.96, 115.52, 111.57, 106.37, 67.29, 52.53, 26.36; HRMS (M+): Calc’d for C\(_{19}\)H\(_{19}\)N\(_2\)O\(_4\): 339.1345. Found: 339.1345.

1-(2-(4-hydroxyphenyl)-2-oxoethoxy)-1-oxo-3-phenylpropan-2-aminium-2,2,2-trifluoroacetate, 149. The precursor, 149e, was obtained from an associate, without
characterization. White ppt., 74%, Mp: 64-66°C; IR (KBr): 3405, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 10.86, (s, 1H), 8.58, (s, 3H), 7.87 (d, J = 8.04, 2H), 7.31 (m, 5H), 6.92 (d, J = 8.15, 2H), 5.61 (dd, J = 16.15, J = 4.00, 2H), 4.45 (t, J = 4.20, 1H), 3.30 (dd, J = 15.88, J = 8.03, 1H), 3.18 (dd, J = 12.10, J = 4.05, 1H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 189.73, 168.75, 163.00, 158.66, 134.66, 130.50, 129.55, 128.60, 127.26, 124.95, 117.95, 115.51, 67.27, 53.01, 36.03; HRMS (M⁺): Calc’d for C₁₇H₁₈NO₄: 300.1230. Found: 300.1244.

(4R)-2-(4-(benzyloxy)phenyl)-2-oxoethyl-4-((3R,5R,8R,10S,12S,13R,17R)-3,12-dihydroxy-10,13-dimethylhexadecahydro-1H-cyclopent[a]phenanthren-17-yl)pentanoate, 150d. The reaction was patterned after the one to synthesize 146d but used DMF instead of CH₃CN as the solvent. White ppt., 84%, Mp: 77-79°C; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 7.90 (d, J = 8.47, 2H), 7.40 (m, 5H), 7.01 (d, J = 9.00, 2H), 5.28 (s, 2H), 5.14 (s, 2H), 3.99 (s, 1H), 3.67 (s, 1H), 2.56 (m, 1H), 2.46 (m, 1H), 2.00-1.00 (m, 26 H), 1.01 (s, 3H), 0.91 (s, 3H), 067 (s, 3H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 197.78, 173.72, 163.16, 135.98, 130.11, 128.75, 128.34, 127.51, 127.48, 114.89, 73.18, 71.88, 70.21, 65.89, 65.62, 53.46, 48.28, 47.37, 46.54, 42.10, 36.46, 36.31, 36.06, 35.23, 35.05, 34.14, 33.69, 30.92, 30.82, 30.54, 28.66, 27.44, 26.14, 23.66, 23.19, 17.39, 17.29, 15.30, 12.79.

(4R)-2-(4-hydroxyphenyl)-2-oxoethyl-4-((3R,5R,8R,10S,12S,13R,17R)-3,12-dihydroxy-10,13-dimethylhexadecahydro-1H-cyclopent[a]phenanthren-17-yl)pentanoate, 150. The same method to produce 130e' was used. White ppt., 93%,
Mp: 97-99°C; $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 7.84 (d, $J = 8.42$, 2H), 6.88 (d, $J = 8.95$, 2H), 5.35 (s, 2H), 4.21, (s, 1H), 3.81, (s, 1H), 2.52 (m, 1H), 2.40 (m, 1H), 2.00-1.00 (m, 26 H), 0.96 (s, 3H), 0.85 (s, 3H), 0.62 (s, 3H); $^{13}$C NMR (125 MHz, d$_6$-DMSO) $\delta$ (ppm), 190.91, 173.97, 160.98, 130.46, 127.41, 115.72, 73.29, 71.98, 65.90, 65.65, 60.46, 53.45, 48.29, 47.10, 42.08, 36.37, 36.02, 35.20, 34.13, 33.70, 30.73, 30.43, 28.61, 27.45, 26.12, 23.65, 23.17, 21.10, 17.37, 15.29, 14.22, 12.75; HRMS (M + H): Calc’d for C$_{32}$H$_{47}$O$_6$: 527.3373. Found: 527.3397. (M + Na): Calc’d for C$_{32}$H$_{46}$O$_6$Na: 549.3192. Found: 549.3191. (M + NH$_4$): Calc’d for C$_{32}$H$_{50}$O$_6$N: 544.3639. Found: 544.3634.

diethyl 2-(4-(benzyloxy)phenyl)-2-oxoethylphosphonate, 151d. The general method of De Borggraeve$^{147}$ was utilized. An Argon purged solution of 1-(4-(benzyloxy)phenyl)-2-bromoethanone (146d, 2.44 g (7.99 mmol)) in 10 ml of neat triethylphosphite was heated to 100°C, where it was maintained with stirring and under Argon for 17h. After cooling to room temperature, most of the surfeit CH$_3$CN/Triethylphosphite was removed under reduced pressure. The mixture was then flash chromatographed on silica with gradients of ethyl acetate/hexanes mixtures, until 151d eluted at 100% ethyl acetate. After 1 day of vacuum, viscous oil was observed, 2.48 g (86%). This compound is known. $^1$H NMR (400 MHz, d$_6$-DMSO) $\delta$ (ppm), 8.00 (d, $J = 9.25$, 2H), 7.40 (m, 5H), 7.13 (d, $J = 9.56$, 2H), 5.22 (s, 2H), 4.00 (m, 4H), 1.24 (t, $J = 8.33$, 6H).

diethyl 2-(4-hydroxyphenyl)-2-oxoethylphosphonate, 151. The same general method to produce 130e’ was used. White ppt., 96%, Mp: 108-110°C; IR (KBr):
3460, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 10.45 (s, 1H), 7.89 (d, \(J = 9.31\), 2H), 6.85 (d, \(J = 9.40\), 2H), 5.09 (s, 2H), 4.00 (m, 4H), 1.24 (t, \(J = 8.47\), 6H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.28, 162.46, 131.89, 127.80, 115.07, 60.88, 37.66, 16.10; HRMS (M + Na): Calc’d for C\(_{12}\)H\(_{17}\)PO\(_5\)Na: 295.0711; Found: 295.0698; (M + H): Calc’d for C\(_{12}\)H\(_{18}\)PO\(_5\): 273.0892. Found: 273.0877.

1-(4-(benzyloxy)phenyl)-2-fluoroethanone, 152d. To a solution of 1-(4-(benzyloxy)phenyl)-2-bromoethanone (analog of 146d, 820 mg (2.69 mmol)) in 100 ml CH\(_3\)CN was added cesium fluoride (1.63 g, 10.7 mmol). The heterogeneous mixture was refluxed for 1.5 h at which time the reaction was deemed complete as indicated by GC/MS. Excess CH\(_3\)CN was removed under reduced pressure, equal volumes of EtOAc/H\(_2\)O added (50 ml), the organic phase retained and flash chromatographed on silica gel (1:1 Et2O, Hex), affording 590 mg of 152d as a white ppt, 90%, Mp: 106-108°C; IR (KBr): 3460, 3310, 3100-2800, 1735, 1710, 1690, 1595, 1520, 1430, 1370, 1248, 1200, 1165, 1050, 818, 760, 710 cm\(^{-1}\); \(^1\)H NMR (400 MHz, CDCl\(_3\)) \(\delta\) (ppm), 7.98 (d, \(J = 8.67\), 2H), 7.40 (m, 5H), 7.06 (d, \(J = 9.10\), 2H), 5.54 (s, 1H), 5.43 (s, 1H), 5.16 (s, 2H); \(^{13}\)C NMR (125 MHz, CDCl\(_3\)) \(\delta\) (ppm), 192.01, 163.37, 135.91, 131.58, 130.30, 128.77, 127.50, 126.45, 114.80, 84.22, 70.24; HRMS (FAB +): Calc’d for C\(_{15}\)H\(_{13}\)FO\(_2\): 245.0978. Found: 245.0978.

2-fluro-1-(4-hydroxyphenyl)ethanone, 152. White ppt., 92%. This compound is known. \(^1\)H NMR (400 MHz, MeOD) \(\delta\) (ppm), 7.91 (d, \(J = 8.50\), 2H), 6.91 (d, \(J =
9.15, 2H), 5.72 (s, 1H), 5.61 (s, 1H); HRMS (M - H): Calc’d for C₈H₆FO₂: 153.0352. Found: 153.0326. (M + Na): Calc’d for C₈H₇FO₂Na: 177.0328. Found: 177.0327.

**benzyl 5-acetyl-2-(benzyloxy)benzoate, 153b.** The reaction conditions to produce 153c were amenable to those to create 135b. White ppt., 93%, Mp: 98-100 °C; IR (KBr): 3325, 3100-2800, 1726, 1697, 1676, 1598, 1500, 1452, 1370, 1269, 1230, 1176, 1060, 1030, 1008, 821, 757, 721, 698 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.46 (d, J = 2.10, 1H), 8.08 (dd, J = 9.00, J = 2.34, 1H), 7.44-7.33 (m, 10H), 7.09 (d, J = 8.85, 2H), 5.38 (s, 2H), 5.25 (s, 2H), 5.28 (s, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 195.42, 165.04, 160.91, 135.11, 134.24, 133.04, 130.28, 129.16, 128.22, 127.81, 119.35, 117.36, 112.49, 70.08, 68.86, 25.74; HRMS (M + Na): Calc’d for C₂₃H₂₀O₄Na: 383.1259. Found: 383.1262.

**benzyl 2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 153c.** The synthetic method to generate 122d was followed. Oil, 95%; IR (KBr): 3325, 3100-2800, 1726, 1697, 1676, 1598, 1500, 1452, 1370, 1269, 1230, 1176, 1060, 1030, 1008, 821, 757, 721, 698 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.50 (d, J = 2.15, 1H), 8.10 (dd, J = 9.00, J = 2.40, 1H), 7.44-7.33 (m, 10H), 7.10 (d, J = 8.90, 2H), 5.38 (s, 2H), 5.25 (s, 2H), 4.41 (s, 2H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 189.39, 165.32, 162.18, 135.68, 135.48, 134.52, 133.49, 128.74, 128.26, 127.11, 126.47, 120.87, 113.44, 70.86, 67.21, 30.48; HRMS (M + Na): Calc’d for C₂₃H₁₉BrO₄Na: 461.0365. Found: 461.0372.

**benzyl-2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)-butanoyloxy)-acetyl)-benzoate, 153d.** The synthetic mode to engender 122e was utilized. White ppt.,
75%, Mp: 75-77°C; IR (KBr): 3271, 3100-2800, 1733, 1701, 1697, 1600, 1577, 1500, 1454, 1415, 1365, 1269, 1251, 1215, 1166, 1056, 1028, 1008, 919, 875, 821, 758, 700 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.26 (d, J = 2.44, 1H), 8.12 (dd, J = 9.00, J = 2.33, 1H), 7.44-7.33 (m, 11H), 6.89 (t, J = 5.51, 2H), 5.45 (s, 2H), 5.33 (s, 2H), 2.97 (t, J = 7.20, 2H), 2.42 (t, J = 7.00, 2H), 1.68 (m, 2H), 1.38 (s, 9H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 190.85, 172.13, 165.28, 161.01, 155.90, 135.82, 133.51, 130.81, 128.42, 128.04, 127.89, 127.33, 126.27, 113.95, 77.78, 70.10, 64.96, 59.73, 39.30, 30.64, 28.22, 25.24; HRMS (M + Na): Calc’d for C₃₂H₃₅NO₈Na: 584.2260. Found: 584.2239.

5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoic acid, 153e. The reaction conditions to produce 130e¹ were abided. Adhesive ppt., 96%; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.37 (d, J = 2.35, 1H), 8.08 (dd, J = 9.15, J = 2.30, 1H), 7.10 (d, J = 8.90, 2H) 6.90 (t, J = 5.51, 2H), 5.43 (s, 2H), 2.98 (t, J = 7.15, 2H), 2.44 (t, J = 7.05, 2H), 1.68 (m, 2H), 1.36 (s, 9H).

4-(2-(3-carboxy-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 153. The method used to produce 130 was followed. Adhesive ppt., 94%; IR (KBr): 3440, 3274, 3100-2800, 1733, 1741, 1692, 1649, 1631, 1596, 1500, 1450, 1416, 1299, 1201, 1184, 1124, 1053, 1006, 823, 800, 760, 707, 680 cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm), 8.36 (s, 1H), 8.00 (d, J = 8.93, 1H), 7.08 (d, J = 8.10), 5.46 (s, 2H), 3.12 (t, J = 7.20, 2H), 2.68 (t, J = 7.00, 2H), 2.06 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm), 193.77, 174.11, 171.62, 165.14, 163.06, 134.87,
131.89, 119.72, 117.89, 115.09, 113.41, 66.60, 38.55, 30.45, 21.97; HRMS (M+): Calc’d for C_{13}H_{16}NO_{6}: 282.0978. Found: 282.0967.

4-(2-(4-hydroxy-3-(methoxycarbonyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 154. The precursor, methyl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoate, was received from an associate. Adhesive ppt., 79%; IR (KBr): 3440, 3274, 3100-2800, 1733, 1741, 1692, 1649, 1631, 1596, 1500, 1450, 1416, 1299, 1201, 1184, 1124, 1053, 1006, 823, 800, 760, 707, 680 cm⁻¹; ¹H NMR (400 MHz, D₂O) δ (ppm), 8.40 (s, 1H), 8.03 (d, J = 9.05, 1H), 7.05 (d, J = 8.00, 1H), 5.48 (s, 2H), 3.96 (s, 3H), 3.10 (t, J = 7.15, 2H), 2.70 (t, J = 7.00, 2H), 2.05 (m, 2H); ¹³C NMR (125 MHz, D₂O) δ (ppm), 193.33, 174.36, 169.84, 165.22, 162.84, 133.99, 132.18, 121.01, 117.44, 115.87, 113.73, 66.88, 51.10, 39.02, 29.97; HRMS (M+): Calc’d for C_{14}H_{18}NO_{6}: 298.1134. Found: 298.1123.

Benzy 2-(benzyloxy)-5-(2-(dodecanoyloxy)acetyl)benzoate, 155d. The same method to produce 153d was used but with dodecanoic acid. Oil, 79%; IR (KBr): 3271, 3100-2800, 1733, 1701, 1697, 1600, 1577, 1500, 1454, 1415, 1365, 1269, 1251, 1215, 1166, 1056, 1028, 1008, 919, 875, 821, 758, 700 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.26 (d, J = 2.30, 1H), 8.14 (dd, J = 8.85, J = 2.25, 1H), 7.44-7.33 (m, 11H), 5.42 (s, 2H), 5.32 (s, 4H), 2.42 (t, J = 7.15, 2H), 1.55 (m, 2H), 1.26-1.15 (m, 16), 0.84 (t, J = 5.95, 3H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.35, 173.29, 165.34, 162.09, 135.67, 133.62, 132.43, 130.80, 128.33, 127.10, 126.76, 120.77, 113.49, 70.82, 67.20, 65.60, 33.94, 31.80, 30.51, 29.64, 29.37, 29.29,

5-(2-(dodecanoyloxy)acetyl)-2-hydroxybenzoic acid, 155. The same method to generate 153e was followed. White ppt., 92%, Mp: 58-60°C; IR (KBr): 3271, 3100-2800, 1733, 1701, 1697, 1600, 1577, 1500, 1454, 1415, 1365, 1269, 1251, 1215, 1166, 1056, 1028, 1008, 919, 875, 821, 758, 700 cm<sup>-1</sup>; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ (ppm), 11.02, (s, 1H), 8.51 (d, J = 2.05, 1H), 8.11 (dd, J = 4.80, J = 4.10, 1H), 7.10 (d, J = 7.95, 1H), 5.32 (s, 2H), 2.25 (t, J = 7.20, 2H), 1.70 (m, 2H), 1.39-1.27 (m, 16H), 0.88 (t, J = 6.00, 3H); <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ (ppm), 190.34, 173.82, 172.53, 166.28, 135.87, 128.91, 126.16, 118.74, 111.25, 108.59, 35.44, 33.99, 31.93, 31.00, 29.63, 29.36, 29.12, 24.93, 22.71, 14.15; HRMS (M - H): Calc’d for C<sub>21</sub>H<sub>29</sub>O<sub>6</sub>: 377.1964. Found: 377.1959.

The same procedures were utilized to synthesize 156.

benzyl 2-(benzyloxy)-5-(2-(tetradecanoyloxy)acetyl)benzoate, 156d. White ppt., 71%, Mp: 55-57°C; IR (KBr): 3271, 3100-2800, 1733, 1701, 1697, 1600, 1577, 1500, 1454, 1415, 1365, 1269, 1251, 1215, 1166, 1056, 1028, 1008, 919, 875, 821, 758, 700 cm<sup>-1</sup>; <sup>1</sup>H NMR (400 MHz, d<sup>6</sup>-DMSO) δ (ppm), 8.26 (d, J = 2.30, 1H), 8.14 (dd, J = 8.85, J = 2.25, 1H), 7.44-7.33 (m, 11H), 5.42 (s, 2H), 5.33 (s, 4H), 2.40 (t, J = 7.10, 2H), 1.56 (m, 2H), 1.26-1.22 (m, 20H), 0.84 (t, J = 6.00, 3H); <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ (ppm), 190.955, 174.49, 165.34, 162.09, 135.67, 133.62, 132.43, 130.80, 128.33, 127.10, 126.76, 120.77, 113.49, 70.82, 67.20, 65.60, 33.94, 31.80,
30.51, 29.64, 29.37, 29.29, 29.13, 28.66, 28.31, 24.93, 22.72, 21.09, 14.23; HRMS (M + Na): Calc’d for C_{37}H_{46}O_{6}Na: 609.3192. Found: 609.3201.

**2-hydroxy-5-(2-(tetradecanoyloxy)acetyl)benzoic acid, 156.** White ppt., 94%, Mp: 60-62°C; IR (KBr): 3271, 3100-2800, 1733, 1701, 1697, 1600, 1577, 1500, 1454, 1415, 1365, 1269, 1251, 1215, 1166, 1056, 1028, 1008, 919, 875, 821, 758, 700 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.30 (d, \(J = 2.30\), 1H), 7.92 (dd, \(J = 8.80\), \(J = 2.20\), 1H), 6.88 (d, \(J = 8.60\), 1H), 5.36 (s, 2H), 2.40 (t, \(J = 7.05\), 2H), 1.56 (m, 2H), 1.26-1.22 (m, 20H), 0.85 (t, \(J = 6.00\), 3H); \(^{13}\)C NMR (125 MHz, CDCl\(_3\)) \(\delta\) (ppm), 190.40, 174.49, 172.42, 170.63, 133.28, 130.85, 123.29, 117.31, 116.06, 65.78, 33.62, 31.27, 29.04, 28.99, 28.95, 28.86, 28.69, 28.51, 28.33, 26.23, 24.47, 22.08, 13.94; HRMS (M + Na): Calc’d for C\(_{23}\)H\(_{34}\)O\(_6\)Na: 429.2253. Found: 429.2250.

**5-acetyl-2-(benzyloxy)benzoic acid, 157b.** The general benzylation method as 122b was followed. To a solution of methyl-5-acetyl-2-hydroxybenzoate (157a, 3.06 g, 15.7 mmol) in 50 ml of DMF was added potassium carbonate (6.53 g, 47.3 mmol) and benzyl bromide (4.68 ml, 39.4 mmol). The reaction was stirred at room temperature for 2 days. At this time, the mixture was diluted with 200 ml of 1:1 EtOAc/H\(_2\)O, the organic phase retained, concentrated, and flash chromatographed on silica gel (4:1 Hex/EtOAc) to obtain an oil, 4.40 g (98%) that was then dissolved in 100 ml of CH\(_3\)OH with 25 ml of aqueous 6 M KOH. The mixture was stirred for 3 h, at which time, conc. HCl was added slowly until an acidic solution was manifest by pH paper. The profuse solid material (KCl) was filtered off, 150 ml of CHCl\(_3\) added, the organic phase extracted, dried with MgSO\(_4\), and inspissated in vacuo to produce
157b as a white precipitate, 3.50 g (92%). This compound is known. $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 8.77 (d, $J = 2.00$, 1H), 8.23 (dd, $J = 8.68$, $J = 2.20$, 1H), 7.45 (m, 5H), 7.22 (d, $J = 8.75$, 1H), 5.38 (s, 2H), 2.63 (s, 3H).

5-acetyl-N-(4-benzoylephynyl)-2-(benzyloxy)benzamide, 157c. The general approach utilized by Fairbanks, et. al.$^{148}$ was applied. To a solution comprised of 157b (614 mg, 2.27 mmol) in 25 ml of dry CH$_2$Cl$_2$ was added DCC (469 mg, 2.27 mmol) and DMAP (69 mg, 0.568 mmol). This admixture was immured in an ice/saturated NaCl solution (~ 0°C). While stirring, a solution of 4-aminophenyl-(phenyl)methanone (448 mg, 2.27 mmol) in 10 ml of CH$_2$Cl$_2$ was slowly added dropwise. Once added, the reaction mixture was removed from the ice bath and persisted at room temperature overnight. At this time, the excess DCU was filtered off and remaining solution flash chromatographed (CH$_2$Cl$_2$/EtOAc) to afford 157c as a white precipitate, 58%. Mp: 185-187 ºC; IR (KBr): 3271, 3100-2800, 1685, 1670, 1649, 1593, 1535, 1500, 1323, 1311, 1200, 1054, 1028, 1008, 927, 823, 760 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 10.10 (s, 1H), 8.94, (d, $J = 2.32$, 1H), 8.24 (dd, $J = 8.94$, $J = 2.32$, 1H), 7.77-7.46 (m, 14H), 6.69 (d, $J = 8.28$, 1H), 5.33 (s, 2H), 2.68 (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) $\delta$ (ppm), 196.50, 195.64, 162.29, 159.96, 150.90, 142.07, 138.86, 135.49, 134.25, 133.63, 132.88, 132.19, 131.30, 129.78, 129.55, 128.98, 128.29, 127.20, 120.98, 118.92, 113.65, 72.48, 26.62; HRMS (M + H): Calc’d for C$_{29}$H$_{24}$NO$_4$: 450.1705. Found: 450.1700.

N-(4-benzoylephynyl)-2-(benzyloxy)-5-(2-bromoacetyl)benzamide, 157d. The synthetic method paralleled that for 122d. Oil, ~92%; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$
(ppm), 10.10 (s, 1H), 8.95, (d, J = 2.32, 1H), 8.24 (dd, J = 8.94, J = 2.32, 1H), 7.77-7.46 (m, 15H), 5.35 (s, 2H), 4.53 (s, 2H).

2-(3-(4-benzoylphenylcarbamoyl)-4-(benzyloxy)phenyl)-2-oxoethyl-4-(tert-butoxycarbonylamino)butanoate, 157e. The same reaction conditions were utilized as those to produce 122e. White ppt., 48%, Mp: 199-201°C; IR (KBr): 3271, 3100-2800, 1685, 1670, 1649, 1593, 1535, 1500, 1323, 1311, 1200, 1054, 1028, 1008, 927, 823, 760 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 10.69 (s, 1H), 8.20, (d, J = 2.10, 1H), 8.14 (dd, J = 9.00, J = 2.40, 1H), 7.77-7.46 (m, 14H), 6.89 (t, J = 5.95, 1H), 5.48 (s, 2H), 5.35 (s, 2H), 2.97 (t, J = 6.80, 2H), 2.40 (t, J = 7.80, 2H), 1.70 (m, 2H), 1.38 (s, 9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 194.56, 191.01, 172.23, 164.29, 159.60, 155.61, 142.90, 137.45, 136.00, 132.32, 132.08, 131.12, 131.73, 131.12, 129.48, 129.41, 128.50, 128.15, 127.71, 126.59, 125.80, 118.68, 113.41, 77.47, 70.35, 66.17, 30.65, 28.22, 24.95; HRMS (M + Na): Calc’d for C₃₈H₃₈N₂O₈Na: 673.2526. Found: 673.2534.

2-(3-(4-benzoylphenylcarbamoyl)-4-hydroxyphenyl)-2-oxoethyl-4-(tert-butoxycarbonylamino)butanoate, 157f. The same reaction conditions were utilized as those to produce 130'. Adhesive ppt., 86%; ¹H NMR (400 MHz, CD₃CN) δ (ppm), 9.21 (s, 1H), 8.45, (d, J = 2.20, 1H), 8.03 (dd, J = 9.00, J = 2.40, 1H), 7.56 (d, J = 8.65, 2H), 7.31-7.24 (m, 7H), 7.05 (d, J = 7.20, 1H), 5.48 (s, 2H), 5.35 (s, 2H), 3.09 (t, J = 7.00, 2H), 2.45 (t, J = 7.80, 2H), 1.78 (m, 2H), 1.40 (s, 9H).

4-(2-(3-(4-benzoylphenylcarbamoyl)-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 157. The same reaction conditions
were utilized as those to generate 130. Adhesive ppt., 74%; IR (KBr): 3271, 3100-2800, 1685, 1670, 1649, 1593, 1535, 1500, 1323, 1311, 1200, 1054, 1028, 1008, 927, 823, 760 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 12.75 (s, 1H), 10.57 (s, 1H), 8.56 (d, J = 2.00, 1H), 8.08 (dd, J = 8.80, J = 2.10, 1H), 7.87 (s, 3H), 7.69 (d, J = 8.20, 2H), 7.35 (t, J = 7.68, 2H), 7.30 (m, 4H), 7.25 (t, J = 7.30, 1H), 7.18 (d, J = 8.69, 1H), 5.56 (s, 2H), 2.96 (t, J = 6.80, 2H), 2.65 (t, J = 7.80, 2H), 1.96 (m, 2H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.68, 171.77, 165.31, 162.69, 157.90, 157.65, 141.35, 137.96, 135.97, 132.78, 130.02, 129.00, 128.61, 128.41, 125.94, 125.12, 121.06, 118.60, 117.61, 116.05, 66.28, 38.11, 30.12, 22.59; HRMS (M+): Calc’d for C₂₆H₂₅N₂O₆: 461.1707. Found: 461.1715.

The same reaction conditions were utilized to produce 159-165.

3,5-difluorophenyl-5-acetyl-2-(benzyloxy)benzoate, 158c. White ppt., 82%, Mp: 115-117 °C; IR (KBr): 3275, 3100-2800, 1751, 1731, 1677, 1606, 1502, 1465, 1363, 1271, 1208, 1122, 1056, 1028, 1008, 822, 760 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.60, (d, J = 2.20, 1H), 8.19 (dd, J = 9.00, J = 2.20, 1H), 7.50 (d, J = 8.10, 2H), 7.38 (m, 3H), 7.18 (d, J = 8.2, 2H), 6.77 (m, 3H), 5.30 (s, 2H), 2.63 (s, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 195.82, 164.05, 162.93, 161.95, 152.16, 135.34, 134.81, 133.67, 132.84, 128.91, 128.19, 127.09, 118.69, 113.33, 106.19, 101.78, 70.63, 26.42; HRMS (M + H): Calc’d for C₂₂H₂₅F₂O₄Na: 405.0914. Found: 405.0918.

3,5-difluorophenyl-2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 158d. Oil, ~93%; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.63, (d, J = 2.20, 1H), 8.20 (dd, J = 9.00, J =
2.20 (1H), 7.50 (d, J = 8.10, 2H), 7.38 (m, 3H), 7.18 (d, J = 8.2, 2H), 6.77 (m, 3H),
5.32 (s, 2H), 4.44 (s, 2H).

3,5-difluorophenyl-2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonylamino)-butanoyloxy)acetyl)benzoate, 158e. Oil, 44%; IR (KBr): 3271, 3100–2800, 1685, 1670, 1649, 1593, 1535, 1500, 1323, 1311, 1200, 1054, 1028, 1008, 927, 823, 760 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.50, (d, J = 2.10, 1H), 8.24 (dd, J = 9.00, J = 2.40, 1H), 7.53–7.49 (m 6H), 7.39 (d, J = 5.60, 1H), 7.13 (dd, J = 8.20, J = 2.10, 2H), 6.89 (t, J = 5.95, 1H), 5.49 (s, 2H), 5.40 (s, 2H), 2.98 (t, J = 6.80, 2H), 2.42 (t, J = 7.80, 2H), 1.70 (m, 2H), 1.37 (s, 9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.88, 172.44, 170.36, 164.36, 162.80, 161.74, 155.55, 151.39, 135.99, 134.59, 131.77, 128.44, 127.65, 126.97, 118.45, 114.22, 106.30, 102.18, 77.44, 70.32, 66.16, 59.75, 39.50, 30.42, 28.71, 24.33; HRMS (M + Na): Calc’d for C₃₁H₃₃F₂NO₈ Na: 606.1916. Found: 606.1899. (M + H): Calc’d for C₃₁H₃₂F₂NO₈: 584.2096. Found: 584.2078.

3,5-difluorophenyl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoate, 158f. Adhesive ppt., 86%; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.56, (d, J = 2.10, 1H), 8.22 (dd, J = 9.00, J = 2.40, 1H), 7.22 (d, J = 9.60, 1H), 7.39 (d, J = 5.60, 1H), 7.08 (dd, J = 8.20, J = 1.90, 1H), 7.02 (m, 2H), 5.49 (s, 2H), 2.97 (t, J = 6.80, 2H), 2.42 (t, J = 7.80, 2H), 1.69 (m, 2H), 1.39 (s, 9H).

4-(2-(3-(3,5-difluorophenoxy)carbonyl)-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 158. Adhesive ppt., 82%; IR (KBr): 3271, 3100-2800, 1685, 1670, 1649, 1593, 1535, 1500, 1323, 1311, 1200, 1054,
1028, 1008, 927, 823, 760 cm$^{-1}$; $^1$H NMR (400 MHz, MeOD) $\delta$ (ppm), 8.68, (d, $J$ = 2.10, 1H), 8.22 (dd, $J$ = 9.10, $J$ = 2.40, 1H), 7.19 (d, $J$ = 9.10, 1H), 7.07 (d, $J$ = 5.42, 1H), 7.00 (m, 2H), 5.51 (s, 2H), 3.08 (t, $J$ = 6.80, 2H), 2.66 (t, $J$ = 7.80, 2H), 2.04 (m, 2H). $^{19}$F NMR (376 MHz, MeOD) $\delta$ (ppm), -109.79; $^{13}$C NMR (125 MHz, MeOD) $\delta$ (ppm), 190.89, 172.05, 166.59, 165.62, 164.08, 162.23, 161.54, 151.63, 135.33, 131.31, 126.05, 118.20, 112.15, 106.07, 101.66, 66.00, 38.80, 28.69, 22.50; HRMS (M+): Calc'd for C$_{19}$H$_{18}$F$_2$NO$_6$: 394.1102. Found: 394.1101.

2-(trifluoromethyl)phenyl-5-acetyl-2-(benzyloxy)benzoate, 159c. White ppt., 51%. Mp: 83-85 °C; IR (KBr): 3271, 3100-2800, 1755, 1681, 1598, 1502, 1494, 1456, 1409, 1361, 1323, 1197, 1168, 1132, 1112, 1056, 1028, 1009, 822, 760, 698 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 8.72, (d, $J$ = 2.20, 1H), 8.20 (dd, $J$ = 9.00, $J$ = 2.20, 1H), 7.75 (d, $J$ = 8.20, 1H), 7.65 (dd, $J$ = 12.05, $J$ = 8.00, 1H), 7.52 (d, $J$ = 8.20, 2H), 7.43-7.32 (m, 5H), 7.17 (dd, $J$ = 8.50, $J$ = 2.20, 1H), 5.35 (s, 2H), 2.63 (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) $\delta$ (ppm), 195.95, 162.59, 162.39, 148.22, 135.63, 134.63, 133.14, 129.88, 128.72, 128.13, 127.04, 126.37, 124.61, 124.20, 123.36, 122.03, 119.86, 118.21, 113.36, 70.76, 26.31; HRMS (M + Na): Calc’d for C$_{23}$H$_{17}$F$_3$O$_4$Na: 437.0977. Found: 437.0966.

2-(trifluoromethyl)phenyl-2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 159d. Oil, ~93%. $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ (ppm), 8.72, (d, $J$ = 2.20, 1H), 8.20 (dd, $J$ = 9.00, $J$ = 2.20, 1H), 7.75 (d, $J$ = 8.20, 1H), 7.65 (dd, $J$ = 12.05, $J$ = 8.00, 1H), 7.51 (d, $J$ = 8.20, 2H), 7.43-7.32 (m, 5H), 7.17 (dd, $J$ = 8.50, $J$ = 2.20, 1H), 5.35 (s, 2H), 4.45 (s, 2H).
2-((trifluoromethyl)phenyl-2-(benzyloxy)-5-(2-((4-(tert-butoxycarbonyl-amino)-butanoyloxy)acetyl)benzoate, 159e. White ppt., 44%, Mp: 117-119 °C; IR (KBr): 3271, 3100-2800, 1753, 1697, 1666, 1600, 1529, 1494, 1456, 1415, 1380, 1367, 1321, 1272, 1195, 1166, 1143, 1112, 1055, 1028, 1009, 821, 760 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.52, (d, \(J = 2.20\), 1H), 8.37 (dd, \(J = 9.00, J = 2.20\), 1H), 7.85 (m, 2H), 7.58-7.50 (m, 5H), 7.37 (t, \(J = 9.00\), 2H), 7.31 (d, \(J = 8.10\), 1H), 6.89 (t, \(J = 5.80\), 1H), 5.48 (s, 2H), 5.42 (s, 2H), 2.98 (t, \(J = 6.80\), 2H), 2.42 (t, \(J = 7.80\), 2H), 1.68 (m, 2H), 1.37 (s, 9H); \(^1^3\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.80, 172.23, 162.09, 155.61, 147.63, 135.99, 134.90, 134.22, 131.55, 128.41, 127.88, 127.06, 126.84, 126.34, 125.10, 124.11, 121.95, 121.20, 119.78, 117.98, 114.35, 77.46, 70.19, 66.09, 39.43, 30.62, 28.20, 20.73; HRMS (M + Na): Calc’d for C\(_{32}\)H\(_{32}\)F\(_3\)O\(_8\)Na: 638.1978. Found: 638.1960.

2-((trifluoromethyl)phenyl-5-(2-((4-(tert-butoxycarbonyl-amino)-butanoyloxy)acetyl)-2-hydroxybenzoate, 159f. Adhesive ppt., 90%; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 11.43, (s, 1H), 8.51, (d, \(J = 2.20\), 1H), 8.17 (dd, \(J = 9.00, J = 2.20\), 1H), 7.62 (d, \(J = 8.40\), 2H), 7.56 (m, 2H), 7.19 (d, \(J = 8.10\), 1H), 6.89 (t, \(J = 5.80\), 1H), 5.44 (s, 2H), 2.98 (t, \(J = 6.80\), 2H), 2.43 (t, \(J = 7.80\), 2H), 1.68 (m, 2H), 1.37 (s, 9H).

4-((2-(4-hydroxy-3-((2-(trifluoromethyl)phenoxy)carbonyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 159. Adhesive ppt., 81%; IR (KBr): 3442, 3271, 3100-2800, 1741, 1691, 1645, 1600, 1494, 1454, 1417, 1380, 1323, 1269, 1199, 1172, 1130, 1053, 1026, 821, 800, 760, 717 cm\(^{-1}\); \(^1\)H NMR (400 MHz,
d6-DMSO) δ (ppm), 11.43 (s, 1H), 8.51, (d, J = 2.20, 1H), 8.17 (dd, J = 9.00, J = 2.20, 1H), 7.88-7.83 (m, 5H), 7.62-7.58 (m, 2H), 7.19 (d, J = 8.20, 1H), 5.49 (s, 2H), 2.90 (t, J = 6.80, 2H), 2.42 (t, J = 7.80, 2H), 1.86 (m, 2H); 19F NMR (376 MHz, d6-DMSO) δ (ppm), -63.07; 13C NMR (125 MHz, d6-DMSO) δ (ppm), 190.46, 171.74, 165.18, 163.73, 159.43, 158.27, 147.48, 134.97, 134.20, 131.86, 128.46, 126.94, 125.21, 124.13, 121.61, 118.11, 115.73, 114.45, 66.16, 38.10, 30.15, 22.32; HRMS (M+): Calc’d for C20H19NF3O6: 426.1165. Found: 426.1140.

2-fluorophenyl 5-acetyl-2-(benzyloxy)benzoate, 160c. White ppt., 57%, Mp: 91-93 °C; IR (KBr): 3271, 3100-2800, 1755, 1681, 1598, 1502, 1494, 1456, 1409, 1361, 1323, 1197, 1168, 1135, 1056, 1028, 1009, 822, 760, 698 cm−1; 1H NMR (400 MHz, CDCl3) δ (ppm), 8.73, (d, J = 2.20, 1H), 8.23 (dd, J = 9.00, J = 2.20, 1H), 7.40 (t, J = 7.50, 2H), 7.38-7.33 (m, 3H), 7.28-7.20 (m, 5H), 5.34 (s, 2H), 2.71 (s, 3H); 19F NMR (376 MHz, CDCl3) δ (ppm), -128.93; 13C NMR (125 MHz, CDCl3) δ (ppm), 190.82, 172.10, 169.24, 161.98, 155.77, 154.92, 151.67, 138.03, 136.08, 134.43, 131.67, 128.55, 127.00, 124.36, 124.87, 118.09, 116.78, 115.42, 77.01, 26.87; HRMS (M + Na): Calc’d for C22H17FO4Na: 387.1009. Found: 387.1015.

2-fluorophenyl-2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 160d. Oil, ~84%; 1H NMR (400 MHz, CDCl3) δ (ppm), 8.78, (d, J = 2.20, 1H), 8.22 (dd, J = 9.00, J = 2.20, 1H), 7.40 (t, J = 7.50, 2H), 7.38-7.33 (m, 3H), 7.28-7.20 (m, 5H), 5.36 (s, 2H), 4.44 (s, 2H).

3-fluorophenyl-2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)-butanoyloxy)acetyl)benzoate, 160e. White ppt., 74%, Mp: 114-116 °C; IR (KBr): 3271, 3100-
2800, 1755, 1681, 1598, 1502, 1494, 1456, 1409, 1361, 1323, 1197, 1168, 1132, 1112, 1056, 1028, 1009, 822, 760, 698 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.51, (d, \(J = 2.20\), 1H), 7.51 (m, 3H), 7.38-7.33 (m, 3H), 7.28-7.20 (m, 5H), 6.88 (t, \(J = 6.90\), 1H), 5.49 (s, 2H), 5.41 (s, 2H), 2.98 (t, \(J = 6.80\), 2H), 2.42 (t, \(J = 7.80\), 2H), 1.68 (m, 2H), 1.37 (s, 9H); \(^19\)F NMR (376 MHz, CDCl\(_3\)) \(\delta\) (ppm), -128.98; \(^13\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.91, 172.24, 170.37, 161.92, 155.62, 154.57, 152.61, 137.66, 136.01, 134.74, 131.67, 128.41, 127.65, 127.06, 126.39, 125.21, 124.23, 118.09, 116.79, 114.37, 77.49, 70.23, 66.18, 59.75, 30.64, 28.21, 24.96; HRMS (M + Na): Calc’d for C\(_{31}\)H\(_{32}\)FNO\(_8\)Na: 588.2010. Found: 588.2021.

**2-fluorophenyl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoate, 160f.** White ppt., 74%, Mp: 114-116 °C; IR (KBr): 3271, 3100-2800, 1755, 1681, 1598, 1502, 1494, 1456, 1409, 1361, 1323, 1197, 1168, 1132, 1112, 1056, 1028, 1009, 822, 760, 698 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.52, (d, \(J = 2.20\), 1H), 8.13 (dd, \(J = 9.00, J = 2.20\), 1H), 7.45-7.29 (m, 5H), 7.19 (d, \(J = 8.10\), 1H), 5.44 (s, 2H), 2.96 (t, \(J = 6.80\), 2H), 2.42 (t, \(J = 7.80\), 2H), 1.68 (m, 2H), 1.37 (s, 9H); \(^19\)F NMR (376 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), -130.36; \(^13\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.67, 172.24, 170.37, 161.92, 155.62, 154.56, 152.59, 137.42, 134.88, 132.08, 127.91, 125.37, 124.35, 118.15, 116.76, 114.20, 77.49, 66.18, 59.75, 30.64, 28.21, 24.96; HRMS (M + Na): Calc’d for C\(_{24}\)H\(_{26}\)FNO\(_8\)Na: 498.1540. Found: 498.1533.

**4-(2-(3-((2-fluorophenoxy)carbonyl)-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 160.** Adhesive ppt., 92%; IR (KBr):
3271, 3100-2800, 1755, 1681, 1598, 1502, 1494, 1456, 1409, 1361, 1323, 1197, 1168, 1132, 1112, 1056, 1028, 1009, 822, 760, 698 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 11.55 (s, 1H), 8.55 (d, J = 2.20, 1H), 8.14 (dd, J = 9.00, J = 2.20, 1H), 7.86 (s, 3H), 7.47-7.39 (m, 4H), 7.21 (d, J = 8.10, 1H), 5.51 (s, 2H), 2.89 (t, J = 6.80, 2H), 2.59 (t, J = 7.80, 2H), 1.86 (m, 2H), 1.37 (s, 9H); ¹⁹F NMR (376 MHz, d⁶-DMSO) δ (ppm), -130.75; ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.58, 171.73, 163.63, 163.14, 158.23, 154.56, 152.60, 134.83, 132.11, 127.88, 125.22, 124.35, 118.17, 116.76, 115.64, 114.34, 66.26, 38.08, 30.10, 22.57; HRMS (M⁺): Calc’d for C₁₉H₁₉FNO₆: 376.1191. Found: 376.1197.

3-nitrophenyl 5-acetyl-2-(benzyloxy)benzoate, 161c. White ppt., 46%; Mp: 106-108°C; IR (KBr): 3271, 3100-2800, 1728, 1712, 1677, 1604, 1502, 1454, 1392, 1353, 1321, 1284, 1209, 1168, 1051, 1024, 1004, 902, 823, 761, 725 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.69 (d, J = 2.20, 1H), 8.17-8.15 (m, 2H), 8.10 (d, J = 8.20, 1H), 7.59-7.49 (m, 4H), 7.41-7.35 (m, 3H), 7.20 (d, J = 8.60, 1H), 5.31 (s, 2H), 2.64 (s, 3H); ¹³C NMR (125 MHz, CDCl₃) δ (ppm), 196.02, 163.13, 162.36, 151.09, 148.85, 135.49, 133.36, 131.40, 130.10, 128.35, 127.20, 121.97, 120.93, 118.48, 117.64, 115.45, 113.39, 112.19, 71.02, 26.63; HRMS (M + Na): Calc’d for C₂₂H₁₇NO₆Na: 414.0954. Found: 414.0966.

3-nitrophenyl 2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 161d. Oil, ~65%; ¹H NMR (400 MHz, CDCl₃) δ (ppm), 8.71 (d, J = 2.20, 1H), 8.17-8.15 (m, 2H), 8.10 (d, J = 8.20, 1H), 7.59-7.49 (m, 4H), 7.41-7.35 (m, 3H), 7.22 (d, J = 8.60, 1H), 5.35 (s, 2H), 4.46 (s, 2H).
3-nitrophenyl-2-(benzylxoy)-5-(2-(4-(tert-butoxycarbonyl-amino)butanoyloxy)-acetyl)benzoate, 161e. Adhesive ppt., 39%; IR (KBr): 3271, 3100-2800, 1728, 1712, 1677, 1604, 1502, 1454, 1392, 1353, 1321, 1284, 1209, 1168, 1051, 1024, 1004, 902, 823, 761, 725 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.56, (d, \(J = 2.30, 1\)H), 8.20 (m, 3H), 7.75 (d, \(J = 8.00, 2\)H), 7.52-7.48 (m, 3H), 7.37-7.34 (m, 3H), 6.89 (t, \(J = 7.10, 1\)H), 5.49 (s, 2H), 5.40 (s, 2H), 2.96 (t, \(J = 6.90, 2\)H), 2.43 (t, \(J = 7.70, 2\)H), 1.67 (m, 2H), 1.36 (s, 9H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.67, 172.34, 164.67, 161.54, 155.80, 150.84, 148.34, 136.36, 134.51, 131.82, 129.85, 128.84, 128.13, 127.96, 127.66, 126.94, 121.11, 118.24, 117.48, 114.27, 78.77, 70.42, 66.13, 39.72, 30.34, 28.14, 24.27; HRMS (M + Na): Calc’d for C\(_{31}\)H\(_{32}\)N\(_2\)O\(_{10}\)Na: 615.1955. Found: 615.1971.

3-nitrophenyl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)-acetyl)-2-hydroxybenzoate, 161f. Adhesive ppt., 87%; \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.52, (d, \(J = 2.30, 1\)H), 8.13 (dd, \(J = 8.40, J = 2.30, 1\)H), 7.70 (d, \(J = 8.00, 1\)H), 7.59-7.51 (m, 2H), 7.19 (dd, \(J = 7.90, J = 1.90, 2\)H), 6.88 (t, \(J = 7.10, 1\)H), 5.49 (s, 2H), 2.97 (t, \(J = 6.90, 2\)H), 2.44 (t, \(J = 7.70, 2\)H), 1.67 (m, 2H), 1.36 (s, 9H).

4-(2-(4-hydroxy-3-((3-nitrophenoxy)carbonyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 161. White ppt., 80%, Mp: 58-60°C; IR (KBr): 3271, 3100-2800, 1728, 1712, 1677, 1604, 1502, 1454, 1392, 1353, 1321, 1284, 1209, 1168, 1051, 1024, 1004, 902, 823, 761, 725 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 8.51, (d, \(J = 2.30, 1\)H), 8.13 (dd, \(J = 8.40, J = 2.30, 1\)H), 7.84 (s, 3H), 7.71 (d, \(J = 8.00, 1\)H), 7.59-7.51 (m, 2H), 7.19 (dd, \(J = 7.90, J = 1.90, 2\)H), 5.52
(s, 2H), 2.91 (t, J = 6.90, 2H), 2.60 (t, J = 7.70, 2H), 1.86 (m, 2H); $^{13}$C NMR (125 MHz, d$^6$-DMSO) δ (ppm), 190.60, 171.74, 164.55, 158.22, 157.77, 154.78, 150.11, 137.46, 134.55, 131.87, 130.01, 125.16, 122.46, 120.44, 118.68, 116.79, 115.89, 114.49, 66.29, 38.09, 30.08, 22.57; HRMS (M+): Calc’d for C$_{19}$H$_{19}$N$_2$O$_8$: 403.1136. Found: 403.1145.

4-nitrophenyl 5-acetyl-2-(benzylloxy)benzoate, 162c. White ppt., 91%. Mp: 93-95°C; IR (KBr): 3250, 3100-2800, 1751, 1728, 1677, 1649, 1598, 1523, 1411, 1348, 1269, 1195, 1165, 1055, 1026, 1007, 905, 860, 823, 760 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 8.64, (d, J = 2.20, 1H), 8.30 (d, J = 9.20, 2H), 8.21 (d, J = 8.40, 1H), 7.50 (dd, J = 8.40, J = 2.10, 2H), 7.41-7.35 (m, 5H), 7.20 (d, J = 8.65, 1H), 5.31 (s, 2H), 2.63 (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm), 196.07, 171.26, 162.79, 155.58, 145.43, 135.47, 133.55, 131.44, 130.17, 128.79, 127.57, 126.76, 125.53, 122.70, 118.48, 115.67, 71.02, 26.44; HRMS (M + Na): Calc’d for C$_{22}$H$_{17}$NO$_6$Na: 414.0954. Found: 414.0969.

4-nitrophenyl-2-(benzylloxy)-5-(2-bromoacetyl)benzoate, 162d. Oil, ~84%; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 8.70, (d, J = 2.20, 1H), 8.33 (d, J = 9.20, 2H), 8.23 (d, J = 8.40, 1H), 7.51 (dd, J = 8.40, J = 2.10, 2H), 7.41-7.35 (m, 5H), 7.24 (d, J = 8.65, 1H), 5.35 (s, 2H), 4.46 (s, 2H).

4-nitrophenyl2-(benzylloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)butanoyloxy)acetyl)benzoate, 162e. White ppt., 57%, Mp: 105-107 °C; IR (KBr): 3275, 3100-2800, 1745, 1697, 1658, 1602, 1523, 1413, 1348, 1271, 1195, 1165, 1051, 1026, 1007, 860, 823, 761 cm$^{-1}$; $^1$H NMR (400 MHz, d$^6$-DMSO) δ (ppm), 8.55, (d, J
$\delta$ (ppm), 190.88, 172.22, 162.33, 161.89, 156.56, 155.31, 145.18, 136.02, 134.64, 131.89, 128.46, 127.66, 127.18, 126.35, 125.34, 123.32, 118.44, 114.28, 77.46, 70.30, 66.19, 40.06, 30.64, 28.22, 24.44; HRMS (M + Na): Calc’d for $C_{31}H_{32}N_2O_{10}$Na: 615.1955. Found: 615.1937.

4-nitrophenyl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoate, 162f. White ppt., 87%, Mp: 144-146°C; IR (KBr): 3275, 3100-2800, 1745, 1697, 1658, 1602, 1523, 1413, 1348, 1271, 1195, 1165, 1051, 1026, 1007, 860, 823, 761 cm$^{-1}$; $^1$H NMR (400 MHz, $d_6$-DMSO) $\delta$ (ppm), 11.30 (s, 1H), 8.54, (d, $J = 2.20$, 2H), 8.38 (d, $J = 9.20$, 2H), 8.18 (d, $J = 8.40$, 1H), 7.65 (dd, $J = 8.40, J = 2.10$, 2H), 7.20 (d, $J = 8.65$, 1H), 6.89 (t, $J = 7.10$, 1H), 5.47 (s, 2H), 2.98 (t, $J = 6.90$, 2H), 2.43 (t, $J = 7.70$, 2H), 1.68 (m, 2H), 1.38 (s, 9H); $^{13}$C NMR (125 MHz, $d_6$-DMSO) $\delta$ (ppm), 190.63, 172.00, 165.08, 156.04, 145.21, 142.34, 134.78, 132.13, 130.99, 126.12, 125.31, 123.44, 118.08, 114.64, 77.45, 66.08, 39.96, 30.64, 28.21, 24.95; HRMS (M + Na): Calc’d for $C_{24}H_{26}N_2O_{10}$Na: 525.1485. Found: 525.1487.

4-(2-(4-hydroxy-3-((4-nitrophenoxy)carbonyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 162. White ppt., 91%, Mp: 88-90°C; IR (KBr): 3275, 3100-2800, 1745, 1697, 1658, 1602, 1523, 1413, 1348, 1271, 1195, 1165, 1051, 1026, 1007, 860, 823, 761 cm$^{-1}$; $^1$H NMR (400 MHz, $d_6$-DMSO) $\delta$ (ppm), 11.48
(s, 1H), 8.54, (d, J = 2.20, 2H), 8.38 (d, J = 9.20, 2H), 8.15 (d, J = 8.40, 1H), 7.80 (s, 3H), 7.63 (dd, J = 8.40, J = 2.10, 2H), 7.19 (d, J = 8.65, 1H), 5.51 (s, 2H), 2.89 (t, J = 6.90, 2H), 2.59 (t, J = 7.70, 2H), 1.86 (m, 2H); 13C NMR (125 MHz, d6-DMSO) δ (ppm), 190.59, 171.78, 163.55, 157.99, 157.75, 155.18, 145.23, 134.76, 132.19, 125.34, 125.19, 123.44, 118.38, 115.99, 114.76; HRMS (M+): Calc’d for C_{19}H_{19}N_{2}O_{8}Na: 403.1141. Found: 403.1131.

3,5-bis(trifluoromethyl)phenyl 5-acetyl-2-(benzyloxy)benzoate, 163c. White ppt., 90%, Mp: 73-75°C; IR (KBr): 3270, 3100-2800, 1753, 1731, 1679, 1629, 1598, 1500, 1413, 1371, 1282, 1205, 1180, 1137, 1056, 1029, 1008, 844, 821, 760, 700 cm⁻¹; 1H NMR (400 MHz, CDCl₃) δ (ppm), 8.65, (d, J = 2.20, 1H), 8.23 (dd, J = 8.80, J = 2.40, 1H), 7.80 (d, J = 2.20, 1H), 7.67 (d, J = 8.80, 2H), 7.50 (dd, J = 8.40, J = 2.10, 2H), 7.42-7.36 (m, 3H), 7.20 (d, J = 8.65, 1H), 5.31 (s, 2H), 2.64 (s, 3H); 13C NMR (125 MHz, CDCl₃) δ (ppm), 195.71, 171.20, 162.94, 162.40, 151.27, 135.28, 133.45, 132.55, 130.01, 128.81, 127.35, 126.03, 123.86, 121.69, 119.84, 118.15, 113.41, 71.13, 26.42; HRMS (M + Na): Calc’d for C_{24}H_{16}F_{6}O_{4}Na: 505.0850. Found: 505.0866.

3,5-bis(trifluoromethyl)phenyl 2-(benzyloxy)-5-propionylbenzoate, 163d. Oil, ~92%. 1H NMR (400 MHz, CDCl₃) δ (ppm), 8.69, (d, J = 2.20, 1H), 8.25 (dd, J = 8.80, J = 2.40, 1H), 7.81 (d, J = 2.20, 1H), 7.68 (d, J = 8.80, 2H), 7.51 (dd, J = 8.40, J = 2.10, 2H), 7.43-7.37 (m, 3H), 7.23 (d, J = 8.65, 1H), 5.33 (s, 2H), 4.45 (s, 2H).

3,5-bis(trifluoromethyl)phenyl 2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)butanoyloxy)acetyl)benzoate, 163e. Adhesive ppt., 58%; IR (KBr): 3270,
3100-2800, 1745, 1697, 1602, 1529, 1500, 1460, 1371, 1282, 1253, 1174, 1139, 1054, 1029, 1008, 823, 760, 680 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 8.60, (d, J = 2.20, 1H), 8.29 (dd, J = 8.80, J = 2.40, 1H), 8.11 (m, 3H), 7.54-7.50 (m, 3H), 7.39-7.34 (m, 3H), 7.20 (d, J = 8.65, 1H), 6.89 (t, J = 7.10, 1H), 5.50 (s, 2H), 5.41 (s, 2H), 2.97 (t, J = 6.90, 2H), 2.44 (t, J = 7.70, 2H), 1.70 (m, 2H), 1.38 (s, 9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.86, 172.24, 162.38, 161.99, 155.60, 151.32, 135.98, 134.72, 132.09, 131.66, 128.40, 128.00, 127.32, 126.30, 123.91, 121.72, 119.98, 118.22, 114.19, 77.46, 70.40, 66.17, 39.63, 30.64, 28.21, 24.97; HRMS (M + Na): Calc’d for C₃₃H₃₁F₆NO₈Na: 706.1852. Found: 706.1844; (M + H): Calc’d for C₃₃H₃₂F₆NO₈: 684.2032. Found: 684.2028.

3,5-bis(trifluoromethyl)phenyl-5-(2-(4-((tert-butoxycarbonyl)amino)butanoyl-oxy)acetyl)-2-hydroxybenzoate, 163f. Adhesive ppt., 35%; ¹H NMR (400 MHz, CD₃CN) δ (ppm), 8.70, (d, J = 2.20, 1H), 8.16 (dd, J = 8.80, J = 2.40, 1H), 8.01 (m, 3H), 7.15 (d, J = 8.65, 1H), 5.36 (s, 2H), 3.08 (t, J = 6.90, 2H), 2.44 (t, J = 7.70, 2H), 1.79 (m, 2H), 1.39 (s, 9H).

4-(2-(3-(3,5-bis(trifluoromethyl)phenoxy)carbonyl)-4-hydroxyphenyl)-2-oxoethoxy)-4-oxobutan-1-aminium-2,2,2-trifluoroacetate, 163. White ppt., 93%, Mp: 77-79°C; IR (KBr): 3438, 3270, 3100-2800, 1741, 1691, 1600, 1460, 1413, 1373, 1282, 1201, 1176, 1130, 1054, 1029, 1008, 823, 761, 717, 680 cm⁻¹; ¹H NMR (400 MHz, d⁶-DMSO) δ (ppm), 11.49, (s, 1H), 8.58, (d, J = 2.20, 1H), 8.21 (d, J = 8.80, 2H), 8.14 (d, J = 6.10, 2H), 7.85 (s, 3H), 7.20 (d, J = 8.65, 1H), 5.51 (s, 2H), 2.88 (t, J = 6.90, 2H), 2.59 (t, J = 7.70, 2H), 1.88 (m, 2H); ¹³C NMR (125 MHz, d⁶-
DMSO) δ (ppm), 190.55, 171.76, 163.68, 157.97, 157.48, 151.23, 134.81, 132.35, 131.34, 125.14, 124.07, 121.75, 119.97, 118.24, 115.88, 114.51, 66.26, 38.08, 30.10, 22.57; HRMS (M+): Calc’d for C_{21}H_{18}F_{6}NO_{6}: 494.1038. Found: 494.1021.

**quinolin-8-yl 5-acetyl-2-(benzyloxy)benzoate, 164c.** White ppt., 86%, Mp: 179-181°C; IR (KBr): 3275, 3100-2800, 1745, 1670, 1598, 1581, 1496, 1467, 1448, 1382, 1276, 1238, 1199, 1163, 1056, 1028, 1007, 873, 821, 760, 692, 680 cm^{-1}; 1H NMR (400 MHz, CDCl₃) δ (ppm), 8.92, (m, 2H), 8.23 (m, 2H), 7.78 (dd, J = 7.20, J = 2.47, 1H), 7.59-7.52 (m, 4H), 7.46 (dd, J = 8.28, J = 4.25, 1H), 7.32-7.28 (m, 3H), 7.18 (d, J = 8.70, 1H), 5.34 (s, 2H), 2.65 (s, 3H); 13C NMR (125 MHz, CDCl₃) δ (ppm), 196.24, 163.86, 162.41, 156.72, 150.62, 147.60, 136.00, 135.87, 134.29, 134.06, 129.92, 129.63, 128.59, 128.42, 128.10, 127.07, 126.82, 126.29, 121.79, 119.54, 113.41, 70.71, 24.96; HRMS (M + H): Calc’d for C_{25}H_{20}NO_{4}: 398.1392. Found: 398.1389.

**quinolin-8-yl 2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 164d.** Oil, ~84%. 1H NMR (400 MHz, CDCl₃) δ (ppm), 8.96, (m, 2H), 8.23 (m, 2H), 7.80 (dd, J = 7.20, J = 2.47, 1H), 7.63-7.58 (m, 4H), 7.55 (dd, J = 8.28, J = 4.25, 2H), 7.49-7.30 (m, 3H), 7.21 (d, J = 8.70, 1H), 5.37 (s, 2H), 4.52 (s, 2H).

**quinolin-8-yl 2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)-butanoyloxy)-acetyl)benzoate, 164e.** Adhesive ppt., 48%; IR (KBr): 3275, 3100-2800, 1745, 1670, 1598, 1581, 1496, 1467, 1448, 1382, 1276, 1238, 1199, 1163, 1056, 1028, 1007, 873, 821, 760, 692, 680 cm^{-1}; 1H NMR (400 MHz, d^{6}-DMSO) δ (ppm), 8.91, (d, J = 8.05, 1H), 8.89 (d, J = 7.60, 1H), 8.70 (d, J = 9.40, 1H), 8.50 (d, J = 8.75, 1H), 7.97 (dd, J
quinolin-8-yl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-
hydroxybenzoate, 164f. White ppt., 92%, Mp: 137-139°C; IR (KBr): 3469, 3298,
3274, 3100-2800, 1743, 1697, 1664, 1631, 1604, 1500, 1421, 1365, 1299, 1249,
1232, 1199, 1166, 1056, 1027, 1009, 875, 821, 760 cm⁻¹; ¹H NMR (400 MHz, d⁶-
DMSO) δ (ppm), 11.34 (s, 1H), 8.92, (d, J = 8.05, 1H), 8.67 (d, J = 7.60, 1H), 8.52
(d, J = 9.40, 1H), 8.21 (d, J = 8.75, 1H), 8.01 (dd, J = 7.50, J = 2.10, 1H), 7.76 (d, J =
8.20, 2H), 7.71 (dd, J = 8.28, J = 4.25, 1H), 7.21 (d, J = 8.40, 1H), 6.89 (d, J = 7.10,
1H), 5.48 (s, 2H), 2.96 (t, J = 6.90, 2H), 2.43 (t, J = 7.70, 2H), 1.68 (m, 2H), 1.37 (s,
9H); ¹³C NMR (125 MHz, d⁶-DMSO) δ (ppm), 190.76, 172.24, 164.92, 163.64,
155.59, 150.85, 146.45, 140.05, 136.53, 134.94, 131.91, 129.14, 128.34, 126.98,
125.60, 122.34, 121.88, 118.31, 114.49, 77.45, 66.10, 39.47, 30.65, 28.21, 24.95;
Calc’d for C₂₇H₂₈N₂O₈Na: 531.1743. Found: 531.1739.
4-(2-(4-hydroxy-3-((quinolin-8-yl)oxy)carbonyl)phenyl)-2-oxoethoxy)-4-hydroxy-1-aminium 2,2,2-trifluoroacetate, 164. Adhesive ppt., 92%; IR (KBr): 3469, 3298, 3274, 3100-2800, 1743, 1697, 1664, 1631, 1604, 1500, 1421, 1365, 1299, 1249, 1232, 1199, 1166, 1056, 1027, 1009, 875, 821, 760 cm$^{-1}$; $^1$H NMR (400 MHz, d$^6$-DMSO) δ (ppm), 11.34 (s, 1H), 8.91, (d, $J = 8.05$, 1H), 8.67 (d, $J = 7.60$, 1H), 8.52 (d, $J = 9.40$, 1H), 8.21 (d, $J = 8.75$, 1H), 8.01 (dd, $J = 7.50$, $J = 2.10$, 1H), 7.80 (s, 3H), 7.76 (d, $J = 8.20$, 2H), 7.71 (dd, $J = 8.28$, $J = 4.25$, 1H), 7.21 (d, $J = 8.40$, 1H), 5.53 (s, 2H), 2.89 (t, $J = 6.90$, 2H), 2.60 (t, $J = 7.70$, 2H), 1.87 (m, 2H); $^{13}$C NMR (125 MHz, d$^6$-DMSO) δ (ppm), 190.70, 171.76, 164.82, 163.69, 158.09, 157.84, 150.83, 146.43, 140.03, 136.57, 134.91, 131.96, 129.14, 126.56, 125.44, 122.35, 121.90, 118.33, 117.98, 115.55, 66.30, 38.09, 30.09, 22.55; HRMS (M+): Calc’d for C$_{22}$H$_{21}$N$_2$O$_6$: 409.1400. Found: 409.1394.

Naphthalen-1-yl 5-acetyl-2-(benzyl)benzoate, 165c. White ppt., 39%, Mp: 134-136°C; IR (KBr): 3275, 3100-2800, 1745, 1670, 1598, 1581, 1496, 1467, 1448, 1382, 1276, 1238, 1199, 1163, 1056, 1028, 1007, 873, 821, 760, 692, 680 cm$^{-1}$; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 8.75, (d, $J = 2.36$, 1H), 8.22 (dd, $J = 8.79$, $J = 2.36$, 1H), 7.97 (d, $J = 8.66$, 1H), 7.91 (d, $J = 8.73$, 1H), 7.81 (d $J = 8.53$, 1H), 7.54-7.49 (m, 4H), 7.41-7.34 (m, 5H), 7.23 (d, $J = 8.93$, 1H), 5.36 (s, 2H), 2.65 (s, 3H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm), 196.03, 164.30, 162.08, 146.86, 135.52, 134.72, 134.39, 134.15, 133.30, 130.03, 128.74, 128.59, 128.24, 127.99, 127.22, 126.48, 126.23, 125.46, 121.57, 119.77, 118.28, 113.37, 70.97, 26.48; HRMS (M + H): Calc’d for C$_{26}$H$_{20}$O$_4$Na: 419.1259. Found: 419.1268.
Naphthalen-1-yl 2-(benzyloxy)-5-(2-bromoacetyl)benzoate, 165d. Oil, ~88%; $^1$H NMR (400 MHz, CDCl$_3$) δ (ppm), 8.77, (d, $J = 2.36$, 1H), 8.25 (dd, $J = 8.79$, $J = 2.36$, 1H), 8.01 (d, $J = 8.66$, 1H), 7.93 (d, $J = 8.73$, 1H), 7.82 (d $J = 8.53$, 1H), 7.56-7.50 (m, 4H), 7.41-7.34 (m, 5H), 7.27 (d, $J = 8.93$, 1H), 5.36 (s, 2H), 4.45 (s, 2H).

Naphthalen-1-yl-2-(benzyloxy)-5-(2-(4-(tert-butoxycarbonyl-amino)-butanoyloxy)acetyl)benzoate, 165e. This compound was manufactured by an associate and was not characterized prior to further reactivity.

Naphthalen-1-yl-5-(2-(4-(tert-butoxycarbonylamino)butanoyloxy)acetyl)-2-hydroxybenzoate, 165f. Yellow, adhesive ppt, 96%; IR (KBr): 3450, 3275, 3100-2800, 1745, 1670, 1598, 1581, 1496, 1467, 1448, 1382, 1276, 1238, 1199, 1163, 1056, 1028, 1007, 873, 821, 760, 692, 680 cm$^{-1}$; $^1$H NMR (400 MHz, d$_6$-DMSO) δ (ppm), 11.54 (s, 1H), 8.62, (d, $J = 2.36$, 1H), 8.14 (dd, $J = 8.79$, $J = 2.36$, 1H), 8.02 (m, 2H), 7.93 (d, $J = 8.73$, 1H), 7.62-7.58 (m, 3H), 7.49 (d, $J = 8.10$, 1H), 7.16 (d, $J = 8.93$, 1H), 6.89 (t, $J = 5.90$, 1H), 5.36 (s, 2H), 2.98 (t, $J = 6.90$, 2H), 2.43 (t, $J = 7.70$, 2H), 1.70 (m, 2H), 1.37 (s, 9H); $^{13}$C NMR (125 MHz, CDCl$_3$) δ (ppm), 190.50, 172.24, 165.18, 164.27, 155.60, 146.19, 134.52, 134.17, 132.18, 129.71, 128.48, 127.99, 126.85, 126.36, 125.75, 124.80, 121.22, 119.75, 118.70, 115.02, 77.45, 66.07 39.47, 30.67, 28.22, 24.97; HRMS (M + Na): Calc’d for C$_{28}$H$_{29}$NO$_8$Na: 530.1791. Found: 530.1794.

4-(2-(4-hydroxy-3-((naphthalen-1-yloxy)carbonyl)phenyl)-2-oxoethoxy)-4-oxobutan-1-aminium 2,2,2-trifluoroacetate, 165. White ppt., 91%, Mp: 63-65°C; IR (KBr): 3450, 3275, 3100-2800, 1745, 1670, 1598, 1581, 1496, 1467, 1448, 1382,
1276, 1238, 1199, 1163, 1056, 1028, 1007, 873, 821, 760, 692, 680 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 11.55, (s, 1H), 8.65, (d, \(J = 2.36\), 1H), 8.17 (d, \(J = 7.60\), 2H), 8.05 (d, \(J = 8.20\), 1H), 7.95 (s, 3H), 7.84 (d, \(J = 8.55\), 1H), 7.63 (m, 3H), 7.51 (d, \(J = 8.10\), 1H), 7.27 (d, \(J = 8.93\), 1H), 6.89 (t, \(J = 5.90\), 1H), 5.56 (s, 2H), 2.90 (t, \(J = 6.90\), 2H), 2.60 (t, \(J = 7.70\), 2H), 1.88 (m, 2H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 190.64, 171.75, 165.04, 163.52, 158.02, 157.54, 146.12, 134.63, 132.21, 129.73, 128.65, 127.67, 126.88, 125.76, 122.10, 121.15, 119.74, 118.59, 118.11, 116.03, 115.17, 66.32, 38.09, 30.11, 22.58; HRMS (M\(^+\)): Calc’d for C\(_{23}\)H\(_{22}\)NO\(_6\)Na: 408.1447. Found: 408.1447.

\((R)-(R)-1-(4-(benzyloxy)phenyl)-1-oxopropan-2-yl)-2-acetoxy-2-phenylacetate + \((R)-(S)-1-(4-(benzyloxy)phenyl)-1-oxopropan-2-yl)-2-acetoxy-2-phenylacetate\)

166a, 167a. A parallel process as described for the synthesis of 122e was utilized. Viscous oil, 94%; IR (KBr): 3275, 3100-2800, 1745, 1697, 1658, 1602, 1523, 1413, 1348, 1271, 1195, 1165, 1051, 1026, 1007, 860, 823, 761 cm\(^{-1}\); \(^1\)H NMR (400 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 7.96 (d, \(J = 8.17\), 2H), 7.88 (d, \(J = 8.95\), 2H), 7.48-7.35 (m, 20H), 7.11 (d, \(J = 8.17\), 2H), 7.07 (d, \(J = 8.20\), 2H), 6.07, (d, \(J = 3.00\), 2H), 6.04 (d, \(J = 2.90\), 2H), 5.22 (s, 2H), 5.21, (s, 2H), 2.13 (s, 3H), 2.10 (s, 3H), 1.39 (d, \(J = 7.00\), 3H), 1.33 (d, \(J = 6.75\), 3H); \(^{13}\)C NMR (125 MHz, d\(^6\)-DMSO) \(\delta\) (ppm), 193.85, 193.83, 170.33, 169.79, 169.70, 167.77, 167.74, 162.69, 162.58, 161.99, 136.49, 136.34, 133.59, 133.11, 130.91, 130.81, 130.70, 130.05, 129.67, 129.26, 129.19, 128.71, 128.58, 128.50, 128.37, 128.11, 128.04, 127.97, 127.80, 127.78, 127.58, 127.43, 127.41, 126.43, 126.38, 115.08, 114.92, 114.84, 114.61, 73.82, 73.68, 72.43,
72.16, 69.54, 69.52, 20.74, 20.32, 16.95, 16.89; HRMS (M + Na): Calc’d for C_{26}H_{24}O_{6}Na: 455.1471. Found: 455.1474. (M + H): Calc’d for C_{26}H_{25}O_{6}: 433.1651. Found: 433.1670.

(R)-(R)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl)-2-acetoxy-2-phenylacetate + (R)-(S)-1-(4-hydroxyphenyl)-1-oxopropan-2-yl)-2-acetoxy-2-phenylacetate 166, 167. A parallel process as described for the synthesis of 130e’ was utilized. Viscous oil, 99%; IR (KBr): 3455, 3100-2800, 1745, 1697, 1658, 1602, 1413, 1348, 1271, 1195, 1165, 1051, 1026, 1007, 860, 823, 761 cm^{-1}; ¹H NMR (400 MHz, d{⁶-DMSO}) δ (ppm), 10.53 (s, 2H), 7.84 (d, J = 8.17, 2H), 7.80 (d, J = 8.95, 2H), 7.53-7.42 (m, 10H), 6.85 (d, J = 8.17, 2H), 6.82 (d, J = 8.20, 2H), 6.07, (d, J = 3.00, 2H), 6.04 (d, J = 2.90, 2H), 2.12 (s, 3H), 2.10 (s, 3H), 1.38 (d, J = 7.00, 3H), 1.32 (d, J = 6.75, 3H); ¹³C NMR (125 MHz, d{⁶-DMSO}) δ (ppm), 193.49, 193.46, 169.78, 169.69, 167.76, 167.73, 162.68, 162.57, 133.64, 133.57, 131.07, 130.96, 130.28, 129.26, 129.19, 129.07, 128.88, 128.70, 128.62, 128.57, 128.13, 127.99, 127.83, 127.74, 124.88, 124.83, 115.62, 115.47, 114.91, 114.43, 74.83, 74.18, 72.34, 72.03, 20.38, 20.32, 17.08, 17.00; HRMS (M + Na): Calc’d for C_{19}H_{18}O_{6}Na: 365.1001. Found: 365.0996.

6-((R)-4-((3R,5R,8R,9S,10S,12S,13R,14S,17R)-3,12-dihydroxy-1,13-dimethyl-hexadecahydro-1H-cyclopenta[a]phenanthren-17-yl)pentan-amido)hexan-1-aminium, 168c. A solution of deoxycholic acid 168a, (DOC, 503 mg, 1.28 mmol), N-hydroxybenzotriazole (HOBT, 173 mg, 1.28 mmol), and 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDCI, 368 mg, 1.92 mmol) in 10 ml of
DMF was stirred for 30 min to ensure complete dissolution. While stirring, tert-butyl 6-aminohexylcarbamate (277 mg, 1.28 mmol) in 5 ml DMF was slowly added over 5 minutes. The reaction proceeded for 22 h under ambient conditions. After this time, the solution was diluted with ethyl acetate and 100 ml of water was added. An extraction of the organic layer then was performed, followed by addition of anhydrous magnesium sulfate. After concentration, 15 ml of 4 M HCl in 1,4-dioxane was added. Stirring of this solution, under ambient conditions, occurred for 1 h. Excess solution was removed under reduced pressure for 2 days to afford 168c as a white, adhesive precipitate, 456 mg (75%). IR (KBr): 3584, 3226, 2950, 2846, 1657, 1569, 1448, 1419, 1390, 1340, 1157, 1100, 1062, 1016; $^1$H NMR (400 MHz, MeOD) δ 7.98 (s, 1H), 3.61 (m, 2H), 3.19 (t, 2H), 2.94 (t, 2H), 2.31 (m, 4H), 1.61-1.33 (m, 31H), 1.23 (s, 3H), 0.99 (s, 3H), 0.88 (s, 3H); $^{13}$C NMR (125 MHz, MeOD), δ 177.97, 74.17, 72.68, 48.83, 48.15, 47.72, 43.75, 40.89, 40.80, 37.61, 37.44, 37.34, 37.13, 36.57, 35.46, 34.98, 33.66, 32.08, 31.22, 30.96, 30.09, 30.01, 28.87, 28.63, 27.49, 27.15, 25.02, 23.86, 17.74, 13.33; HRMS (M+): Calc'd for C$_{30}$H$_{55}$N$_2$O$_3$: 491.4213. Found: 491.4209.

(R)-4-((3R,5R,8R,9S,10S,12S,13R,14S,17R)-3,12-dihydroxy-10,13-dimethyl-hexadecahydro-1H-cyclopenta[a]phenanthren-17-yl)-N-(6-(3-(3',6'-dihydroxy-3-oxo-3H-spiro[isobenzofuran-1,9'-xanthene]-5-yl)thioureido)hexyl)pentanamide, 168. A solution of 168c (238 mg, 0.484 mmol) and fluorescein-4-isothiocyanate (FITC, 226 mg, 0.581 mmol) in 20 ml of MeOH was stirred for 30 min. While stirring at 0°C, triethylamine (TEA, 67 mL (0.484 mmol) was added over a period of
5 minutes. The reaction proceeded under ambient conditions for 16 h. Inspissation under reduced pressure followed to yield a dark red/brown precipitate. Flash column chromatography (100% Ethyl acetate, then 1:1 Ethyl acetate/Acetone, then 1:1 Acetone/Methanol furnished 227 mg of 168 (53%) as a yellowish, waxy precipitate. IR (KBr): 3580, 3278, 2933, 2856, 1757, 1712, 1614, 1587, 1504, 1452, 1365, 1334, 1220, 1180, 1118, 1081, 871, 850, 786, 711, 673; $^1$H NMR (400 MHz, MeOD) δ 8.22 (s, 1H), 7.76 (m, 2H), 7.16 (d, 1H), 6.68 (m, 3H), 6.55 (m, 3H), 3.61 (m, 2H), 3.19 (t, 2H), 2.94 (t, 2H), 2.31 (m, 4H), 1.61-1.33 (m, 31H), 1.23 (s, 3H), 0.99 (s, 3H), 0.88 (s, 3H); $^{13}$C NMR (125 MHz, MeOD), δ 182.84, 176.96, 171.68, 154.70, 142.76, 131.75, 130.66, 126.28, 126.11, 126.02, 120.22, 118.83, 114.35, 112.29, 112.04, 74.17, 72.68, 48.83, 48.15, 47.72, 43.75, 40.89, 40.80, 37.58, 37.34, 36.96, 36.58, 35.45, 34.96, 34.38, 33.60, 31.22, 30.49, 30.05, 30.00, 29.69, 28.88, 28.55, 27.75, 27.62, 25.03, 23.91, 17.83, 13.43, 9.38. HRMS (M + H): Calc'd for C$_{51}$H$_{65}$N$_3$O$_8$S: 880.4571. Found: 880.4578.
References

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Ge, M.; Cline, E.; Yang, L. Tetrahedron Letters, 2006, 47, 5797-5799.


Dr. Gerry Lushington, University of Kansas

IUPAC definition, 1994

Prof. J. Wirz; Dr. D. Heger; Dr. P. Sebej.


Provided in the Chem Draw® 3D software package


Cope, Elizabeth. Dissertation 2008


Dr. Chicheng Ma, unpublished results

Professor Emeritus, Department of Chemistry, UCLA.


Stensrud, K.; Givens, R.S.; Noh, J.; Kandler, K. Manuscript in Progress.


Prof’s Wendy and Bill Picking, Molecular Biosciences, University of Kansas.


Chan Ho Park, Dissertation 1996.


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