Metastable-State Photoacids: Synthesis, Properties, and Applications

Parth Patel
University of Central Florida

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ABSTRACT

Reversible photochromic compounds have the ability to reversibly change its color when it absorbs photons of a particular wavelength. This process of color change is a consequence of structural changes within the compound, such as cis-trans photo-isomerization. Some examples of photochromic compounds are spiropyrans, spirooxazines, diarylethenes and azobenzenes. These compounds have been extensively studied for decades, and are used in various applications such as biomedicine, chemical sensors and harvesting solar energy. However, majority of photochromic compounds are initially activated by ultraviolet (UV) light. The use of UV light is harmful for biological applications and photo-degrade the compound over repeated use. To overcome these limitations, a new class of reversible photochromic compound was introduced, called metastable-state photoacid (mPAH). In brief, mPAH is a photochromic compound which can photo-dissociate its protons under visible light and can thermally re-capture the released protons efficiently in the dark. Based on this unique property, in this research, we (1) synthesized different mPAH, and (2) studied and characterized their physicochemical (acidity, kinetics, and optical) properties. Additionally, we (3) applied different visible light activated mPAHs towards light controllable polymeric-based ion-selective optodes for detection of calcium ions and sodium ions, and modulate fluorescence with pH. The research presented herein opens new avenues towards the synthesis of mPAH derivatives and could be applied to any proton-transfer process related applications which requires wireless controllability with high sensitivity.
To my loving family and lovely wife.

Thanks for providing endless support throughout this “carefree” journey.
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CHAPTER 1 GENERAL INTRODUCTION

1.1 Motivation

Media 1-1 Visual illustration of a colored solution in response to light. See supplemental file to view Media 1-1.

As shown in Media 1-1, a solution contains an organic dye initially exhibiting a yellow color. When this solution was irradiated with blue LED light and just after switching off the light source, the yellow solution turned almost clear. Interestingly, the near clear solution slowly regains the original yellow color after some time. Also, this color change can be switched back forth and it amazed me. Likewise, a bunch of questions arose. What dye was used and why the color changed? How can we apply such a dye to advance today’s technology? Answering these simple
fundamental questions is what inspired me to pursue research about this interesting dye called metastable-state photoacid.

1.2 Reversible Organic Photochromic Compounds

Reversible organic photochromic compounds have a unique property. These compounds can reversibly undergo molecular structural or electronic configuration changes by absorbing photons of a particular wavelength of light.\textsuperscript{1-3} Likewise, structural transformation in response to the non-invasive stimuli (light), these compounds are able to remotely modulate physicochemical properties; such as color, fluorescence, dipole moment, electrical property and/or acidity.\textsuperscript{4-13} Also, the use of light provides spatial and temporal control over the processes involved.\textsuperscript{14,15} To name a few, spiropyrans, spirooxazines, azobenzenes and diarylenethenes are some of the most extensively studied reversible organic photochromic compounds (Figure 1-1).

Figure 1-1 Examples of light responsive organic photochromic compound (A) spiropyran, (B) spirooxazine, (C) diarylethene, and (D) azobenzene.
These compounds have been applied towards designing light-responsive materials for various applications; for instance, chemical sensors, energy conversion, actuators, optoelectronic devices or drug delivery applications.\textsuperscript{16-24} However, most organic-based photochromic compounds use high-energy ultraviolet light (UV) for promoting changes in physicochemical property.\textsuperscript{25} As a result of employing highly energetic light source to activate these compounds, its use in biological applications could prove detrimental.\textsuperscript{26,27} Additionally, these organic photochromic compounds exhibit fatigueness in response of repeated activation by UV light.\textsuperscript{25}

1.3 \textbf{Reversible Organic Photo-acidochromic Compounds}

The physicochemical property that is most intriguing is the reversible modulation of acidity or proton-transfer by light. This is because proton-transfer dependent chemical reaction is one of the most common reaction found in nature.\textsuperscript{28} It is responsible for the production of adenosine triphosphate and controlling the photosynthetic pathway in plants.\textsuperscript{29-31} Moreover, proton-transfer plays a vital role towards regulating many pH-dependent enzymes.\textsuperscript{32-36} Therefore, controlling this important process by light could prove extremely beneficial for any proton-dependent applications. Since light can be applied remotely and non-invasively, is energy efficient, and provides spatiotemporal control.\textsuperscript{37} As so, extensive research has been undertaken to artificially create reversible proton-gradient by light. This can be achieved by using pH and light-sensitive organic compounds, such as photobases (example: sipropyrans) and photoacids (example: excited-state photoacids and metastable-state photoacids).\textsuperscript{13,38-45} A distinction must be made between these pH and light-sensitive organic compounds.
When a spiropyran compound (Sp) absorb UV light, it converts to its more basic merocyanine form (Mc); hence, a photobase (Figure 1-2).\textsuperscript{13,38} Still under UV light, this merocyanine form can capture protons thermodynamically from an external proton source which is governed by the acidity constants and diffusion of protons. Additionally, protonated merocyanine (McH\textsuperscript{+}) is visible light sensitive.\textsuperscript{13,38} Therefore, once UV light ceases and the visible light is exposed, McH\textsuperscript{+} can than photo-release the protons back to the original source. Nonetheless, the use of UV light as mentioned previously could hinder its potential use.\textsuperscript{26,27}

![Diagram of spiropyran reaction](image)

Figure 1-2 Reversible photo-acidochromic behavior of spiropyran.

On the other hand, photoacids in general behave like a strong acid under light activation.\textsuperscript{28} They can photo-release their protons and can be re-captured when the light is switch off. Of the two types of reversible photoacids, excited-state photoacids (Figure 1-3) photo-release their protons from an electronically high-acidity excited state upon light activation.\textsuperscript{39-42} When the light is switched off, the high-acidity excited state undergoes extremely fast relaxation (proton re-capture
process) to the original low-acidity ground state; thereby, hindering large change in proton gradient.\(^{28}\) Hence, this type of photoacid may not be an ideal candidate for applications that rely on slow diffusion mediated proton-transfer reactions.

![Figure 1-3 Examples of excited-state photoacid (A) Pyranine, and (B) 1-Naphthol.](image)

1.4 Metastable-state Photoacids

Contrary to excited-state photoacids,\(^{39-42}\) a new class of visible light activated photoacid, called metastable-state photoacid, was first introduced by Liao and co-workers in 2011.\(^{43}\) This is the type of compound that was used to illustrate color changing behavior (Media 1-1). As shown in Figure 1-4, there are two types of metastable-state photoacids: merocyanine-type (Mer-mPAH) and tricyanofuran-type (TCF-mPAH).\(^{43-45}\) Both these metastable-state photoacid were designed by connecting an electron-accepting moiety to a weakly acidic nucleophilic moiety by a trans-double bond.\(^{28}\) In brief, these compounds are able to change color and photo-release their protons under
visible light activation via trans-cis photoisomerization followed by nucleophilic reaction between the two moieties.\textsuperscript{28} When visible light is switched off, the color is regained and the photo-dissociated metastable-state photoacid displays thermodynamically driven long relaxation (proton re-capture process).\textsuperscript{28}

![Diagram of Merocyanine-type and Tricyanofuran-type metastable-state photoacids]

Figure 1-4 Reversible photo-acidochromic behavior of metastable-state photoacids (A) Merocyanine-type (Mer-mPAH), and (B) Tricyanofuran-type (TCF-mPAH).

This relaxation process is greatly slower than compared to the excited-state photoacids. Due to the improved stability of the photo-dissociated metastable-state photoacid, the photo-released protons can theoretically be used for any proton-transfer based applications. Furthermore, metastable-state photoacids can cause a large pH change with high efficiency and good reversibility. Due to these
unique factors, metastable-state photoacids have found numerous applications in recent years. To name a few, they have been applied towards controlling supramolecular structures, killing bacteria, used as an acid-catalyst, activate ion-channels, release fragrant molecules, ATP production, and manage waste.

Hence, presented in the following chapters, different visible light activated metastable-state photoacids (merocyanine- and tricyanofuran-type) were synthesized to meet the applications’ requirement. These modified metastable-state photoacids were characterized and evaluated in terms of its optical properties, acidity, reversibility, and stability. In accordance to the evaluations, they were then applied towards cation-selective optode sensors, and fluorescence and pH modulations.

1.5 References


45 C. Yang, T. Khalil, Y. Liao, RSC Adv., 2016, 6, 85420.


CHAPTER 2 VISIBLE-LIGHT ACTIVATED METASTABLE-STATE PHOTOACID POLYMER FOR CALCIUM DETECTION UNDER NON-EQUILIRIUM CONDITIONS


2.1 Abstract

Presented here is a sensing membrane consisting of a modified merocyanine photoacid polymer and a calcium ionophore in plasticized poly(vinyl chloride). This membrane is shown to actively exchange protons with calcium ions when switched ON after illumination at 470 nm, and the exchange can be followed by UV−vis spectroscopy. The sensing membrane shows no response in the ON state when calcium ions are absent. The limit of detection of the sensor is $5.0 \times 10^{-4}$ M with an upper detection limit of 1.0 M. Thus, we demonstrate for the first time the use of a visible light activated, lipophilic photoacid polymer in an ion-sensing membrane for calcium ions, which highly discriminates potassium, sodium, and magnesium ions.

2.2 Introduction

A sensor that uses the change in absorption or emission of a dye provides a means for detecting analytes using different methods.\textsuperscript{1-4} When the dye is a photoactive molecule, it promises control
over the sensor’s characteristics using light as a noninvasive stimulus. Readily available monochromatic light sources, advances in photochemistry, and synthetic methods open the door for the use of such photoactive molecules in sensors.

Umezawa’s group pioneered the use of photoswitchable molecules in potentiometric sensing\(^5\) while other researchers have used different types of photoactive species in optical ion sensors.\(^6\)\(^-\)\(^9\)

A typical cation selective sensor consists of a lipophilic polymer membrane such as plasticized poly(vinyl chloride) (PVC) containing an ion-exchanger and two ionophores: one designed to bind to a cation of interest and the other is a pH indicator (chromoionophore) that interacts with a reference ion (H\(^+\)) and changes its optical properties. This change is related to the activity of the ion of interest. These sensors typically function under conditions of mass transfer equilibria and are passive in nature. However, they can be made dynamic by the use of photoactive compounds. For instance, Bakker and co-workers replaced the chromoionophore in a traditional optode with a spiropyran that photoisomerizes to a zwitterionic merocyanine form with increased basicity.\(^10\)\(^,\)\(^11\)

In order to be used as a sensor for cations, they had to introduce an external source of protons to protonate the photoisomerized merocyanine.\(^12\) In contrast, the use of molecules that photorelease protons could eliminate the need for an external source of H\(^+\).

There are two main types of molecules that release protons upon illumination: photoacid generators (PAGs) and photoacids (PAHs). PAGs undergo proton photodissociation irreversibly,\(^13\)\(^,\)\(^14\) while PAHs are molecules that undergo proton photodissociation and thermal reassociation, hence promising reversibility.\(^15\)\(^-\)\(^17\) Shvarev et al. recently used a PAG to perturb the
equilibrium of a conventional ion-sensing optode; however, to the best of our knowledge, no one has used a PAH in such sensing systems.\textsuperscript{18} This ability to switch proton fluxes back and forth using light makes PAHs better candidates for use in reusable optical ion sensors and, at the same time, obviates the need for conditioning prior to use.

Recently, a new PAH was reported that undergoes proton dissociation under visible light and thermal reassociation over multiple cycles with no observable fatigue and has a long-lived high-acidity state.\textsuperscript{19,20} In addition, this PAH was shown to change the volume of a hydrogel upon photoillumination, used to inhibit bacterial growth,\textsuperscript{21} and change the conductivity of a polymer.\textsuperscript{22} The long-lived proton dissociated state allows for this PAH to be used in diffusion controlled processes. In contrast, most known PAHs have short-lived high acidity states,\textsuperscript{15} hence limiting the use of these compounds in diffusion controlled states. We envision the introduction of this photoacid in optical ion sensors that can substitute the chromoionophore as well as serve to switch the sensor ON only when desired. Thus, this work presents our results toward a “first of a kind” cation-selective sensor based on a polymerized PAH reported before.\textsuperscript{23}

2.3 Results and Discussion

The PAH mentioned before\textsuperscript{19} is useful in applications which involve hydrophilic matrices, being incompatible with sensing membranes based on lipophilic components such as in optodes. A previous study described the synthesis of a copolymer between a PAH unit and methyl acrylate.\textsuperscript{23} Acrylates are viable monomers for the synthesis of polymers used in optical sensing membranes.\textsuperscript{24,25} Thus, the covalent attachment of the PAH unit to a lipophilic acrylic polymer
backbone might allow the introduction of this photoacid into a lipophilic sensing membrane. These membranes are typically composed of an inert polymer such as PVC dissolved in a plasticizer such as dioctylsebacate (DOS) or o-nitrophenyl octyl ether (o-NPOE). The proper choice of a plasticizer is very important as it fixes the dielectric constant, improves the selectivity, ensures a fast response, and elongates the lifetime of the sensor.26 With a few modifications, polymer 1 (Scheme 2-1) was synthesized with ~10% PAH component (in blue) and ~90% methyl acrylate (in black) which was optimal for the lipophilic requirements of the PVC/DOS or o-NPOE membrane matrix described in this work. Higher percentages of photoacid polymer were synthesized but were not appropriate for PVC/DOS or PVC/o-NPOE matrix due to solubility problems.

Scheme 2-1 shows the likely mechanism of the sensor. The sensor can be activated (switched ON) upon irradiation at 470 nm for 5 min, where polymer 1 photocyclizes to polymer 2 with the concomitant release of protons. Incorporation of a cation selective ionophore (L) results in an exchange of protons with the cation of choice. The photogenerated protons should actively undergo two processes once the light is switched OFF. One is the recombination of the protons with polymer 2 to form polymer 1, and the second, the exchange of protons with the ion analyte, which should occur to maintain electroneutrality in the membrane. Considering the long-lived proton dissociated state, the exchange of protons with the analyte should be much faster than the protons recombining with polymer 2. The dynamics of this process can be monitored via UV–vis spectroscopy providing a means to detect cations optically.
The sensing membrane used here was prepared with 9.74% (84.56 mmol/kg of PAH unit) of polymer 1, 8.48% (105.70 mmol/kg) of calcium ionophore IV, 27.26% of PVC, and 54.52% of o-NPOE. All components were dissolved in tetrahydrofuran, and the cocktail was spin coated onto glass slides to form the membranes. For measurements, 0.5 M tris(hydroxymethyl)aminomethane buffer at pH 7.40 (adjusted using 6.0 M HCl) was used. Figure 2-1 depicts the spectra of the ion-sensing membrane before irradiation (a), after 5 min of illumination at 470 nm (b), and finally after 35 min in the dark (c), while the sensing membrane was immersed in buffer solution.
Figure 2-1 Absorbance spectra of ion-sensing membrane (a) before illumination, (b) after 5 min at 470 nm, and (c) after 35 min in the dark, in 0.5 M tris(hydroxymethyl)aminomethane buffer at pH 7.40.

The sensing membrane showed a change in absorbance at different concentrations of calcium ions. An increase in the calcium ion concentration in the sample solution favors the ion exchange between calcium ions and the photoreleased protons. Hence, more protons are released to the sample solution. Consequently, the formation of polymer 1 decreased, corresponding to a loss in absorbance (Figure 2-2A). A plot of the change in absorbance vs log of calcium ion activity is show in the Figure 2-2A inset. A blank experiment was performed where the sensing membrane was immersed in a buffer solution that did not contain calcium ions. The sensing membrane was illuminated for 5 min followed by 35 min in the dark. Ten subsequent runs were performed
showing barely any change in absorbance (Figure 2-2B). These results indicate that the sensor is stable over extended periods of illumination.

Figure 2-2 Absorbance spectra of ion-sensing membrane activated (A) after 35 min in the dark at different concentrations of calcium ions (a) 0.0, (b) $1.0 \times 10^{-4}$, (c) $1.0 \times 10^{-3}$, (d) $5.0 \times 10^{-3}$, (e) $1.0 \times 10^{-2}$, (f) $5.0 \times 10^{-2}$, (g) $1.0 \times 10^{-1}$, (h) $5.0 \times 10^{-1}$, (i) 1.0, and (j) 1.25 M. Inset in (A) shows the corresponding calibration curve for calcium ions. Conditions for the experiment were similar to Figure 2-1. (B) Blank experiment where ten subsequent runs were performed, after 5 min of illumination followed by 35 min in the dark when no calcium ions were present.
The reproducibility of the ion-sensing membrane response was evaluated by performing calibration curves with at least three different membranes, as can be seen in Figure 2-3. Concentrations in the range between $1.0 \times 10^{-5}$ and $1.25$ M $\text{Ca}^{2+}$ were tested, and a detection limit of $5.0 \times 10^{-4}$ M calcium ions was obtained. The response range of the photoactivated ion-sensing membrane presented here is similar to that reported in the literature based on the same ionophore$^{27,28}$ and is promising for applications requiring remote control. The change in absorbance for this sensor with an increase in $\text{Ca}^{2+}$ concentration was evaluated after a 5 min activation at different times of the ion-exchange process (5, 10, 20, and 35 min), where the greatest
change was obtained at 35 min (see Appendix A, Figure A-1). For further experiments, 35 min was used.

Figure 2-4 Selectivity for the ion-sensing membrane measured 35 min after activation. Conditions for the experiment are similar to Figure 2-1.

Two different matrices were tested for sensing: PVC/DOS and PVC/o-NPOE. While working with PVC/DOS, the sensing membrane did not discriminate against potassium ions. This was likely due to ion-pair formation of potassium ions with the sulfonate moiety present in polymer 2. The replacement of DOS by o-NPOE provided a polar medium to the membrane, due to its higher dielectric constant disfavoring the formation of such ion-pairs between soft ions such as potassium and the sulfonate moiety. To further explore the selectivity of the ion-sensing membrane, different
ions were tested. Thus, Figure 2-4 shows the response of the calcium ion sensor after it was exposed to different concentrations of sodium, potassium, and magnesium ions in 0.5 M buffer solution for 35 min, which indicates a good selectivity with respect to the change in absorbance. When compared to traditional optodes using the same ionophore, the discrimination of interfering ions was found to be similar.27

2.4 Conclusion

For the first time, a visible light activated, lipophilic PAH polymer was incorporated into an ion-sensing membrane, which can be regenerated. A new dynamic sensing technique was also demonstrated for calcium ions based on phototriggered proton exchange. The importance of this sensor lies in the fact that the sensing membrane can be triggered only when desired by using visible light and needs no conditioning or ion-exchanger.

Work is in progress to improve the limits of detection and extend this sensing mechanism to other cations. It is anticipated that the PAH polymer will replace the pH sensitive dye used in traditional optodes. Such sensors will be especially useful for controlled sensing where the sensor can be turned ON/OFF merely by flipping a switch.
2.5 Acknowledgments

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2.6 References


3.1 Abstract

Herein, we present different functionalized meta-stable photoacids in ion-sensors to tune the equilibrium response time. As a proof of concept, two new meta-stable photoacids were synthesized, one contains an ether group and the other has an ester functional group. These functionalized meta-stable photoacids were elucidated in solution of ethanol and in ion-sensing films via pH and kinetic studies. In comparison from our previous work, the response time was reduced from hours to minutes by utilizing an ether functional group in the meta-stable photoacid.

3.2 Introduction

The last few decades have seen considerable progress in ion sensing electrodes and optodes. Specifically in terms of improving the selectivity and detection limits at equilibrium and non-equilibrium conditions, as discussed by Bobacka et al. Since ion-selective optodes provide “wireless” detection of ions, researchers have shown increased interest in developing these optodes.
sensors.\textsuperscript{5–9} Optodes are made of a plasticized polymer (that serves as a matrix support) containing an ion-exchanger (that maintains electroneutrality), an ionophore (that is selective to the ion of interest), and chromoionophore or pH indicator (that is selective to a reference ion (H\textsuperscript{+})).\textsuperscript{2} The detection of the analyte depends on mass transfer equilibria, which is dictated by the activity of the analyte and the reference ion.\textsuperscript{10–12} The changes in the optical property of the indicator relates to the activity of the ion of interest, and can be followed via absorbance and/or fluorescence spectroscopy.\textsuperscript{13–15}

Most optodes are passive in nature which limits control over the ion-exchange process.\textsuperscript{2,13} Whereas, an active optode would be ideal for ion detection applications done locally. One possible way to make an optode active is by using photoactive compounds. While different research groups have used photoacid generators\textsuperscript{14} and spiropyrans (Sp)\textsuperscript{16,17} to achieve this, we have used meta-stable photoacids (mPAHs).\textsuperscript{18} A distinction must be made between photoacids (PAHs) and mPAHs. While both are reversible, PAHs release protons from an electronically excited state\textsuperscript{19}; hence the proton recapture is too fast (in the order of nanoseconds to seconds) to be useful in diffusion mediated sensing. Alternatively, mPAHs photo-release protons from a meta-stable state.\textsuperscript{20,21} This state is sufficiently long-lived (in the order of minutes to hours) for the ion-exchange process to take place.

In addition, mPAHs-based ion sensors can be remotely activated by visible light, do not suffer photo-fatigue\textsuperscript{20,21}, and enable reversibility of the sensor\textsuperscript{18}. However, in order for those advantages to be realized, further studies are required. One of the major issues we encountered previously was
the fact that the proton re-association took hours to reach the equilibrium, consequently the
detection was performed under non-equilibrium conditions.\textsuperscript{18}

In order to achieve detection under equilibrium conditions, the photo-dissociated state of the
mPAH should have a shorter lifetime (in the order of minutes). The stability of the photo-
dissociated form of the mPAH is likely dictated by electronic effects. As the mPAH we used was
covalently attached to the polymer backbone by an ester linkage,\textsuperscript{18} we hypothesized that the
electron withdrawing nature of the ester group at the meta position from the pyran oxygen could
be extending the lifetime of the photo-dissociated state. We thus decided to study the effect of
electron demand of the linker with respect to the mPAH to determine if the lifetime of the photo-
dissociated state could be shortened.

Here, we present the synthesis of two mPAHs containing different functional groups, one has an
ether group (PA-4Octadec) and the other has an ester group (PA-4Stearoyl). These functionalized
mPAHs are characterized in solution and in the ion-sensing film. In addition, kinetic studies are
presented to relate the effect of the functional group with the proton recapture process.

3.3 Experimental

3.3.1 Reagents

2,4-Dihydroxybenzaldehyde, stearoyl chloride, high molecular weight poly(vinyl chloride) (PVC),
terakis(4-chlorophenyl)borate tetradecylammonium salt (ETH 500), N,N-dicyclohexyl-N,N-
dioctadecyl-3-oxapentanediamide (calcium ionophore IV), triethylamine (Et$_3$N), 1,3-dichloro-1,1,3,3-tetramethyldisiloxane, and 2-nitrophenyl octyl ether (o-NPOE) were obtained from Sigma Aldrich, USA. Potassium iodide (KI), tris(hydroxymethyl) aminomethane (tris), ethanol (EtOH), acetonitrile (MeCN), hydrogen peroxide (H$_2$O$_2$), sulfuric acid (H$_2$SO$_4$), and hydrochloric acid (HCl) were obtained from Fischer Scientific, USA. Dichloromethane, ethyl acetate, hexane, and acetone were obtained from Macron, USA. Calcium chloride, 1-bromooctadecane, tetrahydrofuran (THF), and cyclohexanone were obtained from Acros, Belgium. Potassium carbonate anhydrous (K$_2$CO$_3$), and sodium sulfate anhydrous (Na$_2$SO$_4$) were obtained from Amresco, USA. Chloroform-D (CDCl$_3$) and dimethyl sulfoxide-D$_6$ (DMSO) were obtained from Cambridge Isotope Laboratories, Inc., USA.

3.3.2 Instruments

Absorption spectra were obtained using a UV-Visible spectrophotometer (Cary 50) from Varian, Australia. The light source for the photoreaction was a 470 nm LED array with 120 LEDs from Elixa, USA. NMR spectra were obtained with a NMR spectrometer (Avance III 400) of 400 MHz from Bruker, USA. The pH measurements were obtained with a pH meter (Orion Star A211) from Thermo Scientific, USA. The thickness of the ion-sensing films was measured using a profilometer (Tencor AlphaStep 500) from KLA Tencor, USA. DI water used to prepare solutions was purified by a water purification system with resistance of 18 MΩcm (PURELAB Ultra) from Siemens, USA. All experiments were carried out in the dark.

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3.3.3 Synthesis

Scheme 3-1 shows the steps for the synthesis of the two new mPAHs.

3.3.3.1 Synthesis of 2-Hydroxy-4-(octadecyloxy)benzaldehyde (1)

1.0 g (7.2 mmol) of 2,4-dihydroxybenzaldehyde, 1.0 g (7.2 mmol) of K₂CO₃, 0.072 g (0.42 mmol) of KI, and 3.0 mL of dry acetone were added to a round bottom flask and refluxed under N₂ for 30 min until the solution became cloudy and pink. Then, 2.4 g (7.2 mmol) of 1-bromooctadecane and 3.0 mL of dry acetone were added. The solution was refluxed overnight, quenched with water and filtered. The product was extracted in dichloromethane, dried with Na₂SO₄, and purified by column chromatography using 1% ethyl acetate in hexane as the eluent. Yield (2.1 g, 74.8%). H¹ NMR (400 MHz, CDCl₃), δ = 11.5 (s, 1H), 9.7 (s, 1H), 7.4 (m, 1H), 6.5 (m, 1H), 6.4 (m, 1H), 4.0 (t, 2H), 1.8 (m, 2H), 1.4 (m, 2H), 1.2 (m, 30H), 0.9 (m, 3H).

3.3.3.2 Synthesis of 2,3,3-Trimethyl-1-(3-sulfonatepropyl)-3H-indolium (2)

Compound 2 was synthesized according to the literature procedure.²¹,²²

3.3.3.3 Synthesis of 4-Formyl-3-hydroxyphenyl stearate (3)

1.0 g (7.2 mmol) of 2,4-dihydroxybenzaldehyde, 1.4 mL (9.9 mmol) of dry Et₃N, 5.0 mL of dry acetone, and 25 mL of dry dichloromethane were added to a reaction flask under N₂. The reaction mixture was immersed in an ice-bath and stirred for 10 min. After that, 2.2 mL (6.6 mmol) of
stearoyl chloride was added. The mixture was slowly raised to room temperature and allowed to stir overnight, then quenched with water, washed with brine (saturated salt water) and dried with Na₂SO₄. The product was then chromatographed using 1 % ethyl acetate in hexane as the eluent. Yield (1.37 g, 47.1 %). ¹H NMR (400 MHz, CDCl₃), δ = 11.2 (s, 1H), 9.9 (s, 1H), 7.6 (m, 1H), 6.8 (m, 2H), 2.5 (t, 2H), 1.8 (m, 2H), 1.3 (m, 28H), 0.9 (m, 3H).

3.3.3.4 Synthesis of ((E)-3-(2-(2-Hydroxy-4-(octadecyloxy)styryl)-3,3-dimethyl-3H-indol-1-ium-1-yl)propane-1-sulfonate (PA-4Octadec)

Compound 1 (0.26 mmol) and 2 (0.26 mmol) were added into 2 mL EtOH. The mixture was allowed to reflux overnight under N₂. Orange solids were obtained on cooling, which were separated via filtration followed by washing the solids in cold EtOH, then 2 mL of dichloromethane followed by 10 mL of THF to provide the corresponding mPAH. Yield (0.095 g, 55.9 %). ¹H NMR (400 MHz, DMSO), δ = 11.2 (s, 1H), 8.4 (d, 1H, J = 16.1 Hz), 8.2 (d, 1H, J = 9.0 Hz), 7.9 (d, 1H, J = 7.9 Hz), 7.7 (d, 1H, J = 7.2 Hz), 7.6 (d, 1H, J = 16.2 Hz), 7.5 (m, 2H), 6.5–6.4 (m, 2H), 4.6 (t, 2H), 4.0 (t, 2H), 2.5 (t, 2H), 2.0 (m, 2H), 1.7 (s, 6H), 1.3–1.1 (m, 32H), 0.8 (m, 3H).

3.3.3.5 Synthesis of (E)-3-(2-(2-Hydroxy-4-(stearoyloxy)styryl)-3,3-dimethyl-3H-indol-1-ium-1-yl)propane-1-sulfonate (PA-4Stearoyl)

Compound 2 (0.26 mmol) and 3 (0.26 mmol) were added into 5 mL MeCN. The mixture was allowed to reflux overnight under N₂. Orange solids were obtained on cooling which were separated by filtration followed by washing the solids in copious amounts of hexane to provide the corresponding mPAH. Yield (0.050 g, 28.8 %). ¹H NMR (400 MHz, DMSO), δ = 11.4 (s, 1H), 8.5
(d, 1H, J = 16.3 Hz), 8.3 (d, 1H, J = 8.7 Hz), 8.0 (m, 1H), 7.9 (m, 2H), 7.6 (m, 2H), 6.8 (m, 2H),
4.8 (t, 2H), 2.6 (m, 4H), 2.2 (m, 2H), 1.8 (s, 6H), 1.6 (m, 2H), 1.2 (m, 28H), 0.8 (m, 3H).

Scheme 3-1 Steps for the synthesis of mPAHs (PA-4Octadec and PA-4Stearoyl).

3.3.4 Preparation of Ion-Sensing Films

3.3.4.1 Silanization of Glass Slides

The silanization procedure is similar to the previously reported literature with few modifications.23

The glass slides were previously cut to fit the cuvette, which were washed in piranha solution for
30 minutes at 90 °C (safety note: the piranha solution must be handled with caution). The glass
slides were then thoroughly rinsed with deionized (DI) water and dried in a vacuum oven for 30
minutes. The clean glass slides were then silanized in a mixture of 1,3-dichloro-1,1,3,3-tetramethyldisiloxane and dichloromethane for 3 hours at 50 rpm on an orbital shaker. The silanized glass slides were thoroughly rinsed with isopropanol and DI water, then dried again in the vacuum oven, and stored in a vacuum desiccator until use.

3.3.4.2 Ion-Sensor Containing PA-4Octadec

The sensor contains 0.49 mg of PA-4Octadec (15.0 mmol/kg), 1.80 mg of calcium ionophore (45 mmol/kg), 0.86 mg of ETH 500 (15 mmol/kg), PVC (66.0 wt%), and o-NPOE (33 wt%). Cyclohexanone (800 µL) was added to dissolve all the components, and the mixture was vortexed for 30 minutes to obtain a homogeneous cocktail solution. Subsequently, 20 µL of cocktail solution was dropped onto silanized glass slides and stored in vacuum overnight. The thickness of the film formed was 9.07±0.22 µm.

3.3.4.3 Ion-Sensor Containing PA-4Stearoyl

The sensor contains 0.50 mg of PA-4Stearoyl (15 mmol/kg), 1.80 mg of calcium ionophore (45 mmol/kg), 0.86 mg of ETH 500 (15 mmol/kg), PVC (66 wt%), and o-NPOE (33 wt%). Cyclohexanone was used to dissolve all the components, and the following procedure was the same as in Section 3.3.4.2. The thickness of the film formed was 7.10±0.52 µm.
3.3.5 Characterization of mPAHs

3.3.5.1 pH Measurements

To measure the pH for both mPAHs, PA-4Octadec (1.04 mg) and PA-4Stearoyl (0.97 mg) were dissolved in a mixture of 90% EtOH and 10% DI water, separately. Prior to irradiation with visible light (470 nm), the pH of the solution was recorded. Subsequently, the dissolved mPAHs were irradiated for 2 minutes and the pH was registered. From the data acquired, the pKa’s were calculated before irradiation for each mPAH.

3.3.5.2 Kinetics Measurements

Kinetics studies were performed by dissolving the mPAHs in EtOH (PA-4Octadec at the concentration of $21.6 \times 10^{-6}$ M, and PA-4Stearoyl at $23.2 \times 10^{-6}$ M) and irradiating for 2 minutes with visible light (470 nm). The rate of proton re-association was monitored by measuring the absorbance (of the reprotonated form) every 15 seconds for PA-4Octadec and every 15 minutes for PA-4Stearoyl, respectively. The molar absorptivities for both mPAHs in EtOH were calculated from the maximum absorbance obtained by acidifying the solution with 6.0 M HCl.

The above procedure was also used to determine the time it takes for proton re-association for each mPAH in the ion-sensing film using a 0.025 M tris buffer at pH 7.40 (adjusted by 6 M HCl).

All kinetic data were plotted as the maximum absorbance vs. time.
3.4 Results and Discussion

Figure 3-1 $^1$H NMR of PA-4Stearoyl for (A) before irradiation, and (B) after 2 minutes irradiation (470 nm).

PA-4Octadec and PA-4Stearoyl were synthesized according to Scheme 3-1. PA-4Octadec was linked to a lipophilic alkyl chain via an ether functional group, whereas PA-4Stearoyl uses an ester as the linker. Initially, the mPAHs were studied by NMR spectroscopy to understand the photo-induced structural change and the effect of functional groups upon irradiation. Thus, spectra of PA-4Octadec in deuterated DMSO was collected before irradiation and after 2 min of irradiation.
The resulting spectral properties for both before and after irradiation showed no change. However, when PA-4Stearoyl was evaluated, a drastic change was observed (Figure 3-1A). Comparing the aromatic region of the spectra before and after irradiation showed the disappearance of the peak at 11.4 ppm (loss of the proton from the –OH group). In addition, the bridging hydrogens at 8.55 and 7.85 ppm disappeared and reappeared as multiplets between 7.19 to 6.43 ppm. This strong downfield shift is also indicative of the changes in structural configuration from a trans to a cis, which is further evidenced by a reduction of coupling constants from 16.3 Hz to 10.3 Hz. Moreover, the aromatic peaks shifted upfield after irradiation indicating a loss in conjugation. The spectrum (Figure 3-1B) after irradiation corresponds well with the structure of an anionic spiropyran.20

The stability of the spiropyran could be dictated by the electron density at the pyran oxygen which influences the proton recapture rate of the mPAHs after irradiation. The NMR study indicates that the PA-4Stearoyl photo-dissociates into a stable anionic spiropyran, while PA-4Octadec may or may not be photoactive. To determine whether the PA-4Octadec photo-dissociates, a pH study was conducted before and after irradiation.

The mPAHs (PA-4Octadec and PA-4Stearoyl) were dissolved in a solution of 90 % EtOH and 10 % DI water, showed pHs of 5.95±0.25 and 5.17±0.15, respectively. The corresponding pKa’s were calculated to be 8.41±0.52 and 6.82±0.32, respectively. When the solutions were irradiated, the pH of the mPAHs dropped to 4.37±0.04 and 4.42±0.15 for PA-4Octadec and PA-4Stearoyl
respectively. This change in pH indicates that both mPAHs release their protons upon irradiation, thereby confirming photoactivity.

Figure 3-2 Kinetics of mPAHs in EtOH after 2 minutes of irradiation (470 nm) for (A) PA-4Octadecc, and (B) PA-4Stearoyl. Left Figures are the response curve over time for both mPAHs obtained from its corresponding absorption spectra (right). (Inset: Plot of fitting the kinetic data in a second-order rate equation for both mPAHs)
Furthermore, kinetic studies were performed to understand the proton re-association rate of the mPAHs after irradiation. If the proton recapture process is the rate limiting step for the respective mPAHs, i.e. protonation of $\text{Sp}^-$ ($\text{Sp}^- + \text{H}^+ \rightarrow \text{MeH}$) ($\text{MeH}$: protonated neutral merocyanine state; $\text{Sp}^-$: photo-dissociated anionic spiropyran state), by the majority of the protons in the solution produced from the high-acidity state $\text{MeH}$ (i.e. $[\text{H}^+] = [\text{Sp}^-]$). Then the kinetics should follow a 2nd order rate equation\textsuperscript{20}

$$\frac{d[\text{MeH}]}{dt} = k[\text{Sp}^-][\text{H}^+] = k[\text{Sp}^-]^2$$  \hspace{1cm} [3.1]

If no other long-lived species are involved during this process, then $[\text{Sp}^-]=[\text{MeH}]_T-[\text{MeH}]$ ($[\text{MeH}]_T$ is the total concentration of mPAH), and Equation 3.1 can be integrated to give Equation 3.2.\textsuperscript{20}

$$\frac{1}{([\text{MeH}]_T-[\text{MeH}])} = kt + \frac{1}{[\text{MeH}]_T}$$  \hspace{1cm} [3.2]

Thus, the proton recapture process was monitored by UV-Vis spectroscopy after the mPAHs were irradiated for 2 min. The kinetics for both mPAHs in EtOH is shown in Figure 3-2, where the absorbance of PA-4Octadec (Figure 3-2A) and PA-4Stearoyl (Figure 3-2B) stabilized at 5 min and 225 min, respectively. This evidence indicates that the time of proton re-association is much faster for PA-4Octadec than for PA-4Stearoyl. This could be the reason why the NMR spectra for PA-4Octadec were similar before and after irradiation. The data provided from the kinetic experiment was fitted into Equation 3.2. And, as shown in the inset of Figure 3-2, the kinetics fits well into the 2nd order rate equation. The rate constant was calculated to be $2811.50 \text{ M}^{-1}\text{s}^{-1}$ with $R^2$ of 38
0.9996 for PA-4Octadec (Figure 3-2A Inset) and molar absorptivity of 41158.2 M$^{-1}$cm$^{-1}$. On the other hand, PA-4Stearoyl exhibited 1971.70 M$^{-1}$s$^{-1}$ as the rate constant with R$^2$ of 0.9994 (Figure 3-2B Inset) and molar absorptivity of 23567.16 M$^{-1}$cm$^{-1}$. The fact that the kinetic data fits well into Equation 3.2 indicates that the protonation of Sp$^-$ is indeed involved during the rate-limiting step.

Figure 3-3 Kinetics of mPAHs in ion-sensing film after 2 minutes of irradiation (470 nm) in 0.025 M tris buffer solution at pH 7.40 for (A) PA-4Octadec (scans every 6 seconds), and (B) PA-4Stearoyl (scans every 15 minutes).

Likewise, the time for protons to re-associate for PA-4Octadec and PA-4Stearoyl in the ion-sensing film was evaluated. Figure 3-3 shows the change in absorbance arising from the protonation of Sp$^-$ over time after the sensors were irradiated. The equilibrium response times were 8 min and 150 min for PA-4Octadec and PA-4Stearoyl in the ion-sensing film, respectively.
As mentioned before, the proton re-association time was reduced from hours to minutes by the use of an ether functional group (PA-4Octadec) in place of the ester group (PA-4Stearoyl). This is likely because PA-4Stearoyl in its Sp⁻ state is more stable after irradiation. The ester group withdraws electron density from the pyran oxygen (Scheme 3-2) and consequently reduces the tendency for the proton recapture by the pyran oxygen. On the other hand, PA-4Octadec is less stable in its deprotonated form because the ether group donates electron density to the pyran oxygen, hence speeding up the proton recapture process. Thus, PA-4Octadec has a short-lived photo-dissociated state. It is therefore clear that the lifetime of the photo-dissociated state is
dependent on the nature of the substituent (–R) meta to the pyran oxygen as represented in Scheme 3-2.

3.5 Conclusion

For the first time, we tuned the proton re-association rate of mPAHs by appropriate functionalization. When compared to our previous work\textsuperscript{18}, we were able to improve the response time from hours to minutes at pH 7.40. We envision that, by using mPAHs in ion-sensors, we can control the ion-exchange process that would ultimately lead to improved detection limits.

3.6 Acknowledgements

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3.7 References


CHAPTER 4 VISIBLE-LIGHT ACTIVATED METASTABLE-STATE PHOTOACID INION-SELECTIVE OPTODE SENSORS FOR SODIUM AND CALCIUM IONS: EXPERIMENTAL AND THEORETICAL APPROACH


4.1 Abstract

A new platform of ion-selective optodes is presented here to detect cations under thermodynamic equilibrium via ratiometric analysis. This novel platform utilizes a ‘one of a kind’ visible light-induced metastable photoacid as a reference ion indicator to achieve activatable and controllable sensors. These ion-selective optodes were studied in terms of their stability, sensitivity, selectivity, and theoretical aspects.

4.2 Introduction

Ion selective optodes (ISOs) have been extensively studied to monitor different cations for biomedical and environmental applications. In recent years, there has been considerable interest to study ISOs that are activatable, reversible and controllable for their use in ion sensing applications that require localized detection of free ions without perturbing their surroundings. Attempts have been made to convert the traditional passive mode ISOs into active mode by the
use of photoactive compounds such as spiropyrans and photoacid generators.\textsuperscript{16–19} However, the use of spiropyrans in an ISO requires UV light to activate the sensor, which can result in cellular damage when applied in biomedical applications.\textsuperscript{20} In addition, UV light photodegrades the active compound in the ISO, shortening the sensor's lifetime.\textsuperscript{13–17} Moreover, photoacid generators in an ISO undergo photolysis, making the sensor irreversible.\textsuperscript{18,19} To overcome photodegradation and irreversibility for such ISOs, we propose to use a ‘one of a kind’ visible light activatable and controllable ISO based on a metastable photoacid (mPAH) to detect cations under equilibrium conditions by ratiometric analysis.

Scheme 4-1 Cation-exchange process of ISOs that utilize a (A) neutral basic indicator, and a (B) neutral acidic indicator. (In: neutral basic indicator; L: cation selective ionophore; R:\textsuperscript{−}: anionic additive; InH: neutral acidic indicator).

\begin{align*}
A) & \quad a_{M^{2+}}(aq) + xL(org) + zInH^+(org) \rightleftharpoons L_xM^{2+}(org) + zIn(org) + za_{H+}(aq) \\
B) & \quad a_{M^{2+}}(aq) + xL(org) + zInH(org) \rightleftharpoons L_xM^{2+}(org) + zIn^{-}(org) + za_{H+}(aq)
\end{align*}
A typical ISO consists of a neutral basic indicator (In) that is selective to a reference ion (H⁺), an ionophore (L) that is selective to the cation of interest, and anionic additives (R⁻) that maintain electroneutrality in the plasticized polymer matrix (Scheme 4-1A). The response mechanism of these ISOs is dictated by mass transfer equilibria (cation-exchange process) between an organic (ISO) and an aqueous phase. Consequently, the detection of the cation of interest, after achieving thermodynamic equilibrium, is made by measuring the changes in the indicators’ optical properties, either by absorption or fluorescence spectroscopy. Nevertheless, prior to cation detection, these ISOs need an external source of protons for the cation-exchange process to proceed, resulting in a response mechanism with multiple steps. In contrast, the proposed tricyanofuran based mPAH (CF₃PhTCF-PAH) acts as a neutral acidic indicator (InH), providing protons for the cation-exchange process. Considering that mPAH is a photoactive compound, it photodissociates its proton upon irradiation and thermally undergoes proton reassociation with a dissociated state that is sufficiently long-lived, thus eliminating the need for an external source of protons, while the dissociated state of the mPAH acts as an anionic additive to maintain electroneutrality within the ISO as shown in Scheme 4-1B. Hence, an ISO based on the proposed mPAH exhibits a one-step response mechanism.

We have recently shown that merocyanine based mPAH linked to an acrylate polymer backbone can be utilized in an ISO to detect calcium ions. This merocyanine based mPAH is a neutrally charged acidic indicator which exhibits a longer equilibrium response time (in the order of hours). Consequently, the detection of calcium ions was performed under non-equilibrium conditions. Moreover, we have shown that by modifying the merocyanine based mPAH with an appropriate
functional group (electron donating group), the equilibrium response time can be shortened to the order of minutes.\textsuperscript{30} However, the charged nature of the merocyanine based mPAH was optimal only with a polar plasticizer, such as 2-nitrophenyl octyl ether (o-NPOE), to reduce ion-pair formation that affects selectivity.\textsuperscript{29} o-NPOE absorbs under 400 nm, interfering with the absorbance peak of the deprotonated state merocyanine based mPAH, and inhibited ratiometric analysis. In contrast, the non-charged CF\textsubscript{3}PhTCF-PAH has shown ideal compatibility with a non-polar plasticizer, such as bis(2-ethylhexyl)sebacate (DOS) which does not interfere optically, allowing ratiometric analysis to further increase the sensor sensitivity and signal reproducibility.\textsuperscript{31}

4.3 Results and Discussion

Scheme 4-2 Photoresponsive behaviour of CF\textsubscript{3}PhTCF-PAH when incorporated in an ISO.
As shown in Scheme 4-2, it is expected that CF$_3$PhTCF-PAH within the ISO would undergo oxidative photoreaction under visible light irradiation, resulting in a stable carbanion state (CF$_3$PhTCF-PA$^-$) of the mPAH along with its photodissociated proton.$^{32}$ Likewise, the photodissociated protons would be exchanged when exposed to cations of interest, as the CF$_3$PhTCF-PA$^-$ state is sufficiently long-lived to allow for the diffusion mediated cation-exchange process.

![Figure 4-1](Image)

Figure 4-1 Stability of CF$_3$PhTCF-PAH in ISOs for (A) calcium sensor in 0.5 M formate buffer at pH 4.5, and (B) sodium sensor in 0.3 M magnesium acetate buffer at pH 5.5. Absorbance was recorded at 470 nm. (ON state: after 1 minute irradiation with 470 nm; OFF state: after 25 minutes in the dark).

CF$_3$PhTCF-PAH was synthesized according to a literature procedure.$^{32-34}$ The calcium and sodium ISOs proposed here contain CF$_3$PhTCF-PAH (7.5 mmol kg$^{-1}$), and calcium ionophore IV (22.5...
mmol kg\(^{-1}\)) or sodium ionophore X (7.5 mmol kg\(^{-1}\)) within poly(vinyl chloride) (33 wt\%) and DOS (66 wt\%).

At first, these ISOs were exposed to their respective buffer solutions without any cation of interest. As shown in Figure 4-1, the ISOs based on CF\(_3\)PhTCF-PAH were stable over repeated activation cycles between the ON (deprotonated form) and OFF states (protonated form) without any loss of the absorbance signal, indicating no observable photodegradation.

![Figure 4-2](image)

Figure 4-2 Absorption spectra of ISOs for 25 minutes in the dark after activation for different concentrations of (A) calcium ions in 0.5 M formate buffer at pH 4.5; concentrations (a) 0, (b) 1.0 \(\times 10^{-8}\), (c) 1.0 \(\times 10^{-7}\), (d) 1.0 \(\times 10^{-6}\), (e) 1.0 \(\times 10^{-5}\), (f) 1.0 \(\times 10^{-4}\), and (g) 1.0 \(\times 10^{-3}\) M, and (B) sodium ions in 0.3 M magnesium acetate buffer at pH 5.5; concentrations (a) 0, (b) 1.0 \(\times 10^{-6}\), (c) 1.0 \(\times 10^{-5}\), (d) 1.0 \(\times 10^{-4}\), (e) 1.0 \(\times 10^{-3}\), (f) 1.0 \(\times 10^{-2}\), (g) 1.0 \(\times 10^{-1}\), and (h) 1.0 M.
Figure 4-2 shows the absorption spectra of the ISOs based on CF₃PhTCF-PAH towards different concentrations of calcium (Figure 4-2A) and sodium (Figure 4-2B) ions at thermodynamic equilibrium (25 minutes in the dark) once the ISOs were activated by visible light (470 nm) for 1 minute. From both absorption spectra, it is observed that as the concentration of the cation of interest increases, there is a gradual decrease in the CF₃PhTCF-PAH peak (470 nm) and a gradual increase in the CF₃PhTCF-PA⁻ peak (318 nm). These absorbance changes allow for ratiometric analysis (protonated and deprotonated absorbance peaks of the CF₃PhTCF-PAH) to indirectly correlate the activity of the cation of interest.

The experimental data obtained were then compared to the theoretical response function. The theoretical response function (eqn [4.1]), in terms of the activity of protons and the cation of interest, was generated from the ion-exchange equilibria (Scheme 4-1B). This was derived by utilizing mass balance (eqn [4-2] and eqn [4-3]), charge balance (eqn [4-4]), degree of deprotonation using the ratio of absorbance at 470 nm and 318 nm (eqn [4-5]), and the ion-exchange equilibrium constant (eqn [4-6]) equations. This is analogous to that of traditional ISO theory which was established in the early 1990s.²¹⁻²³

Theoretical response function for cation:

\[
\alpha_{M^{z+}} = \left(K_{exch}\right)^{-\frac{1}{z}} \left(\frac{(1-\alpha)a_{H^+}}{\alpha}\right)^{z} \frac{[InH]_{T(1-\alpha)}}{z[L]_{T}^{-\frac{1}{z}}[InH]_{T(1-\alpha)}}^{x}
\]  

[4.1]

Mass balance equations:
\[ [\text{InH}]_T = [\text{InH}] + [\text{In}^-] \quad [4.2] \]

\[ [L]_T = [L] + x[L_xM^{z+}] \quad [4.3] \]

**Charge balance equation:**

\[ [\text{In}^-] = z[L_xM^{z+}] \quad [4.4] \]

**Degree of deprotonation (\( \alpha \)):**

\[ \alpha = \frac{[\text{InH}]}{[\text{InH}]_T} = \frac{A - A_D}{A_P - A_D} \quad [4.5] \]

**Cation-exchange equilibrium constant (\( K_{exch} \)) for the neutral acidic indicator:**

\[ K_{exch} = \left( \frac{[\text{In}^-]a_{H^+}^x}{[\text{InH}]a_{M^{z+}}^z} \right)^x \frac{[L_xM^{z+}]}{a_{M^{z+}}[L]^x} \quad [4.6] \]

The subscript “T” indicates the total concentration of CF3PhTCF-PAH ([\text{InH}]_T) and the ionophore ([L]_T). The activity of the cation of interest and the proton is denoted as \( a_{M^{z+}} \) and \( a_{H^+} \). Also, “x” and “z” denote the value of the ionophore chelating with the cation of interest and charge of the cation of interest, respectively. The ratio of absorbance for CF3PhTCF-PAH is denoted as \( A \), the protonated state of CF3PhTCF-PAH in 1 M hydrochloric acid as \( A_P \), and the deprotonated state of CF3PhTCF-PAH in 1 M sodium hydroxide as \( A_D \); for 25 minutes in the dark after activation.

As shown in Figure 4-3, the experimental data present a strong correlation with the theoretical response curve. From the resulting cation response curves, the experimental limit of detection
(LOD) for calcium and sodium ions was $2.6 \times 10^{-6}$ M and $2.3 \times 10^{-3}$ M, respectively. These values were obtained by intersecting two extrapolated segments of the response curve as indicated in the literature.\textsuperscript{35} Furthermore, the cation-exchange constant ($\log K_{\text{exch}}$) for calcium and sodium ISOs was $-9.3 \pm 0.3$ and $-5.3 \pm 0.1$, respectively. The ionophore–cation complex ratio for calcium ISO was 3 to 1 and for sodium ISOs was 1 to 1.\textsuperscript{36,37}

Figure 4-3 Experimental (and) and theoretical (lines) responses of ISOs based on CF$_3$PhTCF-PAH for the detection of (A) calcium, and (B) sodium ions for 25 minutes in the dark after activation ($n = 3$).

It was hypothesized that ISOs which contain no anionic additive, cannot maintain constant ionic strength within the ISO.\textsuperscript{23} As a result, the cation-exchange constant cannot be retained, due to each
change in the activity of protons and the activity of the cation of interest.\textsuperscript{23} However, the changes in the ionic strength within the ISO based on CF\textsubscript{3}PhTCF-PAH, containing no additional anionic additive, were negligible as observed in the kinetic data for different concentrations of sodium ions (Figure 4-4), where stable responses over time were obtained at thermodynamic equilibrium. It is noteworthy that the interactions between the negatively and positively charged components for such an ISO do occur, after a certain threshold ($1 - \alpha > 0.5$), as the ionic strength within the ISO changes drastically. Accordingly, higher concentrations of cations are not shown in Fig. 3 because the thermodynamic equilibrium was not maintained, and thus the upper detection limit was $6.3 \times 10^{-4}$ M and 0.6 M for calcium and sodium ions, respectively (at $\alpha = 0.5$).
Figure 4-4 Kinetic study for ISOs based on CF₃PhTCF-PAH at different concentrations of sodium ions in 0.3 M magnesium acetate buffer at pH 5.5. Scans every 30 second after activation; concentrations (a) $1.0 \times 10^{-6}$, (b) $1.0 \times 10^{-5}$, (c) $1.0 \times 10^{-4}$, (d) $1.0 \times 10^{-3}$, (e) $1.0 \times 10^{-2}$, (f) $1.0 \times 10^{-1}$, and (g) 1.0 M.
Figure 4-5 Selectivity response for (A) calcium (buffer: 0.5 M formate at pH 4.5), and (B) sodium (buffer: 0.3 M magnesium acetate at pH 5.5) ISOs towards interfering cations for 25 minutes in the dark after activation (n = 3).

Furthermore, these ISOs based on CF$_3$PhTCF-PAH were studied in terms of selectivity towards interfering cations (J) by a separate solution method.$^{25,38}$ As shown in Figure 4-5, all interfering ions were highly discriminated for both ISOs, calcium (Figure 4-5A) and sodium (Figure 4-5B). Thus, the selectivity coefficient values for the calcium ISO towards magnesium, sodium and potassium ions were $-3.8$, $-3.1$ and $-3.3$, respectively. When compared to other ISOs, containing the same ionophore and spiropyran as the reference indicators, the selectivity coefficients were $-2.9$ (magnesium ion), $-6.6$ (sodium ion) and $-8.9$ (potassium ion).$^{16}$ The log $K_{M,J}$ values for the sodium ISO towards potassium, calcium and magnesium ions were $-4.5$, $-6.6$ and $-2.7$, respectively. These values are comparable to the selectivity coefficient of sodium against the potassium ion ($-2.4$), calcium ion ($-4.0$) and magnesium ion ($-4.1$) containing spiropyran as a
reference indicator. The selectivity coefficient \( \log K_{M,J} \) values were calculated using eqn [4.7], and is analogous to that of ISO theory at \( \alpha = 0.5 \).

Selectivity coefficient for interfering cations:

\[
K_{M,J}^{opt} = \frac{K_{exch}^{J}}{K_{exch}^{M}} \left[ \frac{(1-\alpha)\alpha_{H^+}}{\alpha} \right]^{z-w} \frac{w(\{L\}_T - \frac{p}{w}\{lnH\}_T(1-\alpha))}{z(\{L\}_T - \frac{z}{w}\{lnH\}_T(1-\alpha))}^{x}
\]

[4.7]

The high discrimination observed may be due to the proton affinity towards \( \text{CF}_3\text{PhTCF-PA}^- \) (deprotonated form) to be greater than the binding energy of interfering cations by the respective ionophores. Consequently, these ISOs inhibit the competitive behaviour between the interfering cations and protons during the cation-exchange process, even at high concentrations. As a result, an ISO based on \( \text{CF}_3\text{PhTCF-PAH} \) may also be utilized for practical purposes which demands negligible interactions towards interfering cations.

4.4 Conclusion

In conclusion, this novel visible light activatable and controllable ISO based on \( \text{CF}_3\text{PhTCF-PAH} \) that does not exhibit photodegradation, showed good stability, selectivity and reproducibility. Furthermore, the responses of these ISOs towards calcium and sodium ions were standardized following the cation-exchange equilibria. Similarly, other cations can be detected using this platform by interchanging the ionophore within the polymer matrix. Also, we aim to optimize this type of ISO in terms of size reduction and sensitivity. Likewise, we expect that this novel mPAH
(CF$_3$PhTCF-PAH) may act as a substitute to neutral basic indicators for their use in cation sensing applications that provide control using visible light as needed.

4.5 Acknowledgements

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4.6 References


CHAPTER 5 SWITCHING FLUORESCENCE AND ACIDITY BY VISIBLE-LIGHT ACTIVATED METASTABLE-STATE PHOTOACID AND BODIPY IN SOLUTION


5.1 Abstract

Small changes in the pH gradient play a critical role in numerous biological and chemical pathways. Systems capable of monitoring and regulating these changes with high sensitivity and minimum photo-fatigue are in demand. Herein, we propose a visible light-triggered molecular system that allows for reversible regulation of acidity and fluorescence. This robust bi-functional system opens a new horizon towards novel studies that rely on small changes in acid-mediated controlled processes with high sensitivity. The two photosensitive compounds employed, a metastable-state photoacid (mPAH) and a boron-dipyrrromethene (BODIPY) derivative, allow for consistent modulation of both fluorescence (based on the working principle of the inner filter effect) and pH (around a magnitude) over multiple cycles.
5.2 Introduction

Organic photochromic molecules such as spiropyrans, azobenzenes or diarylethenes are compounds that undergo reversible structural changes when exposed to light.\textsuperscript{1–3} The physicochemical changes associated with the structural transformations of these compounds include variations in their optical properties, dipole moment, and/or conductivity.\textsuperscript{4–8} As so, pH sensitive photochromic compounds are attractive for applications concerning proton-transfer processes. Traditionally, spiropyrans have been used for this purpose.\textsuperscript{9} Nonetheless, these molecules require acidification using an external source of protons while activated with UV light to photoisomerize them from the spiropyran form (SP) to the merocyanine form (MC). The protonated merocyanine species obtained by this procedure (MCH\textsuperscript{+}) is then sensitive to visible light (Scheme 5-1A). In practice, these conditions could prove challenging in terms of bio-applicability and robustness. To circumvent these obstacles, a new class of visible light-triggered molecular switches called metastable-state photoacids (mPAHs) has been proposed.\textsuperscript{10}

In contrast to classical super photoacids,\textsuperscript{11} this new class of visible light-triggered mPAHs also allows for a large pH change with high efficiency and good reversibility. These novel mPAHs are designed by linking an electron-accepting (EA) moiety and a weakly acidic nucleophilic moiety (NuH) with a double bond (Scheme 5-1B).\textsuperscript{10} Under irradiation, the trans–cis photoisomerization, followed by a nucleophilic reaction between the two moieties, results in an efficient proton release. In the dark, the metastable-state relaxes to the initial conformation within a few seconds to hours, thus making it essential to study proton-transfer processes due to the long-lived nature of the metastable state.\textsuperscript{12–16} Due to their acidity-modulating capabilities, mPAHs have found applications
in hydrogels,\textsuperscript{17} ion-selective optodes,\textsuperscript{18,19} polymerization reactions,\textsuperscript{20} photo-activated ion-channels,\textsuperscript{21} and waste management systems.\textsuperscript{22}

Scheme 5-1 (A) Photochromic equilibrium of spiropyran and external acidification to produce light triggered proton releasing merocyanine. (B) General scheme for mPAHs. (C) Acido-basic and photochromic equilibria between Mer-mPAH (protonated-open form), Mer-mPA\(^{-}\) (conjugate basic-open form) and SP-mPA\(^{-}\) (conjugate basic-closed form).

Recently, a new class of multi-stimulus-responsive fluorescent molecular switches based on mPAHs has been shown.\textsuperscript{23} The authors provided a unique approach to covalently link a UV light activated fluorophore to the mPAH moiety. As mentioned before, the need of a UV light source to
induce fluorescence may likely compromise the robustness of the photonic device and its use in biological applications.

Herein, we present a simple and robust pH and fluorescence modulating system that can be activated with visible light. This system is composed of a high quantum yield pH insensitive fluorophore (a boron-dipyrrromethene (BODIPY) derivative\textsuperscript{24–26} and a novel merocyanine-type mPAH (Mer-mPAH; Scheme 5-1C) acting as a molecular switch. By utilizing the concept of the inner filter effect,\textsuperscript{27} both the absorption and emission of BODIPY are modulated by the state of Mer-mPAH. The approach of using the inner filter effect has found success in improving pH-dependent fluorescence sensitivity for other systems.\textsuperscript{28,29} Nonetheless, these systems do not exploit the benefits of mPAHs.

5.3 Synthesis

A lipophilic Mer-mPAH was synthesized according to the pioneering work of Shi et al. with few modifications to make the product more stable and compatible with organic solvents (Scheme C-1 in Appendix C).\textsuperscript{10}
5.4 Acido-basic equilibrium

Figure 5-1 (A) pH dependent absorption spectra of Mer-mPAH (50 µM) in 0.1 M of different buffers in co-solvent (ethanol/water – 50/50) system. (B) Time-dependent absorbance spectra of Mer-mPAH (65 µM) in pure ethanol recorded every 0.1 minute after 470 nm light irradiation (ON) and in the dark (OFF). Inset: Changes in major absorption peaks between protonated and deprotonated forms of Mer-mPAH with respect to time.
First, the acido-basic equilibrium of Mer-mPAH (50 μM) was investigated by acquiring absorption spectra in 0.1 M Britton–Robinson (pH 2 to 7), 0.1 M Tris–HCl (pH 8 to 9) and 0.1 M Glycine–NaOH (pH 10) buffers. Different buffers were used due to solubility issues at high pH in the co-solvent (ethanol/water – 50/50) system. This experiment was performed to identify the spectral signatures of three acido-basic species (Scheme 5-1C). As shown in Figure 5-1A, a strong absorption band with a peak at 474 nm was predominant in acidic media (pH 2.18) with a distinct yellow colour. This intense and broad spectral feature was assigned to the unique protonated species Mer-mPAH. TD-DFT computations (Section C.3 in Appendix C) on optimized geometries (Figure C-7 in Appendix C) confirmed this statement. Although the theoretical transition energy seemed to be slightly overestimated (0.22 eV), the general shape of the absorption band was well reproduced (Figure C-8A in Appendix C).

When the pH was increased, the absorption band peak of the protonated-open form decreased, and the two absorption band peaks at 284 nm and 556 nm increased (colour of the solution turned to light pink). These two absorption bands correspond to the two different conjugate basic forms of Mer-mPAH. The red shifted peak centered at 556 nm corresponds to the conjugate basic open-form (Mer-mPA\(^-\)), due to the electronic enhancement by the resonance of Mer-mPAH. The blue shifted peak centered at 284 nm is assigned to the conjugate basic-closed form (SP-mPA\(^-\)), given that the resonance is broken at the chiral sp\(^3\) carbon center. Theoretical calculations allowed confirming the identity of the two species that give rise to the peaks in basic media (Figure C-8B and C in Appendix C). The red-shift that occurs upon deprotonation is well reproduced for Mer-
mPA$^-$ (experimental: 0.39 eV, theoretical: 0.31 eV) and the shape of the absorption spectra of SP-mPA$^-$ is well reproduced.

With complete rationalization of acido-basic species, the thermodynamic acidity constant of Mer-mPAH was then estimated to be 6.15 by following a literature protocol (Section C.4 in Appendix C).  

5.5 *Photochromism*

Next, the visible light-triggered photochromic behaviour of Mer-mPAH (65 μM) was examined in pure ethanol solution. It is evident from Figure 5-1B that in ethanol solution, the starting material is exclusively in its protonated-open form with a strong absorption band peak at 478 nm. Upon irradiation with visible light (470 nm), the yellow coloured solution turns colourless within a few seconds. This is the result due to the drastic decrease in the protonated-open form and reaching a photo-stationary state (Figure 5-1B, inset, blue region). Only the conjugate basic SP-mPA$^-$ form was observed, as attested by the increase of the 278 nm absorption band (analogous to the 284 nm band reported before) and the lack of the 556 nm shoulder. When irradiation of visible light was ceased, the peak at 478 nm appeared again, and the solution recovered its yellow colour. The exponential time constant for this recovery kinetic is $1.56 \pm 0.01$ minutes, which is indicative of the fast thermal back reaction via carbon–oxygen bond breaking followed by unfolding merocyanine isomerization with proton assistance.  

Using the ground state structures obtained theoretically, Mer-mPAH was found to be more stable than SP-mPA$^-$ (Section C.3 in Appendix C). Accordingly, the changes in Gibbs free energy between the photo-stationary (SP-mPA$^-$ form)
and ground (Mer-mPAH form) states were 87.8 kJ mol$^{-1}$ in ethanol and 83.0 kJ mol$^{-1}$ in aqueous solutions. To the best of our knowledge, this is the first theoretical report on the thermodynamics of mPAH. For comparison, based on theoretical thermodynamics reported for spiropyans, the changes in Gibbs free energy between the photo-stationary (MC form, see Scheme 5-1A) and ground (SP form, see Scheme 5-1A) states are also positive values.$^{31-36}$ Likewise, due to the more stable nature of Mer-mPAH, this difference in energy may likely assist the thermodynamic reverse process in darkness.

To corroborate that protons have been released from the Mer-mPAH by visible light, pH measurements under irradiation were performed (Fig. C-10 in Appendix C). A 65 μM Mer-mPAH solution in ethanol/water (90/10) yielded an initial pH of 5.5. When visible light was switched ON, the pH of the solution decreased to 4.3, with a 1.20 ± 0.02 change. This is an important performance compared to previously reported mPAHs.$^{10,12-15}$ These observations confirmed that the Mer-mPAH has the ability to concomitantly release its protons while converting to its conjugate basic form using visible light. Likewise, when the visible light was switched OFF, the pH increased to its initial state, confirming the thermodynamic reversibility of the process. Moreover, a consistent magnitude of pH change can be modulated over multiple cycles with no observable photo-degradation.
5.6 Visible light-triggered pH/fluorescence bi-molecular system

Figure 5-2 (A) Absorption spectra for the optimized chromophore mixture (2.6 µM of BODIPY with 65 µM of Mer-mPAH) in pure ethanol without (black curve) and under visible light irradiation (red curve). Arrows indicate possible fluorescence from BODIPY (inner filter effect illustration). (B) Emission spectra (excitation at 495 nm) for BODIPY (2.6 µM) in pure ethanol with increasing Mer-mPAH concentration (a - i: 0, 1.3, 2.6, 5.2, 13, 26, 39, 52, 65 µM). Inset: Stern–Volmer plot for the ratio of fluorescence lifetime decays, where ‘‘τ₀’’ and ‘‘τ’’ represent the lifetimes of BODIPY only and the mixture, respectively. (C) pH modulation by 470 nm light for an ethanol/water (90/10) solution containing the chromophore mixture (2.6 µM of BODIPY with 65 µM Mer-mPAH) with respect to time. (D) Fluorescence modulation for the optimized chromophore mixture in pure ethanol: green curve refers to BODIPY only emission (I_{\text{max}}); black curve refers prior to visible light irradiation; red curve refers after 0.25 minute of visible light irradiation (I_{\text{ON}}); and blue dashed curve refers after 10 minutes in the dark (I_{\text{OFF}}). Inset: Repeatability cycles.

The objective of this study was to obtain an efficient visible light-triggered BODIPY fluorescence in parallel to pH modulation using Mer-mPAH through the well-known inner-filter effect.²⁷
better understand the visible light-triggering of fluorescence in the chromophore mixture, we measured the absorption spectra for a blend of BODIPY (2.6 μM) and Mer-mPAH (65 μM) (Figure 5-2A). In the dark, Mer-mPAH absorption is predominant, thus avoiding the BODIPY molecules to absorb the excitation light (495 nm) and consequently resulting in negligible fluorescence (Mer-mPAH is poorly fluorescent, Figure C-11 in Appendix C). In this state, the fluorescence of BODIPY is OFF. Reversely, when the blend was exposed to visible radiation, Mer-mPAH converts to its conjugate basic forms with reduced absorbance in the visible region, thus allowing the BODIPY molecules to absorb the excitation light. In this state, the fluorescence of BODIPY is ON.

As a matter of fact, the key points to design such a system are (i) a good overlap between Mer-mPAH and BODIPY absorption bands and (ii) significant differences between extinction coefficients between acidic and basic species of Mer-mPAH. Consequently, we decided to investigate the best choice of concentrations for both molecules in the blend. Photochemical parameters of BODIPY are well known, and for ethanol solution, the absorption maximum is typically observed at approximately 500 nm (Figure C-12A in Appendix C). After testing different BODIPY concentrations in ethanol (excitation at 495 nm), we found that 2.6 μM yields a maximum emission intensity within the linear dynamic range at 509 nm (Fig. C-12B, inset, in Appendix C). Furthermore, as shown in Fig. 2B, ~99% of BODIPY (2.6 μM) emission was inhibited when 25 equivalents of Mer-mPAH (65 μM) were used. In fact, the optimized choice of concentrations was contemplated by the differences in molar extinction coefficients of BODIPY.
(98040 M$^{-1}$cm$^{-1}$) and Mer-mPAH (40138 M$^{-1}$cm$^{-1}$) (Figure C-13 in Appendix C). Therefore, excess Mer-mPAH was required to maximize fluorescence inhibition.

To explore possible dynamic quenching of BODIPY fluorescence by Mer-mPAH, and to validate that the inner filter effect is the only responsible criterion for fluorescence inhibition, fluorescence lifetime measurements were performed for BODIPY (2.6 μM) with different concentrations of Mer-mPAH in ethanol solution (Figure 5-2B, inset; and Figure C-14, Table C-1 in Appendix C). The results showed no observable differences, which indicates that the fluorescence inhibition is intrinsically governed by the inner filter effect.$^{27}$

With the optimized concentrations of the chromophore mixture (BODIPY – 2.6 μM and Mer-mPAH – 65 μM), the repetitive modulation of pH and fluorescence was investigated. Like the pH modulation study shown earlier for Mer-mPAH using visible light, this chromophore mixture in ethanol/water (90/10) solution displayed an identical pH change of 1.20 ± 0.01 over multiple cycles (Figure 5-2C). Moreover, this chromophore mixture in ethanol solution exhibits similar photochromic response when monitored by absorption with an exponential time constant for the recovery kinetic of 1.41 ± 0.01 minutes (Fig. C-15 in Appendix C). No observable changes were observed for solution containing BODIPY only. Finally, upon examination of fluorescence switching for this chromophore mixture in ethanol solution, ~67% of emission after visible light irradiation was observed (Figure 5-2D) compared to the emission of BODIPY only. This is due to a part of excitation light (495 nm) being absorbed by the remaining protonated-open form of Mer-mPAH after irradiation with visible light. This outcome underlines the fact that for future
development and the design of a mPAH, one must consider a strong absorption overlap between the initial form of mPAH and the fluorophore for the best inner filter effect. Likewise, upon visible light irradiation of mPAH, the inner filter effect will be eliminated. Thereby, allowing the fluorophore to absorb excitation light and emit as fluorescence. Moreover, the observed emission intensity was completely inhibited to its initial state after 10 minutes in the dark. Likewise, no photo-degradation was observed while performing fluorescence switching of this chromophore mixture by visible light over multiple times for more than two hours (Figure 5-2D, inset). With these cumulative results, Mer-mPAH has demonstrated its ability to modulate both pH and fluorescence by incorporating BODIPY in solution.

5.7 Conclusions

We have shown a novel visible light-triggered bimolecular system capable of simultaneously modulating pH and fluorescence. This system exhibits good photostability, and consistent pH and fluorescence changes over multiple cycles. This controllable system is expected to find applications in the study of pH dependent enzymes, pH sensitive drug tracking and release near cancerous sites, and improvement of the sensitivity of light stimulated sensing platforms for ions.

The fluorescence modulation shown in the Mer-mPAH and BODIPY blend is the result of the inner filter effect. However, the switching of fluorescence is not as effective as the dynamic quenching mechanism. A better understanding of this family of chromophores and further investigation of ultrafast mechanisms in these systems are still necessary for more emissive mPAHs. This possibility is currently being explored in our laboratory.
5.8 Acknowledgements

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5.9 References


APPENDIX A: SUPPORTING INFORMATION OF CHAPTER 2
A.1 Materials
Calcium Ionophore IV (ETH 5234), poly (vinyl chloride) high molecular weight (PVC), o-nitrophenyl octyl ether (o-NPOE), bis(2-ethylhexyl) sebacate (DOS) and tetrahydrofuran (THF) were provided from Sigma Aldrich (St. Louis, MO). Calcium, potassium sodium, magnesium, and cadmium chlorides, tris(hydroxymethyl)aminomethane and hydrochloric acid were purchased from Fisher Chemicals (Fair Lawn, NJ).

A.2 Instrumentation
A 470 nm LED array with 56 10,000 mcd LEDs (Elixa, Ltd.) was used for irradiation. A UV-Vis spectrophotometer (Cary 50 Bio UV/Visible) from Varian, Australia; a spin coater (WS-650MZ-23NPP) from Laurell North Wales, PA; and a deionized water laboratory purification system (PURELAB Ultra) from Siemens, Orlando, FL.

A.3 Preparation of sensing membranes
The ion-sensing optode membrane contains 9.74% (84.56 mmol/kg of photoacid unit) of polymer 1, 8.48% (105.70 mmol/kg) of calcium ionophore IV, 27.26% PVC and 54.52% o-NPOE. THF was added and the mixture was vigorously shaken for 30 minutes to dissolve all the components. Subsequently, a 20 µL of the cocktail was added onto clean glass slides and spin coated for 2 minutes at 600 rpm to form a thin optode membrane for UV-Vis analysis. The glass slides were previously cut to fit the cuvette, cleaned in an ethanol bath while being scrubbed with cotton swabs and dried under mild heat using a heat gun.

A.4 UV-Vis spectrophotometer measurements

UV-Vis measurements were done in 0.5 M Tris(hydroxymethyl)aminomethane solution at pH 7.40 (pH was adjusted using 6.0 M HCl). Calibration curve was performed using different concentrations of calcium chloride solutions in the range of 1.0 x 10^{-5} to 1.25 M. First, a baseline was corrected using glass slide in buffer solution. Prior to irradiation, the sensing membrane was scanned in the buffer to obtain the absorbance maxima. Immediately, the sensing membrane was irradiated for 5 minutes at 470 nm to turn the system ON, and subsequently scanned. Then the sensing membrane was kept in dark for 35 minutes and scanned, the resulting spectra was referred as in the dark. The last two steps were repeated subsequently with increasing calcium concentrations. The absorbance registered was plotted against the log of the activity of calcium ions. Figure A-1 shows the change in absorbance for different concentrations of calcium ions after 5 min activation at different times of ion-exchange process (5, 10, 20 and 35 min).
Figure A-1 Absorbance response of calcium-sensing optode membrane activated at 5 min and allowed for different ion-exchange times at a) 5, b) 10, c) 20 and d) 35 minutes in the dark at concentrations of calcium ions ranging from $1.0 \times 10^{-5}$ to 1.25 M. In 0.5 M of tris(hydroxymethyl)aminomethane buffer at pH 7.40.

A.5 Selectivity measurements

For selectivity measurements a similar process was followed as described above. First, the signal was obtained in the buffer solution and then in the corresponding analyte ($\text{Ca}^{2+}$, $\text{Na}^+$, $\text{K}^+$ and $\text{Mg}^{2+}$) at different concentrations.
APPENDIX B: SUPPLEMENTARY INFORMATION OF CHAPTER 4

B.1 Experimental Methods

B.1.1 Reagents

N,N-Dicyclohexyl-N’,N’-dioctadecyl-3-oxapentanediamide (Calcium ionophore IV), 4-tert-butylcalix[4]arene-tetraacetic acid tetraethyl ester (Sodium Ionophore X), high molecular weight poly(vinyl chloride) (PVC), bis(2-ethylhexyl) sebacate (DOS), tetrahydrofuran (THF), sodium chloride, potassium chloride, calcium chloride, magnesium chloride, magnesium acetate tetrahydrate and sodium hydroxide were purchased from Sigma Aldrich, USA. Formic acid, hydrochloric acid and glacial acetic acid were purchased from Fisher Scientific, USA. Potassium hydroxide was purchased from Amresco, USA. Tricyanofuran based metastable photoacid (CF₃PhTCF-PAH) was synthesized according to literature procedures.¹⁻³

B.1.2 Instrumentation

The absorbance measurements were performed using a UV-Vis spectrophotometer (Cary 50 Bio UV/Visible) from Varian, Australia. The source of visible light irradiation was a 470 nm LED array with 120 LEDs from Elixa, USA. The thickness of the cation-selective optode membranes was measured using a profilometer (Tencor AlphaStep 500) from KLA Tencor, USA. Deionized
water used to prepare solutions for analysis was purified by a water purification system with resistance of 18 MΩcm (PURELAB Ultra) from Siemens, USA. The pH of the solutions was obtained with a pH meter (Orion Start A211) from Thermo Scientific, USA. All experiments were carried out in the dark.

**B.1.3 Preparation of Cation-selective optodes**

The calcium and sodium ion-selective optodes contain CF$_3$PhTCF-PAH (7.5 mmol/kg), calcium ionophore IV (22.5 mmol/kg) and sodium ionophore X (7.5 mmol/kg), respectively, PVC (33 wt%), and DOS (66 wt%). These components were dissolved in THF (0.8 mL) and shaken until the solution was homogenous. Subsequently, 20 μL of the cocktail solution was added onto clean silanized glass slides. The resultant ISO membranes were dried first in air for 1 hour (in the dark) and then stored in vacuum until use. The membrane thicknesses were between 35 – 45 μm.

**B.1.4 Response measurements**

The absorbance measurements were performed in 0.5 M formate buffer at pH 4.5 (pH was adjusted with 2.0 M KOH) for calcium-selective optode and 0.3 M magnesium acetate buffer at pH 5.5 (pH was adjusted with dilute acetic acid) for sodium-selective optode. First, a baseline was corrected using a clean silanized glass slide in buffer solution. Prior to irradiation, the cation-selective optode membrane was scanned in the buffer. Immediately, the sensing membrane was turned ON by irradiating with 470 nm visible light for 1 minute and subsequently scanned. Then, the sensing
membrane was kept in dark for 25 minutes (OFF state) and scanned. Afterwards, the buffer solution was changed and the last two steps were repeated four more times.

The response for both cation-selective optode membranes towards its cation of interest and the kinetic experiment for sodium-selective optode membrane were performed similarly as mentioned above. First, the response (ON and OFF state) of the cation sensing membrane was obtained in buffer solution. Then, the last two steps were repeated with increasing cation concentrations (calcium chloride: 1.0x10^{-8} M to 1.0x10^{-3} M, and sodium chloride: 1.0x10^{-6} M to 1.0 M) in their respective buffer. This experiment was performed in triplicates. However, for the kinetic experiment, the response of the CF₃PhTCF-PAH was obtained by scanning the sensing membrane every 30 seconds for 25 minutes after 1 minute irradiation (470 nm).

**B.1.5 Selectivity measurements**

These measurements were performed similarly as mentioned above in triplicates. The interfering cations used were magnesium, sodium and potassium for calcium-selective optode, and potassium, calcium and magnesium for sodium-selective optode.

**B.2 Supplementary References**


C.1 Experimental Section

C.1.1 Materials and Instrumentation

Acetone, dichloromethane (DCM), hexanes, ethyl acetate, methanol, ethanol, glacial acetic acid, hydrochloric acid (HCl), sodium hydroxide (NaOH) and sodium bicarbonate were purchased from Fisher, USA. Benzoyl chloride, 2,4-dimethylpyrrole, boron trifluoroetherate, glycine, acetonitrile, and 2-ethylhexyl bromide were purchased from Acros, USA. 2,4-dihydroxybenzaldehyde and cyclohexyl methyl ketone were purchased from Alfa Aesar, USA. Phenyl hydrazine hydrochloride was purchased from TCI, USA. Potassium carbonate (K2CO3), tris(hydroxymethyl)aminomethane (Tris) and sodium sulfate were purchased from Amresco, USA. Potassium iodide (KI), sodium chloride (NaCl), boric acid, phosphoric acid, 1,3-propanesultone and triethylamine were purchased from Sigma-Aldrich, USA. Deuterated NMR solvents were purchased from Cambridge Isotope Laboratories, Inc., USA. BODIPY was synthesized according to literature procedure.¹

¹H and ¹³C NMR spectroscopy were carried out with NMR spectrometer (Avance III 400) from Bruker, USA. Absorbance based experiments were performed using a UV-Vis spectrophotometer (Cary 50 Bio UV/Visible) from Varian, Australia. The source of visible light activation was a 470
nm 120 LED array (maximum average power of 1800 mW) from Elixa, USA. Deionized water used to prepare solutions for pH studies was purified by a water purification system with resistance of 18 MΩcm (Milli-Q Academic) from EMD Millipore Corporation, USA. The pH of the solutions was obtained with a pH meter (Orion Start A211) from Thermo Scientific, USA. Steady-state fluorescence experiments were performed using a spectrofluorometer (FluoroMax-4 Spectrofluorometer) from Horiba Scientific, USA. Time-resolved fluorescence (lifetime decay profiles) studies were performed using a spectrofluorometer (FLS980 Spectrometer) equipped with a 470 nm ± 10 nm picosecond pulsed diode laser as the excitation source (EPL – 470 with a maximum average power of 5 mW) from Edinburgh Instruments, UK. The excitation wavelength of diode laser (EPL – 470) was 470 nm (Δλ = 1 nm), pulse duration was 92 ps, pulse spacing was fixed at 200 ps, and pulse repetition rate was 5 MHz. The emission wavelength selected was 509 nm (Δλ = 1 nm) based on the steady-state fluorescence experiment. The fluorescence lifetime decay profiles were recorded until 2000 peak counts were reached for all ethanolic solutions (see Figure C-14 and Table C-1 for concentrations used) at room temperature. The decay profiles were corrected with background (instrument response function) and processed to determine the lifetime values obtained by global analysis (reconvolution) fitting using FAST Version 3.4.2. Software (Edinburgh Instruments, UK). All experiments were carried out in the dark unless mentioned otherwise.

C.1.2 Synthesis
Scheme C-1 Synthetic route used to prepare Mer-mPAH.

2-hydroxy-4-(2-ethylhexoxy)benzaldehyde (1): The synthesis of 1 was modified following literature procedure. In a 50 mL two-neck flask, equipped with a stir bar and a reflux condenser, was charged with K₂CO₃ (0.506 g, 3.656 mmol) followed by flame drying and purging with nitrogen gas (three times) the equipment setup. Under nitrogen gas, 2,4-dihydroxybenzaldehyde (1.000 g, 7.240 mmol) and dry acetone (8 mL) were added to the reaction flask, followed by raising the temperature to 65°C. After 20 minutes, a solution of KI (10%) and 2-ethylhexyl bromide (2.438 g, 7.312 mmol) in dry acetone (12 mL) was added dropwise to the reaction flask. After 15 hours, the reaction mixture was concentrated in vacuo, and excess 2-ethyl hexyl bromide was removed by distillation. The remaining organic residue was dissolved in minimal DCM and neutralized with
0.1 M HCl. The acid washes were extracted with DCM. Then, the organic phase was washed with brine (saturated NaCl), separated and passed through a drying column. After concentrating the organics in vacuo, the crude was purified by column chromatography (silica) in hexanes and eluted in 50% ethyl acetate in hexanes to afford a colorless oil. Yield (49%). $^1$H NMR (400 MHz, CDCl$_3$): $\delta$ (ppm) = 11.48 (s, 1H), 9.70 (s, 1H), 7.42-7.40 (d, $J = 8.67$ Hz, 1H), 6.55-6.52 (dd, $J = 2.32$ Hz, $J = 8.66$ Hz, 1H), 6.42 (d, $J = 2.29$ Hz, 1H), 3.90-3.89 (d, $J = 5.83$ Hz, 2H), 1.77-1.71 (m, 1H), 1.53-1.30 (m, 10H), 0.95-0.91 (m, 6H). $^{13}$C NMR (400 MHz, CDCl$_3$): $\delta$ (ppm) = 194.24, 166.69, 164.53, 135.14, 114.98, 108.80, 101.07, 71.05, 39.13, 30.40, 29.00, 23.77, 22.99, 11.04.

2'-methylspiro[cyclohexane-1,3'-[3H]indole] (2): The synthesis of 2 was modified following literature procedure.$^3$ In a 30 mL microwave reaction flask, equipped with a stir bar, was charged with cyclohexyl methyl ketone (1.003 g, 7.949 mmol) and glacial acetic acid (13.93 mL). This reaction mixture was sonicated for few minutes before adding phenyl hydrazine hydrochloride (1.138 g, 7.870 mmol). After 20 mins of microwave irradiation (150 psi, 200W), glacial acetic acid was removed in vacuo to result an orange residue. This residue was dissolved in DCM and neutralized with sodium bicarbonate. The organic phase was separated and washed three times with brine. Then, the organic phase was collected, dried with anhydrous sodium sulfate and the DCM was removed in vacuo to afford a red brown oil. Yield (77%). $^1$H NMR (400 MHz, CDCl$_3$): $\delta$ (ppm) = 7.71-7.69 (d, $J = 7.36$ Hz, 1H), 7.56-7.54 (d, $J = 7.54$ Hz, 1H), 7.34-7.30 (td, $J = 7.57$ Hz, $J = 15.21$ Hz 1H), 7.17-7.13 (td, $J = 7.48$ Hz, $J = 14.98$Hz, 1H), 2.27 (s, 3H), 1.98-1.73 (m, 8H), 1.31-1.26 (m, 2H). $^{13}$C NMR (400 MHz, CDCl$_3$): $\delta$ (ppm) = 187.65, 154.12, 144.48, 127.39, 124.14, 124.13, 120.00, 57.77, 31.06, 25.21, 21.51, 16.02.
2’-methyl-1’-(3-sulfopropyl)spiro[cyclohexane-1,3’-[3H]indolium] inner salt (3): The synthesis of 3 was modified following literature procedure. In a 125 mL round bottom flask equipped with a stir bar, 2 (1.10 g, 5.517 mmol) dissolved in acetonitrile (55 mL) was added under nitrogen gas. Then, 1,3-propanesultone (0.741 g, 6.068 mmol) was added dropwise over a period of 30 minutes to the reaction mixture at room temperature under nitrogen gas. Afterwards, the reaction mixture was refluxed under nitrogen gas for 20 hours. Acetonitrile was then removed under vacuo and the crude was purified by column chromatography (silica) using 7% (v/v) methanol in DCM to remove unreacted 1,3-propanesultone. Both purple and brown fractions were combined and concentrated under vacuo, and used without further purification. NMR spectra could not be obtained. Crude yield (60%).

Mer-mPAH: The synthesis of Mer-mPAH was modified following literature procedure. In a 125 mL round bottom flask, crude 3 (0.200 g, 0.622 mmol) was dissolved in anhydrous ethanol (25 mL). Prior to immediate addition of 1 (0.267 g, 0.684 mmol) to the reaction mixture, it was first placed under high vacuum overnight. After addition, the reaction mixture was heated to reflux under nitrogen for 4 hours. Ethanol was removed in vacuo, and the residue was purified by column chromatography (silica, washed with 1% acetic acid in mobile phase) using 6% (v/v) methanol in DCM. All orange and pink fractions were collected and washed with aqueous 1.0 M HCl and extracted with DCM. The organic phase was separated and dried using anhydrous sodium sulfate. The clear reddish orange organic solution was concentrated in vacuo to afford a red solid. Yield (20%). $^1$H NMR (400 MHz, DMSO-d$_6$): δ (ppm) = 11.34 (br. s, 1H), 8.76–8.72 (d, J = 16.16 Hz, 1H), 8.32–8.29 (d, J = 9.61 Hz, 1H), 8.13–8.11 (d, J = 7.43 Hz, 1H), 8.02–8.00 (d, J = 8.07 Hz, 1H),
7.66-7.60 (m, 2H), 7.53-7.49 (t, J = 7.48 Hz, J = 15.00 Hz, 1H), 6.59 (m, 2H), 4.78-4.74 (br. t, J = 7.36 Hz, J = 14.99 Hz, 2H), 3.95-3.93 (d, J = 5.46 Hz, 2H), 2.69-2.66 (br. t, J = 5.95 Hz, J = 12.15 Hz, 2H), 2.22-1.88 (m, 10H), 1.69-1.64 (m, 3H), 1.33-1.24 (m, 6H), 0.90-0.87 (m = 8H). ¹³C NMR (400 MHz, DMSO-d₆): δ (ppm) = 165.77, 161.66, 148.25, 141.74, 141.35, 128.89, 127.43, 125.46, 115.04, 114.80, 108.64, 106.97, 101.07, 70.56, 55.53, 47.31, 44.91, 38.37, 34.26, 29.72, 28.31, 24.27, 24.20, 23.12, 22.40, 20.60, 13.85, 10.79
C.2 $^1$H and $^{13}$C NMR spectra for 1, 2 and Mer-mPAH

Figure C-1 $^1$H NMR spectra of 1.
Figure C-2 $^{13}$C NMR spectra of 1.
Figure C-3 $^1$H NMR spectra of 2.
Figure C-4 $^{13}$C NMR spectra of 2.
Figure C-5 $^1$H NMR spectra of Mer-mPAH.
Figure C-6 $^{13}$C NMR spectra of Mer-mPAH.
C.3 DFT/TD-DFT calculations

All calculations were performed using the Gaussian 09 package. The visualizations were obtained using the Avogadro 1.2.0n software. All calculations including the geometry optimizations, the frequency calculations, and the time-dependent single point energy calculations were carried out with the B3LYP global hybrid functional using the 6-311++G (2d,p) basis set. Ethanol was accounted as an implicit solvent using the IEFPCM solvation model.

![Optimized geometries for ground state structure](image)

Figure C-7 Optimized geometries for ground state structure of (A) Mer-mPAH, (B) Mer-\( \text{mPA}^- \), and (C) SP-\( \text{mPA}^- \) forms.

C.3.1 Gibbs-free energy calculations

The Zero-Point Energy (ZPE) corresponds to the energy of a system where the molecular vibrations that persist at 0K are taken into consideration. To obtain the Gibbs-free energies of mPAH in its different states, we performed frequency calculations on the optimized structures of
Mer-mPAH, Mer-mPA\textsuperscript{-}, and SP-mPA\textsuperscript{-}. Then, the energy of the system in its different states was computed as follows:

\[ \Delta G_{\text{Mer-mPAH}} = ZPE_{\text{Mer-mPAH}} = -5446010.5 \text{ kJ/mol} \]  

\[ \Delta G_{\text{Mer-mPA}^-} = ZPE_{\text{Mer-mPA}^-} + \Delta G_{\text{Solvated } H^+} = (-5444831.6 \frac{\text{kJ}}{\text{mol}}) + (-1104.5 \frac{\text{kJ}}{\text{mol}}) = -5445936.1 \text{ kJ/mol} \]  

\[ \Delta G_{\text{SP-mPA}^-} = ZPE_{\text{SP-mPA}^-} + \Delta G_{\text{Solvated } H^+} = (-5444818.3 \frac{\text{kJ}}{\text{mol}}) + (-1104.5 \frac{\text{kJ}}{\text{mol}}) = -54445922.761 \text{ kJ/mol} \]

All the \( \Delta G \) were normalized to \( \Delta G \) Mer-mPAH as follows:

\[ \text{Normalized } \Delta G_{\text{Mer-mPAH}} = \Delta G_{\text{Mer-mPAH}} - \Delta G_{\text{Mer-mPAH}} = 0 \text{ kJ/mol} \]  

\[ \text{Normalized } \Delta G_{\text{Mer-mPA}^-} = \Delta G_{\text{Mer-mPA}^-} - \Delta G_{\text{Mer-mPAH}} = 74.4 \text{ kJ/mol} \]  

\[ \text{Normalized } \Delta G_{\text{SP-mPA}^-} = \Delta G_{\text{SP-mPA}^-} - \Delta G_{\text{Mer-mPAH}} = 87.8 \text{ kJ/mol} \]

Note that the energy of a solvated proton must be added to the Mer-mPA\textsuperscript{-} and SP-mPA\textsuperscript{-} forms in order to make all the Gibbs free energies comparable. The value for the Gibbs free energy for a solvated proton in aqueous media was obtained from Tissander et al seminal work on solvated ions\textsuperscript{11}. Unfortunately, there is no paper that follows Tissander et al procedure on calculating the Gibbs free energy of a solvated proton in ethanol via cluster calculations. Nonetheless, Markovic et al provide Gibbs free energy calculations for a solvated proton in ethanol (and many other...
solvents) using accurate calculations.\(^{12}\) Although their work does not feature a fully comprehensive solvation model for the proton, it does go on to show that the Gibbs free energies for solvated protons are not so far in water and in ethanol (-1055.8 kJ/mol and -1064.9 kJ/mol, respectively). Hence, we choose to approximate the solvation Gibbs free energy for a proton in ethanol, or in an ethanol-water mixture, by the highly accurate reported value by Tissander et al. It is important to recognize that the predicted energy for the Mer-mPA\(^-\) might have a significant error, since the model does not seem to be able to fully account for its electronic structure (refer to Figure C-8B).

For completion, we repeated the same process for an aqueous solution. All structures were optimized with B3LYP/6-311++G(2d,p) accounting for water as a solvent using the IEFPCM model. A frequency calculation using those same parameters was performed to obtain the ZPE-corrected Gibbs-free energies. The energy of the system (in water) in its different states was computed as follows:

\[
\Delta G_{\text{Mer-mpAH}} = ZPE_{\text{Mer-mpAH}} = -5446016.3 \text{ kJ/mol} \tag{C.1}
\]

\[
\Delta G_{\text{Mer-mpA}} = ZPE_{\text{Mer-mpA}} + \Delta G_{\text{Solvated H}^+} = \left(-5444833.3 \frac{\text{kJ}}{\text{mol}}\right) + \left(-1104.5 \frac{\text{kJ}}{\text{mol}}\right) = -5445937.8 \text{ kJ/mol} \tag{C.2}
\]

\[
\Delta G_{\text{SP-mpA}} = ZPE_{\text{SP-mpA}} + \Delta G_{\text{Solvated H}^+} = \left(-5444828.8 \frac{\text{kJ}}{\text{mol}}\right) + \left(-1104.5 \frac{\text{kJ}}{\text{mol}}\right) = -5445933.3 \text{ kJ/mol} \tag{C.3}
\]

All the \(\Delta G\) were normalized to \(\Delta G\) Mer-mPAH as follows:
Normalized $\Delta G_{Mer-mPAH} = \Delta G_{Mer-mPAH} - \Delta G_{Mer-mPAH} = 0 \text{ kJ/mol}$ \hfill [C.4]

Normalized $\Delta G_{Mer-mPA^-} = \Delta G_{Mer-mPA^-} - \Delta G_{Mer-mPAH} = 78.5 \text{ kJ/mol}$ \hfill [C.5]

Normalized $\Delta G_{SP-mPA^-} = \Delta G_{SP-mPA^-} - \Delta G_{Mer-mPAH} = 83.0 \text{ kJ/mol}$ \hfill [C.6]

C.3.2 Correlation between experimental and theoretical absorbance spectra

Figure C-8 Overlap of the normalized experimental absorption spectra (black line) and the shifter computed spectra (blue line; more details provided below) for (A) Mer-mPAH, (B) Mer-mPA⁻, and (C) SP-mPA⁻ forms.

The first 40 excited states for Mer-mPAH, Mer-mPA⁻, and SP-mPA⁻ forms were calculated using time-dependent DFT on the previously optimized structures. The scales for the theoretical Mer-mPA⁻ spectra seem disproportionate because there is a significant population of the mPAH in its SP-mPA⁻ conformer (evidenced by the peak at 287 nm in the Mer-mPA⁻ experimental spectra). If the experimental spectra contained solely Mer-mPA⁻, the scales would be in better agreement. For example, refer to the spectra of the SP-mPA⁻. The experimental spectrum shows no mPAH in its
Mer-mPA$^-$ conformer (peak at 554 nm) and a negligible amount of mPAH in its Mer-mPAH conformer (peak at 474 nm). As a result, the scales of the theoretical and experimental spectra are much closer.

The spectra in Figure C-8A were normalized to the peak corresponding to the Mer-mPAH form in the experimental spectrum in 100% ethanol. The theoretical spectra required a shift of -0.221 eV to match the experimental first electronic transition (from 437.91 nm to 474.91 nm). A FWHM of 0.21 eV was required to provide the best possible overlap.

The spectra in Figure C-8B were normalized to the peak corresponding to the Mer-mPAH form in the experimental spectrum in ethanol/water (50/50). The theoretical spectra required a shift of -0.282 eV to match the experimental first electronic transition (from 492.21 nm to 554.21 nm). A FWHM of 0.10 eV was required to provide the best possible overlap.

The spectra in Figure C-8C were normalized to the lowest energy electronic transition of the experimental spectrum in 100% ethanol. The theoretical spectra for SP-mPA$^-$ required a shift of 0.178 eV to match the experimental first electronic transition (from 300.24 nm to 287.23 nm). A FWHM of 0.33 eV was required to provide the best possible overlap.

The overlaps are good except in the case of Mer-mPA$^-$ form (Figure C-8B), where the transition corresponding to a shoulder in the experimental spectra is predicted to be at a considerable distance from the transition of lowest energy. As a result, the theoretical spectrum creates a new peak instead of a shoulder. This could be linked to the FWHM that is quite small in this case. A larger
FWMH should merge these two peaks and lead to a better agreement. The inversion of the relative intensities of the two lowest transitions could also be an artifact due to a too crude model (B3LYP/6-311++G (2d,p)) or to the use of ethanol as the only solvent. Further investigations are required to give more insight on these issues.

C.4 Determination of thermodynamic acidity constant

The ground state acidity constant \( pK_a \) of metastable-state photoacid (mPAH) is determined by the following thermodynamic reaction provided below.

\[
\text{mPAH} \quad \xrightarrow{\text{Base}} \quad \text{mPA}^- + \text{H}^+
\]

Likewise, Henderson-Hasselbalch equation \([C.7]\) was correlated with degree of protonation \([C.8]\). Accordingly, mass-balance equation \([C.9]\) was used to modify Equation \([C.7]\) to \([C.10]\).

Henderson-Hasselbalch equation based of mPAH equilibrium reaction:

\[
pH = pK_a + \log\frac{[m^-]}{[mH]}
\]

Degree of protonation, “\(\alpha\)”: \[C.8\]

\[
\alpha = \frac{[mPAH]}{[mPAH]_T} = \frac{A-A_D}{A_P-A_D}
\]
Degree of protonation is defined as the ratio of acidic-open form mPAH, where subscript “T” signifies total concentration of mPAH. Likewise, the concentrations of acidic-open form mPAH can be related to absorbance as denoted by “A”. The symbols “AD” and “AP” represents absorbance values of acidic-open form mPAH absorption peak when it is fully deprotonated and fully protonated.

**Mass-balance equation:**

\[
[mPAH]_T = [mPAH] + [mPA^-] \quad \text{[C.9]}
\]

**Modified Henderson-Hasselbalch equation:**

\[
pH = pK_a + \log \left( \frac{(1-\alpha)}{\alpha} \right) \quad \text{[C.10]}
\]

This theoretical equation was derived by rearranging [C.7] by incorporation of both [C.8] and [C.9], and was used to fit the experimental for estimating the pKa of Mer-mPAH.
Figure C-9 Correlation between theoretical fitting and the obtained experimental data to estimate the thermodynamic acidity constant.
C.5 Additional Figures and Table

Figure C-10 Modulation of pH by 470 nm light over multiple cycles with respect to time for an ethanol/water (90/10) solution containing 65 µM of Mer-mPAH.
Figure C-11 Emission spectra for ethanol solution containing Mer-mPAH (6.50x10^{-5} M) only.

Excitation wavelength 478 nm was used.
Figure C-12 Optical characteristics for different concentrations of BODIPY in ethanol. A) Absorbance spectra. B) Emission spectra (excitation wavelength was 495 nm) with calibration plot to illustrate self-quenching behaviour (Inset).
Figure C-13 Absorbance-based calibration plot for ethanol solutions containing different concentrations of A) Mer-mPAH (1.30x10^{-6} M, 2.60x10^{-6} M, 5.21x10^{-6} M, 1.30x10^{-5} M, 2.60x10^{-5} M, 3.90x10^{-5} M, 5.20x10^{-5} M and 6.50x10^{-5} M), and B) BODIPY (1.04x10^{-9} M, 1.04x10^{-8} M, 5.21x10^{-8} M, 1.04x10^{-7} M, 5.21x10^{-7} M, 1.04x10^{-6} M, 1.04x10^{-6} M, 1.56x10^{-6} M, 2.08x10^{-6} M, 2.60x10^{-6} M, 5.21x10^{-6} M, 1.04x10^{-5} M, 1.56x10^{-5} M and 2.08x10^{-5} M).
Figure C-14 Lifetime decay profiles for ethanol solutions containing fixed concentration of BODIPY (2.60x10^{-6} M) with different concentrations of Mer-mPAH.
Table C-1 Lifetime decay data for different concentration of Mer-mPAH with fixed concentration of BODIPY.

<table>
<thead>
<tr>
<th>[Mer-\textit{m}PAH] + 2.6x10^{-6} M BODIPY</th>
<th>( \tau ) (ns)</th>
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<tr>
<td>0 M</td>
<td>3.317 [a]</td>
</tr>
<tr>
<td>1.3x10^{-6} M</td>
<td>3.328</td>
</tr>
<tr>
<td>2.6x10^{-6} M</td>
<td>3.321</td>
</tr>
<tr>
<td>5.2x10^{-6} M</td>
<td>3.320</td>
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<tr>
<td>6.5x10^{-5} M</td>
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</table>

[a] \( \tau_0 \)
Figure C-15 Absorbance-based experiments for ethanol solution containing A) $6.50 \times 10^{-5}$ M of Mer-mPAH, B) $6.50 \times 10^{-5}$ M of Mer-mPAH with $2.60 \times 10^{-6}$ M of BODIPY (in green: absorbance spectra of $2.60 \times 10^{-6}$ M of BODIPY), and C) Kinetic plots for a solution of $6.50 \times 10^{-5}$ M Mer-mPAH only (in black, $\lambda_{\text{abs}} = 478$ nm), $2.60 \times 10^{-6}$ M BODIPY with $6.50 \times 10^{-5}$ M Mer-mPAH (in red, $\lambda_{\text{abs}} = 478$ nm), and $2.60 \times 10^{-6}$ M BODIPY only (in green, $\lambda_{\text{abs}} = 500$ nm) (ON region signifies visible light activation by 470 nm light, while the OFF region signifies in the dark).

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