Design and Synthesis of
Organic Molecular Models of Artificial Photosynthetic Reaction Center

by

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A Dissertation Presented in Partial Fulfillment
of the Requirements for the Degree
Doctor of Philosophy

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ABSTRACT

A clean and sustainable alternative to fossil fuels is solar energy. For efficient use of solar energy to be realized, artificial systems that can effectively capture and convert sunlight into a usable form of energy have to be developed. In natural photosynthesis, antenna chlorophylls and carotenoids capture sunlight and transfer the resulting excitation energy to the photosynthetic reaction center (PRC). Small reorganization energy, $\lambda$ and well-balanced electronic coupling between donors and acceptors in the PRC favor formation of a highly efficient charge-separated (CS) state.

By covalently linking electron/energy donors to acceptors, organic molecular dyads and triads that mimic natural photosynthesis were synthesized and studied. Peripherally linked free base phthalocyanine (Pc)-fullerene ($C_{60}$) and a zinc (Zn) phthalocyanine-$C_{60}$ dyads were synthesized. Photoexcitation of the Pc moiety resulted in singlet-singlet energy transfer to the attached $C_{60}$, followed by electron transfer. The lifetime of the CS state was 94 ps. Linking $C_{60}$ axially to silicon (Si) Pc, a lifetime of the CS state of 4.5 ns was realized. The exceptionally long-lived CS state of the SiPc-$C_{60}$ dyad qualifies it for applications in solar energy conversion devices. A secondary electron donor was linked to the dyad to obtain a carotenoid (Car)-SiPc-$C_{60}$ triad and ferrocene (Fc)-SiPc-$C_{60}$ triad. Excitation of the SiPc moiety resulted in fast electron transfer from the Car or Fc secondary electron donors to the $C_{60}$. The lifetime of the CS state was 17 ps and 1.2 ps in Car-SiPc-$C_{60}$ and Fc-SiPc-$C_{60}$, respectively.
In Chapter 3, an efficient synthetic route that yielded regioselective oxidative porphyrin dimerization is presented. Using Cu$^{2+}$ as the oxidant, meso-β doubly-connected fused porphyrin dimers were obtained in very high yields. Removal of the copper from the macrocycle affords a free base porphyrin dimer. This allows for exchange of metals and provides a route to a wider range of metallporphyrin dimers.

In Chapter 4, the development of an efficient and an expedient route to bacteriopurpurin synthesis is discussed. Meso-10,20- diformylation of porphyrin was achieved and one-pot porphyrin diacrylate synthesis and cyclization to afford bacteriopurpurin was realized. The bacteriopurpurin had a reduction potential of – 0.85 V vs SCE and $\lambda_{\text{max}}$, 845 nm.
DEDICATION

This dissertation is dedicated to my loving mum who despite all the challenges faced by an uneducated single mother in rural Northern Kenya ensured that I stayed in school and got a decent education.
ACKNOWLEDGMENTS

My research work and education would not have been possible without the support of a number of people. My first thanks goes to my advisor Prof. Devens Gust and collaborators Profs. Ana and Tom Moore. They kindly accepted me into their research group without reservation and guided me through the years of my graduate education and research. Also my research work would not have been possible if not for the talented and helpful members of Gust Moore\textsuperscript{2} research group. In particular I would like to thank: Dr. Paul Liddell for providing me with porphyrin starting material for the bacteriopurpurin project and much invaluable advice and suggestions on synthetic procedures, Dr. Smitha Pillai for her advice on phthalocyanine synthesis and for providing me with the Wittig reagents for the carotene-phthalocyanine-fullerene triad project, Dr. Gerdenis Kodis and Robert Schmitz for their photophysical expertise, Dalvin Mendez, Dr. Ben Sherman and Dr. Maxime Fournier for their electrochemical work, Dr. Mathieu Koepf, Dr. Yuichi Terrazono and Dr. Graeme Copley for many useful suggestions on synthesis and advice during my candidacy oral exam, Katie WongCarter for proof reading and editing my dissertation and many great suggestions. Many thanks to DOE and DARPA for funding.

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<td>BAHA</td>
<td>Tris(4-bromophenyl)aminium hexachloroantimonate</td>
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<td>BBr$_3$</td>
<td>Boron tribromide</td>
</tr>
<tr>
<td>BF$_3$·OEt$_2$</td>
<td>Boron trifluoride diethyl etherate</td>
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<td>BINAP</td>
<td>2,2'-bis(diphenylphosphino)-1,1'-binaphthyl</td>
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<td>C$_{60}$</td>
<td>Fullerene</td>
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<td>2,3-Dichloro-5,6-dicyano-1,4-benzoquinone</td>
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List of Abbreviations

DMAE 2-(dimethylamino)ethanol
DMF N, N-dimethylformamide
DMSO Dimethylsulfoxide
DSCC Dye Sensitized Solar Cells
Et$_3$N Triethylamine
EtOAc Ethyl acetate
Fc Ferrrocene
fs femtosecond
H$_2$Pc Metal-free phthalocyanine
H$_2$SO$_4$ Sulfuric acid
HBr Hydrobromic acid
HCl Hydrochloric acid
HOMO Highest Occupied Molecular Orbital
IBX Iodoxybenzoic acid
InCl$_3$ Indium trichloride
K$_3$PO$_4$ Tripotassium phosphate
KOH Potassium hydroxide
LUMO Lowest Unoccupied Molecular Orbital
MALDI-TOF Matrix Assisted Laser desorption Ionization- Time of flight
MeOH Methanol
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<td>milligram</td>
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<td>mL</td>
<td>milliliter</td>
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<td>NADPH⁺</td>
<td>Nicotinamide adenine dinucleotide phosphate</td>
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<td>NaOMe</td>
<td>Sodium methoxide</td>
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<td>NBS</td>
<td>N-Bromosuccinimide</td>
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<td>NIR</td>
<td>Near infrared</td>
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<tr>
<td>nm</td>
<td>nanometer</td>
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<td>NMR</td>
<td>Nuclear Magnetic Resonance</td>
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<tr>
<td>o-DCB</td>
<td>o-Dichlorobenzene</td>
</tr>
<tr>
<td>Pc</td>
<td>Phthalocyanine</td>
</tr>
<tr>
<td>Pd(PPh₃)₂Cl</td>
<td>Bis(triphenylphosphine)palladium(II) dichloride</td>
</tr>
<tr>
<td>POCl₃</td>
<td>Phosphorous oxychloride</td>
</tr>
<tr>
<td>ppm</td>
<td>parts per million</td>
</tr>
<tr>
<td>PRC</td>
<td>Photosynthetic Reaction Center</td>
</tr>
<tr>
<td>PS II</td>
<td>Photosystem II</td>
</tr>
<tr>
<td>Pt</td>
<td>Platinum</td>
</tr>
<tr>
<td>Q</td>
<td>Quinone</td>
</tr>
<tr>
<td>rpm</td>
<td>revolutions per minute</td>
</tr>
<tr>
<td>Ru</td>
<td>Ruthenium</td>
</tr>
<tr>
<td>SCE</td>
<td>Saturated calomel electrode</td>
</tr>
</tbody>
</table>
### List of Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>SiO₂</td>
<td>Silica</td>
</tr>
<tr>
<td>TBAPF₆</td>
<td>Tetrabutylammonium hexafluorophosphate</td>
</tr>
<tr>
<td>TBPPM</td>
<td>Trans-2-[3-(4-tert-butylphenyl)-2-methyl-2-propenylidene]-malonitrile.</td>
</tr>
<tr>
<td>TFA</td>
<td>Trifluoroacetic acid</td>
</tr>
<tr>
<td>TiO₂</td>
<td>Titanium dioxide</td>
</tr>
<tr>
<td>TLC</td>
<td>Thin Layer Chromatography</td>
</tr>
<tr>
<td>TW</td>
<td>Terrawatt</td>
</tr>
<tr>
<td>UV</td>
<td>Ultraviolet</td>
</tr>
<tr>
<td>ΔG</td>
<td>Change in free energy</td>
</tr>
<tr>
<td>λ</td>
<td>Reorganization energy</td>
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1.1 INTRODUCTION

One of the biggest challenges facing mankind in this century is access to affordable and clean energy. With a very high population growth expected to reach 9.6 billion by 2050 and rapid industrialization being experienced in most parts of the world, the demand for energy will be astronomical and finding a reliable source of energy will be a matter of urgency. In the year 2012, the rate of global energy consumption was 16.3 TW. Going by the current rate of energy consumption, the global annual energy demand will be at least 20 TW by 2030, doubled by mid-century, and tripled by the turn of the century.

Currently 85% of the total global energy consumed comes from burning fossil fuels; for developing countries 90% of their energy comes from fossil fuels. Over-reliance on fossil fuels presents energy security, national security, environmental security, and economic security issues which need to be addressed within the next 10-20 years. At the current rate of fossil fuel consumption, the atmospheric CO$_2$ concentrations within the 21st century will be more than double the pre-anthropogenic values. Against this backdrop of challenges associated with this energy source, there is an urgent need for energy rapid diversification away from fossil fuels and switching to carbon neutral energy sources.

Nuclear fission is a carbon-neutral alternative. It is estimated that existing terrestrial uranium (U) resources are capable of producing 100 TW of electricity using the conventional once-through U reactor technology. At the present global energy consumption rate of 16.3 TW/year, these terrestrial U resources will be exhausted in six years. Besides, use of nuclear energy has associated safety concerns after accidents such as Fukushima in 2011 and Chernobyl
in 1986; in addition nuclear energy presents an additional challenge of waste disposal of used nuclear fuels.

A promising alternative to the fossil fuels that can meet a rapidly increasing global demand for energy is the nearly infinite and inexhaustible solar energy. The energy from the sun that reaches our planet is 100,000 TW/yr. This means that the energy from one hour of sunlight is equivalent to all the energy currently used by mankind in a year.\(^6\) Photovoltaic technologies have been utilized to capture sunlight and produce electricity, and their efficiency and robustness are rapidly improving.\(^8,9\) However, in terms of price, the photovoltaic system currently in use is more costly than fossil based fuels.\(^10\) One approach to realizing a long-term solution to addressing the energy challenge lies in the development of solar conversion systems that are modeled on the principles of the natural photosynthetic systems.

**Natural Photosynthesis**

Natural photosynthesis is a process where sunlight energy is converted into organic molecules of biomass such as carbohydrates.\(^6\)
Figure 1 shows how solar energy is utilized in nature. At the light reactions part of photosynthesis, light absorption, charge separation, water splitting, and electron/proton transfer take place. The dark reaction uses these reducing equivalents from the light reactions in the form of energized electrons and a proton gradient to convert carbon dioxide into carbohydrates (CH$_2$O) and useful organic molecules which makes up biomass, including those that provide humanity with food. Fossil fuels that we burn to power our technology are a product of photosynthetic reactions that happened millions of years ago. Respiration and combustion of biomass are the reverse of photosynthesis, releasing CO$_2$, H$_2$O and energy. It can be concluded that photosynthesis is a way of storing solar energy in the form of chemical bonds.

Natural photosynthesis utilizes two photosystems per electron to drive water splitting and carbon dioxide fixation. The energy of two ‘red’ photons is required for every electron/proton extracted from H$_2$O and used to reduce CO$_2$. The two photosystems are: photosystem II (PSII)
where light is used to split water and release electrons and protons, and photosystem I (PSI), where light is used to provide additional reductive potential energy to the electrons from PSII so as to reduce carbon dioxide via NADPH.

Figure 2 is a schematic representation of how photosynthesis uses PSII and PSI to capture sunlight and convert it to chemical energy. PSII is a very powerful oxidant; in fact, P680$^+$ is the most oxidizing species in living systems and is capable of extracting electrons from water due to its potential at 1.2 V Vs SCE. $^{12}$ To produce one molecule of oxygen, PSII accumulates four oxidizing equivalents that are needed to abstract four reducing equivalents (4e/4H$^+$) from two water molecules (scheme 1).

$$2 \text{H}_2\text{O} + 4h\nu \rightarrow 4\text{H}^+ + 4\text{e}^- + \text{O}_2$$

Scheme 1: Overall water oxidation reaction by P680
Figure 2: A simplified Z-scheme of the light reactions of photosynthesis.\textsuperscript{11}

On the reducing side, PSII has pheophytin (Pheo) and quinone molecules as electron acceptors. The redox potential of Pheo\textsuperscript{2-} (-0.5 V\textsuperscript{6}) is sufficiently negative and in turn it passes the reducing equivalent along an electron transfer chain to PSI (figure 2). In PSI, a second ‘red’ photon is absorbed by a chlorophyll molecule, P700, to attain a reducing potential of -1 V vs SCE or more.\textsuperscript{6} The electron donated by the excited P700 (P700\textsuperscript{*}) gets accepted by a 4Fe-4S center. The low-potential 4Fe-4S center reduces NADP\textsuperscript{+} to NADPH through ferredoxin. The reduced ‘hydrogen carrier’ NADPH is the cofactor used by enzymes to drive fixation of CO\textsubscript{2}. Natural photosynthesis can be used as a blueprint to design and develop highly efficient, artificial, molecular-based conversion of solar energy to chemical energy.

\textbf{Artificial photosynthesis}

Artificial photosynthesis refers to use of the underlying scientific principles of natural photosynthesis to harvest solar energy for electricity or fuel production. There are many
approaches that have been used to convert sunlight to another form of energy. For the purpose of this dissertation, we will restrict artificial photosynthesis to the use of molecular species to store energy from the sun in a charge separated state and eventually a useful fuel.\textsuperscript{13}

Carotenoid (Car)-porphyrin (P)-fullerene (C\textsubscript{60}) molecular triad (figure 3)\textsuperscript{14} dissolved in 2-methyltetrahydrofuran yielded a long-lived charge separated state after excitation of the porphyrin followed by electron transfer and charge shift.\textsuperscript{15}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{structure.png}
\caption{Structure of carotenoid-porphyrin-fullerene artificial reaction center.\textsuperscript{14}}
\end{figure}

On excitation, the porphyrin moiety yields the first excited state Car-\textsuperscript{1}P-C\textsubscript{60}, this decays by transferring its electron to the fullerene with a lifetime \(\tau\) of 32 ps and a quantum yield of 0.99 for Car-P\textsuperscript{**}-C\textsubscript{60}\. Here the molecule has converted the absorbed photon to electrochemical potential, but the back electron transfer to yield the ground state leading to energy wastage occurs with \(\tau = 3.3\) ns. This is a very short time making it difficult to accessing the redox potential. However, in the triad, a fast charge shift with rate constant of 125 ps competes favorably with charge recombination, yielding the long-lived charge separated state Car\textsuperscript{**}-P-C\textsubscript{60}\. with a lifetime of 57 ns and overall quantum yield of 0.95.\textsuperscript{15} A lifetime of 170 ns at ambient temperature and \(\Box 1\) \(\mu\)s at 77 K has been observed for a similar Car-P-C\textsubscript{60} triad.\textsuperscript{16} In the
the electron transfer step is fast enough to compete with other alternative deexcitation pathways. The forward electron transfer is favored by sufficient thermodynamic driving force and a relatively strong electronic coupling between the initial and final states.\textsuperscript{15} In the final state of the triad, Car\textsuperscript{**}-P-C\textsubscript{60}\textsuperscript{•–}, the charges are well separated and the coupling weak. This retards recombination even when the driving force is high. On the other hand, stepwise recombination is slow due to the endergonicity of the first step. Natural photosynthetic reaction centers also use a long-lived, energetic charge-separated state with a high quantum yield achieved by step-wise, short-range electron transfers.

Photosynthetic organisms use antennas to capture sunlight and transfer the resulting singlet excitation energy to reaction centres. Most natural systems use chlorophyll \textit{a} (figure 4) to efficiently capture the photons from sunlight and funnel it to the special pair.

![Molecular structure of chlorophyll \textit{a}](image)

Figure 4: Molecular structure of chlorophyll \textit{a}.\textsuperscript{17}
The chlorin ring is the chromophore that absorbs light, and is responsible for the oxidation-reduction activity of chlorophyll \( a \) pigment. The side chain referred to as phytol tail of the molecule is used to properly orient the pigment molecule when it binds protein. About 99\% of chlorophyll molecules in most photosynthetic membrane are involved in excitation energy-transfer reactions, serving the antennae role. Only 1\% or less of the remaining chlorophyll molecules are involved in the redox-active catalytic site where electron transfer reactions are initiated.\(^{18}\)

In artificial systems, porphyrins, analogues of chlorophyll, have been used as artificial antennae. Porphyrins are naturally occurring macrocyclic heteroaromatic conjugated compounds and they are synthetically easy to prepare and also more photostable than chlorophylls. They exhibit very intense absorption in the visible region, with a Soret band at 400-500 nm and weak Q bands at 550-650 nm spectral regions.\(^{19}\)

Chlorophyll \( a \) has very strong light absorption around 425 nm and 660 nm but exhibits weak absorption bands in the regions between these wavelengths. Natural photosynthetic systems use accessory antenna chromophores to harvest energy in regions where chlorophyll has poor light absorption. Carotenoid polyenes are examples of the some of the commonly used molecules.\(^{15}\)
Figure 5: Absorption spectrum of Chlorophyll $a$ and carotenoid species.\textsuperscript{20}

As shown in figure 5, carotenoids absorb sunlight around 470 and 500 nm, the blue and the green part of the solar spectrum where chlorophyll doesn’t absorb. Use of antennae can also benefit artificial photosynthetic systems. The primary electron donors in artificial reaction centers such as porphyrins and phthalocyanines do not absorb equally at all wavelengths and the overall efficiency of the solar radiation can be increased by incorporating antennas. Carotenoid antennas offer an additional benefit of photoprotection\textsuperscript{21,22} to the artificial system just as it does in the natural systems.

An effective way of using artificial photosynthesis to generate sustainable energy is by efficient use of the renewable solar energy to produce H$_2$ and fuels; particularly those fuels obtained by conversion of carbon dioxide to safe and easily storable liquid fuels are the most useful.\textsuperscript{23,24} A bio-inspired efficient and cost-effective artificial system such as solar fuel making
device should have the key steps of natural photosynthesis; capture of sunlight, long-lived charge separated state and energy transduction into H₂ or another fuel. Figure 6 shows the simplified general scheme of a solar fuel system.

![Simplified diagram of a general solar fuel system.](image)

The solar fuel assembly absorbs sunlight and a charge separated state is created. The resulting electrochemical potential is used by the photocatalyst at the anode to oxidize water. The produced electrons and protons are transported to the cathodic side. At the cathode, proton and electron recombination to form H₂ takes place in presence of a reduction catalyst. An advanced system will have a cathode that can chemsorb CO₂ and in the presence of electrons and protons catalytically covert it to fuels or valuable chemicals.

In a nutshell, an efficient artificial solar energy converting system must meet the following criteria: (i) light captured by antenna molecules and/or photosensitizers should result into excited state formation, (ii) the absorption of the sunlight must result in a fast transfer of electrons to the acceptor component, (iii) the transfer of electrons must be directional, (iv) the charge separated state should be long-lived enough for charge-shift or electrochemical reaction.
to occur, (v) photoprotection capability and/or repair mechanism should be included for long term usage.

1.2 Artificial Photosynthetic Reaction center models

When covalently linked, supramolecular systems such as donor-linker-acceptor (D-L-A) are often used as photosynthetic models. In these systems, D is the light absorbing molecular unit, L is a covalent linker, and A is the component of the supramolecule that accepts light or electrons from D in the light induced process. After the donor absorbs light (scheme 2.1), the singlet excited state of the donor ($^1$D) is formed, and this excited state can relax back to the ground state through different relaxation pathways. Among them is the transfer of energy to the acceptor (scheme 2.2) or transfer of the excited electron to the acceptor to form a charge separated state (scheme 2.3). A schematic representation of these relaxation pathways is shown in Scheme 2.

\[
\begin{align*}
D-L-A + h\nu \rightarrow & \quad ^1D-L-A \quad \text{photoexcitation} \quad (2.1) \\
^1D-L-A \rightarrow & \quad D-L-^1A \quad \text{electronic energy transfer} \quad (2.2) \\
^1D-L-A \rightarrow & \quad D^{**}-L-A^{**} \quad \text{electron transfer} \quad (2.3)
\end{align*}
\]

Scheme 2: Energy transfer and electron transfer pathways of de-excitation of the excited state.

**Energy transfer processes**

There are two energy transfer mechanisms through which transfer of excitation energy from the donor to the acceptor can occur.
Coulombic Mechanism

The coulombic mechanism also referred to as resonance, Förster or through space mechanism is a long range mechanism that can occur without presence of physical contact between donor and acceptor. This mechanism of energy transfer is efficient in systems in which the radiative transitions connecting the ground state and the excited state of each partner have high oscillator strength. The rate of energy transfer ($k_{\text{EnT}}$) in coulombic mechanism is given by the Förster equation (1)\(^{31}\)

$$k_{\text{EnT}} = 0.529\kappa^2 \Phi_D J / n^4 N \tau_D R^6$$  \hspace{1cm} (1)

Where $\kappa$ is the orientation factor which takes into account the directional nature of the dipole-dipole interaction, and $\kappa^2$ is 2/3 for random orientation, $\Phi$ is the quantum yield of fluorescence of the donor, $\tau$ is the lifetime of the excited state of the donor, $R$ is the distance in cm between donor and the acceptor, $J$ is the Förster overlap integral between the emission spectrum of the donor and the absorption spectrum of the acceptor, $n$ is the dielectric constant of the medium, and $N$ is the Avogadro’s number.

When a good spectral overlap integral is present and the photophysical properties are right, the energy transfer by this mechanism occurs efficiently over distances that exceed the molecular diameter.\(^{30}\) Large aromatic molecules can exhibit an efficient singlet-singlet energy transfer; this process is analogous to the one used by nature in the photosynthetic antenna system.\(^{32}\)
Exchange Mechanism

The rate constant for energy transfer by the exchange mechanism, also referred to as Dexter mechanism, is given by equation (2).

\[ k_{\text{EnT}} = K J \exp(-2R/L) \]  

Where \( K \) is related to the specific orbital interactions between the donor and the acceptor, \( J \) is the overlap integral between the emission spectrum of the donor and the absorption spectrum of the acceptor, and \( R \) is the distance between the donor and the acceptor relative to their Van der Waals radii, \( L \).

The exchange mechanism is a double electron transfer process; it involves the transfer of one-electron from the LUMO of the excited donor to the LUMO of the acceptor, and the remaining electron moves from the acceptor HOMO to the donor HOMO. This mechanism allows electrons to be transferred in the cases where excited states involved are spin forbidden in the usual spectroscopic sense. An example of an efficient exchange mechanism is that of triplet-triplet energy transfer in metal complexes from the lowest spin state.

Electron transfer processes

Photoinduced electron transfer is one of the pathways through which deexcitation of the excited state donor occurs via transfer of electrons to the acceptor. It’s in competition with other deactivation pathways such as fluorescence, intersystem crossing (ISC) to the triplet state, and internal conversion to the ground state. Electron transfer leads to a decrease in the lifetime of the excited singlet state of the donor and the fluorescence quantum yield.
Electron transfer processes is described by Marcus, Levich, Hush and Jortner theory. The rate of the electron transfer process is given by equation (3)

\[ k_{eT} = \left( \frac{\pi}{\hbar^2} \lambda \kappa_B T \right)^{1/2} |V|^2 \exp\left(-\frac{(\Delta G^0 + \lambda)^2}{4 \lambda \kappa_B T}\right) \]  

(3)

where \( \hbar \) is the Planck’s constant, \( \lambda \) is the nuclear reorganization energy, \( \kappa_B \) is the Boltzmann’s constant, \( V \) is the electronic coupling matrix element between the donor and the acceptor of the supramolecule, and \( \Delta G^0 \) is the standard free energy of the reaction. For a homogenous series of reactions a log \( k_{eT} \) versus \( \Delta G^0 \) forms a bell-shaped curve. Figure 7 shows the relationship between log \( k_{eT} \) versus \( \Delta G \).

![Figure 7: An example of a Marcus curve showing the normal and the inverted region](image)

As shown in figure 7 above, there are three distinct scenarios in the bell-shaped curve. In the normal region, \( -\Delta G \leq \lambda \), the rate of electron transfer increases with increasing driving force.
At the activationless region, the rate of electron transfer is optimum (\(- \lambda = \Delta G\)), in this region increase or decrease in the driving force does not cause much changes in the rate of the electron transfer. In the “inverted region,” the reaction is strongly exergonic (\(-\Delta G > \lambda\)) and increasing the driving forces leads to a decrease in the electron transfer rate.\(^{41}\)

The reorganization energy, \(\lambda\), is the sum of two independent reorganization energies: there are the “inner” (bond lengths and angles with the donor and acceptor partners) and “outer” (solvent reorientation around the reacting pair) nuclear modes. The outer reorganization energy is the predominant term in the electron transfer processes.\(^{30}\)

From figure 7 it can be deduced that the electron transfer rate in donor-acceptor systems with smaller \(\lambda\) is faster (small \(-\Delta G^0\)) and slower recombination (large \(-\Delta G^0\)) compared to those having larger \(\lambda\) values.\(^{42}\) The early events of natural photosynthetic centers (PRC) have near 100% efficiency and reduced energy-wasting recombination. This is achieved by shuttling electrons away a long distance from its point of origin and kinetically by employing a low reorganization energy (\(\lambda \sim 0.3\) eV) which forces back electron transfer into the Marcus “inverted” region.\(^{43}\) Artificial donor-acceptor systems that replicate these important properties has been the focus of research to design mimics of PRC.\(^{44}\) In artificial systems, fullerene has been used as an electron acceptor due to its superior electron accepting ability and low reorganization energy.

**Fullerenes**

Buckminsterfullerene, \(C_{60}\) was discovered by Kroto and co-workers in 1985.\(^{45}\) Owing to the high degree of symmetry and the arrangement of its molecular orbitals, \(C_{60}\) has a number of
interesting chemical and physical properties.\textsuperscript{46} $C_{60}$ is a superior electron acceptor, and its triply-degenerate low-lying LUMOs (1.5-2.0 eV above the HOMO) allow it to reversibly accept up to six electrons.\textsuperscript{47} Due to the remarkably small reorganization energies associated with reductions of $C_{60}$ and its derivatives, donor-acceptor systems containing fullerenes have been proposed as models for photosynthesis and as energy conversion systems.\textsuperscript{48-50}

Imahori and co-workers compared the electron transfer rate ($k_s$) and the charge recombination rate ($k_{CR}$) of the dyads shown in structure 1a and structure 1b. In structure 1a, benzoquinone (Q), a commonly used acceptor, was replaced with the $C_{60}$ as both of them have similar reduction potentials, the reduction potential for Q = -0.45 V vs. Ag/AgCl while that of $C_{60}$ = -0.65 V vs. Ag/AgCl. Also the semi-rigid amide bond between the electron acceptor and the porphyrin ensures a similar separation distance and nature of the linker between the redox pair.

Structures 1a and 1b above shows molecular structures of two D-L-A systems with a zinc porphyrin as the donor and it was observed that the rate of charge separation was faster using a fullerene than Q by a factor of six. However, the charge recombination rate in 1a was smaller by a factor of less than 1/25 compared with that of 1b. This superior property of $C_{60}$ over the two-dimensional acceptors has been attributed to the smaller reorganization energy of the $C_{60}$
compared to those of their two-dimensional counterparts. The small reorganization energy of $C_{60}$ pushes the photoinduced charge separated state to the top of the Marcus parabolic curve (figure 7) in the normal region, but forces the charge recombination downward into the inverted region. This phenomenon favors photoinduced electron transfer while retarding charge recombination.\textsuperscript{48}

2.1 Design and synthesis of phenylethynyl-bridged phthalocyanine-fullerene dyads

Phthalocyanines (Pcs) are structural analogues of porphyrins which have strong absorption in the visible and near-infrared regions. The stronger absorption at longer wavelengths is as a result of extended conjugation afforded by the peripheral benzene rings. Figure 8 shows the structure of a simple unsubstituted metal free phthalocyanine ($H_2Pc$) and the nomenclature of the positions on the fused benzene ring.

![Figure 8: Phthalocyanine (Pc) core structure and position nomenclature](image)

Pcs have very versatile macrocycles which makes it possible to place seventy different elemental ions in the central cavity, which can be used to tune the physical properties.\textsuperscript{51} Phthalocyanine solutions have a characteristic strong absorption between 670 and 690 nm (Q-
band) (Figure 9). There is also a strong absorption band in the UV region between 320 and 370 nm (B-band or Soret band).

Figure 9: UV/Vis spectrum of H$_2$Pc in chloronaphalene.$^{52}$

It is the Q-band that is responsible for the characteristic intense blue or sometimes blue-green color of the compound and also it is the one that is susceptible to substitution and to the Pc macrocycle environment.$^{53}$ The Q-band absorption is due to $\pi-\pi^*$ transition from the HOMO of $a_{1u}$ symmetry, to the LUMO of $e_g$ symmetry.$^{54}$ This leads to a doubly degenerate first excited state of $^1E_u$ symmetry. The split in the Q-band usually observed in H$_2$Pc is as a result of lower symmetry (D$_{2h}$) compared with those of planar metal Pcs (D$_{4h}$) which leads to the loss of degeneracy of the LUMO orbital to produce $Q_y$ and $Q_x$.$^{53}$ Also, the position of the Q-band in the absorption spectrum of Pcs is sensitive to the central metal ion. Distinct metal-ligand charge
transfer (MLCT) bands arising from excitation from the atomic orbitals of the metal ion to the ligand’s molecular orbitals or *vica versa* have been observed. The effect on the optical properties of Pcs by substitution on the fused benzene ring depends on the position of substituents. α- Substitution greatly influences the energy levels of the molecular orbitals and hence the absorption spectrum. For example, it has been shown that electron-donating groups such as alkoxy moieties cause a 70 nm red-shift of the Q-band of CuPc substituted at the α-position by raising the energy level of the HOMO. On the other hand, with exception of substitutions that results in an extension of the π-orbital, most β-substitution has very little effect on the Q-band’s position. Due to these interesting properties phthalocyanines have been extensively investigated for their promising electrical, optoelectronics, photochemical and catalytic properties.

Pc-fullerene dyads with the electron donor and the electron acceptor covalently linked is an important approach to artificial photosynthesis and molecular electronic devices. Donor-fullerene dyad systems that exhibit a fast photoinduced charge separation (CS) with a fairly long lifetime of the charge separated state was initially observed in porphyrin-fullerene dyads and several of those types of dyads has been prepared. However, connecting Pc donor and fullerene acceptor has several advantages over porphyrin-fullerene dyads. Unlike porphyrins, which have poor absorption in the red part of the visible light, Pcs have strong light absorption in the red region of the solar spectrum (figure 9) and this allows for efficient conversion of solar energy in organic solar cells by collecting a larger section of the available sunlight. In addition, the
relatively large driving force for photoinduced electron transfer favors charge-separated state even in non-polar solvents.\textsuperscript{59}

A number of peripherally connected Pc-fullerene dyads have been synthesized in the last few years.\textsuperscript{60-63} These dyads are known to have an open conformation where the fullerene is located aside the Pc macrocycle rather than on top of it \textsuperscript{59} and it has been concluded that the electron transfer in these dyads take place through space rather than through bonds.\textsuperscript{60,61} The general trend in these type of donor-acceptor units are that the alignment is edge-to-edge and the charge recombination is faster than those that are face-to-face.\textsuperscript{59} The shorter the donor-acceptor linkage, the shorter the recombination time. Quintiliani and co-workers observed that the lifetime of charge-separated state is shorter in the dyads that lack a linker between the ZnPc and the C\textsubscript{60} moiety (figure 10).\textsuperscript{64}

![Figure 10: ZnPc-C\textsubscript{60} dyad](image-url)
The role of the donor-acceptor spacer in dyads are not only structural, it is also involved in the electronic communication between the terminal units. The modular composition of the spacer is also important as it allows one to alter the separation without affecting the electronic nature of the linkage. Of the linkers studied, acetylene linkers have shown the longest lifetime of the radical ion-pair state. It is postulated that a longer linker that has phenyl-ethynyl spacer can lead to fast charge separation and slower recombination. Towards this goal phthalocyanine-C₆₀ dyads separated by phenyl-ethynyl linkers were synthesized and the electrochemical and photochemical properties studied.

Results and Discussion

Synthesis

Phthalonitrile 2 was synthesized by Sonogashira coupling of 4-iodophthalonitrile with 4-ethynylbenzyl alcohol in the presence of [Pd(PPh₃)₂Cl] and CuI as the catalyst (scheme 3). Silica column purification of the crude using ethyl acetate/dichloromethane (1:9) solvent afforded compound 2 in 93% yields. It was observed that higher yield is favored by use of increased amount of catalyst.

![Scheme 3: Synthesis of 4-(2-(4-(Hydroxymethyl)phenyl)ethynylphthalonitrile 2](image-url)
As shown in scheme 4 below, unsymmetrical Pc 3 was obtained by a statistical condensation reaction between phthalonitrile 2 and a commercially obtained 4-tert-butylphthalonitrile in N, N-dimethylaminoethanol (DMAE) heated to reflux in the presence of Zn(OAc)$_2$.2H$_2$O. Thin-layer chromatography (TLC) of the crude product showed two main products: the less polar symmetrically substituted tetra(tert)butylphthalocyaninatozinc (II) side product and the desired asymmetrically substituted Pc 3. Due to the significant polarity difference between the two products separation was easily achieved by using silica column chromatography to yield Pc 3, 17%. This approach was preferred over the alternative method where tri-tert-butyl-iodophthalocyanine get reacted in a Sonogashira coupling reaction with 4-ethynylbenzyl alcohol to afford compound 3, because the condensation reaction to prepare tri-tert-butyl-iodophthalocyanine yields three products which are all very similar in polarity hence difficult to separate, and also the target product was recovered at a lower yield of 10%.

![Scheme 4: Synthesis of Tri(tert)butyl(benzylalcohol-4-ethynyl)phthalocyaninatozinc (II) 3](image)

The hydroxyl functionality on the unsymmetrically substituted Pc 3 was oxidized to a formyl group by use of 2-iodoxybenzoic acid (IBX) in dimethyl sulfoxide/tetrahydrofuran
mixture as the reaction solvent at room temperature to yield Pc 4, 79% (scheme 5). MALDI-TOF (terthiophene): m/z: 874.49

Scheme 5: Tri (tert)butyl(benzaldehyde-4-ethynyl)phthalocyaninato (II) 4

Synthesis of ZnPc-C₆₀ dyad 5 from ZnPc 4 was achieved by a 1, 3-dipolar cycloaddition of azomethine ylide generated in situ in a Prato reaction. Scheme 6 shows the reaction between ZnPc 4 and fullerene, C₆₀ in the presence of sarcosine in refluxing toluene to yield a 53% of dyad 5 after silica column purification with toluene/THF (49:1). MALDI-TOF MS (dithranol): [M]⁺: m/z: 1620.48.
Scheme 6: Synthesis of ZnPhthalocyanine-C$_{60}$ dyad 5

Scheme 7 below shows the synthesis of H$_2$Pc-C$_{60}$ dyad 7 and its precursor, H$_2$Pc 6. Heating Pc 4 in HCl-Pyridine/pyridine mixture at 120 °C for 24 hours afforded H$_2$Pc 6 in a quantitative yield of 98% without the need for a column purification.
Scheme 7: Synthesis of Tri (tert)butyl(benzaldehyde-4-ethynyl)phthalocyanine 6 and H₂Phthalocyanine-C₆₀ dyad 7

Then, a 1,3-dipolar cycloaddition (Prato reaction) between H₂Pc 6 and fullerene C₆₀ in the presence of sarcosine was carried out to obtain H₂Pc-C₆₀ dyad 7, 69% yield, after silica gel column purification with toluene/carbon disulphide (1/1) as the solvent. MALDI-TOF (dithranol matrix) showed a single peak, [M]^+: m/z: 1558.8.
De-zincation of hydroxyl substituted unsymmetrical ZnPc 3 model was achieved by heating it in pyridine-HCl/Pyridine at 120 °C for 21 hours (scheme 8) followed by adding water to the reaction mixture, then cooling, centrifugation and filtration to obtain a pure hydroxyl substituted unsymmetrical H₂Pc model 8, 38% yield. The yield here was far lower than the one obtained during dezincation of ZnPc 4; this may be attributed to the presence of the hydroxyl functionality on ZnPc 3 which may be interfering with the effectiveness of the pyridine-HCl reagent.

Scheme 8: Synthesis of Tri (tert)butyl(benzylalcohol-4-ethynyl)phthalocyanine 8

Scheme 9 shows the synthesis of fulleropyrrolidine phenol model 9 and its precursor, fulleropyrrolidine methoxyphenyl 10. A Prato reaction between 4-anisaldehyde and fullerene C₆₀ in the presence of sarcosine in refluxing toluene afforded compound 9, 60% after silica column purification with toluene/ethyl acetate solvent.
Scheme 9: Synthesis of N-methyl-2-(p-methoxyphenyl)-3, 4-fulleropyrrolidine 9 and N-methyl-2-(p-phenylalcohol)-3, 4-fulleropyrrolidine 10

Demethylation of compound 9 was done by reacting it with boron tribromide (BBr$_3$) at room temperature and the reaction crude was quenched by washing with water. The crude was purified by silica column chromatography with toluene/methanol (19/1) solvent system to afford N-methyl-2-(p-phenylalcohol)-3, 4-fulleropyrrolidine 10, 65% yield.

The structures of the molecules were confirmed by $^1$H NMR (CDCl$_3$), the precursor, hydroxyl substituted phthalonitrile 2 showed all the expected signals. All the 10 protons were accounted for, the single aromatic proton between cyano group and the alkynyl group appeared down field at 7.83 ppm, the two benzylic protons appeared at 4.68 ppm while the hydroxyl proton was a broad signal at 1.85 ppm. Due to the poor solubility of the phthalocyanines in chloroform, all the NMR measurements for all the phthalocyanine molecules were taken in d$_8$-THF. On condensation to form phthalocyanine 3, the hydroxyl proton was down field shifted to 4.35 ppm an indication that it is experiencing deshielding from the phthalocyanine aromatic system. Due to the presence of the regioisomers all the Pc aromatic protons appeared as broad signals. The single proton between the Pc macrocycle and alkynyl group was further shifted
downfield to 10.02 ppm, an indication that being sandwiched between the macrocycle and the triple bond deshields it more than the other Pc aromatic protons. The tert-butyl protons appeared up field between 1.73 and 1.85 ppm. Oxidizing Pc 3 to formyl-derivative phthalocyanine 4, a slight up field shift of most protons was observed. Formyl proton could be seen as a sharp singlet at 10.00 ppm. All the Pc and the phenyl aromatic protons appeared as poorly resolved multiplets. In the ZnPc-C₆₀ dyad 5, the signals of the Pc protons were not significantly impacted, while those on the functionalized fullerene were when compared to those of fullerene model compound 10. The pyrolidine protons signals were shifted down field from 4.97-4.23 ppm in the model compound 10 to 5.06-5.01 ppm in dyad 5, while the N-CH₃ protons in the dyad 5 were up field shifted from 2.97 ppm in the model compound 10 to 2.86 ppm in the dyad 5. Generally the signals in the dyad were broad and poorly resolved which could be attributed to poor solubility of the dyad in d₈-THF, addition of few drops of carbon disulphide to the NMR solvent did not yield much improvement in signal resolutions.

In the free base dyad, H₂Pc-C₆₀ dyad 7, the same pattern as dyad 5 was observed where the Pc signals did not change much from those observed in the model compounds. However, notable differences were observed in the functionalized fullerene components, in H₂Pc-C₆₀ dyad 7, the N-CH₃ signals broadened and were shifted upfield to 2.29 ppm from 2.97 ppm in the model fullerene 10.
Steady-state Absorption

Figure 11 shows the UV/vis spectrum in THF of H$_2$Pc-C$_{60}$ dyad 7 and shows Q-band at 694, 672, 644 and 613 nm and a Soret band at 331 nm, while the Q-band of the H$_2$Pc model compound 8 was at 694, 667, 644, 611 nm and Soret band at 344 nm. For the ZnPc-C$_{60}$ dyad 5 the most red Q-band appeared at 685 nm and the Soret band at 350 nm and the Q-band of the ZnPc model compound 3 was at 683 nm and its Soret band at 350 nm (figure 12). These observations where the Q-bands in both Zn and metal free dyads were not significantly shifted compared to those of the respective models is an indicator that there is no significant ground state interactions between the electronic systems of the two chromophores.

![Figure 11: UV/Vis spectra of H$_2$Pc-C$_{60}$ 7 (solid line) and H$_2$Pc model 8 (dashed line), normalized at Q-band in THF.](image-url)
Figure 12: UV/Vis spectra of ZnPc model 3 (dashed line, - - -), ZnPc aldehyde 4 (dotted line, . . .), and ZnPc-C_{60} 5 (solid line, —) normalized at Pc Q-band in THF.

As shown in figure 12 above, the dyad 5 shows a weak absorption at 432 nm and this is a characteristic of the [6, 6] mono-adduct of C_{60}. The bands for H_{2}Pc 3 model and the H_{2}Pc-C_{60} dyad 7 are broader than those of their zinc counterparts, this can be attributed to the poor solubility of the metal free compounds and this causes aggregation. The better solubility of the zinc containing molecules can be attributed to the ability of the central metal to bind a fifth ligand\(^{66}\), such as the solvent. This binding affects both the optical and electronic transfer properties\(^{67}\), and could be useful in sensor applications.\(^{66}\) Both the dyads show higher absorption in the UV region than their respective models; this is because of the additional absorption by the fullerene moiety. The fulleropyrrolidine component has a high extinction coefficient at 310 nm, seen from the fullerene model 10 (spectrum not shown). The weak absorption at 703 nm of the
fullerene component was not seen in the dyads as it gets eclipsed by the strong Q-bands of the Pcs.

The Q-band of the metal free compounds are red-shifted by 7-8 nm when compared to those of their Zn counterparts, this observation is in agreement with HOMO-LUMO gap differences in these molecules as supported by the electrochemistry data (table 1). Using the oxidation and the reduction potential values the HOMO-LUMO gap were found to be 1.4 eV and 1.6 eV for the H₂Pc model 8 and ZnPc model 3 respectively. Oxidizing Pc alcohol 3 to the corresponding aldehyde 4, the Q-band was shifted from 683 nm to 687 nm, this red-shift can be attributed to the extended conjugation due to the additional formyl double bond which raises the HOMO and consequently reduce the HOMO-LUMO gap.

**Electrochemistry**

Table 1: Redox potentials of Pc-fullerene dyads and their respective models (Electrochemistry measurements were done by Dalvin Mendez)

<table>
<thead>
<tr>
<th>Molecule</th>
<th>Oxidation (V vs SCE)</th>
<th>Reduction (V vs SCE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>H₂Pc model 8</td>
<td>0.62 (160)</td>
<td>-0.78 (54)</td>
</tr>
<tr>
<td>H₂Pc-C₆₀ 7</td>
<td>0.74 (irr)</td>
<td>-0.46 (90)</td>
</tr>
<tr>
<td>C₆₀ model 10</td>
<td>-</td>
<td>-0.63 (80)</td>
</tr>
<tr>
<td>ZnPc-C₆₀ 5</td>
<td>0.54 (210)</td>
<td>-0.66 (74)</td>
</tr>
<tr>
<td>ZnPc model 3</td>
<td>0.59 (145)</td>
<td>-0.99 (90)</td>
</tr>
</tbody>
</table>

The first oxidation potential of the free base Pc model 8 is 0.62 V vs SCE as measured by cyclic voltammetry in benzonitrile solution. A model for ZnPc model 3 is oxidized at 0.54 V vs
SCE whereas the first reduction potential for a model fullerene 10 is -0.63 V vs SCE (table 1). Using these data, the energies of the charge separated states of dyads 5 and 7 in benzonitrile are as follows: $\text{H}_2\text{Pc}^{\bullet^+}\text{-C}_{60}^{-\bullet^+}$ and $\text{ZnPc}^{\bullet^+}\text{-C}_{60}^{-\bullet^+}$ are 1.25 eV and 1.17 eV, respectively. The free base Pc dyad retains higher energy of the absorbed photons.

The electrochemistry data are approximately a linear combination of the oxidation and reduction potentials of the model phthalocyanines and fullerene. This observation agrees with the absorption spectra and these are evidences that there is no strong perturbation of the chromophores due to the covalent linkages.

The energies of the excited singlet states are calculated using the frequency-domain average of the most red wavelength absorption and most blue wavelength emission. They were calculated for the free base phthalocyanine 8 (1.79 eV), the zinc phthalocyanine 3 (1.82 eV), and the fullerene 10 (1.76 eV) (figure 13).

![Figure 13: High energy states and possible interconversion pathways for ZnPc-C$_{60}$ dyad 5 following photoexcitation of the ZnPc chromophore.](image-url)
The energy of the charge-separated state is estimated from the cyclic voltammetric data of model compounds in benzonitrile. Solid arrow represents the main pathway while the dashed arrow represents a minor pathway.

From the spectroscopic and cyclic voltammetric measurements in benzonitrile, a polar solvent, the thermodynamic driving forces for photoinduced electron transfer from $\text{H}_2\text{I}^1\text{Pc-C}_{60}$ to $\text{H}_2\text{Pc}^{++}-\text{C}_{60}^{--}$ and $\text{ZnI}^1\text{Pc-C}_{60}$ to $\text{ZnPc}^{++}-\text{C}_{60}^{--}$ are -0.54 eV and -0.65 eV respectively. The rate constant for most electron transfer processes such as this one is expected to be dependent on the thermodynamic driving force, the electronic coupling, and the nature of the linker. For the ZnPc-C$_{60}$ dyad 5 and the free base Pc-C$_{60}$ dyad 7 under study here, the last two factors are not different. However, from the thermodynamic argument, it is expected that the rate of electron transfer is faster in dyad 5 as long as the process is in the normal region of the Marcus curve.

Figure 14: DFT calculated (gaseous phase) molecular orbitals of the ZnPc-C$_{60}$ dyad 5, the HOMO is localized on the Pc (top) moiety and the LUMO on the C$_{60}$ (bottom).
Figure 15: DFT-calculated (gaseous phase) molecular orbitals of the H$_2$Pc-C$_{60}$ dyad 7, the HOMO is localized on the Pc moiety (top) and the LUMO on the C$_{60}$ (bottom).

As shown in figures 14 and 15 above, the HOMO is exclusively on the Pc and the LUMO is on the C$_{60}$ in both dyads, this gives a further support to the conclusion that in the Pc-C$_{60}$ conjugates under study, Pc is the electron donor and the C$_{60}$ is the electron acceptor and that the molecular orbitals of the components of the dyads retain their individual properties.
Stead-state emission

Figure 16: Steady state emission spectra for H₂Pc model 7 (figure 16A) and ZnPc model 3 (figure 16B), excitation at 330 nm.

The emission wavelength for the models are, H₂Pc (figure 16A): 705, 719 and 781 nm (toluene) and 709, 721 and 784 nm (benzonitrile) and ZnPc (figure 16B): 704 and 773 nm (toluene) and 700 and 770 (benzonitrile). Just like as been observed in the absorption spectra, the emission spectra of the models also show that the free base Pc model emits at a longer wavelength.
Supporting the conclusion that free base Pc model has a smaller HOMO-LUMO gap than the ZnPc model. The emission spectra are mirror images of the absorption spectra, an indication that fluorescence is the major deactivation pathway for the excited singlet state Pc.

In figure 17 the fluorescence intensity is quenched several orders of magnitude in the polar solvent, benzonitrile, compared to the fluorescence intensity in the non-polar solvent.

Figure 17: Emission spectra of the H$_2$Pc-C$_{60}$ dyad 7 (top) and ZnPc-C$_{60}$ dyad 5 (bottom) in toluene and benzonitrile after excitation at 345 nm wavelength.
toluene. The solvent dependent quenching in the dyads means that electron-donor acceptor interactions are involved in the excited-state deactivation of the photoexcited singlet state Pc. This observation is a preliminary evidence that electron transfer from the excited state Pc to C\textsubscript{60} is the predominant deexcitation pathway in polar solvents. Polar solvents are able to stabilize the charge separated state and hence lowering its energy, thus, thermodynamically favoring charge separation as can be seen from the respective $\Delta G_{CS}$ values of the dyads in table 2.

**Energetics**

Table 2: Energies (eV) of the dyads in benzonitrile

<table>
<thead>
<tr>
<th>Feature</th>
<th>ZnPc-C\textsubscript{60}</th>
<th>H\textsubscript{2}Pc-C\textsubscript{60}</th>
</tr>
</thead>
<tbody>
<tr>
<td>$1^*\text{Pc}$</td>
<td>1.82</td>
<td>1.79</td>
</tr>
<tr>
<td>$\text{Pc}^{••}-\text{C}_{60}^{•-}$</td>
<td>1.17</td>
<td>1.25</td>
</tr>
<tr>
<td>$-\Delta G_{CS}$</td>
<td>0.65</td>
<td>0.54</td>
</tr>
<tr>
<td>$-\Delta G_{CR}$</td>
<td>1.17</td>
<td>1.25</td>
</tr>
</tbody>
</table>

As shown in the table 2 above, the energies of the charge-separated states lie well below those of the three singlet excited states in benzonitrile, $1^*\text{ZnPc}$ (3) (1.82 eV), $1^*\text{H}_2\text{Pc}$ (8) (1.79 eV) and $1^*\text{C}_{60}$ (1.76 eV). This favorable driving force ($-\Delta G_{CS}$) suggests a strongly exothermic electron transfer in both cases to form $\text{ZnPc}^{+}-\text{C}_{60}^{•-}$ and $\text{H}_2\text{Pc}^{••}-\text{C}_{60}^{•-}$. 


**Time-resolved studies**

To investigate the dominant pathways of the deactivations in the Pc-C₆₀ as a result of electron-donor-acceptor interactions time-resolved studies were conducted.

**Transient Emission**

Table 3: Fluorescence lifetimes (ns) measurements of Pc models and Pc-C₆₀ dyads after excitation at 330 in toluene and benzonitrile. (Measurements were taken by Robert Schimtz)

<table>
<thead>
<tr>
<th>Molecule</th>
<th>Toluene</th>
<th>Benzonitrile</th>
</tr>
</thead>
<tbody>
<tr>
<td>ZnPc (3)</td>
<td>2.4</td>
<td>2.3</td>
</tr>
<tr>
<td>H₂Pc (8)</td>
<td>6.5</td>
<td>6.3</td>
</tr>
<tr>
<td>ZnPc-C₆₀ (5)</td>
<td>0.11 (51%)</td>
<td>□ 0.02 (90%)</td>
</tr>
<tr>
<td></td>
<td>1.7 (22%)</td>
<td>0.1 (9%)</td>
</tr>
<tr>
<td></td>
<td>2.18 (27%)</td>
<td>2.3 (1%)</td>
</tr>
<tr>
<td>H₂Pc-C₆₀ (7)</td>
<td>0.22 (59%)</td>
<td>0.08 (55%)</td>
</tr>
<tr>
<td></td>
<td>2.0 (39%)</td>
<td>0.2 (44%)</td>
</tr>
<tr>
<td></td>
<td>6.5 (2%)</td>
<td>2.3 (1%)</td>
</tr>
</tbody>
</table>

As shown by data in table 3 above, both ZnPc 3 and H₂Pc 8 models have solvent independent fluorescence lifetime. The lifetimes of Pc excited state in both dyads were significantly reduced, although the reduction is much greater in benzonitrile than in toluene. In the non-polar solvent, toluene, there is a presence of a characteristic emission of the \(^1C₆₀\), in dyad 5 it is 1.7 ns (22 %) and in 7 it is 2.0 ns (39 %). In benzonitrile, for
both dyads there is no evidence of $^1C_{60}$ emission, it appears if does exist, it get quenched very fast.

In both solvents, the lifetime of the excited state Pc is shorter in ZnPc-C$_{60}$ 5 than in H$_2$Pc-C$_{60}$ 7. These observations in addition to the solvent dependent quenching of dyads observed in steady fluorescence can be used to conclude the existence of energetically favorable fast intramolecular electron transfer processes from locally excited Pc to the C$_{60}$. The long-lived minor component in the free base dyad 7 which makes up 2% and 1% of the decay in toluene and benzonitrile respectively is most likely an impurity

**Transient absorption**

To test the nature and dynamics of the charge-separated radical ion-pair state of ZnPc-C$_{60}$ conjugate, femtosecond-resolved transient absorption spectroscopy was utilized. Due to the observed aggregation phenomenon H$_2$Pc-C$_{60}$ dyad 7 was not studied. In figure 18 transient absorption spectra for ZnPc-C$_{60}$ 5 are gathered. There is a competing energy transfer from Pc singlet excited state to C$_{60}$ singlet excited state and formation of ZnPc$^{\bullet\bullet}$-C$_{60}$$^{\bullet}$ CS state with 1.3 ps. After that there is slower (with 10 ps) formation of ZnPc$^{\bullet\bullet}$-C$_{60}$$^{\bullet}$ CS state. The CS state due to ZnPc$^{\bullet\bullet}$-C$_{60}$$^{\bullet}$ decays in 94 ps. The longer-lived CS state is attributed to the presence of the phenylethynyl linker.
Figure 18: differential absorption spectra (visible) obtained upon femtosecond flash photolysis (695 nm) of 5 in benzonitrile with several time delays at RT (Experiment was done by Dr. Gerdenis Kodis)

Torres and co-workers reported shorter CS state lifetime for ZnPc-C_{60} conjugates with no linker (figure 10) or with just an ethynyl linker (figure 19).

Figure 19: Ethynyl-linked ZnPc-C_{60} dyad
The lifetime (τ) of the CS state for the structure in figure 10 was reported to be 32 ps and that of the structure in figure 19 was 36 ps. The driving force (-ΔG\textsubscript{CR}) for the CS for 5 and for the structure in figure 10 is 1.19 eV and 1.17 eV respectively, therefore it cannot be the main factor favoring the longer lifetime of the CS state of 5. It can be concluded that for ZnPc-C\textsubscript{60} conjugates with similar charge-recombination (CR) driving force, the lifetime of CS state is significantly influenced by the linker.

**Experimental section**

**Synthesis**

4-(2-(4-(Hydroxymethyl)phenyl)ethynylphthalonitrile 2

Thirty mL of Et\textsubscript{3}N was placed in a round-bottom flask immersed in ice-water and degassed with argon for 40 min. A mixture of 4-ethynylbenzyl alcohol (126 mg, 0.953 mmol) and 4-iodophthalonitrile (219 mg, 0.862 mmol) was dissolved in the degassed and cold Et\textsubscript{3}N (30 mL). This was followed by addition of [Pd(PPh\textsubscript{3})\textsubscript{2}Cl\textsubscript{2}] (63 mg, 0.0898 mmol) and CuI (21 mg, 0.11 mmol). The reaction mixture was stirred for 15 hrs at 70°C under argon atmosphere. The solvent was removed under reduced pressured and the crude product purified by flash column chromatography on silica gel CH\textsubscript{2}Cl\textsubscript{2}/EtOAc (9:1) to yield 207 mg (93%) of 2 as a white solid.

\(^1\)H-NMR (CDCl\textsubscript{3}, 400 MHz) δ\textsubscript{H} 7.83 (s, 1H), 7.75-7.69 (m, 2H), 7.48-7.46 (d, J=8, 2H), 7.35-7.33 (d, J=8, 2H), 4.68 (s, 2H), 1.85 (br s, 1H)
**Tri (tert)butyl(benzylalcohol-4-ethynyl)phthalocyaninatozinc (II) 3**

A mixture of 4-tertbutylphthalonitrile (715 mg, 3.88 mmol), phthalonitrile 2 (200 mg, 0.775 mmol), and Zn(OAc)$_2$.2H$_2$O (825 mg, 3.76 mmol) in 30 mL DMAE was stirred and heated at 130°C for 24 hrs. Then the mixture was cooled to room temperature, the solvent was removed and the residue was washed with MeOH/H$_2$O (5:1) and the dark green crude obtained was purified by flash column chromatography on silica gel, toluene/THF (9:1). The blue symmetrical Pc was eluted first, followed by the greenish blue Pc 3 119 mg, 17%

$^1$H-NMR (d$_8$-THF, 400 MHz) $\delta$H, ppm 10.02 (s, 1H), 9.45-9.20 (m, 8H), 8.33-8.27 (m, 3H), 7.86-7.72 (m, 4H), 4.71 (s, 2H), 4.35 (br s, 1H), 1.85-1.73 (m, 27H)

UV/Vis (THF): $\lambda_{\text{max}}$ 683, 611 and 350 nm

MALDI-TOF (terthiophene): calc. for C$_{53}$H$_{46}$N$_8$OZn: [M]$^+$: m/z: 874.37, found 874.75.

**Tri (tert)butyl(benzaldehyde-4-ethynyl)phthalocyaninato (II) 4**

A mixture of Pc 3 (20 mg, 0.0229 mmol), IBX (20 mg, 0.0714 mmol) in THF (3 mL)/DMSO (3 mL) was stirred under argon atmosphere at room temperature for 48 hrs. Stirring was stopped and 10 mL of brine was added to the mixture followed by extraction with ether. Solvent was removed from the organic layer under reduced pressure and crude purified by column chromatography toluene/THF (19:1) to yield a green Pc 4, 16 mg, 79%.

$^1$H-NMR (d$_8$-THF, 400 MHz) $\delta$H, ppm 9.99 (s, 1H), 9.40-9.28 (m, 4H), 9.22-911 (m, 4H), 8.22-8.15 (m, 4H), 7.96-7.86 (m, 4H), 1.75-1.62 (m, 27H)

UV/Vis (THF): $\lambda_{\text{max}}$ 687, 672, 611, 352 nm

MALDI-TOF (terthiophene): calc. for C$_{53}$H$_{44}$N$_8$OZn: [M]$^+$: m/z: 872.29, found 874.49.
**Phthalocyanine-C$_{60}$ dyad 5**

A mixture of Pc 4 (15 mg, 0.0172 mmol), C$_{60}$ (41 mg, 0.0569 mmol), sarcosine (11 mg, 0.123 mmol) in an anhydrous toluene (25 mL) was stirred at reflux temperature for 22 hrs under argon atmosphere. After this time the mixture was cooled and solvent removed under reduced pressure. The crude was purified by a column chromatography in silica gel, toluene/THF (49:1) to obtain the ZnPc-C$_{60}$ dyad 5 as a dark-blue powder 15 mg, 53%.

$^1$H-NMR (d$_8$-THF, 400 MHz) $\delta_H$ , ppm: 9.41-9.49 (m, 4H), 9.31-9.23 (m, 4H), 8.31-8.22 (m, 4H), 7.98 (br s, 2H), 7.87 (br s, 2H), 5.06-5.01 (m, 3H), 4.28-4.25 (d, $J=12$, 2H), 2.86 (s, 3H, N-CH$_3$), 1.94-1.76 (m, 27H)

UV/Vis (THF): $\lambda_{max}$ 685, 672, 632,610, 350 nm

MALDI-TOF MS (dithranol): calc. for C$_{115}$H$_{49}$N$_9$Zn: [M]$^+$: m/z: 1620.34, found 1620.48

**Tri (tert)butyl(benzaldehyde-4-ethynyl)phthalocyanine 6**

Tri(tert)butyl(benzaldehyde-4-ethynyl) phthalocyaninatozinc(II) 4 (100 mg, 0.115 mmol), pyridine (10 mL) and pyridine-HCl (0.3 g) were stirred under argon for 24 hrs. After this time, stirring and heating was stopped and to the hot mixture H$_2$O (10 mL) was added. The mixture was allowed to cool and resulting precipitate was collected by centrifugation at 3,250 rpm. The bluish green precipitate was washed several times with water, MeOH followed by drying on high vacuum to obtain 91 mg of Pc 5, a 98% yield.

$^1$H-NMR (d$_8$-THF, 400 MHz) $\delta_H$ , ppm: 10. 05 (s, 1H), 8.70-7.19 (m, 16H), 1.71-1.88 (m, 27H)

UV/Vis (CH$_2$Cl$_2$): $\lambda_{max}$ 693, 677, 646, 620, 338 nm

MALDI-TOF MS (terthiophene): calc. for C$_{53}$H$_{46}$N$_8$: [M]$^+$: m/z: 810.38, found 810.55
Phthalocyanine-C$_{60}$ dyad 7

A solution of tri(tert)butyl(benzaldehyde-4-ethynyl) Pc 6 (20 mg, 0.0247 mmol), C$_{60}$ (59 mg, 0.0819 mmol), sarcosine (15.8 mg, 0.177 mmol) in an anhydrous toluene (25 mL) was heated to reflux with stirring under argon atmosphere for 22 hrs. After this time, solvent was removed under reduced pressure and the crude product purified by column chromatography toluene/CS$_2$ (1:1) to obtain a green product 26.7 mg, 69 %.

$^1$H-NMR (d$_8$-toluene, 400 MHz) $\delta$H , ppm 9.14-8.65 (m, 9H), 7.96-7.68 (m, 7H), 4.77 (s, 1H), 4.67-4.64 (d, $J$=12, 1H), 3.99-3.97 (d, $J$=8, 1H), 2.62 (s, 3H, N-CH$_3$), 1.66-1.63 (m, 27 H), 2.29 (br s, 2H)

UV/Vis (THF): $\lambda_{max}$ 694, 672, 644, 613, 343 nm

MALDI-TOF MS (dithranol): calc. for C$_{115}$H$_{51}$N$_9$: [M]$^+$: m/z: 1558.43 found 1558.79

Tri(tert)butyl(benzylalcohol-4-ethynyl)phthalocyaninato 8

Tri(tert)butyl(benzylalcohol-4-ethynyl) phthalocyaninatozinc(II) 3 (36 mg, 0.412 mmol), pyridine (10 ml) and pyridine-HCl (0.089 g) were heated at 120 °C while stirring under argon for 21 hrs after which to the hot mixture H$_2$O (5 mL) was added. The mixture was allowed to cool and resulting precipitate was collected by centrifugation at 3,250 rpm. The green precipitate was washed several times with water, MeOH followed by drying on high vacuum to obtain 12.7 mg of Pc 6, a 38% yield.

$^1$H-NMR (d$_8$-toluene, 400 MHz) $\delta$H , ppm

UV/Vis (THF): $\lambda_{max}$ 694, 667, 644, 611, 344 nm

MALDI-TOF MS (terthiophene): calc. for C$_{53}$H$_{48}$N$_8$O: [M]$^+$: m/z: 812.39 found 812.44
**N-methyl-2-(p-methoxyphenyl)-3, 4-fulleropyrrolidine 9**

A mixture of C\(_{60}\) (216 mg, 0.3 mmol), 4-anisaldehyde (20 mg, 0.15 mmol), and sarcosine (136 mg, 1.5 mmol) in toluene (60 mL) was warmed to reflux under argon environment while stirring for 26 hrs. Solvent was removed under reduced pressure. The crude was purified on silica column chromatography, toluene/EtOAc (99:1) to obtain a brown powder fullerene 9 (78.8 mg, 60% yield).

\(^1\)H-NMR (CDCl\(_3\), 400 MHz) \(\delta_H, \) ppm 7.69 (br, s, 2H), 6.95-6.93 (d, \(J=8, 2H\)), 4.97-4.95 (d, \(J=8, 2H\)), 4.87 (s, 1H), 4.25-4.21 (d, \(J=12, 2H\)), 3.80 (s, 3H, N-CH\(_3\)), 2.77 (s, 3H, O-CH\(_3\))

MALDI-TOF MS (terthiophene): calc. for C\(_{53}\)H\(_{48}\)N\(_8\)O: [M]\(^+\): m/z: 883.10 found 883.18

**N-methyl-2-(p-benzylalcohol)-3, 4-fulleropyrrolidine 10**

To a solution of fullerene 9 (32 mg, 0.0362 mmol) in toluene (20 mL) immersed in ice-cold water, 1.5 mL of 1M BBr\(_3\) (1.5 mmol) in dichloromethane was added drop wise. The mixture was stirred at room temperature for 24 hrs under an argon atmosphere. The mixture was transferred to a separating funnel and washed with water (30 mL×3) and then the solvent was removed under a reduced pressure. The crude was purified by a column chromatography on silica gel, toluene/MeOH (19:1) to obtain a brown fullerene 10 (20 mg, 65%). \(^1\)H-NMR (CDCl\(_3\), 400 MHz) \(\delta_H, \) ppm 7.65 (br, s, 2H), 6.85-6.83 (d, \(J=8, 2H\)), 4.97-4.95 (d, \(J=8, 1H\)), 4.86 (s, 1H), 4.25-4.23 (d, \(J=8, 2H\)), 2.973 (s, 3H)

UV/Vis (Toluene): \(\lambda_{max}\) 704, 433, 310 nm

MALDI-TOF MS (terthiophene): calc. for C\(_{69}\)H\(_{11}\)NO: [M]\(^+\): m/z: 869.08 found 869.15
2.2 **Synthesis and photophysical studies of a novel carotenoid silicon Pc-C\textsubscript{60} triad**

Due to the presence of 18 \(\pi\)-electron aromatic cloud, most Pcs have cofacial intermolecular \(\pi\)-\(\pi\) interactions, even at low concentrations, and this causes aggregation which leads to fluorescence quenching and poor solubility.\textsuperscript{70} The steric effect of the axial substituents on silicon phthalocyanines (SiPc) have been shown to exhibit a non-aggregation characteristics and increased solubility in common organic solvents.\textsuperscript{71}

Phthalocyanines with two symmetrical fullerene substituents connected axially through central silicon have been reported in the last few years.\textsuperscript{72-75} In addition to improving the solubility by diminishing intermolecular interactions, fullerenes bring additional properties to the phthalocyanine molecule. As mentioned earlier, fullerenes (C\textsubscript{60}) are good electron acceptors and have low reorganization energy; \(\lambda\).\textsuperscript{46}

C\textsubscript{60}-SiPc-C\textsubscript{60} triads are good models for artificial photosynthetic reaction centers. However, to develop a better mimic of natural photosynthesis, artificial photosynthesis will require a system with two electron donors and one acceptor where a long-lived charge separation can be achieved via a step wise electron transfer.\textsuperscript{76}

In natural photosynthesis the roles of carotenoids includes acting as an antennae by transferring excitation energy to the photosynthetic reaction center\textsuperscript{77} and protection by quenching chlorophyll triplet states to prevent generation of the cell damaging reactive singlet oxygen species.\textsuperscript{49,77} Excited state carotene has a high energy (~1.9 eV)\textsuperscript{78} and this allows it to act as antenna via singlet-singlet energy transfer, but the triplet state is much lower in energy (0.63 eV) than singlet oxygen (0.98 eV)\textsuperscript{79} and triplet chlorophyll (~1.3 eV)\textsuperscript{80} and these properties allow it
to be a good energy acceptor as well. Figure 20 shows some of the naturally occurring carotenoids that play a light-harvesting role in photosynthesis.

![Molecular structures of four important carotenoids.](image)

Figure 20: Molecular structures of four important carotenoids.\(^{81}\)

In addition, under stress conditions, for example in Mn-depleted PSII or at low temperature, β-carotene has been observed to be an electron donor to the highly oxidizing P680\(^+\).\(^{82}\)
Incorporating carotenoids into artificial photosynthetic reaction center may lead to a long-lived charge separated state by charge shift and provide photoprotection. Also use of carotenoids and phthalocyanines together will ensure access to a wider spectral window of solar radiation. Hence an asymmetrical axial linkage on SiPc with potential electron donor on one side and a potential electron acceptor on the other side will be a good mimic for natural photosynthesis. To this end, synthesis of carotene and fullerene axially connected to a SiPc was achieved. Electron transfer and energy transfer processes in the triad were investigated by use of spectroscopic and electrochemical methods.

**Results and Discussions**

**Synthesis**

Axially substituted phthalocyanine 11 was afforded by a reaction between a commercially available silicon phthalocyanine dihydroxide with 4-iodobenzoate and 4-formylbenzoate (1:1) as shown in scheme 10. The crude product contained a statistical mixture of diformylbenzoate, diiodobenzoate, and the target asymmetrically linked 4-formylbenzoate-4-iodobenzoate SiPc products. By column purification, the bluish green phthalocyanine 11 was obtained in 30% yield. MALDI-TOF (terthiophene matrix) showed the molecular ion peak at m/z=1160.3 as the 100 % peak, fragments were also observed at m/z=1011.33 (target molecule with the formylbenzoate cleaved off) and m/z=913.45 (target molecule with the iodobenzoate cleaved off). Jesse Bergkamp and co-workers reported a similar kind of fragmentation pattern with axial ether linkages on SiPc.83

Heating a mixture of phthalocyanine 11, sarcosine and C$_{60}$ in o-dichlorobenzene at 140°C afforded SiPc-C$_{60}$ dyad 12 as shown in scheme 11. Column purification gave a non-fluorescent green compound dyad 2, 40 % yield. MALDI-TOF (TBPPM matrix, mode-reflector 6000) showed the molecular ion peak at m/z=1909.2 as the only peak, no fragments beside the peak at m/z=720 for the unfunctionalized C$_{60}$ was observed.
Scheme 11: (p-iodobenzoate)-[p-(N-methyl-3′, 4′-fulleropyrrodin-2′-yl) benzoate] (2, 9, 16, 23-tetra-tert-butylphthalocyaninato) silicon 12.

Compounds 13 and 14 shown in scheme 12 were synthesized following a literature method. Reacting commercially obtained β-apocarotenal with 4-(N-acetylamino)benzyltriphenylphosphonium bromide (kindly donated by Dr. Smitha Pillai) in a Wittig type reaction in presence of sodium methoxide in dimethyl sulfoxide solvent gave 4-amidophenyl-β-carotene 13, 30% yield after column purification and crystallization over hexane/dichloromethane to remove the cis isomer.
Schemes 12: synthesis of \( (p' \text{-} 7' \text{-apo-7'} \text{-}(4\text{-aminophenyl}) \text{-}\beta \text{-carotene} \) 14.

4-Aminophenyl-\( \beta \)-carotene 13 was hydrolyzed using methanolic potassium hydroxide in THF at 65 °C to obtain 4-aminophenyl-\( \beta \)-carotene 14 in near quantitative yield of 94% after column purification.

Synthesis of carotene-SiPc-C\text{60} triad 15 was attempted by stirring a mixture of SiPc-C\text{60} dyad 12, 4-amino-\( \beta \)-carotene 14 in presence of a new commercially acquired tris(dibenzylidene acetone) dipalladium (0)/BINAP/sodium tert-butoxide (1:3:4) in freshly distilled and degassed toluene and heated at 80°C under argon environment (scheme 13). No product was realized after 7 hours of stirring. Increasing the temperature to 110 °C, addition of more catalysts and increase of the reaction time to 45 hours did not help to form a product. After stirring was stopped and the mixture cooled to room temperature, the reactants were partially recovered. It appears that the presence of C\text{60} may have interfered with the effectiveness of the palladium (0) catalyst.
Scheme 13: Attempted synthesis of \([p-7’-apo-7’-(4-aminophenyl)\-\(\beta\)-carotenobenzoate]-[\(p\)--(N-methyl-3’, 4’-fulleropyrrolo-2’-yl) benzoate] (2, 9, 16, 23-tetra-\(\text{tert}\)-butylphthalocyaninato) silicon 15.

Carotene-SiPc dyad 16 was synthesized by reacting SiPc 11 with 4-aminophenyl-\(\beta\)-carotene 14 in presence of a new commercially obtained catalyst/ligand/base mixture of tris(dibenzylidene acetone) dipalladium (0)/BINAP/sodium \(\text{tert}\)-butoxide-1:3:4 in freshly distilled and degassed toluene at 80 \(\text{C}\) (scheme 14), on column purification a yellowish green product, compound 16 was obtained, 33% yield. It was observed the yield was inversely...
dependent on the temperature; lowering the temperature increases the yield. Increasing the
temperature to 90°C, the yield decreased to 12%, at 95°C and the yield was 4% at 100°C there
was no detectable product formed. This is thought to be as a result of decomposition of the
reaction intermediates, this conclusion is supported by the presence of multiple unaccounted for
spots on the analytical TLC of the reaction crude and it was also observed that the number of
spots increased with increasing temperature.

MALDI-TOF (Tertiophene matrix) showed only a tiny peak of the molecular ion,
m/z=1537, the major peaks observed were the fragments, m/z=913.6, 100% intensity (target
product with 4-aminophenyl-β-carotenobenzoate cleaved off) and m/z=1389.0, 20% intensity
(target product with 4-formylbenzoate cleaved off).
Scheme 14: Synthesis of (p-7'-apo-7'-(4-aminophenyl)-β-carotenobenzoate)-(p-formylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 16.

As shown by scheme 15, carotene-SiPc-C_{60} triad 17 was afforded by a single Prato reaction from (p-7'-apo-7'-(4-aminophenyl)-β-carotenobenzoate)-(p-formylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 16 in 62% yield after purification by silica column chromatography.
Scheme 15: Synthesis of \([p-7'-apo-7'-(4-aminophenyl)-\beta\text{-}\text{carotenobenzoate}]-[p--(N\text{-}methyl\text{-}3', 4'\text{-}fulleropyrrodin\text{-}2'-yl) \text{ benzoate}]\) (2, 9, 16, 23-tetra-tert-butylphthalocyaninato) silicon 17.

MALDI-TOF (TBPPM matrix) showed the molecular ion peak at \(m/z=2284.5\) as the only peak, no fragments beside the peak at \(m/z=720\) for the unfunctionalized \(C_{60}\) was observed.

The model compound, SiPc 18 and compound 19 were prepared as shown in scheme 16. Reacting commercially obtained silicon phthalocyanine dihydroxide with 4-toluic acid/4-formyl benzoic acid mixture (1:1) gave SiPc 18 and SiPc 19 after column purification in 17% and 27% yield, respectively. MALDI-TOF (Tertiophene matrix) showed both the molecular ion peaks and the fragments peaks. For Pc 18, peaks were observed at \(m/z=1048.5\), which was the molecular ion peak, \(m/z=913.4\) (target molecule with the 4-methylbenzoate cleaved off) and \(m/z=899.5\) (target molecule with the 4-formylbenzoate cleaved off). For Pc 19, peaks were observed at...
m/z=1034.5, which was the molecular ion peak and m/z=899.4 (target molecule with the 4-methylbenzoate cleaved off).

Scheme 16: Bis(p-methylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 18 and (p-formylbenzoate)-(p-methylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 19.

To obtain SiPc-\(C_{60}\) dyad 20, Pc 19 was reacted with \(C_{60}\) and sarcosine in \(o\)-dichlorobenzene (scheme 17). Column purification afforded a near quantitative yield of dyad 10, 94%.

MALDI-TOF (terthiophene matrix) showed the molecular ion peak at m/z=1795.4 as the only peak, no fragments beside the peak at m/z=720 for the unfunctionalised C₆₀ was observed.

As shown in scheme 18, C₆₀ model 21 was prepared by a Prato type reaction of 4-methylester benzaldehyde, C₆₀ and sarcosine in refluxing toluene. On column purification a yield of 14% of C₆₀ 21 was obtained. It was observed that no detectable product was obtained when the reaction was carried out in o-dichlorobenzene (o-DCB), this may be attributed to solubility difference of the reactants in toluene and o-DCB. Better solubility in the former may be favoring product formation.

Scheme 18: N-Methyl-2-(p-methylbenzoate)-3,4-fulleropyrrolidine 21.

The ¹H NMR (CDCl₃) spectrum of phthalocyanine 11 showed an up field shift of the signals of benzoate protons to the range 6.67-4.83 ppm as opposed to the expected region of around 8 ppm for phenyl aromatic protons. This observation confirms the influence of the shielding effect of the phthalocyanine π cloud. In addition, the presence of four distinct benzoate signals at 6.73, 4.58, 5.26 and 4.83 ppm gives further support to the presence of an asymmetric axial linkage on the phthalocyanine. On the other hand, attachment of the carotene to the
phthalocyanine to form dyad 16 had no effect on the resonance of either of the components, i.e. phthalocyanine and carotene. The $^1$H NMR (CDCl$_3$) spectrum of carotene-phthalocyanine dyad 16 was a sum of the spectrum of the two parts. This could be attributed to the presence of the two phenyl rings that separate the two components and hence protect the carotene from the shielding effect of the phthalocyanine aromatic electron cloud. In contrast, earlier work by our group showed that when a carotenoid is directly, axially attached to SiPc there was a very significant shift in the resonance of the carotene protons. On attaching the C$_{60}$ to form the triad 17, the up field shift of the benzoate protons and those protons of the pyrrolidine linker between the C$_{60}$ and the phthalocyanine were similar to the one reported by Martin-Gomis and co-workers. The signal of the benzoate protons adjacent to C$_{60}$, pyrrolidine protons and N-CH$_3$ protons were all shifted up field when compared to the corresponding protons of the C$_{60}$ model 21. Benzoate protons were shifted from 7.84 ppm to 5.17 ppm, the pyrrolidine protons from 4.97, 4.96, and 4.25 ppm were moved to 4.63, 4.28, and 3.84 ppm; and the N-CH$_3$ protons were shifted from 2.78 to 2.17 ppm. This observation suggests that the attached C$_{60}$ is in the range of the strong aromatic current of the Pc.

**Steady-state Absorption**

The UV/Vis absorption spectrum of the triad 17 and model compounds were taken in dichloromethane. It was observed that the absorption spectrum of triad 17 showed that the Q-band of the phthalocyanine is unperturbed on formation of the triad (figure 21); the addition of the carotenoid pigment causes a larger increase in absorption in the middle of the spectrum. Figure 22 shows that the carotenoid moiety band at 480 nm in carotene-Si Pc dyad 16 is red-
shifted compared to model 4-aminophenyl-β-carotene, which has a maximum absorbance at 475 nm. The carotene moiety absorption maximum is at 479 nm in triad 17, not a significant change from that observed in the carotene-Pc 16 dyad. However, the absorption spectrum of the SiPc-C_{60} dyad 20 in dichloromethane is a close superposition of the spectra of the component chromophores making up the dyad. This is an indication that there is not much electronic interaction between the individual chromophores at the ground state. In addition, the sharp Q-bands are indications that the silicon phthalocyanine is not aggregating. C_{60} has its intense absorption at 220, 265 and 330 nm\(^{42}\) and these bands are buried under the stronger Soret transition of the phthalocyanine and are only seen as an overall intensification of the absorption in the UV/Vis region.

![Normalized UV/Vis absorption spectrum of triad 17 (—), carotene-SiPc dyad 16 (- - - -) and SiPc model 18 (…..).](image)

Figure 21: Normalized UV/Vis absorption spectrum of triad 17 (—), carotene-SiPc dyad 16 (- - - -) and SiPc model 18 (…..).
The UV/Vis spectrum of SiPc 18 and SiPc 19 were exactly the same and yet at the para-position of the phenyl moiety on SiPc 18, is the electron donating methyl group while on one of the para-position of the phenyl moiety of SiPc 19 there is the electron withdrawing formyl group. Based on this observation it can be concluded that the functional group on the para-position of the axial phenyl substituent does not have any influence on the conjugation of the Pc core.

**Electrochemistry**

Electrochemistry measurements were performed by Dr. Maxime Fournier

**Table 4:** First reduction potentials of C₆₀ model 21 and C₆₀-SiPc dyad 20.

<table>
<thead>
<tr>
<th>Molecule</th>
<th>E_{red}^-</th>
</tr>
</thead>
<tbody>
<tr>
<td>18</td>
<td>-0.47</td>
</tr>
<tr>
<td>21</td>
<td>-0.56</td>
</tr>
<tr>
<td>20</td>
<td>-0.64</td>
</tr>
</tbody>
</table>
Table 5: Oxidation potentials of carotene model, carotene-SiPc dyad 16, SiPc model 18, C_{60}-SiPc dyad 20.

<table>
<thead>
<tr>
<th>Molecule</th>
<th>$E_{ox}^1$</th>
<th>$E_{ox}^2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carotene model</td>
<td>0.49</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>0.47</td>
<td>0.93</td>
</tr>
<tr>
<td>18</td>
<td>0.93</td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>1.10</td>
<td></td>
</tr>
</tbody>
</table>

The first charge separated state, Car-SiPc^•+^-C_{60}^•, lies at 1.48 eV (figure 25) above the ground state, therefore the charge recombination to reform the ground state is likely to lie in the inverted region of the Marcus curve, where an increase in the driving force past the optimum leads to a decrease in electron transfer rate.\textsuperscript{35,86,87} This phenomenon is expected to favor the charge shift towards the carotene moiety.

Electron transfer rates are determined by thermodynamics, steric and electronic effects. In the present triad, an initial electron transfer from excited singlet SiPc to C_{60}, followed by a charge shift to the carotenoid moiety to form a long-living charge separated state (C^•+^-SiPc-C_{60}^•) is thermodynamically favorable.
Steady-State Emission
Figure 23 shows the steady-state emission spectra of Car-SiPc-C_{60} triad 17 taken in toluene and benzonitrile after excitation at 363 nm.

![Emission spectra of Car-SiPc-C_{60} triad 17 in toluene and benzonitrile after excitation at 363 nm](image)

Figure 23: Emission spectra of Car-SiPc-C_{60} triad 17 in toluene and benzonitrile after excitation at 363 nm

Time-resolved Emission

The steady-state emission spectra of triad 17 shown in figure 23 indicate that there is solvent-dependence quenching, an indirect indication of existence of electron transfer in polar solvent. To get an insight into the nature of fluorescence, time resolved fluorescence experiments were carried out using single-photon-counting technique (Experiments were done by Robert Schimtz). The results are summarized in Table 6
<table>
<thead>
<tr>
<th>Compound</th>
<th>Toluene</th>
<th>Benzonitrile</th>
</tr>
</thead>
<tbody>
<tr>
<td>SiPc 18</td>
<td>7.4</td>
<td>6.43</td>
</tr>
<tr>
<td>SiPc-C_{60} 20</td>
<td>0.035 (65%)</td>
<td>~0.035 (52%)</td>
</tr>
<tr>
<td></td>
<td>1.69 (26%)</td>
<td>0.490 (46%)</td>
</tr>
<tr>
<td></td>
<td>2.62 (9%)</td>
<td>4.84 (2%)</td>
</tr>
<tr>
<td>Car-SiPc-C_{60} 17</td>
<td>0.030 (74%)</td>
<td>0.020 (84%)</td>
</tr>
<tr>
<td></td>
<td>0.303 (8%)</td>
<td>0.430 (13%)</td>
</tr>
<tr>
<td></td>
<td>1.32 (14%)</td>
<td>5.56 (3%)</td>
</tr>
<tr>
<td></td>
<td>3.66 (4%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 6: Lifetimes (ns) of excited singlet states of Pc moieties as determined from time-resolved fluorescence studies in toluene and benzonitrile. The excitation wavelength was 635 nm.

As was observed with peripheral ZnPc 3 and H_{2}Pc 8 models, SiPc 18 model also shows similar lifetime of excited state in both toluene and benzonitrile. The fluorescence lifetimes of model SiPc 18 in toluene and benzonitrile were 7.4 and 6.43 ns respectively. The lifetime of excited Pc is significantly reduced in dyad 20 in both solvents but roughly by the same magnitude, this solvent independent quenching may be due to energy transfer from the locally excited SiPc to C_{60}. In toluene for dyad 20, there is presence of characteristic {1}C_{60} emission (1.69 ns, 26 %) and this is missing in benzonitrile. The same phenomenon was observed in the triad

63
where the characteristic $^1\text{C}_{60}$ emission occurs with 1.32 ns (14%) lifetime and this is again absent in benzonitrile, and it can be concluded that in benzonitrile, either $^1\text{C}_{60}$ doesn’t form or it get quenched very fast as soon as it forms.

**Time-resolved absorption studies**

Pump-probe transient absorption experiments were carried out to characterize the various excited species in the dyad models and the triad and to allow detection of nonemissive species such as charge-separated states. Experiments were performed by Dr. Gerdenis Kodis.

The occurrence of electron transfer from the excited state Pc to the C$_{60}$ acceptor in the SiPc-C$_{60}$ dyad 20 was confirmed by femtosecond laser flash photolysis experiment. A degassed benzonitrile solution containing SiPc-C$_{60}$ dyad 20 was subjected to 100 fs at 800 nm laser pulses. The resulting transients were monitored in the visible and the near-IR region (figure 24). Distinct characteristic absorption bands due to SiPc$^{\bullet⁺}$ (555 nm and 880 nm)$^{88,89}$ and C$_{60}^{\bullet⁻}$ (1000 nm)$^{90,91}$ were observed. This data is in agreement with that reported by Martin-Gomis and co-workers for C$_{60}$-SiPc-C$_{60}$.$^{73}$ This is a clear evidence of formation of charge-separated (CS) state. A long-lived band at $\lambda$=530 nm is a characteristic of $^3\text{SiPc}^\bullet$. $^{88}$

The presence of the long-lived $^3\text{SiPc}^\bullet$ is an evidence that the back electron transfer from the CS state forms the excited SiPc triplet state instead of the ground state because of the lower triplet energy (1.26 eV)$^{70}$ compared to the CS state (1.48 eV) determined from the redox potentials of the model compounds as shown in figure 25.
Figure 24: Differential absorption spectra (visible and near-infrared) of SiPc-C$_{60}$ dyad 20 upon femtosecond photolysis (695 nm) in deaerated benzonitrile.

Overall, there is energy transfer from Pc singlet excited state to form C$_{60}$ singlet excited state with ~24 ps. After that there is slow (with ~540 ps) formation of Pc$^+$-C$_{60}^-$ CS state, which is expected due to closeness in energy between Pc-1C$_{60}$ (1.75 eV) and Pc$^+$-C$_{60}^-$ (1.48 eV) (figure 25).
Figure 25: High energy states and possible interconversion pathways for SiPc-C$_{60}$ dyad following photoexcitation of the SiPc chromophore.

The energy of the charge-separated state is estimated from the cyclic voltammetric data of model compounds in benzonitrile.

The CS state decays in ~4.5 ns. Because of the slow formation of CS state there is substantial formation of C$_{60}$ triplet excited state which decays to Pc triplet excited state.

The charge recombination (CR) of the CS state to directly regenerate the ground state has a driving force of $\Delta G_{CR} = 1.48$ eV and this must be lying in the inverted region of the Marcus curve and this explains the long lifetime of the CS state and the decay first to the long-lived $^3$SiPc$^*$ and then to the ground state.
The decay associated spectrum of Car-SiPc dyad 16 dissolved in benzonitrile and excited at 695 nm is shown in figure 26. The excitation light at 695 nm is exclusively absorbed by the Pc moiety leading to a fast (with 3.3 ps) formation of Car\(^{+}\)-Pc\(^{-}\) CS state. This is evidenced by the presence of a near-IR (~980 nm) band, a characteristic absorption of a carotenoid radical species. Previous work from our group on carotene-SiPc-carotene triad where the carotene lacked the aminophenyl linker showed a carotenoid radical at 860 nm.\(^9\) The bathochromic shift in the carotenoid radical transient of the current may be attributed to the extended conjugation which makes it a better electron donor by raising the HOMO.

The transient in the 550-640 nm range also rises and decays with same time constant as the carotenoid radical. The huge negative amplitude at ~690 nm also recovers in the same time constant and is associated with the ground state bleaching of SiPc moiety. The match of the rise of the carotenoid radical cation transient with the decay of SiPc S\(_1\) level (3.3 ps) is an evidence of electron transfer from the carotenoid to the excited \(^1\)SiPc\(^{+}\) state as a dominant fluorescence quenching mechanism.
Figure 26: Decay associated spectra of the transient absorption of Car-SiPc dyad \textbf{16} in benzonitrile after a 100 fs laser pulse at 695 nm.

As shown on figure 26, the CS state forms after 3.3 ps and decays in 13 ps. There is some minor dyad conformation which shows much slower CS state recombination lifetime of \sim 940 ps. There is small impurity of SiPc that has a singlet excited state that does not decay on time scale measured. The fast CS and CR is in agreement with the large driving force for both processes, $\Delta G_{\text{CS}} = 0.81$ and the $\Delta G_{\text{CR}} = 0.96$ (figure 27).
Figure 27: Deexcitation pathways for Car-SiPc dyad 16 following photoexcitation of the SiPc chromophore.

The energy of the charge-separated state is estimated from the cyclic voltammetric data of model compounds in benzonitrile. The ultrafast formation of $\text{Car}^\bullet^\text{-Pc}^\bullet^\text{-}$ CS state is in agreement with previous studies on Car-SiPc-Car triad which gave a value of 2.5 ps and 10.5 ps as the rate of CS formation and the lifetime of the CS respectively.\textsuperscript{85}
Figure 28 shows the decay associated spectrum (DAS) of Car-SiPc-C\(_{60}\) triad 17 dissolved in benzonitrile and excited at 695 nm.

![Graph of Car-Pc-C\(_{60}\) triad in BZN with excitation at 695 nm]

Figure 28: Decay-associated spectra of the transient absorption of Car-SiPc-C\(_{60}\) 17 in benzonitrile of 100 fs laser pulse at 695 nm

There is a fast (with ~2.5 ps) dominating formation of Car\(^{**}\)-Pc\(^{*}\) CS state. There is also possible minor pathway of energy transfer from Pc singlet excited state to C\(_{60}\) singlet excited which competes with formation of Car\(^{**}\)-Pc\(^{*}\) CS. The Car\(^{**}\)-Pc\(^{*}\) CS state decays in ~17 ps. There is some minor triad conformation which shows much slower CS state recombination lifetime of ~945 ps. Since a C\(_{60}\)\(^{*}\) transient has not been observed, this slowly decaying CS state is not likely to be Car\(^{**}\)-SiPc-C\(_{60}\)\(^{*}\). There is small impurity of Pc whose singlet excited state does
not decay on time scale measured. The DAS spectra are mixed; it possibly contains a few transient species and corresponds to a few processes. There is no long living Car radical cation signal in the IR, so there is no significant Car$^{\bullet^+}$-Pc$^-$$^\ddag$C$_{60}$$^\bullet$ final CS state forming in benzonitrile.

![Diagram](image)

Figure 29: Deexcitation pathways for Car-SiPc-C$_{60}$ dyad 17 following photoexcitation of the SiPc chromophore.

Dominant pathways are represented by full arrows and minor pathways are represented by broken arrows. The energy of the charge-separated state is estimated from the cyclic voltammetric data of model compounds in benzonitrile.

As shown in figure 29, the fast CS and CR involving Car$^{\bullet^+}$-SiPc$^\bullet$-C$_{60}$ is the most favored thermodynamically and this supports the data obtained from the transient absorption. The other possible explanation for the fast intramolecular transfer in the Car-SiPc components of the triad is the enhanced coupling that exists between the carotenoid and the phthalocyanine as evidenced
by the 5 nm red shift observed in $\lambda_{\text{max}}$ of carotene moiety after it was attached to the phthalocyanine.

![Graph showing HOMO-LUMO energies](image)

Figure 30: DFT calculated HOMO-LUMO energies of the components of the Car-SiPc-C$_{60}$ triad 17.

Figure 30 shows the HOMO-LUMO energies (calculation done by Dalvin Mendez) of the components making up the triad 17, the HOMO-LUMO gaps are: C$_{60}$ (1.86 eV), SiPc (2.11 eV) and carotene (2.16 eV). The calculations are in agreement with the experimental observations. Because of the relatively lowest level of the HOMO of the C$_{60}$, an alternative approach to realizing a long-lived Car$^\bullet^+$-SiPc-C$_{60}^\bullet$ CS state is by exclusive excitation of the C$_{60}$ chromophore which will in turn result into a flow of electrons by gradient.
Experimental Section

Synthesis of (p-iodobenzoate)-(p-formylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 11

4-Iodobenzoic acid (242 mg, 1.0 mmol), 4-formylbenzoic acid (150 mg, 1.0 mmol), and \((t\text{Bu})_4\text{SiPc(OH)}_2\) (92 mg, 0.115 mmol) in 25 mL anhydrous DMF were stirred at 165 °C under argon atmosphere for 10 hrs. The mixture was cooled to room temperature and solvent removed under reduced pressure and the crude purified by flash chromatography (SiO$_2$, dichloromethane/hexane 7:3) to yield 39 mg (30%) of SiPc 11 as a blue-green solid. $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ 9.74-9.56 (8H, m, Pc-Ar-H), 9.37 (1H, s, CHO), 8.45-8.43 (4H, m, Pc-Ar-H), 6.73 (2H, d, J=8 Hz, Ph-Ar-H), 6.58 (2H, d, J=8 Hz, Ph-Ar-H), 5.26 (2H, d, J=8 Hz, Ph-Ar-H), 4.83 (2H, d, J=8 Hz, Ph-Ar-H), 1.80-1.79 (36H, m, 4×(CH$_3$)$_3$). UV/Vis ($\lambda_{max}$nm, CH$_2$Cl$_2$): 694, 662, 624, 360. MALDI-TOF-MS (Terthiophene matrix) m/z calcd for C$_{63}$H$_{57}$IN$_8$O$_5$Si 1160.33 obsd 1160.35

Synthesis of (p-7'-apo-7'-(4-aminophenyl)-β-carotenobenzoate)-(p-formylbenzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 16

SiPc 11 (53 mg, 0.053 mmol), p-7'-apo-7'-(4-aminophenyl)-β-carotene (31 mg, 0.0608 mmol) 14, and 41 mg of a commercially obtained mixture of tris(dibenzylidene acetone) dipalladium (0): BINAP: sodium tert-butoxide (1:3:4) were stirred in a freshly distilled and degassed toluene (28 mL) at 80°C under argon atmosphere for 48 hrs until almost all of the starting material have been consumed. The solution was then allowed to cool to room temperature. The solvent was evaporated at reduced pressure and the crude purified by column
chromatography (SiO\textsubscript{2}, hexane/dichloromethane/EtOAc 8:1.5:0.5) to obtain 17.6 mg (31%) of the dyad 16. \textsuperscript{1}H NMR (400 MHz, CDCl\textsubscript{3}) \(\delta\) 9.68-9.51 (8H, m, Pc-Ar-H), 9.32 (IH, s, CHO), 8.39-8.37 (4H, m, Pc-Ar-H), 7.01-6.99 (2H, m, Ph-Ar-H), 6.68-6.66 (2H, m, Ph-Ar-H), 6.62-6.44 (8H, m), 6.30-6.25 (2H, m), 6.19-6.08 (6H, m), 5.75 (2H, d, J=8 Hz, Ph-Ar-H), 5.20 (2H, d, J=8 Hz, Ph-Ar-H), 5.13 (1H, br s, NH), 4.98 (2H, d, J=9 Hz, Ph-Ar-H), 1.97-1.87 (m, 14H), 1.74-1.64 (m, 36H), 1.58-1.53 (m, 4H), 1.41-1.38 (m, 3H), 0.97-0.96 (m, 6H). UV/Vis (\(\lambda_{\text{max}}/\text{nm, CH}_2\text{Cl}_2\)): 694, 661, 623, 508 (sh), 480, 450 (sh), 341. MALDI-TOF-MS (Terthiophene matrix) m/z calcd for C\textsubscript{100}H\textsubscript{103}N\textsubscript{9}O\textsubscript{5}Si 1538.79 obsd 1538.60

\textbf{Bis(\textit{p}-methylbenzoate)(2,9,16,23-tetra-\textit{tert}-butylphthalocyaninato)silicon 18 and (\textit{p}-formylbenzoate)-(\textit{p}-methylbenzoate)(2,9,16,23-tetra-\textit{tert}-butylphthalocyaninato)silicon 19}

A portion of 4-toluic acid (136 mg, 1.0 mmol), 4-formylbenzoic acid (150 mg, 1.0 mmol), and \((\text{t-Bu})_4\text{SiPc(OH)}_2\) (92 mg, 0.115 mmol) in 25 mL anhydrous DMF were stirred at 170 °C under argon atmosphere for 10 hrs. The mixture was cooled to room temperature and solvent removed under reduced pressure and the crude purified by flash chromatography (SiO\textsubscript{2}, dichloromethane/hexane 8:2). The symmetrical SiPc 18 was eluted first, yielding 20.5 mg (17%), followed by asymmetrical SiPc 19 (32. 7 mg, 27 %).

SiPc 18 \textsuperscript{1}H NMR (400 MHz, CDCl\textsubscript{3}) \(\delta\) 9.75-9.48 (8H, m, Pc-Ar-H), 8.35-8.33 (4H, m, Pc-Ar-H), 5.95 (4H, d, J= 8 Hz, Ph-Ar-H), 4.96 (4H, d, J= 8 Hz, Ph-Ar-H), 1.73-1.72 (36H, m, 4\(\times\) (CH\textsubscript{3}))\), 1.60 (3H, s, 2\(\times\)CH\textsubscript{3}). UV/Vis (\(\lambda_{\text{max}}/\text{nm, CH}_2\text{Cl}_2\)): 693, 662, 623, 361. MALDI-TOF-MS (Terthiophene) m/z calcd for C\textsubscript{64}H\textsubscript{62}N\textsubscript{8}O\textsubscript{4}Si 1034.47 obsd 1034.45.
SiPc 19 $^1$H NMR (400 MHz, CDCl$_3$) δ 9.69-9.50 (8H, m, Pc-Ar-H), 9.30 (1H, s, CHO), 8.37-8.35 (4H, m, Pc-Ar-H), 6.67 (2H, d, J= 8 Hz, Ph-Ar-H), 5.95 (2H, d, J= 8 Hz, Ph-Ar-H), 5.20 (2H, d, J= 8 Hz, Ph-Ar-H), 4.97 (2H, d, J= 8 Hz, Ph-Ar-H), 1.73-1.72 (36H, m, 4×(CH$_3$)$_3$), 1.60 (3H, s, CH$_3$). UV/Vis ($\lambda_{\text{max}}$/nm, CH$_2$Cl$_2$): 693, 662, 623, 361. MALDI-TOF-MS (Terthiophene) m/z calcd for C$_{64}$H$_{60}$N$_8$O$_5$Si 1048.45 obsd 1048.46

(p-methylbenzoate)-(p-(N-methyl-3’,4’-fulleropyrrolidin-2’-yl)benzoate)(2,9,16,23-tetra-tert-butylphthalocyaninato)silicon 20

SiPc 19 (32 mg, 0.0305 mmol), C$_{60}$ (55 mg, 0.0764 mmol), and sarcosine (30 mg, 0.337 mmol) in $o$-dichlorobenzene (12 mL) were stirred at 140 ºC under argon atmosphere for 2 hrs. The mixture was cooled and solvent removed under reduced pressure. The crude was purified on silica gel (toluene) yielding 52 mg (94%) of SiPc-C$_{60}$ dyad 20 as green solid. $^1$H NMR (400 MHz, CDCl$_3$) δ 9.73-9.55 (8H, m, Pc-Ar-H), 8.42-8.40 (4H, m, Pc-Ar-H), 6.68 (2H, br s, Ph-Ar-H ), 6.01 (2H, d, J=8 Hz, Ph-Ar-H), 5.23 (2H, br s, Ph-Ar-H), 4.59 (2H, d, J=8 Hz, Ph-Ar-H), 4.59 (1H, d, J=8 Hz, CHN), 4.25 (1H, s, CHN), 3.83 (1H, d, J=8 Hz, CHN), 2.17 (3H, s, N-CH$_3$), 1.79-1.78 (36H, m, 4×(CH$_3$)$_3$), 1.66 (3H, s, CH$_3$). ($\lambda_{\text{max}}$/nm, CH$_2$Cl$_2$):694, 663, 623, 360, 331. MALDI-TOF-MS (Terthiophene) m/z calcd for C$_{126}$H$_{65}$N$_9$O$_4$Si 1795.49 obsd 1795.35

N-Methyl-2-(p-methylbenzoate)-3, 4-fulleropyrrolidine 21

A mixture of C$_{60}$ (50 mg, 0.0694 mmol), methyl-4-formylbenzoate (30 mg, 0.183 mmol), and sarcosine (28 mg, 0.314 mmol) in toluene (10 mL) were heated to reflux under argon atmosphere for 3 hrs. The mixture was then cooled to room temperature and solvent removed and crude purified by flash chromatography (SiO$_2$, toluene:/EtOAc-19:1) yielding 9 mg (14%) of 21.
21 as a brown solid. $^1$H NMR (400 MHz, CDCl$_3$/CS$_2$ 1:1) δ 8.03 (2H, d, J=8 Hz, Ar-H), 7.83 (2H, br s, Ar-H), 4.96-4.94 (2H, m, CHHN & CHN), 4.25 (1H, d, J=9.6 Hz, CHHN), 3.84 (3H, s, N-CH$_3$), 2.77 (3H, s, CH$_3$). (λ$_{max}$/ nm, toluene): 703, 432, 326. MALDI-TOF-MS (Terthiophene) m/z calcd for C$_{71}$H$_{13}$NO$_2$ 911.09 obsd 910.61

Synthesis of [p-7’-apo-7’-(4-aminophenyl)-β-carotenebenzoate]-[p--(N-methyl-3’,4’-fulleropyrrodin-2’-yl)benzoate](2,9,16,23-tetra-tert-butylphalocyaninato)silicon 17

Dyad 16 (23 mg, 0.015 mmol), C$_{60}$ (54 mg, 0.07 mmol), and sarcosine (42 mg, 0.471 mmol) in anhydrous toluene (20 mL) were stirred at 105 °C under argon atmosphere for 24 hrs. The mixture was then cooled and the solvent removed under reduced pressure. The crude was purified on silica gel (toluene) yielding 21 mg (62%) of triad 17.

$^1$H NMR (400 MHz, CDCl$_3$) δ 9.68 (m, 8H, Pc-Ar-H), 8.36-8.34 (m, 4H, Pc-Ar-H), 7.00-6.98 (d, 2H, J=8, Ph-Ar-H), 6.72-6.43 (m, 10H, vinyl H, Ph-Ar-H), 6.30-6.03 (m, 8H, vinyl H, Ph-Ar-H), 5.75-5.72 (d, 2H, J=12, Ph-Ar-H), 5.18-5.16 (br, d, 2H, Ph-Ar-H), 4.96-4.94 (d, 2H, J=8, Ph-Ar-H), 4.63-4.62 (d, 1H, J=4, CHHN), 4.28 (br s, 1H, CHN), 3.85-3.83 (d, 1H, J=8, CHHN), 2.17 (s, 3H, N-CH$_3$), 1.97-1.88 (m, 14H, CH$_3$-18C, CH$_3$-20C, CH$_3$-19’C, CH$_3$-20’C, CH$_2$-4C), 1.75-1.65 (m, 36H, 4× (CH$_3$)$_3$), 1.57-1.53 (m, 3H, CH$_3$-19C), 1.41-1.38 (m, 4H, CH$_2$-2C, CH$_2$-3C), 0.97-0.96 (m, 6H, CH$_3$-16 and CH$_3$-17C).

UV/Vis (λ$_{max}$/nm, CH$_2$Cl$_2$): 693, 663, 623, 508 (sh), 479, 455 (sh), 360, 329.

MALDI-TOF-MS (Terthiophene) m/z calcd for C$_{162}$H$_{108}$N$_{10}$O$_4$Si 2284.83 obsd 2284.50
2.3 Design and Synthesis of Ferrocene-Silicon Phthalocyanine-Fullerene triad

The design, synthesis and study of artificial photosynthetic reaction center molecular models utilizing SiPc-C\textsubscript{60} dyad was investigated further by incorporating an easily oxidizable molecule, ferrocene, as the secondary electron donor in the triad in place of the carotenoid moiety. Ferrocene (Fc)-porphyrin (P)-C\textsubscript{60} triads\textsuperscript{90,93-95} have been shown to have a long-lived charge separated (Fc\textsuperscript{**}-P-C\textsubscript{60}\textsuperscript{**}) state. A Fc-P-C\textsubscript{60} (figure 31) in which the chromophores are separated by short linkers was reported to have the longest charge-separated lifetime, 630 \(\mu\text{s}\), ever reported for triads at room temperature.\textsuperscript{95}

Figure 31: Ferrocene-Zinc porphyrin-fullerene triad.\textsuperscript{95}

Due to the superior light harvesting and optical properties of axially substituted silicon phthalocyanine over porphyrins, it was thought that attaching ferrocene as a secondary donor to SiPc-C\textsubscript{60} dyad will afford fast forward electron transfer and a long-lived CS state, towards this goal, a ferrocene (Fc)-silicon phthalocyanine (SiPc)-fullerene (C\textsubscript{60}) triad (figure 32) was synthesized and the photophysical properties were studied.
The short linkers between the components are expected to provide better coupling between adjacent chromophores which will enhance a fast forward electron transfer. The excellent electron accepting ability of C$_{60}$ is expected to favor forward electron transfer from the excited state SiPc and on the other hand its low reorganization, $\lambda$ is expected to retard CR and consequently favoring hole transfer to the Fc moiety. Sequential electron transfer is expected to be achieved by initial electron transfer from an excited phthalocyanine to the C$_{60}$ (Fc-$^{1*}$SiPc-C$_{60}$ → Fc-SiPc$^{+*}$-C$_{60}^{*}$) and followed by charge shift to obtain the final charge separated state (Fc-SiPc$^{+*}$-C$_{60}^{*}$ → Fe$^{+*}$-SiPc-C$_{60}^{*}$).

**Results and Discussions**

**Synthesis**

Reaction between the commercially available tetra-tert-butylphthalocyanine silicon dihydroxide, ferrocene carboxylic acid and 4-formylbenzoic acid (scheme 19) yielded a
statistical mixture of products. Silica column purification (hexane/dichloromethane 3:7) afforded Fc-SiPc dyad 22, a 22% yield

Scheme 19: Synthesis of Fc-SiPc dyad 22.

A Prato-type reaction between compound 22, sarcosine and C₆₀ in o-DCB as the solvent was used to obtain the triad 23. The reaction crude was purified on silica column with toluene:EtOAc (49:1) as the solvent to obtain ferrocene-silicon phthalocyanine-fullerene triad 23, a 65% yield. MALDI-TOF (Terthiophene matrix) showed a single peak at m/z: 1889.47 which matched the target triad. No peaks representing fragments were observed.
Scheme 20: Synthesis of ferrocene-silicon phthalocyanine-fullerene triad 23

$^{1}$H NMR spectrum was measured in CDCl$_3$ solvent. On attachment of the ferrocene carboxylic acid, the resonance signals of the phthalocyanine moiety did not change significantly from that of phthalocyanine model 18. The SiPc signals shifted from 9.75-9.55 ppm and 8.43-8.41 ppm in the model SiPc 18 to 9.70-9.52 ppm and 8.40-8.37 ppm in the Fc-SiPc dyad 22. This is an indicator of some slight shielding of the Pc protons by the Fc. The formyl proton of Fc-SiPc dyad 22 appears as a singlet at 9.32 ppm which is also slightly up field from 9.37 ppm of SiPc 19. The nine Fc protons appeared as three distinct broad peaks at 3.17 ppm (2H), 2.75 ppm (5H) and 2.17 ppm (2H).

In the final Fc-SiPc-C$_{60}$ triad 23, no shift in the resonance of the Pc protons were observed from that of Fc-SiPc dyad 22. The phenyl aromatic protons between the SiPc and the C$_{60}$ became broader due to the effect of the latter. The protons of the fulleryrrolidine moiety in the triad 23 were shifted upfield when compared to those of the C$_{60}$ model 21. The pyrrolidine
and the N-CH$_3$ protons were shifted from 4.97, 4.95, 4.25 and 2.78 ppm in C$_{60}$ model 21 to 4.57, 4.21, 3.80 and 2.13 ppm in the triad 23, respectively. This means that the fulleropyrrolidine is close enough to have its protons shielded by the π-cloud of the phthalocyanine moiety.

**Steady-state Absorption**

![Normalized UV/Vis spectra](image)

Figure 33: Normalized UV/Vis spectra (dichloromethane) of SiPc model 18 (solid line), Fe-SiPc dyad 22 (- - dashed line), and Fe-SiPc-C$_{60}$ triad 23 (·· dotted line).

From figure 33, it can be seen that no shift in the Q-band of the SiPc chromophores occurs after the attachment of ferrocene and C$_{60}$ which supports the notion that there are no ground state interactions among the chromophores. The small absorption at 431 nm and the band at 343 nm in the spectrum of triad 23 is a characteristic peak for [6, 6] fulleropyrrolidine and Soret band of fullerene, respectively.
**Electrochemistry**

Table 7: First reduction potentials of SiPc model 18, C₆₀ model 21, and C₆₀-SiPc-Fc triad 23

<table>
<thead>
<tr>
<th>Molecule</th>
<th>$E_{\text{red}}$^1</th>
</tr>
</thead>
<tbody>
<tr>
<td>18</td>
<td>-0.47</td>
</tr>
<tr>
<td>21</td>
<td>-0.56</td>
</tr>
<tr>
<td>23</td>
<td>-0.55</td>
</tr>
</tbody>
</table>

Table 8: Oxidation potentials of SiPc model 18, C₆₀ model 21, SiPc-Fc dyad 22 and C₆₀-SiPc-Fc triad 23

<table>
<thead>
<tr>
<th>Molecule</th>
<th>$E_{\text{oxy}}^1$</th>
<th>$E_{\text{oxy}}^2$</th>
<th>$E_{\text{oxy}}^3$</th>
</tr>
</thead>
<tbody>
<tr>
<td>18</td>
<td>0.93</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td></td>
<td>1.48 (irr)</td>
<td></td>
</tr>
<tr>
<td>22</td>
<td>0.74</td>
<td>1.19</td>
<td></td>
</tr>
<tr>
<td>23</td>
<td>0.71</td>
<td>1.14</td>
<td>1.54 (irr)</td>
</tr>
</tbody>
</table>

The reduction potential of the C₆₀ moiety in the triad 23 did not change significantly from that of the model C₆₀ 21, this confirms that C₆₀ is a superior electron acceptor, a property it retains even in the triad. The first oxidation potential of the SiPc-Fc dyad 22 did not change much after coupling to form triad 23; however the oxidation potential of SiPc changed from 0.93 V vs SCE in the model SiPc 18 to 1.19 V vs. SCE and 1.14 V vs SCE in the SiPc-Fc dyad 22 and triad 23, respectively. This anodic shift in the oxidation potential of the SiPc moiety in the dyad and the triad can be attributed to the fact that, in these two molecules, SiPc oxidation is the 2nd
oxidation potential of the molecule and the presence of the neighboring oxidized ferrocene may make the Pc macrocycle less electron rich hence the observed reduced ease of oxidation. The observed irreversible oxidation peak at 1.48 V in the model C_{60} 21 and 1.54 V in the triad 23 has been reported in similar molecules and is due to the oxidation of the nitrogen atom in the fulleropyrrolidine.\textsuperscript{96}

The electrochemistry data above supports the conclusion made from the absorption spectra data that at ambient condition there is no much ground state chromophoric interactions between the components of the triad.

**Time-resolved absorption**

The decay-associated spectra of Fc-SiPc dyad 22 and Fc-SiPc-C_{60} triad 23 dissolved in benzonitrile and subjected to 100 fs laser pulse at 695 nm is shown in figure 34. There is a very fast (with ~200 fs) formation of Fc^{+}-Pc^{-} CS state in both dyad and triad. The CS state decays in 1.2 ps.
Figure 34: Femtosecond transient absorption spectra of Fc-SiPc dyad 22 (top) and Fc-SiPc-C\textsubscript{60} triad 23 (bottom) in deaerated benzonitrile after laser excitation at 695 nm
There is some impurity or most likely decomposition product [fluorescence SPC does not show impurity, <5% (data not shown)] of Pc whose singlet excited state decays in 6.4 ns forming triplet excited state. The 24 ps component is most likely solvation of Pc excited singlet state. Absence of characteristic transients of SiPc\(^{**}\) (~880 nm) and C\(_{60}\)\(^-\) (1000 nm) rules out formation of Fc-SiPc\(^{**}\)-C\(_{60}\)\(^-\) CS state.

The dominant factor that determines formation of respective CS state is the thermodynamic driving force, and this is what seems to be favoring the formation of Fc\(^{**}\)-SiPc\(^-\)-C\(_{60}\) (\(-\Delta G_{CS}=0.56 \text{ eV}\)) over Fc-SiPc\(^{**}\)-C\(_{60}\)\(^-\) (\(-\Delta G_{CS}=0.29 \text{ eV}\)). The shorter distance between the SiPc and the Fc could also explain the ultra fast charge separation and recombination between these two moieties. The ultrafast charge-recombination in Fc\(^{**}\)-SiPc\(^-\)-C\(_{60}\) is also due to the enhanced coupling between the ion radical pair as a result of a short linker and short distance.

**Experimental Section**

**Synthesis of ferrocene-silicon phthalocyanine dyad 22**

A mixture of \((t\text{Bu})_4\text{SiPc(OH)}_2\) (46 mg, 0.0576 mmol), 4-formylbenzoic acid (75 mg, 0.5 mmol), and ferrocene carboxylic acid (115 mg, 0.5 mmol) in DMF (10 mL) was stirred and warmed to 160 °C under argon atmosphere for 3 hrs. Stirring was stopped, the mixture was cooled to room temperature and the solvent removed under reduced pressure. The crude was purified by silica column chromatography (hexane: DCM 3:7) and three greenish blue bands were observed. The middle band which was the target dyad 22 was isolated and solvent removed to obtain a greenish blue solid Fc-SiPc dyad 22, with a 22% yield. \(^1\text{H NMR}\) (400 MHz, CDCl\(_3\)) \(\delta\) 9.70-9.52 (m, 8H, Pc-Ar-H), 9.32 (s, 1H, CHO), 8.39-8.37 (m, 4H, Pc-Ar-H), 6.69-6.67 (d, 2H,
\[ J=8, \text{ Ph-Ar-H}) , 5.23-5.20 \text{ (d, 2H, J}=8.4, \text{ Ph-Ar-H}), 3.17 \text{ (br s, 2H, Fc-H), 2.75 (br s, 5H, Fc-H), 2.20-2.10 \text{ (m, 2H, Fc-H}), 1.75-1.74 \text{ (m, 36H, 4× (CH}_3)_3\text{).} \]

UV/Vis (\(\lambda_{\text{max/\text{nm, dichloromethane}}}: 694, 663, 624, 361\))

MALDI-TOF-MS (Terthiophene) m/z calcd for C\(_{67}\)H\(_{62}\)FeN\(_8\)O\(_5\)Si 1142.43 obsd 1142.31.

**Synthesis of ferrocene-silicon phthalocyanine-fullerene triad 23**

To a mixture of ferrocene-silicon phthalocyanine dyad 22 (14 mg, 0.0123 mmol), C\(_{60}\) (22 mg, 0.0305 mmol), and sarcosine (12 mg, 0.135 mmol), o-DCB (5 mL) anhydrous was added. The resulting mixture was stirred and heated to reflux under argon atmosphere. The reaction progress was monitored by TLC and after three hrs all the dyad 22 starting material has been consumed and the stirring and refluxing was stopped and the reaction mixture cooled to room temperature. The solvent was removed under reduced pressure and the crude was purified by silica gel column chromatography (toluene: EtOAc 49:1) to obtain a blue compound, triad 23, 65% yield.

\(^1\)H NMR (400 MHz, CDCl\(_3\)) \(\delta 9.70-9.50 \text{ (m, 8H, Pc-Ar-H), 8.36-8.34 \text{ (m, 4H, Pc-Ar-H), 6.63 (br s, 2H, Ph-Ar-H), 5.16-5.14 \text{ (d, 2H, } J=6.8 \text{ Hz, Ph-Ar-H), 4.58-4.56 \text{ (d, 1H, } J=9.2 \text{ Hz, CHHN), 4.21(br s, 1H, CHN ), 3.81-3.79 \text{ (d, 1H, } J=9.2 \text{ Hz, CHHN), 3.26 (br s, 4H, Fc-H), 2.88-2.75 (m, 5H, Fc-H), 2.13 (s, 3H, NCH}_3\text{), 1.74-1.73 (m, 36H, 4× (CH}_3)_3\text{ ).}}\)

UV/Vis (\(\lambda_{\text{max/\text{nm, CH}_2\text{Cl}_2}}: 694, 664, 624, 431, 360, 343\))

MALDI-TOF-MS (Terthiophene) m/z calcd for C\(_{129}\)H\(_{67}\)FeN\(_9\)O\(_4\)Si 1889.48 obsd 1889.47.
3. Regioselective oxidative synthesis of meso-β fused diporphyrin

Introduction
Extending the conjugated aromatic system of porphyrins leads to a significant change in optical and electrochemical properties, and such changes find use in a wide range of fields\textsuperscript{97} that includes technology and medicine\textsuperscript{98} which utilizes the linear and non-linear optical properties of the π-extended porphyrinoids systems. Also, owing to their rigid shape and conjugated electronic system as evidenced by their absorption bands that extend into the NIR, these arrays have potential use as a molecular wires.\textsuperscript{99} There are several methods of extending porphyrin and one of them is the oxidative dimerization.\textsuperscript{100} Oxidative aromatic coupling is a favored method for the extension of the π system of the porphyrin chromophores as it gives the most significant red shift of absorption.\textsuperscript{97} Osuka and coworkers have reported a number of agents for one electron oxidation of porphyrin rings. The most popular oxidants are tris(4-bromophenyl)aminium hexachloroantimonate (BAHA)\textsuperscript{99}, AgPF\textsubscript{6}\textsuperscript{101}, and combinations of 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and scandium trifluoromethanesulfonate (Sc(OTf)\textsubscript{3})\textsuperscript{102}. Although these reagents have been shown to be effective in generating extended porphyrins, BAHA has been observed to cause β chlorination of porphyrins\textsuperscript{99,102} while the other reagents are expensive. In an effort to suppress halogenation, our group has utilized copper (II) perchlorate and tetrafluoroborate salts in acetonitrile\textsuperscript{103}. This method was effective because no chlorination of porphyrins were observed, however it lacked regioselectivity, the products were mixtures of β,β triply-connected and meso-β doubly connected dimers. These posed a purification challenge such that a series of chromatographic column separations were required.
A method that can give a regioselective oxidative porphyrin dimerization without causing unwanted halogenation and a high yield is desirable. In pursuit of this, we utilized Cu$^{2+}$ as an oxidizing agent in the solvent nitromethane to selectively form meso-β doubly-connected fused porphyrin dimer in almost quantitative yield with only a short flash column required for purification$^{100}$. Just like in acetonitrile, Cu$^{2+}$ is a strong oxidizer in nitromethane, with an oxidation potential of 1.0 V vs SCE.$^{104,105}$ Unlike the other oxidation reactions, using Cu$^{2+}$ in acetonitrile allowed us to obtain the pure meso-β doubly connected dimer$^{100}$. Removal of the copper metal from the macrocycle to obtain the free base dimer allows for exchange of metals and provides a route to a wider range of metalloporphyrin dimers.

![Schematic representation of the two HOMO orbitals of the D$_{4h}$ porphyrin ring.](image)

Figure 35: Schematic representation of the two HOMO orbitals of the D$_{4h}$ porphyrin ring.$^{99}$

The regioselectivity of Cu porphyrin to give exclusively a meso-β doubly linked diporphyrins could be due to the $A_{1u}$ HOMO orbital characters of Cu$^{II}$ which has nodal planes through the meso positions and a higher electron density at the β position ($A_{1u}$) $^{106}$. 

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Results and Discussions

Synthesis

Dimerization of pyrrole was achieved by using a well-known protocol that employs paraformaldehyde, InCl$_3$ and NaOH$^{107}$. The unreacted pyrrole during the first reaction was distilled off and taken through the reaction process one more time, the crude from the two reactions were combined and purified over flash column to yield an overall 3% yield.


Porphyrin 25 was synthesized by a BF$_3$•OEt$_2$ catalyzed condensation of dipyrromethane 24 with mesityl aldehyde followed by DDQ oxidation and neutralization by triethyl amine.$^{108}$ Silica column purification afforded 26% of pure compound 25.

Scheme 22: Synthesis of porphyrin 25
The brominating agent N-bromosuccinimide was used to carry out monobromination of the \textit{meso} position of porphyrin 25\textsuperscript{109}. The HBr generated was neutralized with pyridine. The reaction was very fast and it was monitored by TLC and MALDI-TOF and it was stopped as soon as the dibromo product appeared. Lower temperature favored the monobromo product and to optimize the yield of the monobrominated porphyrin the reaction was carried out at below 0\textdegree C temperature.

Scheme 23: Synthesis of porphyrin 26

A yield of 50\% was obtained and an unfragmented molecular ion peak, m/z = 624.3 that correspond to the target monobrominated porphyrin were observed.

Porphyin 27 (scheme 24) was synthesized by Suzuki coupling reaction between toyl boronic acid and monobromo pophyrin 26. Under a reflux condition in presence of heterogeneous solvent mixture of toluene/water/methanol, a high yield of 72\% of porphyrin 27 was realized. In MALDI-TOF a single peak at m/z = 636.6 was observed.
Scheme 24: Synthesis of porphyrin 27

As shown in scheme 25, stirring of porphyrin 27 with Cu(II) acetate monohydrate in methanol at room temperature gave a quantitative yield of Cu(II) porphyrin 28 which was easily isolated by washing with water and brine without need for a chromatographic purification. MALDI-TOF gave a single peak at m/z = 697.5 indicating that the Cu (II) is tightly held in the macrocycle under the MALDI condition.
Scheme 25: Synthesis of porphyrin 28

The porphyrin dimer 29 was obtained by a selective one electron oxidative coupling of porphyrin 28 using Cu(II) tetrafluoroborate in nitromethane solvent. By treating porphyrin 28 with 3.3 equivalents of Cu(II) tetrafluoroborate in nitromethane at room temperature for 3 hours all starting materials were consumed and the reaction mixture was quenched with aqueous potassium ferrocyanide so as to reduce any remaining Cu(II) and unreacted oxidized porphyrin (scheme 26).
A short and a quick silica gel column were used to obtain a pure compound 29 with almost quantitative yield of 90%. MALDI-TOF result of the crude mixture also showed only a single peak at m/z = 1393.0 which matches the calculated mass of the target porphyrin dimer 29.

Stirring compound 29 in a mixture of trifluoroacetic acid and concentrated sulfuric acid (1:10 v/v) at ambient temperature yielded compound 30 after only 20 minutes. Only a single product was obtained as evidenced by a single peak observed in MALDI-TOF, m/z = 1267.0 and a single spot on TLC. A short silica column was run to obtain a pure compound 30, 85% yield; there was no trace of the starting material on the column.
Scheme 27: Synthesis of compound 30

The ease of demetallating compound 29 to produce a free base dimer 30 opens up a possibility of producing a wider range of metallodiporphyrins.

$^1$H NMR (CDCl$_3$) spectroscopy was used to characterize the copper free intermediates and the final product. The aromaticity of the electronic $\pi$ system of porphyrins can be deduced from the chemical shift of the $\beta$-protons and the inner N-H. In the monomer porphyrin 27, the $\beta$-protons appear at 8.71-9.26 ppm and the inner N-H protons appear at -2.89 ppm in CDCl$_3$. The $^1$H NMR spectrum of the *meso*-β doubling linked fused diporphyrin 30 shows upfield shifts of the $\beta$-protons to 8.16-9.24 ppm and downfield shift of the inner N-H protons at 0.72 ppm. The upfield shifts of the $\beta$-protons are an indicator that the aromatic ring current is decreased probably as a result of increased conjugation over the diporphyrin. The dramatic downshift of the inner N-H chemical resonance of the diporphyrin 30 relative to that of the monomer 28 is a further evidence that aromaticity, as measured by ring-current effects, is reduced in the doubly fused porphyrin as opposed to the monoporphyrins.
**Steady-state Absorption**

The UV/Vis/NIR absorption spectra shows a broad and bathochromically shifted three major bands (bands I, II and III) as shown in figure 36. The shift in the major bands towards red is understood to be as a result of extensive conjugation over the diporphyrin π-electronic systems. The positions of band I (414-423 nm) of the diporphyrins are similar to those of the Soret bands of the monomers, only slight red-shift were observed, Soret band of Cu porphyrin dimer 29 is at 414 nm while the Soret of the corresponding monomer 27 is at 410 nm. On the other hand, the Soret band of free base porphyrin dimer 30 is at 423 nm and that of the corresponding monomer 27 is at 412 nm.

![UV-visible-NIR spectrum](image)

**Figure 36:** UV-visible-NIR of 27 (broken), 28 (dots), 29 (solid) and 30 (dashed) in dichloromethane.

The spectra was normalized at the highest absorbance peak.

Band II are not only red-shifted and their intensities magnified, but they are also more broad and complicated. This is an indication of presence of interactions in the ground state and
first excited state. The most red-shifted band is band III and it is dramatically intensified when compared to the Q-bands of the porphyrin monomers. There is a difference in the position of bands, particularly band III between the free-base diporphyrin 30 and Cu\textsuperscript{II} diporphyrin 29. Band III of Cu\textsuperscript{II} diporphyrin 29 is observed at high-energy 770 nm and free-base diporphyrin 30 is observed at low-energy position 814 nm., this mean that the HOMO-LUMO gap is narrower in the latter diporphyrin when compared to the former.

**Experimental Section**

**General procedure for metallation of porphyrin.**

Metalation of porphyrins was performed by adding porphyrin and 5-10 equivalents of Cu(II) acetate to a 5:1 solution of dichloromethane / methanol. The solution was heated for 30 min and then allowed to stir overnight under a nitrogen atmosphere. The organic layer was washed with distilled water, saturated aqueous sodium bicarbonate, and again with distilled water, and the solvent removed under reduced pressure. All reactions went to completion.

**Synthesis of di(1H-pyrrol-2-yl)methane 24**

Undistilled pyrrole was passed through a short alumina column to obtain a clear pyrrole. To the clear pyrrole (205 g, 3.06 mol), paraformaldehyde (0.92 g, 30.6 mmol) was added, and the mixture degassed for 30 min while stirring. The mixture was then warmed at 50 °C in an oil bath for 1 hr followed by addition of InCl\textsubscript{3} (0.68 g, 3.07 mmol), and then the resulting mixture was stirred for another 3 hr after which NaOH (3.63 g, 90.75 mmol) was added and stirring continued for another 1.5 hr at room temperature. The crude was passed through celite and the filtrate was
taken through distillation. The dark oily residue was stored while the pyrrole (169 g, 2.5 mol) distillate was taken through the procedure described above with paraformaldehyde (0.76 g, 25.3 mmol), InCl$_3$ (0.56 g, 2.53 mmol), and NaOH (0.297 g, 7.43 mmol). The crude from the two reactions were combined and silica column chromatography was run with dichloromethane/hexane 7:3 to obtain a snow white fluffy solid dipyrromethane 24, 3.2 g, 3%.

$^1$H NMR (400 MHz, CDCl$_3$) $\delta$ ppm 7.80 (2H, br s, N-H), 6.64-6.63 (2H, m, $\beta$-H), 6.15-6.13 (2H, m, $\beta$-H), 6.03 (2H, br s, $\beta$-H), 3.96 (2H, s, meso).

Synthesis of 5, 15-dimesitylporphyrin 25

A solution of dipyrromethane 24 (0.2 g, 1.37 mmol) and mesityl aldehyde (0.23 g, 1.54 mmol) in CHCl$_3$ (140 mL) was purged with argon for 15 minutes, and then BF$_3$ • OEt$_2$ (0.14 g, 0.972 mmol) was added and the mixture stirred at room temperature. After 6 hr, DDQ (0.467, 2.06 mmol) was added. After a total of 8 hr, the reaction mixture was neutralized by adding 2 mL of triethylamine. The crude mixture was filtered through a pad of basic alumina and the filtrate was evaporated under a reduced pressure. Purification was carried out by silica column chromatography (hexane: CH$_2$Cl$_2$- 3:7) to afford 5, 15-dimesitylporphyrin 25, 98.5 mg, 26% yield.

$^1$H NMR (400 MHz, CDCl$_3$) $\delta$ ppm 10.21 (2H, s, meso), 9.32 (4H, d, $J$=4 Hz, $\beta$-H), 8.88 (4H, d, $J$=5 Hz, $\beta$-H), 7.33 (4H, s, Ar-H), 2.66 (6H, s, Ar-CH$_3$), 1.85 (12H, s, Ar-CH$_3$), $-3.05$ (2H, s, N-H). MALDI-TOF-MS m/z calcd for C$_{38}$H$_{34}$N$_4$ 546.3 obsd 546.4.
Synthesis of 10-Bromo-5, 15-dimesitylporphyrin 26

85 mg (0.156 mmol) of 5, 15-dimesitylporphyrin 25 was dissolved in CHCl₃ (40 mL) and the mixture cooled to below -20 °C by immersing the round-bottom flask in dry ice bath cooled with isopropyl alcohol. To the mixture, NBS (29 mg, 0.163 mmol) and pyridine (0.196 g, 2.48 mmol) were added and the resulting mixture was stirred under argon atmosphere for 20 minutes after which stirring was stopped and acetone was added to the crude mixture. Solvent was removed under a reduced pressure and the resulting crude was purified on a short flash silica gel column (hexane/toluene 1:1) to obtain 10-Bromo-5, 15-dimesitylporphyrin 26, 48.5 mg, 50% yield.

1H NMR (400 MHz, CDCl₃) δ 10.07 (1H, s, meso), 9.65 (2H, d, J=4 Hz, β-H), 9.21 (2H, d, J=4 Hz, β-H), 8.77 (4H, d, J=5 Hz, β-H), 7.29 (4H, s, Ar-H), 7.64 (6H, s, Ar-CH₃), 1.84 (12H, s, Ar-CH₃), -2.88 (2H, s, N-H). MALDI-TOF-MS m/z calcd for C₃₈H₃₃BrN₄ 624.2 obsd 624.3

Synthesis of 5,15-dimesityl-10-tolylporphyrin 27.

A solution of 48 mg (7.69 × 10⁻⁵ mol) of 26, 30 mg (2.24 × 10⁻⁴ mol) of tolyl boronic acid, and 54 mg of potassium phosphate hydrate in 11 mL of toluene/water/methanol (27:1.5:1.5) was degassed for 40 minutes and then 14.4 mg (1.21 × 10⁻⁵ mol) of tetrakis (triphenyl phosphine) palladium (0) was added and the resulting mixture refluxed under argon atmosphere for 10 hrs. The crude mixture was filtered through celite and washed with 50 mL diethyl ether. The filtrate was successively washed with saturated sodium bicarbonate, water and brine. The organic layer was dried over anhydrous magnesium sulfate and the solvent removed. The purplish residue was purified on a silica gel column using a mixture of hexane/dichloromethane (2:1), to obtain a pure
porphyrin 26 (35mg, 72%). $^1$H NMR (400 MHz, CDCl$_3$) δ 10.10 (1H, s, meso), 9.26 (2H, d, J=4 Hz, β-H), 8.85-8.81 (4H, m, β-H), 8.72 (2H, d, J=5 Hz, β-H), 8.10 (2H, d, J= 8Hz, Ar-H), 7.53 (2H, d, J= 8Hz, Ar-H), 7.30 (4H, s, Ar-H), 2.67 (3H, s, Ar-CH$_3$), 2.64 (6H, s, Ar-H), 1.84 (12H, s, Ar-H), -2.89 (2H, s, N-H). MALDI-TOF-MS m/z calcd for C$_{45}$H$_{40}$N$_4$ 636.3 obsd 636.6; UV-visible ($\lambda_{max}$, CH$_2$Cl$_2$) 412, 507, 538, 582, 637 nm.

Synthesis of Cu (II) 5,15-dimesityl-10-tolylporphyrin porphyrin 28

This material was prepared from 27 as specified by the general procedure for porphyrin metalation to obtain a quantitative yield. MALDI-TOF-MS m/z calcd for C$_{45}$H$_{38}$CuN$_4$ 697.2, obsd 697.5; UV-visible-NIR ($\lambda_{max}$, CH$_2$Cl$_2$) 533, 410 nm

Synthesis of Cu$^{II}$ diporphyrin 29

Thirty-nine mg (5.59× 10$^{-5}$ mol) of 28 in 20 mL nitromethane was sonicated for 30 minutes and then degassed and then transferred to a round bottom flask containing 63 mg (1.83× 10$^{-4}$ mol) of Cu(BF$_4$)$_2$•6 H$_2$O in 15 mL nitromethane which has previously been sonicated until all salt has dissolved followed by degassing. The color of the mixture immediately turned to dark purple. Stirring at room temperature under argon atmosphere for 2 hrs was continued. The reaction mixture was quenched with potassium ferrocyanide in 50 mL water followed by addition of 50 mL chloroform. Organic layer was dried over anhydrous sodium sulfate and filtered. Solvent was removed to obtain purplish crude. The purplish residue was purified on a silica column using a mixture of hexane/dichloromethane (4:1), to obtain copper (II) diporphyrin 29 (35 mg, 90%). MALDI-TOF-MS m/z calcd for C$_{90}$H$_{76}$Cu$_2$N$_8$ 1392.5, obsd 1392.7;
UV/Visible/NIR (λ_max, CH₂Cl₂): 414, 487 (sh), 552, 628, 702 (sh), 770 nm.

**Synthesis of free base diporphyrin 30**

A solution of 35 mg (2.51× 10⁻⁵ mol) of copper (II) diporphyrin 29 in 15 mL H₂SO₄/TFA (1:10 v/v) was stirred at room temperature under argon atmosphere for 20 min when all the starting material were gone by TLC using hexane/dichloromethane (1:2) mixture. The reaction mixture was quenched by adding excess ice. The product was extracted with dichloromethane and the organic phase was washed with aqueous NaOH (20%, w/w) water, brine and dried over anhydrous Na₂SO₄. The resulting crude was purified on a short silica gel column using hexane/dichloromethane (1:2) to obtain free base diporphyrin 30 (27 mg, 85%). ¹H NMR (400 MHz, CDCl₃) δH 9.23 (2H, d, J= 5Hz, β-H), 8.90 (2H, s, β-H), 8.42 (2H, d, J=5 Hz, β-H), 8.36 (2H, d, J=5 Hz, β-H), 8.31 (2H, d, J=4 Hz, β-H), 8.20 (2H, d, J= 5Hz, β-H), 8.16 (2H, d, J= 5Hz, β-H), 7.95 (4H, d, J= 8Hz, Ar-H), 7.49 (4H, d, J=13 Hz, Ar-H), 7.29 (4H, s, Ar-H), 7.23 (4H, s, Ar-H), 2.64 (12H, d, J= 4Hz, Ar-CH₃), 2.60 (6H, s, Ar-CH₃), 2.07 (12H, s, Ar-CH₃), 1.94 (12H, s, Ar-CH₃), 0.72 (4H, br, s, N-H). MALDI-TOF-MS m/z calcd for C₉₀H₇₆N₈ 1269.6, obsd 1484.2; UV-visible-NIR (λ_max, CH₂Cl₂) 423, 499, 560, 611, 737(sh), 814 nm. ⁸⁹
4. Expedient synthesis of bacteriopurpurin: A red-absorbing dye for potential application in hydrogen production

Introduction

Dye-sensitized solar cells (DSCC) based on TiO$_2$ using Ruthenium (Ru) dyes with efficiencies of up to 11% has been reported. However, the cost and the environmental issues associated with ruthenium dyes demands for discovery of cheaper and safer organic dyes. Some of the advantages of organic dyes include, diversity, ease of modification of their molecular structures and their intense absorption.

Porphyrin (figure 37) sensitizers make good candidates for this purpose owing to their intense absorption in Soret and Q bands to harvest solar energy efficiently in a broad spectral region. However, porphyrin-sensitized TiO$_2$ cells have poor light harvesting properties relative to Ru dyes at 450-500 nm and 650-900 nm due to the existence of a gap between the Soret and Q bands and this is a major limitation. One possible way to overcome this problem is to modulate the electronic structures of porphyrins so that we can match the light-harvesting properties with the energy distribution on the earth particularly in the visible and NIR regions.

Figure 37: Porphyrin 1-24 numbering system
These types of dyes can be used to replace Ru dyes as TiO$_2$ sensitizers and coupled to water splitting catalytic center to produce hydrogen. Figure 38 shows a working of water splitting DSSC.

![Figure 38: Schematic diagram of the water splitting dye sensitized solar cell.](image)

The electrons injected by the sensitizer are conducted through the wire to the cathode, where protons are reduced to hydrogen. The net result is solar production of hydrogen gas and molecular oxygen. Towards the goal of developing a dye with the right absorption spectra and redox properties, bacteriopurpurin 31 was synthesized.

![Bacteriopurpurin 31](image)
Bacteriopurpurins possesses two annelated cyclopentyl rings which are opposite to each other. The reduction of the pyrrolic rings in the porphyrin macrocycle has a significant effect on the absorption spectrum of the reduced compounds.\textsuperscript{115}

**Results and Discussions**

**Synthesis**

Synthesis of bacteriopurpurin 31 was started from metalation of Porphyrin 32 (obtained from Dr. Paul Liddell) as shown in scheme 28.\textsuperscript{116}

\[
\begin{align*}
\text{Scheme 28: Nicklyation of porphyrin 32} \\
\text{The yield was not very good, and also it was also observed that the percent yield decreased when scaling up was attempted. This may have been due to the HCl formed displacing Ni from the product and reforming the starting material. Different reagents and reaction conditions were used to obtain a better yield of 33. Reaction between compound 32 and Ni (II) acetylacetone hydrate in refluxing toluene for 24 hours afforded compound 33 in a near quantitative yield, 92% yield (scheme 29) }
\end{align*}
\]
Scheme 29: Nickel 5, 15-bis(tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 33

Monoformylation of 33 using Vilsmeier formylation protocol with (chloromethylene)dimethyl ammonium chloride yielded a deep green compound 34 as shown in scheme 30.\textsuperscript{115} The diformyl product was a tiny slow moving band on the silica gel column and separation was easily achieved.

Scheme 30: Nickel 10-meso-formyl- 5, 15-bis(tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 34

A Wittig type reaction between compound 34 and (carboethoxymethylene) triphenyl phosphorane afforded compound 35 (scheme 31). Compound 35 was purified by flash chromatography on silica column (dichloromethane: hexane 3:1) followed by recrystallization in methanol: dichloromethane to obtain a small dark green crystals\textsuperscript{117} to obtain high yield, 84\%. 

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Scheme 31: Synthesis of porphyrin 35

Several attempts to formylate position 20 (figure 37) of compound 35 using the method used by Robinson\textsuperscript{115} were not successful. Although it has been reported by Morgan et al\textsuperscript{118} that nickel porphyrin undergo only monoformylation, diformylation of Ni (II) porphyrin 33 was realized by modifying the method used in scheme 30. Using an excess of a freshly prepared Vilsmeier reagent obtained by drop wise addition of phosphorous oxychloride to DMF and running the reaction longer at higher temperature, the diformyl product, compound 36 was obtained as the major product. The diformyl product was separated from the monoformyl product on a silica flash column (dichloromethane: hexane 3:1) and obtained in 61\% yield. Scheme 32 shows conversion of nickel porphyrin 33 to 10, 20-diformyl nickel porphyrin 36.

Scheme 32: Nickel 10,20-bis(meso-formyl)- 5, 15-bis(tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 36
Compound 36 was converted to compound 37 as shown in scheme 33, column purification afforded porphyrin 37, 43% yield. The side product had one of the formyl group unreacted and it was separated from compound 37 on a silica flash column with dichloromethane as the mobile phase.

Scheme 33: Synthesis of nickel Porphyrin 37

Demetalation of 37 was attempted following the method used by Robinson\textsuperscript{115} which employed the use of concentrated sulfuric acid followed by neutralization with saturated sodium bicarbonate as shown in scheme 34.

Scheme 34: Attempted synthesis of porphyrin 38
Compound 38 was not obtained; from the MALDI-TOF results it was observed that the concentrated sulfuric acid used hydrolyzed the vinylic esters to the corresponding vinylic acids without removing nickel from the macrocycle.

We thought of a way of avoiding reactions involving concentrated sulfuric acid in presence of esters, an easily hydrolysable group, the only way to do that was to remove the nickel from 10, 20-diformyl nickel porphyrin 34, generally aldehydes are known to be less susceptible to acid hydrolysis than esters. After all, the role of nickel was to facilitate formylation, a role it has served diligently and it is no longer needed in further steps.

Free base 10, 20-diformyl porphyrin 39 was obtained by reacting Ni (II) 10, 20-diformyl porphyrin 36 with concentrated sulfuric acid as shown in scheme 35. The progress of the reaction was monitored by TLC, and in less than 20 minutes all the starting material has been consumed. After neutralization with sodium bicarbonate the crude was purified on silica column to afford free base porphyrin 39, 21% yield

Scheme 35: 10,20-bis(meso-formyl)-5,15-bis(tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 39
The low yield was attributed to high polarity of the product which made it stick to the silica flash column used. 4% methanol: dichloromethane was used to elute the compound from a short column.

Synthesis of 5,15-(3,7,13,17-tetraethyl-2,8,12,18-tetramethyl-10,20-ditolyld) bacteriopurpurin 31 was achieved by reacting free base porphyrin 39 with (carboethoxymethylene) triphenyl phosphorane in a refluxing xylene. Scheme 36 shows synthesis of bacteriopurpurin 31. Column purification with dichloromethane as the eluent afforded bacteriopurpurpurin 31 as a lime green product, 31% yield.

Scheme 36: 5,15-(3,7,13,17-tetraethyl-2,8,12,18-tetramethyl-10,20-ditolyld) bacteriopurpurin 31

It important to note that the formation of meso diacrylate free base porphyrin and cyclization to bacteriopurpurin 31 occurred in one step without the use of non-nucleophic base such as DBU as used in the literature.

Steady-state Absorption

Bacteriopurpurin 31 dye shows a characteristic Soret band at 435 and Q-bands at 578, 611, 696 and 845 nm. Figure 39 shows UV-Vis-NIR absorption of bacteriopurpurin 31.
Figure 39: UV/Vis/NIR spectra absorption of Bacteriopurpurin 31 in dichloromethane.

The ability to absorb into the NIR makes it a good harvester of light across the electromagnetic spectrum.
**Electrochemistry**

Figure 40 show the CV scan of bacteriopurpurin 31 (Electrochemistry measurement was done by Dr. Ben Sherman)

![CV scan of bacteriopurpurin 31](image)

Figure 37 : CV scans of bacteriopurpurin 31

The experiment was performed in dichloromethane with 0.1 M TBAPF$_6$, Pt working electrode, Ag$^+$/Ag quasi-reference calibrated to SCE with ferrocene (Fc$^+$/Fc 0.45 V vs SCE). Scans shown were measured at a rate of 100 mV/s.

As shown in figure 39 bacteriopurpurin 31 undergoes successive oxidations. First reversible single oxidation occurs at 0.38 V vs SCE and the second irreversible oxidation occurs at 0.92 V vs SCE. The first reduction potential of the excited state is -1.07 V vs SCE. Therefore, the synthesized bacteriopurpurin 31 is likely to be a good donor to TiO$_2$ (E$^{\text{red}}$ = -0.85 V vs SCE) when incorporated onto a photocathode catalyst of a photoelectrochemical device for light driven H$_2$O splitting by reducing hydrogen ions (E$^{\text{red}}$ = -0.6 5V vs SCE) $^{19}$ to produce hydrogen gas.
In conclusion, an expedient route to the synthesis of bacteriopurpurin has been developed. CV studies have shown that this dye has promising redox properties to be used as a TiO$_2$ sensitizer in a dye sensitized solar cells for the production of hydrogen. The molecule was found to be unstable in air due to oxidative decomposition.

**Experimental Section**

**Nickel 5, 15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 33**

A solution of 5, 15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 32 0.208 g (0.313 mmol) and nickel (II) acetylacetone hydrate 1.241 g (4.8 mmol) in toluene (50 mL) was heated under reflux for 24 hrs. The resulting solution was cooled and the solvent was removed under reduced pressure. The resulting crude was dissolved in dichloromethane and washed with water several times. The organic layers were combined and solvent removed. Silica column chromatography of the crude product on silica with dichloromethane as eluent gave only one band, which was collected to yield 33 (0.206 g, 92 %) as a bright red solid.

$^1$H-NMR (CDCl$_3$, 400 MHz) δ$_H$ 9.41 (s, 2H, meso-H), 7.69 (d, 4H, $J$=7.6 Hz, Ar-H), 7.41 (d, 4H, $J$=7.6 Hz, Ar-H), 3.71-3.66 (m, 8H, 4×C$_H_2$), 2.63 (s, 6H, 2×Ar-CH$_3$), 2.26 (s, 12H, 4×CH$_3$), 1.62-1.58 (t, 12H, 4×CH$_2$CH$_3$)

UV/Vis (CH$_2$Cl$_2$): $\lambda_{max}$ nm 562, 520, 400

**Nickel 10,20-bis(meso-formyl) - 5, 15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 36**

A 20-mL portion of Vilsmier reagent, prepared by a drop wise addition of POCl$_3$ (14 mL) to DMF (11 mL) at 0 °C and allowed to stand at room temperature for 1 hr during which the
solution turned pink was transferred to a round-bottom flask immersed in an oil bath heated to 70 °C. To this solution, nickel 15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 33 (107 mg, 0.15 mmol) dissolved in anhydrous 1,2-dichloroethane (62 mL) was added drop wise. The reaction mixture was stirred for an additional 9 hr while maintaining at around 70 °C. A saturated solution of sodium acetate (200 mL) was then added, and stirring and heating were continued for 2 more hours. The organic and the aqueous phase were separated and the former was washed with dichloromethane three times. Organic layers were combined and the solvent removed under reduced pressure. The resulting crude was purified on silica flash column, the first red band was eluted with CH₂Cl₂/hexane 4:1 and the major green band eluted with CH₂Cl₂. On drying, the green band gave a bright green solid nickel 10,20-bis(meso-formyl)-5,15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 36 (95 mg, 82%).

\[ ^1H-NMR \text{(CDCl}_3, 400 \text{ MHz}) \delta \text{H} 11.53 \text{ (s, 2H, 2×CHO)}, 7.55 \text{ (d, 4H, } J= 8 \text{ Hz, Ar-} \text{H}), 7.39 \text{ (d, 4H, } J= 8 \text{ Hz, Ar-} \text{H}), 3.39-3.35 \text{ (m, 8H, 4×CH}_2), 2.58 \text{ (s, 6H, 2×Ar-CH}_3), 1.96 \text{ (s, 12H, 4×CH}_3), 1.48-1.45 \text{ (t, 12H, 4×CH}_2 \text{CH}_3) \]

MALDI-TOF-MS m/z calcd for C₄₈H₄₈N₄NiO₂ 770.3 obsd 770.6

10,20-bis(meso-formyl)-5,15-bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylporphyrin 39

To a solution of nickel porphyrin 36 (110 mg, 0.1429 mmol) in dichloromethane (30 mL), concentrated sulfuric acid (0.4 mL) was added drop wise and the mixture stirred at room temperature for 6 minutes. The mixture was neutralized by addition of saturated aqueous sodium...
hydrogen carbonate (30 mL) and the organic layer was collected, washed and dried. The solvent was removed under reduced pressure, and the crude product purified on a silica gel (CH$_2$Cl$_2$/MeOH 9.5:0.5) to yield a dark green solid 39 (61.7 mg, 60%).

$^1$H-NMR (CDCl$_3$, 400 MHz) $\delta$H 12.14 (s, 2H, 2×CHO), 7.98 (d, 4H, J= 8 Hz, Ar-$\text{H}$), 7.541 (d, 4H, J= 8 Hz, Ar-$\text{H}$), 3.02-2.99 35 (m, 8H, 4×C$_2$H$_2$), 2.66 (s, 6H, 2×Ar-CH$_3$), 1.85 (s, 12H, 4×CH$_3$), 1.28-1.25 (t, 12H, 4×CH$_2$CH$_3$).

MALDI-TOF-MS m/z calcd for C$_{48}$H$_{50}$N$_4$O$_2$ 714.9 obsd 715.4

5, 15-Bis(β-tolyl)-2,8,12,18-tetraethyl-3,7,13,17-tetramethylbacteriopurpurin 31

(Carboethoxymethylene)triphenyl phosphorane (0.5955 g, 1.71 mmol) was added to a solution of porphyrin 39 (13 mg, 0.0182 mmol) in xylene (20 mL) and the mixture heated to reflux while stirring under argon atmosphere for 24 hrs. The mixture was cooled and solvent removed under reduced pressure. The crude was purified on silica gel chromatography (dichloromethane) and the only lime green band observed was collected to yield bacteriopurpurin 31 (4.8 mg, 31%).

$^1$H-NMR (CDCl$_3$, 400 MHz) $\delta$H 9.22 (s, 2H, 2× isocyclic ring H), 7.70-7.68 (m, 4H, Ar-H), 7.52-7.50 (m, 4H, Ar-H), 4.39-4.37 (m, 4H, 2×CO$_2$CH$_2$), 4.24-4.16 (m, 2×CHCH$_3$, 2×CH$_2$CH$_3$, 2×CHCH$_3$), 2.55 (s, 6H, 2×CH$_3$), 2.34-2.69 (m, 4H, 2×CHCH$_2$), 2.02 (s, 6H, 2×CH$_3$), 1.71-1.62 (m, 6H, 2×CO$_2$CH$_2$CH$_3$), -0.26-0.3 (t, 6H, 2×CHCH$_2$CH$_3$).

MALDI-TOF-MS: M/z calcd for C$_{56}$H$_{92}$N$_4$O$_4$ 854.5, obsd. 854.9.
Conclusions

An efficient synthesis of several organic molecular models of artificial photosynthetic reaction centers were realised and interesting and promising photophysical properties were observed. Both free base (H\textsubscript{2}) phthalocyanine(Pc)-fullerene (C\textsubscript{60}) and Zinc (Zn) Pc-C\textsubscript{60} where the Pc and the C\textsubscript{60} moieties are separated by a phenylethynyl linker was realised in high yields. While H\textsubscript{2}Pc-C\textsubscript{60} exhibited aggregation characteristics and was insoluble in most common organic solvents, ZnPc-C\textsubscript{60} was soluble. While in non-polar solvents the fluorescence of the excited Pc chromophore is quenched mainly by singlet-singlet energy transfer to C\textsubscript{60} in polar solvents both dyads exhibited a ultrafast electron transfer to the covalently-linked C\textsubscript{60}. Transient absorption studies on ZnPc-C\textsubscript{60} showed formation of a charge-separated (CS) with quantum yield of unit that had a lifetime of 94 ps. This is the longest life time reported for similir dyads with similar electron transfer driving force, it can be concluded that the unusually long-lived CS state achieved with the current dyad is as result of the phenylethynyl linker.

A novel non-aggregating silicon (Si)-Pc dyad where the Pc and the C\textsubscript{60} are linked axially via the central Si was synthezied and lifetime of CS state of ~5 ns was observed. The lifetime for axially linked dyad’s CS state is about 50 times that of the the peripherally linked dyads, an observation that can be used to infer that the lifetime of CS state is not only dependent on the type of linker separating the donor and the acceptor but also on the orientation, pherIPHERAL or axial. Assymetrical diaxial substitution on the central Si in SiPc was achieved, and this presents a possibility of attaching different kind of ligands such as donor-acceptor, push-pull systems on the opposite plane of the SiPc chromophore through the central Si without the occurrence of the
photoinactive aggregates in solution and this allow tuning of the optical and electrochemical properties of the phthalocyanine for efficient solar energy capture and other photochemical applications.

A carotenoid (Car) and a ferrocene (Fc) secondary electron donors were covalently attached to the SiPc-C$_{60}$ through axial linkage to form Car-SiPc-C$_{60}$ and Fc-SiPc-C$_{60}$ triads respectively. A very fast charge separation and charge-recombination (CR) within Car-SiPc and Fc-SiPc dyads were observed. Lifetime of the CS states were 1.2 ps and 200 fs for Car-SiPc and Fc-SiPc dyads respectively. The ultra fast charge separation observed is favored by enhanced coupling afforded by the short linkers and a large driving force in the Car-SiPc and Fc-SiPc dyads. The driving force (-\(\Delta G_{CS}\)) for the formation of the CS state for Car$^{*+}$-SiPc$^-$ and Fc$^{*+}$-SiPc$^-$ are 0.81 eV and 0.56 eV respectively. In the Car-SiPc-C$_{60}$ triad, the transfer of electrons to the C$_{60}$ to form Car-SiPc$^{*+}$-C$_{60}^-\cdot$ is a minor pathway for quenching of the SiPc fluorescence. This is because the slower electron transfer from SiPc to C$_{60}$ gets outcompeted by the very fast electron transfer from the ground state carotenoid to the hole in the HOMO of the excited SiPc. Molecular orbital calculations has showed that the highest occupied molecular orbital (HOMO) of the C$_{60}$ moeity is below those of both car and SiPc, and transfer of electrons from SiPc to C$_{60}$ has been observed to occur after singlet-singlet energy transfer from excited state SiPc to the C$_{60}$ followed by a hole transfer from the HOMO of the C$_{60}$ to the energetically higher HOMO of SiPc. Currently, exclusive excitation of C$_{60}$ chromophore so as to realise step wise hole transfer to the Car moeity to afford a long-lived Car$^{*+}$-SiPc-C$_{60}^-\cdot$ is under investigation. Alternative approach that may favor formation of Car$^{*+}$-SiPc-C$_{60}^-\cdot$ as the final state includes: Use of shorter
carotenoid to lower the HOMO of the Car moiety, so as to make it a poor donor to the excited state Pc, decoupling of the Car and SiPc dyad will also slow the rate of electron transfer between the Car to the SiPc, while introducing a shorter linker between the SiPc and the C_{60} will make the intramolecular electron transfer from the SiPc to the C_{60} faster hence enabling this step to be kinetically competitive to afford a charge-shift to the Car moiety. The above modifications between the respective dyads can also be applied to Fc-SiPc-C_{60} to give a long-lived Fc^{**}-SiPc-C_{60}^{●+} CS that will enable harnessing of the stored energy in the CS state.

An efficient route to meso-β doubly connected porphyrin dimer was developed. A near quantitative yield with minimal purification was obtained. Use of cheap Cu^{2+} oxidants and chloride free solvent ensured absence of β- chlorinated porphyrin impurities as was reported with use of tris(4-bromophenyl)amine hexachloroantimonate (BAHA), 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and trifluoromethanesulfonate. Unlike the lack of regioselectivity reported with use of copper (II) perchlorate and tetrafluoroborate salts in acetonitrile, use of Cu^{3+} salt as the oxidant in nitromethane afforded meso-β doubly-connected fused porphyrin dimer exclusively. Stirring the copper (II) porphyrin dimer in H_{2}SO_{4}/TFA mixture easily converted it into free base porphyrin dimer in 20 mins with a high yield of 85 %. The optical and the electrochemical properties of the porphyrin dimer can be tuned by inserting different metals or combinations of metals into the macrocycles. The interesting photophysical properties and the low band gap of the porphyrin dimer find applications not only in solar energy conversion but in other fields of technology and medicine.
An efficient and expedient route to bacteriopurpurin, a red-absorbing dye was developed. Due to the presence of stereocenters and two annelated cyclopentyl rings, synthesis of bacteriopurpurin has been reported to be a synthetically challenging task. Starting with a quantitative nicklyation of porphyrin 32 by nickel (II) acetylacetone hydrate, one-step 10,20-diformylation of porphyrin 33 was achieved by using a freshly prepared Vilsmaer reagent as opposed to the commercially available one which only yields monoformyl product. Use of (Carboethoxymethylene)triphenyl phosphorane, Wittig reagent, to synthesize meso-diacrylate product and stereoselective cyclization to afford bacteriopurpurin 31 occurred in a single step without the need for additional reagents. Overall the seven steps from porphyrin to bacteriopurpurin reported in literature were reduced to four and still with a higher yield of the final product.

The bacteriopurpurin obtained was observed to have interesting electrochemical and optical properties, reduction potential (-0.85 V vs SCE) and $\lambda_{\text{max}}$: 845 nm. These properties makes it an ideal candidate for use as a dye to sensitize TiO$_2$ which can be coupled to the H$^+$ reduction side of the water splitting catalytic center to produce H$_2$, a clean source of energy.

**General Experimental Methods**

**Materials:** Unless noted all reagents and solvents were obtained from commercials sources which are referenced where applicable. Toluene, dichloromethane and hexanes were distilled prior to use. Silica gel chromatography was carried out using 230-400 silica gel (Silicyale, Siliflas F60).
General analytical methods

Stead-state spectroscopy

UV-Visible-NIR spectroscopy was performed using a Shimadzu UV-3101PC UV-Visible spectrophotometer or Shimadzu UV-2550 UV-Visible spectrophotometer. Excitation was provided by a 75 W xenon-arc lamp and single grating monochromator. Fluorescence was detected at 90° to the excitation beam via a single grating monochromator and an R928 photomultiplier tube having S-20 spectral response and operating in the single photon counting mode.

Time-resolved fluorescence

Fluorescence decay measurements were performed on optically dilute (ca. $1 \times 10^{-5}$ M) samples by the time-correlated single-photon-counting method. Two different systems were employed. Excitation source for the first system was a mode-locked Ti:Sapphire laser (Spectra Physics, Millennia-pumped Tsunami) with a 130-fs pulse duration operating at 80 MHz. The laser output was sent through a frequency doubler and pulse selector (Spectra Physics Model 3980) to obtain 370-450 nm pulses at 4 MHz. Fluorescence emissions was detected at the magic angle using a double grating monochromator (Jobin Yvon Gemini-180) and a microchannel plate photomultiplier tube (Hamamatsu R3809U-50). The instrument response function was 35-55 ps. The spectrometer was controlled by software based on the LabView programming language and data acquisition was done using a single photon counting card (Becker-Hickl, SPC-830).

Excitation source for the second system was a fiber supercontinuum laser based on a passive modelocked fiber laser and a high-nonlinearity photonic crystal fiber supercontinuum
The laser provides 6-ps pulses at a repetition rate variable between 0.1 – 40 MHz. The laser output was sent through an Acousto-Optical Tunable Filer (Fianium AOTF) to obtain excitation pulses at desired wavelength. Fluorescence emission was detected at the magic angle using a double grating monochromator (Jobin Yvon Gemini-180) and a microchannel plate photomultiplier tube (Hamamatsu R3809U-50). The instrument response function was 35-55 ps. The spectrometer was controlled by software based on the LabView programming language and data acquisition was done using a single photon counting card (Becker-Hickl, SPC-830).

**Transient absorption**

The femtosecond transient absorption apparatus consisted of a kilohertz pulsed laser source and a pump-probe optical setup. Laser pulses of 100 fs at 800 nm were generated from an amplified, mode-locked Titanium Sapphire kilohertz laser system (Millennia/Tsunami/Spitfire, Spectra Physics). Part of the laser pulse energy was sent through an optical delay line and focused on to a 3 mm sapphire plate to generate a white light continuum for the probe beam. The remainder of the pulse energy was used to pump an optical parametric amplifier (Spectra Physics) to generate excitation pulses, which were selected using a mechanical chopper. The white light generated was then compressed by prism pairs (CVI) before passing through the sample. The polarization of pump beam was set to the magic angle (54.7°) relative to the probe beam and its intensity adjusted using a continuously variable neutral density filter. The white light probe is dispersed by a spectrograph (300 line grating) onto a charge-coupled device (CCD)
camera (DU420, Andor Tech.). The final spectral resolution was about 2.3 nm for over a nearly 300 nm spectral region. The instrument response function was ca. 150 fs.

The nanosecond-millisecond transient absorption measurements were made with excitation from an optical parametric oscillator driven by the third harmonic of a Nd:YAG laser (Ekspla NT342B). The pulse width was ~4-5 ns, and the repetition rate was 10 Hz. The detection portion of the spectrometer (Proteus) was manufactured by Ultrafast Systems. The instrument response function was ca. 4.8 ns.

Data analysis

Data analysis was carried out using locally written software (ASUFIT) developed under a MATLAB environment (Mathworks Inc.). Decay-associated spectra (DAS) were obtained by fitting the transient absorption or fluorescence change curves over a selected wavelength region simultaneously as described by Eq 1 (parallel kinetic model),

$$
\Delta A(\lambda, t) = \sum_{i=1}^{n} A_i(\lambda) \exp\left(-t / \tau_i\right)
$$

where $\Delta A(\lambda, t)$ is the observed absorption (or fluorescence) change at a given wavelength at time delay $t$ and $n$ is the number of kinetic components used in the fitting. A plot of $A_i(\lambda)$ versus wavelength is called a decay-associated spectrum, and represents the amplitude spectrum of the $i^{th}$ kinetic component, which has a lifetime of $\tau_i$.

The global analysis procedures described here have been extensively reviewed in literature. Random errors associated with the reported lifetimes obtained from fluorescence and transient absorption measurements were typically $\leq 5\%$. 

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Mass spectrometry was performed using the MALDI-TOF method using a Voyager DE STR from Applied Biosystems in reflector mode for all molecules. Commonly used matrices were terthiophene, dithranol and trans-2-[3-(4-tert-butylphenyl)-2-methyl-2-propenylidene]-malononitrile.

$^1$H NMR spectra for the molecules were obtained in deuterated solvents using a Varian 400 MHz instrument with tetramethysilane as internal reference. Proton assignments were by the use of COSY and NOESY experiments. 

Electrochemical measurements were performed using a CH instruments 650C or 760D electrochemical workstation using a standard 3-electrode cell setup. Counter and reference electrodes were platinum gauze and pseudo Ag/AgCl. Working glassy electrodes and electrolytes were specific for each experiment. Potentials were referenced to SCE using the ferrocene/ferrocenium redox couple as an internal standard, with values depending on the solvent.\textsuperscript{120}
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