Silicon phthalocyanines: synthesis and resurgent applications

Koushambi Mitra (✉) and Matthew C. T. Hartman (✉)

The intense far-red absorption and emission features have made silicon phthalocyanines (SiPcs) distinct from the structurally related porphyrin analogues. Unlike most other phthalocyanines, SiPcs possess two additional axial bonds which reduce aggregation in solution and can be synthetically tailored, thereby creating further scope for modulation of optical, chemical and electronic properties. Multiple synthetic strategies have been employed for facile construction of symmetrical or unsymmetrical SiPc variants bearing desired substituents at the axial and the aromatic ring positions. The overarching motive of this concise review article is to highlight and summarize the key synthetic routes and the fast-emerging applications of SiPcs in photouncaging techniques, photothermal and photoimmunotherapy, photovoltaics, optoelectronics and photocatalysis.

1. Introduction

The mid-twentieth century witnessed a wave of enthusiasm among researchers in developing new classes of macroheterocyclic compounds by inserting different elements in the core of phthalocyanines (Pcs).1–4 Like their closely related por-
These axial ligands serve as pivotal handles for fine-tuning of the properties of SiPcs as desired. Furthermore, bulky or polar substitutions in the axial positions perturb planarity and are more effective than the \( \alpha \) or \( \beta \) phthalocyanine ring substitutions in reducing aggregation and improving solubility. Extensive synthetic methodologies have been developed for preparation of symmetrical and unsymmetrical SiPcs, often containing both tetravalent (axial) and hexavalent (core) silicon in the same structure. Remarkably, axial Si–O and Si–C bonds exhibit NIR light induced dissociation rendering novel applications of SiPcs in medicinal biology, beyond the context of photodynamic therapy. These unique properties coupled with the inexpensive cost of synthesis and low toxicity have resulted in fascinating applications of SiPcs in major fields of research including cancer phototherapy, NIR imaging, organic photovoltaics, organic electronics and photocatalysis.

In this condensed review article, we present an overview of the broad synthetic methodologies adapted to improve the design of symmetrical and non-symmetrical SiPcs, since their first successful isolation in around 1960. Here, we also account for the interesting properties and the related applications of SiPcs over the past decade, emphasizing mostly on the emerging and novel applications. In the concluding section, we have included a brief perspective to provide guidance for rationally designing an envelope of new SiPc derivatives, to discern areas of impending developments, and to anticipate the future directions of this interesting field.

2. Syntheses of silicon phthalocyanines

Facile, short and high yielding synthetic strategies comprising of non-hazardous and easily accessible starting materials are an essential prerequisite to realize the complete potential of any material. Various synthetic approaches have been developed to increase the structural flexibility of SiPcs. These approaches offer easy modulation of the physico-chemical properties as desired and help to rapidly generate a vast library of SiPc derivatives with diverse functionalities and applications. For example, successful applications of SiPcs in biological fields can be achieved by incorporation of water soluble or bio-compatible groups. On the other hand, insertion of electron donating and electron withdrawing moieties can alter the opto-electronic properties of SiPc molecules. Likewise, availability of electrophilic or nucleophilic reactive handles in the SiPc structure will allow to append desired functionalities for versatile applications.

Such fine-tuning of SiPc structures is mainly obtained by reactions at two major sites: (i) the axial and (ii) the phthalocyanine ring \((\alpha)\) and \((\beta)\) positions (Fig. 1C). While the axial handles are exploited for imparting variants without considerably affecting the absorption and emissive properties of SiPc, the \(\alpha/\beta\) substitutions result in a prominent change of spectral features. Below, we have elaborated on the discovery of SiPc
and the various synthetic methods developed henceforth resulting in the formation of several symmetrical and unsymmetrical SiPcs.

2.1 Synthesis of SiPc precursors

The synthesis of most SiPcs is based on two main precursors, the dichloro SiPc (SiPcCl₂), 1 (Schemes 1 and 2) and the bis-hydroxy adduct [SiPc(OH)₂], 2, (Scheme 3). In this section, we have described the synthetic strategies for these important building blocks.

2.1.1 Synthesis of dichloro SiPc (SiPcCl₂). Linstead and co-workers attempted but failed to synthesize SiPc using a direct reaction of Pc with silicon tetrachloride (SiCl₄).¹ Later, in a seminal report, Joyner and Kenney envisioned a different synthetic route involving 1,2-dicyanobenzene or phthalonitrile (a precursor extensively used for synthesis of Pcs) instead of a metal free Pc as the starting material.²⁴ They exposed phthalonitrile to repeated treatments with SiCl₄ for 2 h using dry quinoline as a solvent and a reaction temperature of >200 °C (Route A, Scheme 1). Further purification using vacuum sublimation techniques resulted in formation of blue-green crystals which were presumed to be the desired dichloro SiPc analogue (SiPcCl₂), 1 (Scheme 1). Use of hexachlorodisiloxane instead of SiCl₄ as the source of silicon also resulted in improved yields of 1 under similar conditions, which was purified by Soxhlet extraction process (Route B, Scheme 1).¹³ However, similar attempts to synthesize 1 by reacting organo chlorosilanes such as methyltrichlorosilane and phenyltrichlorosilane with phthalonitrile were unsuccessful.¹³ Interestingly, the Si–N bonds exhibited unprecedented stability under extreme acidic or basic conditions.

Though Kenney and co-workers successfully isolated this extraordinary class of compounds, the initial synthetic routes were experimentally inconvenient due to poor yields and laborious purification methods. This entailed exploration of refined strategies with alternate starting materials and/or inclusion of additional reaction steps. Reaction of 2-cyanobenzamide (obtained from hydrolysis of phthalimide with concentrated ammonium hydroxide) with SiCl₄ in quinoline at elevated temperatures (~205 °C) resulted in the formation of a blue solid (Route A, Scheme 1).²³ The crude precipitate was washed with quinoline, benzene, pyridine, acetic acid, ethanol and ether to obtain the pure 1 (yield ~35%). Another two-step synthetic route comprised of ammonolysis of phthalonitrile in methanol and sodium methoxide to form 1,3-diminoisoindoline (DII) in the initial step, followed by reflux of DII with SiCl₄ in quinoline to afford compound 1 (yield ~71%) (Route B, Scheme 2).²⁵

Brusso and co-workers recently proposed an alternate route to synthesize DII from phthalonitrile by using air-stable lithium bis(trimethylsilyl)amide etherate, Li[N(TMS)]₂·Et₂O in dry toluene at room temperature for 5 h, followed by exposure to gaseous hydrochloric acid at ice bath temperature.²⁶ The dichloride salt, H₂DIICl₂, thus formed when refluxed with SiCl₄ in quinoline afforded 1 (yield ~51%). Among all these synthetic pathways, route B of Scheme 2 remains the most convenient and preferred choice for the construction of SiPcs.

The axial chloro groups of SiPc 1 can be replaced with other halogens. The chlorides are replaced by fluoride atoms on treat-
2.2 Synthesis of SiPCs with axial substitutions

2.2.1 Synthesis of symmetrical SiPCs with axial Si–OR bonds. Compounds 1 and 2 can be further fabricated by substituting the unique axial chloro and hydroxyl groups by alcohols, phenols, silanols, chlorosilanes and carboxylic acids to generate the compounds 3–5 (Scheme 4). The axial Si–Cl bonds in 1 undergo smooth reactions with alcohols and phenols to generate the corresponding symmetrically substituted bis-alkoxy derivatives, 3. Such reactions are reported with phenols (such as 2-naphthol, 1-naphthol, phenol, 4-hydroxybenzaldehyde, 4-aminophenol, 4-nitrophenol, 4-hydroxyproline and 3-hydroxypropridine) as well as other hydroxyl compounds (such as solketal, polyethylene glycols, 1,3-bis(dimethylamino)-2-propanol and mono-6-hydroxy permethylated β-cyclodextrin). In a typical reaction, 1 was refluxed with 3–7 times excess ROH (R = alkyl, aryl groups) at ∼120 °C in dry solvents like toluene or dichlorobenzene for about 24–96 h.14,20,27,31–38 Reactions proceeded under argon atmosphere in presence of ∼10 times excess of a base such as sodium hydride or dry pyridine (Scheme 4). Generally, small volumes of solvent were used to accelerate the precipitation of the blue coloured bis-alkoxy SiPC products. The yields of these reactions varied from 20–40% depending on the nature of the R groups.

In order to form a siloxane bond in the axial position, precursor 2 is refluxed with an excess of silanols or monochlorosilanes to obtain 4 (yields ∼40–70%).13,16,17,39–42 Reactions were usually carried out for short durations ranging from 3–6 h in non-polar solvents (e.g. dichlorobenzene and toluene) in presence of a mild base (e.g. pyridine) (Scheme 4).6,16,39 Interestingly, the axial Si–O–R linkages in certain bis-alkoxy SiPC derivatives, 3, can react with silanols under reflux conditions in dry pyridine to generate siloxane (Si–O–Si) bonds thereby forming SiPC derivatives.4,13,43 Alternatively, 4 were obtained by directly refluxing 1 with trialkylchlorosilanes in sodium hydroxide and chlorobenzene in presence of Aliquat HTA-1, a high temperature phase transfer catalyst (Scheme 4).14

Oligomeric and polymeric SiPC derivatives having axial Si–O and Si–O–Si linkages have been reported as well. Such μ-oxo bridged SiPC polymers were formed when compound 1 was allowed to react with 2 and the resulting hydroxy-capped oligomers were reacted with a chlorosilane.6 The desired polymer was isolated by column chromatography. Another way to obtain oligomers is from reaction of 1 or 2 with dendritic hydroxyl compounds (e.g. polyethyleneglycols and triethanolamines) or silanols containing multiple hydroxyl moieties in the axial position. The properties of these polymers were investigated in both solid and solvent phases and some of these derivatives were even processed into thin polymeric films.45–50

Attempts to react 1 with alternate ligand functionalities such as carboxylic acids were successful and expanded the domain of axial functionalization of SiPCs (Scheme 4).27 Esterification reactions of 1 occurred under reflux conditions in polar solvents such as 2-methoxyethyl ether (∼160 °C), diglyme (∼160 °C) or 1,4-dioxane (∼100 °C), DMF (∼160 °C) to produce various SiPC esters, 5. Unlike etherification, esterification reactions proceeded easily in the absence of a base. The reaction completion times ranged from as short as 30 min to as long as 48 h.20,51–56 The SiPC bis-esters, 5 were obtained as blue solids in moderate (∼30%) to high (∼90%) yields.
2.2.2 Synthesis of unsymmetrical SiPcs with axial Si–OR bonds. The strategies described in Scheme 4 have led to a wide array of axially functionalized symmetrical SiPcs, 3–5; however, in comparison to their symmetrical analogues, only a few SiPc derivatives with unsymmetrical axial Si–O bonds have been reported to date.

Kenney and co-workers noticed that only one of the two axial Si–O–Si linkages hydrolysed when 4a \( (R = [(CH_3)_2Si(CH_2CH_2CH_2N(CH_3)_2]) \) in 4) was stirred with trichloroacetic acid in dichloromethane at room temperature for 5 h (Scheme 5A).\(^{22,57}\) This hydrolysed product, 6 was then reacted with methanol in dry pyridine to form unsymmetrical SiPc, 7 in high yields of \( \sim 80\% \) (Scheme 5A). Another synthetic route to obtain 6 is strategized on exploiting the photolabile nature of Si–C bonds which will be discussed in the proceeding sections of organo SiPcs.\(^{58}\)

In an endeavour to synthesize and explore amphiphilic SiPc derivatives as PDT agents, Ng and co-workers serendipitously discovered a facile route to prepare unsymmetrical SiPc derivatives having axial Si–O linkages (Scheme 5B).\(^{59}\) In an attempt to purify a bis-alkoxy symmetrical SiPc, 3a \( (R = 1,3\text{-bis(di-methylamino)-2-propanol in 3}) \) by crystallization from methanol-chloroform solvent at room temperature, they observed the formation of an unsymmetrical SiPc, 8a in high reaction yields (\( \sim 73\% \)). Similar displacements reactions of 3a occurred with slight excess of ethanol and \( n \)-hexanol, but only under reflux conditions (\( \sim 70\; ^\circ\text{C} \)) in chloroform for 48 h to form 8b and 8c (yields \( \sim 29\%\text{–}37\% \)). In contrast, no mono-substituted unsymmetrical SiPc could be isolated when 3a was refluxed with dodecanol for 48 h. All these observations led to the conclusion that the length of the alkyl group of the alcohols played a crucial role in determining the ratio of the mono and di-substituted SiPc products, however their different reactivities were unaccounted. Recently, Schnemann and co-workers utilized this method and displaced one of the bulky axial ligands from 3a with biologically relevant phenols such as coumarins and combretastatin to obtain SiPcs, 8d and 8e (Scheme 5B).\(^{19}\) The use of phenols and prior neutralization of chloroform over anhydrous potassium carbonate possibly led to improved reaction yields of \( \sim 80\% \).

A recent report also utilized a similar displacement methodology to synthesize SiPc with unsymmetrical axial Si–OR bonds.\(^{60}\) Oligomeric ethylene glycols of various lengths were incorporated in one of the axial positions by refluxing a symmetrical amino-containing bis-phenoxy SiPcs with large excess (40\%) of desired glycols in toluene in presence of sodium hydride. Moderate yields of 23–31\% were achieved using this strategy.

In another report, Ng and co-workers used a slightly different strategy to synthesize SiPcs with unsymmetrical axial

![Scheme 5](image-url)
substituents, 9 (Scheme 6).\(^{61}\) Compound 1 was initially refluxed with various alkyl alcohols bearing sugar or diamino moieties in dry toluene and sodium hydride. After an hour of reaction, mono-6-hydroxy permethylated \(\beta\)-cyclodextrin and sodium hydride were added to the mixture and the reaction was continued for another 48 h. In these reactions, 1 was always taken in 1.5-folds molar excess of that of the alcohols to prevent exclusive formation of axially symmetrical SiPcs. However, the reaction yields of 9 were comparatively low (10–14\%).

SiPcs with unsymmetrical axial Si–O bonds have also been synthesized by selectively functionalizing one of the axial groups of a symmetrical SiPc. In a recent example, only one of the axial carboxylic acid groups in a symmetrical SiPc was subjected to a typical amide coupling reaction to form a series of unsymmetrical SiPc–peptide conjugates.\(^{62}\)

### 2.2.3 Synthesis of SiPcs with axial Si–C bonds

#### Symmetrical and unsymmetrical SiPcs with axial Si–C bonds

The formation and isolation of SiPcs with direct Si–C axial bonds have been comparatively difficult because of two prime reasons: (i) cationic nature of hexa-coordinated silicon which instead prefers bonding with electronegative elements such as oxygen and chloride and (ii) photo-instability of Si–C bonds.\(^{28}\) In a similar fashion, compounds having an axial C bond containing SiPcs were synthesized and isolated crude or pure 10 as green solids with yields of ~5–52\%. Use of allyl and benzyl trichlorosilanes in these strategies, resulted in formation of only bis-silyloxy ether SiPcs, 4 therefore, indicating the limited scope of axial Si–C ligands that can be installed in SiPcs.\(^{63}\)

Symmetrical SiPcs, 11 containing trans-bis Si–C linkages were obtained in high yields (~90\%) by refluxing 1 with 2–3 fold excess of Grignard reagents of choice (RMgI) in dry THF for several hours under inert and dark conditions (Scheme 7).\(^{49,64}\) However, only primary alkyl, aryl or alkyne groups (such as methyl, ethyl and phenyl) were successfully inserted using the Grignard reactions. Kumada et al. aptly combined the above two strategies and successfully prepared unsymmetrical SiPcs, 12 with direct Si–C bonds in axial positions.\(^{65}\) They reacted 10 with 6-fold excess of the corresponding Grignard reagent (R'MgI) in dry diethyl ether and stirred at room temperature overnight (Scheme 7). Hydrolysis of the intermediate magnesium adducts was carefully accomplished using aqueous hydrochloric acid at 0 °C. The pure products, 12 were obtained as dark green solids in high yields of ~85%. In contrast to the SiPc analogues with axial Si–O bonds, certain organo SiPcs exhibited better solubility in organic solvents such as chloroform and dichloromethane.

SiPcs with mixed axial Si–O and Si–C bonds. The axial chloride of 10 was hydrolysed with concentrated sulfuric acid, followed by addition of ice, to produce monoorgano monohydroxy SiPc derivatives, 13 [R'SiPc(OH)] (Scheme 8A).\(^{28}\) These hydroxyl derivatives were subsequently refluxed with phenols (e.g. phenol, 4-chlorophenol) in 1,2,4-trimethyl benzene for 30–90 min to yield SiPcs, [R'SiPc(OR)] with trans axial Si–C and Si–O bonds.\(^{28}\) In a similar fashion, compounds having an axial silyl group, 14 (where R' = silyl) was obtained by directly exposing crude 10 to siloxy ethers (R'SiOCH\(_3\)).\(^{22}\)

Though organo SiPcs were long discovered, appendage of suitable reactive handles in the Si–C bonds was not achieved until recently. Kenney et al. synthesized and isolated crude organo SiPc derivative, 10c (R = ClCH\(_2\)CH\(_2\)CH\(_3\)) in 10 where the axial Si–C ligand contained a reactive alkyl halide substituent (Scheme 7).\(^{22}\) Compound 10c was obtained by refluxing equimolar amounts of DII and 3-chloropropyltrichlorosilane in solvents like quinoline, 2,6-lutidine or DII–cyclodextrin (shown as cylindrical structure), sodium hydride, toluene, refluxed with various alkyl alcohols bearing sugar or diamino moieties, 63

![Scheme 6](image)

**Scheme 6** Synthesis of axially unsymmetrical Si–O bond containing SiPcs. Reagents and conditions: (i) (a) ROH, sodium hydride, toluene, r.t., 30 min, (b) 1, reflux, 1 h, (c) mono-6-hydroxy permethylated \(\beta\)-cyclodextrin (shown as cylindrical structure), sodium hydride, toluene, reflux, 48 h. R represents various alkyl moieties.

![Scheme 7](image)

**Scheme 7** Synthesis of axially symmetrical and unsymmetrical Si–C bond containing SiPcs. Reagents and conditions: (i) RSiCl\(_3\), dry quinoline, reflux, 90 min; (ii) (a) RMgI, dry diethyl ether, N\(_2\), r.t., overnight, (b) 6 M HCl, 0 °C. R denotes alkyl groups.
Room temperature rendered the hydroxyl SiPc analogues, of interesting synthetic routes for the preparation of unsymmetrically modified α modified at axial substitutions, the peripheral rings of SiPcs can be substituted to form SiPcs, by a siloxy ether, [(CH3)2Si(OCH3)(CH2CH2CH2N(CH3)2)] (Scheme 8A). Next, using the chloride group, [ClSiR3], was isolated in yield of 5% (Scheme 8A). Next, using the chloride group, 14a which was isolated in yield of ~5% (Scheme 8A). The presence of a nucleophilic thiol moiety dangling at the end of the Si–C axial bond created the possibility of functionalization of the organo SiPcs.

The photolabile axial Si–C bonds triggered the development of interesting synthetic routes for the preparation of unsymmetrical SiPcs with axial bis Si–O bonds. For example, compounds 14 (R' = silyl) on UV or visible light irradiation in trimethylamine saturated water and tolune solvent system at room temperature rendered the hydroxyl SiPc analogues, in high yields which were further derivatized with hydroxyl groups to form SiPcs, 7.

### 2.3 Synthesis of SiPcs with phthalocyanine ring substitutions

Like axial substitutions, the peripheral rings of SiPcs can be modified at α and β positions to perturb planarity, reduce aggregation, and increase hydrophilicity or to insert reactive handles for attachment of desired functionalities (Fig. 1C). More importantly, α/β substitutions are exploited to modulate the spectral properties of SiPcs. Various synthetic strategies have been developed for incorporation of halides, alkyl, alkoxy, phenoxy, phenylthio, sulphonate, glycols and vinyl groups at the peripheral and non-peripheral positions and often result in the formation of tetra- or octa-substituted symmetrical SiPcs. A few routes have been also been proposed for synthesis of SiPcs with unsymmetrical peripheral substitutions. However, the latter reactions are synthetically more challenging as they usually result in low synthetic yields and formation of mixtures of products which are difficult to separate.

#### 2.3.1 SiPcs with symmetrical phthalocyanine ring substitutions

The α/β decorated symmetrical SiPcs are primarily obtained from three major synthetic approaches: (i) direct reactions of the aromatic rings in SiPc, (ii) initial synthesis of substituted phthalonitrile precursors, followed by conversion to SiPcs via the corresponding DII derivatives and (iii) initial formation of a substituted Pc ring, followed by direct insertion of the core Si atom. These three strategies are discussed in detail below.

Direct reaction of the aromatic rings in SiPc. By means of a direct reaction, the aromatic rings of SiPc can be sulphonated, to obtain enhanced photostability as well as aqueous solubility. In a typical reaction, 1 was heated with excess chlorosulphonic acid (ClSO3H) at 130 °C for 5 h and then cooled to 85 °C. Thionyl chloride (30-fold excess) was slowly added to the solution and the reaction was continued for another 4 h. The resulting sulphonyl chloride SiPc adduct was isolated in pure form (yield ~68%) and then hydrolysed in methanol–water (1 : 5 v/v mixture) under reflux conditions for 3 h to obtain the tetrasulphonated SiPc, in high reaction yields of ~87% (Scheme 9). Importantly, the varying degrees of sulphonation depend on the SiPc: ClSO3H molar ratio and thus can be controlled. For example, use of SiPc and ClSO3H in 1 : 4 molar ratio resulted in isomeric mixtures of α or β substituted (Fig. 1) monosulphonated unsymmetrical SiPcs.

Using substituted phthalonitriles as precursors. The choice of substituted phthalonitriles, or 17 as precursors varied according to the nature and position of the desired substituents (Scheme 10). For synthesis of symmetrical β, β substituted SiPcs (22, where X = H), 4,5-disubstituted phthalonitriles, as precursors varied according to the nature and position of the desired substituents (Scheme 10). For synthesis of symmetrical β, β substituted SiPcs (22, where X = H), 4,5-disubstituted phthalonitriles, (16, where X = H) were used as precursors. Depending on the nature of Y groups, compounds 16 were synthesized from either 4,5-dibromo catechol or 4,5-dichloro dicyanobenzene. The subsequent steps for formation of SiPcs, 22 were common in both the cases and proceeded via the formation of the corresponding DII derivatives, 18 (Scheme 10). In some cases, purification of the SiPc chloro analogues were difficult and required further transformation of the axial chloro group to a hydroxyl or siloxy group.
Diether dicyano phthalonitriles (16a, where X = H and Y = OR) were obtained from 4,5-dibromo catechol by initial base mediated nucleophilic reaction of the hydroxyl groups and subsequent conversion of the bromides to cyanides via Rosemund-von Braun reaction. Lieberman et al. used this approach to obtain a crude product mixture containing octapentenyloxy SiPc derivatives, 22a (X = H; Y = pentenyloxy; Z = Cl in 22) via the DII derivative 18a. A similar approach was also adopted to obtain pentyloxy groups in the peripheral positions of SiPc (yield ∼45%).

The other route involved 4,5-dichloro dicyanobenzene as the starting material. The two chloro groups were readily substituted with nucleophiles in presence of a base in polar solvents such as DMF and DMSO and afforded phthalonitriles, 16 (where X = H) in a single step reaction. This approach has been utilized for the synthesis of SiPc variants bearing phenoxy (∼yield ∼67%). In another report, 4,5-dihalo dicyanobenzenes (Y = Cl/Br), as such, was converted to phthalonitriles, 16b and further reacted to form DII 18b and insert halogens in the peripheral positions of SiPcs, 22b (where X = H, Y = Cl/Br; Z = Cl in 22; yields ∼72–80%).

In another example, eight peripheral pentyl groups were incorporated in a SiPc structure (22c, where X = H; Y = pentyl; Z = CH3 in 22) using 1,2-dichlorobenzene as the starting material. In the first step, Grignard coupling of 1,2-dichlorobenzene with pentylbromomagnesium was performed to obtain the 1,2-dipentyl benzene. This product was initially subjected to bromination and next, the aromatic bromides were converted to cyanides using CuCN. Finally, the resulting phthalonitrile 16c was converted into DII derivative, 18c and subsequently to 22c with a reaction yield of ∼42%. Similarly, use of naphthalonitrile (2,3-naphthalenedicarbonitrile) instead of a phthalonitrile, generated silicon naphthalocyanines (SiNcs) with a reaction yield of ∼82%. The SiNcs, a subclass of SiPcs, absorb and emit in the NIR region owing to the extended π-conjugation of the aromatic rings.

SiPcs bearing α,α substituents (23, where Y = H in Scheme 10) were generated through the use of 3,6-disubstituted phthalonitriles (17, where Y = H). One such example has been reported by Nakagawa and co-workers. The synthesis involved prior conversion of 3,6-hydroxyphthalonitrile to 3,6-(bisphenylthio)phthalonitrile via tosylation of the hydroxyl groups and base-mediated displacement by thiophenol. The phthalonitrile 17, thus obtained was first converted to corresponding DII derivative, 19 and eventually to α,α-octaphenylthio SiPc derivative, 23a (X = phenylthio; Y = H; Z = OH in 23; yield ∼84%).

Direct insertion of the core silicon to a pre-formed Pc. A more flexible synthetic methodology to form SiPc variants at α/β positions involved initial formation of core element-free Pcs, followed by insertion of core silicon atom (Scheme 11). In a
3. Applications of silicon phthalocyanines

SiPc derivatives display interesting optical, redox, electronic properties and high thermal stability. In addition, availability of multiple synthetic strategies have resulted in enormous flexibility of SiPc frameworks. Easy modulation of solubility, optical and redox properties, bio-compatibility, bioactivity or solid-state structural arrangements has been achieved through decorating both axial and peripheral positions. This led to a multitudinous array of SiPc derivatives that have been crafted according to the demands for particular applications. The past decade has experienced an upsurge of novel applications of SiPcs in medicinal chemistry, photovoltaics, optoelectronics and photocatalysis. Selected examples of these emerging applications are discussed below.

3.1 Applications of SiPcs in medicinal chemistry

Silicon phthalocyanines have been extensively pursued in light-based medical applications, primarily as anticancer, antibacterial and antifungal agents.\(^{23,76-78}\) Owing to the strong NIR emission, SiPcs have been potentially applied for \textit{in vitro} and \textit{in vivo} bioimaging.\(^{32,52,79-81}\) Two fluorescent SiPc derivatives, namely La Jolla Blue® and IRD700DX®, are commercially available as labelling agents for biomolecules.\(^{82-84}\) Likewise, due to the intense far red and NIR Q-band absorptions, several photosensitive SiPc molecules have been deployed in photodynamic therapy (PDT), photounescing techniques, photoimmunotherapy (PIT), and photothermal therapy (PTT). A few recent noteworthy examples are highlighted in the subsequent sections.

3.1.1 Incorporation water soluble axial or peripheral ligands. The most sought after modification of SiPcs has been to reduce aggregation tendencies and overcome the stinted solubility, a factor that restrains the biological applications of most Pcs. One extensive methodology pursued to improve the aqueous solubility is the incorporation of nitrogen rich substituents in the axial or peripheral positions of SiPcs, which can be readily alkylated to give quaternary ammonium salts using methyl iodide, dimethyl sulfate or 1,3-propanedialates to obtain cationic water soluble SiPc derivatives, mostly suitable for biological applications (Scheme 12).\(^{15,17,19,31,59,60,85,86}\)

3.1.2 SiPcs in photodynamic therapy (PDT) of cancer. Photodynamic therapy is a minimally-invasive anticancer modality which typically involves a local or systemic administration of a photosensitizer followed by activation in the cancerous tissues with light irradiation.\(^{87,88}\) SiPc based compounds are second generation photosensitizers which display moderate to high quantum yields of NIR light induced generation of singlet oxygen, a short-lived and highly cytotoxic species. The promising activity of Pc-4 (6) in clinical trials\(^{89}\) has sparked renewed interest in researchers to improve efficacy of Pc-4.\(^{90-94}\) In parallel, several approaches were adopted to increase the aqueous solubility and bio-compatibility leading to development of new SiPc photosensitizers. For example, a typical reaction, phthalonitriles (16 or 17) were refluxed with lithium carbonate (~4-fold excess) in 1-pentanol for 66 h to obtain Pcs (20 or 21) in ~30% reaction yields.\(^{71}\) This conversion proceeded faster (~48 h) and efficiently (high reaction yields ~55–72%) in presence of non-nucleophilic strong bases like 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU).\(^{68}\) In the final step, these Pcs (20 and 21) were reacted with 10-30-folds excess of trichlorosilane (HSiCl\(_3\)) to form the desired SiPcs with symmetrical \(\alpha/\beta\) substituents and hydroxyl axial ligands, 23 and 24 (Scheme 11). Reactions were conventionally carried out in anhydrous and inert atmosphere using organic solvents such as DCM, tetrahydrofuran, acetonitrile or toluene at room temperature or under reflux conditions.\(^{70}\) The nature of the base (such as tripropylamine, tributylamine and DBU) employed affected both reaction completion times and yields (~12–53%) of desired SiPcs (23 or 24).\(^{72,73}\)

2.3.2 SiPcs with unsymmetrical phthalocyanine ring substituitions. Mono or di-substituted unsymmetrical SiPcs are synthesized either directly by condensing two different phthalonitriles or indirectly by initial conversion of these phthalonitriles to the corresponding DII derivatives followed by condensation. In the former strategy, an unsymmetrical Pc is first obtained which is then subjected to direct insertion method to insert a silicon atom in the unsymmetrical Pc core. In almost all the cases, the axial chlorides were replaced by hydroxyl or siloxy groups for successful isolation of the peripherally unsymmetrical SiPcs in pure form. The final reaction yields of pure SiPcs varied from 3–27%. This technique led to the synthesis and isolation of SiPcs bearing a reactive handle only in one of the four benzene rings (Fig. 2). This enabled incorporation of functional groups at the periphery and led to the development of SiPc molecules with remarkable biological properties.\(^{41,74,75}\)

In conclusion to this section, all the aforementioned strategies provided diverse reaction schemes to fabricate SiPcs with ligands of choice such as acidic, basic, neutral, hydrophilic, lipophilic, electron rich or electron withdrawing substituents. This led to exploitation of the NIR optical properties of SiPc in various contexts.

![Fig. 2 Chemical structure of a SiPc polymeric derivative, La Jolla Blue with unsymmetrical peripheral (●) substitutions.](image-url)
SiPc derivative with two bulky permethylated cyclodextrin moieties in the axial positions displayed reduced self-aggregation in the cell culture media and remarkable *in vitro* and *in vivo* PDT effects. Likewise, SiPcs conjugated axially with biomolecules such as nucleosides and serum albumin exhibited enhanced cellular uptake. Furthermore, incorporation of tumor targeting moieties such as erlotinib, hyaluronic acid, and biotin in the SiPc framework resulted in excellent tumor specific uptake and PDT effects. In other approaches, SiPc molecules have been covalently linked with chemotherapeutic drugs to generate chemo-photodynamic conjugates having dual mode of cytotoxic action.

Recently, You’s research group developed SiPc pseudoprodrugs to execute multipurpose objectives such as (i) fluorescence imaging, (ii) far red light-mediated uncaging, and (iii) combined effects of singlet-oxygen induced PDT and local chemotherapy. They utilized singlet oxygen-labile amionoacrylate bonds to append clinical chemotherapeutic drugs such as combretastatin and paclitaxel in the axial positions of SiPcs (3b and 3c in Fig. 3A). The caged forms 3b and 3c exhibited significantly reduced dark cytotoxicity as evident from 19 and 190-folds lower IC50 values when compared to the respective free drugs. On light irradiation, SiPc produced singlet oxygen which led to an immediate PDT effect and simultaneous cleavage of the aminoacrylate linker to un cage the intact chemotherapeutic drugs. This bystander or local chemotherapeutic effect after the illumination was demonstrated by exposing only one-half of each well containing cancer cells treated with vehicle only, 3b or a structurally equivalent non-pho tocleavable SiPc conjugate, NCL-3b (Fig. 3B). The PDT effect of NCL-3b was restricted only in the light exposed half of the well due to the short half-life (10–320 ns) and limited diffusion distance (20–200 nm) of singlet oxygen (panel c of Fig. 3B). However, the 3b-treated wells showed prominent bystander effects. The photorelease combretastatin diffused throughout the well and caused similar toxicity in the light unexposed cells as compared to light-exposed cells (panel b of Fig. 3B).

Compound 3b was further modified to incorporate a tumor-targeting folic acid at the second axial position via a hydrophilic PEG linker. It was observed that the longer length of the PEG spacer enhanced specific uptake and localization in folate receptor overexpressing colon 26 tumors. The SiPc conjugate having PEG linker length of ~45 showed the most impressive light induced cytotoxicity giving a IC50 value of 1.65×10-8 M (690 nm, 10 J cm-2 for 15 min). At these concentrations, 3b and NCL-3b did not produce any significant dark toxicity. Reproduced with permission from ref. 101. Further permissions related to the material excerpted should be directed to the ACS (https://pubs.acs.org/doi/10.1021/jm5000722).

3.1.3 NIR dissociation of axial ligands of SiPcs. An interesting feature of SiPc is the photolability of unique axial Si–C and Si–O bonds. Though this photochemical degradation of SiPcs was long-established, it has been only recently exploited in medicinal chemistry to photorelease PDT and chemotherapeutic anticancer agents selectively in a cancer microenvironment.

3.1.4 SiPcs as photocages. Burda et al. utilized the photosensitive Si–C axial bond for NIR light-controlled delivery of Pc-4. They judiciously chose PEGylated gold nanoparticles (Au NPs) as drug-delivery systems due to their excellent biocompatibility and superior drug loading capacity. Facile syn-
thesis of stable covalent assemblies of SiPc and AuNP, Au NP-14b was executed by taking advantage of the thiophilic nature of gold and the free thiol group of 14b. On red light exposure, the Si–C bond in PEGylated Au NP-14b rapidly dissociated to release Pc-4 molecules (Fig. 4). Interestingly, the drug can only be photo-released in lipophilic environments. The Au NPs displayed only ~1% cellular uptake and insignificant cellular toxicity. Therefore, as expected, Au NP-14b conjugates caused no change in viability of HeLa (cervical cancer) cells in the dark. In contrast, almost 80% of the population were killed in Au NP-14b (1 μM) treated HeLa cells which were illuminated (>600 nm, 1 J cm⁻², 12 min). This toxicity profile was comparable to that of unformulated Pc-4, thereby suggesting the photorelease of Pc-4 from the nanoparticle conjugates.

The research groups of Kenney and Burda performed theoretical and experimental measurements to identify the preferred mechanistic pathway of the light triggered Si–C bond cleavage process. Characterization of the photoproducts using various techniques such as steady state spectroscopy, electrospray ionization mass spectrometry, liquid chromatography mass spectrometry, nuclear magnetic resonance and electron paramagnetic resonance hinted towards formation of a relatively long-lived SiPc radical as the key intermediate which consequently undergoes solvolysis. Computational studies additionally confirmed homolytic bond cleavage process involving radical intermediates. Bond-dissociation energy (BDE) calculations indicated that homolytic cleavage (~50 kcal mol⁻¹) is energetically more favoured than heterolytic cleavage (~190 kcal mol⁻¹). This is due to the fact that the unpaired electron is largely delocalized over the SiPc aromatic rings as implied by natural bond orbital (NBO) analysis and spin density calculations.

In another recent report, Schnermann et al. demonstrated the preferential photocleavage of Si–O bonds and release of phenols in axially unsymmetrical SiPcs molecules. On NIR photoexposure at the SiPc Q-band (690 nm), the axial Si–O bonds in 8d and 8e (Scheme 5, above) instantly dissociated in oxygen-deprived physiological conditions and in presence of reducing glutathione (GSH, 5 mM). Thus, photo uncaging of small phenolic molecules such as fluorescent 4-methyl umbelliferone or clinical anticancer drug combretastatin was achieved (Fig. 5A). In contrast, compounds 8d and 8e were fairly stable on light irradiation under normoxic conditions and produced singlet oxygen as the major reactive oxygen species (Fig. 5A). Thus, compound 8e resulted in NIR mediated cytotoxicity in cancer cells which was a manifestation of either PDT or chemotherapy depending on the availability of tissue oxygen. Compound 8e (200 nM) showed NIR (690 nm, 50 J cm⁻²) induced phototoxicity by a PDT mechanism in presence of oxygen leading to 48% cell growth inhibition. In hypoxic conditions, the cell growth inhibition increased to 80% which is comparable to that of free combretastatin suggesting photo-release of the chemotherapeutic agent from the SiPc molecule triggered by NIR light and cytosolic glutathione (Fig. 5A).

Detailed spectroscopic and computational studies were performed to understand the photoredox uncaging reactions. The proposed mechanistic pathway involved a key radical anion intermediate formed through photoinduced electron transfer from the SiPc triplet state (Fig. 5B). This explained the role of reducing agent, glutathione in accelerating the cleavage phenomenon. The transient radical anion was proposed to undergo solvolytic ligand exchange through a heptacoordinate silicon transition state.

3.1.5 SiPcs in photoimmunotherapy. In their seminal paper, Kobayashi and co-workers employed monoclonal antibody–SiPc conjugates, mAb-IR700 (25) in photoimmunotherapy (PIT), a novel molecular-targeted cancer modality that combines PDT and immunotherapy (Fig. 6A). The conjugates...
25 comprised of a cell-impermeable, hydrophilic, NIR absorbing SiPc photosensitizer (IR700 dye) which was covalently tethered to monoclonal antibodies targeting human epidermal growth factor receptors (HER or EGFR). It is important to note that one such SiPc derivative, where IR700 is conjugated to cetuximab, recently gained approval as a NIR PIT drug by the Japanese government for treatment of head and neck cancers. The cell membrane-bound conjugates showed tumor-specific cytotoxicity in HER expressing tumor xenografts only when exposed to NIR light. Interestingly, unlike PDT, internalization of the photosensitizer was not required in PIT and the light induced cytotoxicity was only partially derived from reactive oxygen species. In another report, Mitsunaga et al. combined an antibody (Trastuzumab, T), a chemotherapeutic drug (maytansinoid, DM1) and NIR absorbing SiPc photosensitizer (IR700) to generate bifunctional antibody-drug-photoabsorber conjugates (T-DM1-IR700). In vivo studies with these conjugates showed HER-2 targeted tumor cytotoxicity via dual action NIR-PIT and chemoinmunotherapy. Remarkably, T-DM1-IR700 conjugates exhibited cytotoxicity in deep-seated tumors where NIR-PIT by itself fails due to insufficient NIR light penetration.

Recently, Kobayashi and co-workers unravelled the distinct mechanism of PIT for SiPc-antibody conjugates. They observed NIR light induced dissociation of the axial Si–O–Si linkages in 25 in presence of an electron donor such as ascorbic acid sodium salt, NaAA (Fig. 6B). The NIR light triggered detachment of hydrophilic sulphonated axial ligands led to an increase in hydrophobicity of and consequently resulted in aggregate formation and quenching of NIR fluorescence. Atomic force microscopy (AFM) revealed alteration in the protein conformation such as enlargement of the conjugates and loss of normal Y-shape of the antibody. Thus, the photoinduced degradation not only affected the physical
shape and solubility of the covalently-conjugated antibody, but also influenced the antibody–antigen complexes. These profound morphological changes were speculated to enhance the transmembrane (TM) water flow which created physical stress within the cellular membrane and eventually led to cell bursting and necrotic/immunogenic cell death (ICD) (Fig. 6C).

3.1.6 SiPc in photothermal therapy. Photothermal therapy (PTT) is an oxygen-independent anticancer treatment in which appropriate light energy is absorbed by photosensitive molecules and converted into thermal energy, ultimately leading to hyperthermia mediated cancer cell death.87 Huang and co-workers designed a SiPc derivative, PCB1 or 3d and studied its photothermal behaviour (Fig. 6D).110 The two nitrogen-rich axial substituents effectively fostered photoinduced electron transfer (PET) mechanism and lowered the fluorescence and singlet oxygen generation quantum yields of SiPc, 3d. Temperature escalated by 28 °C on laser irradiation (730 nm) of aqueous solutions of 3d. However, such photothermal characteristics were lost upon protonation of the axial ligands highlighting the crucial roles of nitrogen lone pairs in the PET process. NIR exposure of 3d-treated human liver cancer cells (HepG2) led to ~70% decrease in cell viability indicating its ability to act as a unique PTT agent (Fig. 6E). Similar viability assays performed under ice-bath conditions showed only ~30% loss in cell viability which might be attributed to the PDT effect of 3d (Fig. 6E).

3.2 Applications in organic photovoltaic solar cells

SiPcs have recently generated interest for multifunctional applications in dye-sensitized (DSSCs), bulk heterojunction (BHJ) and planar heterojunction (PHJ) solar cells as additives, acceptors, donors and crosslinking components.111 The intense absorption of chemically and thermally robust SiPcs and SiNcs in the red and far-red spectral region has been utilized to expand the light harvesting window and boost the overall photoconversion efficiency (PCE). The structural tunability has led to easy integration and multi-functional behaviour. Finally, the low toxicity and the elemental abundance of silicon in nature have particularly advocated for their candidacy in organic photowatts and optoelectronics.

3.2.1 SiPc as additives in solar cells. Honda et al. pioneered the use of a bis(tri-n-hexylsilyl oxide) substituted SiPc derivative, 4b as an ternary additive in a P3HT/PCBM [(poly(3-hexylthiophene))/(phenyl-C66-butyric acid methyl ester)] BHJ solar cell and observed a positive influence of the dye in overall device performance (Fig. 7).112 The values for short-circuit current ($J_{sc}$), open-circuit voltage ($V_{oc}$) and the fill factor (FF) of 10.3 mA cm$^{-2}$, 0.57 V and 0.69 respectively were achieved at optimized loading content of 4.8 wt% for 4b (Table 1, Fig. 7). The presence of the dye boosted PCE to 4.1. Transient absorption spectroscopy and surface energy measurements revealed the presence of a cascaded photoconversion mechanism in which facile charge transfer occurred from P3HT to PCBM via SiPc molecules aided by their intermediate energy levels and selective localization in a disordered P3HT domain and at the polymer–fullerene interface.112–114 Hence, the overall device performance was governed by the blend morphology which in turn depended on both dye structure and dye loading content.

Encouraged by these inferences, Bender and co-workers varied the alkyl spacer of the axial silyl groups of dye 4b and observed that use of smaller molecular fragments such as tri-n-butyl silyl (4c) and tri-isopropyl silyl (4d) in the axial positions led to ~10% increase in $f_{sc}$ values when compared to 4b (Fig. 7, Table 1).40 These experimental outcomes led to the conclusion that the SiPc dyes with contrasting properties of high solubility and extreme tendency to crystallize favoured charge transfer and augmented the overall device efficiency. A more rigorous study using eight SiPc variants with axial alkylsilyl groups of different chain lengths has been recently reported.44 It was observed that SiPc variants displayed similar optical, electrochemical and hydrophobicity properties and frontier orbital energy levels. However, the alkyl chain length had a pronounced effect in molecular packing and solubility of the SiPc dyes and thus governed the overall device performance. Best PCE values were achieved in cases where solubility of the ternary additive SiPc dye matched closely to that of P3HT.

In parallel, researchers explored SiPc structural variants with axial functionalities other than siloxy ethers, such as alkyl/aryl ethers and esters, as alternate additives in BHJ solar cells. Bender and co-workers investigated the device performance of bis(3-pentadecylphenoxy)-SiPc, 3e (Fig. 8).30 Albeit having similar spectral features like 4b, the dye 3e failed to significantly increase the $J_{sc}$ and the efficiency ($\eta_{ef}$) of the polymer–fullerene based ternary cascade BHJ solar cell (Table 1). Compound 3e exhibited lower tendencies to crystallize and migrate to the P3HT/PCBM interface as compared to 4a. The authors speculated that these factors might be responsible for its inferior performance.

In another approach, Ameri et al. synthesized SiPc derivatives with bulky t-butyl groups in the periphery and flexible axial pyrene carboxylic acid groups to fabricate ternary BHJ
P3HT/PCBM solar cell devices.\textsuperscript{115} The ternary devices with these variants displayed an increase of over 20% in $J_{SC}$ and over 50% in PCE values compared to the baseline binary device. Among the variants, best fill factor and charge carrier mobility were obtained for the device constructed with 5a (Fig. 9, Table 1).

In an attempt to further increase the light harvesting window up to 800 nm, SiNcs, having appropriate energy levels, were deployed alone or with SiPcs (Fig. 10).\textsuperscript{112,116} The research group of Sellinger used a symmetrical 2,3-SiNc derivative, 26, with peripheral t-butyl groups and two axial trihexylsilyloxide groups as the only light harvesting additive in a ternary P3HT/PMBC solar cell (Fig. 10A).\textsuperscript{116} The presence of bulky t-butyl functional groups conferred higher solubility to 26 and the BHJ morphology of P3HT/PCBM was preserved even with excess dye incorporation. Thus, higher optimal dye loading contents up to 20% was achieved without causing a decrease in $J_{SC}$. The blending of 26 resulted in $\sim$40% enhancement of $J_{SC}$ values and a PCE of up to 4.9% was reported (Table 1).

In another example, Honda et al. blended a SiNc dye, 27 along with 4b, as a fourth component in the polymer–fullerene-SiPc solar cell (Fig. 7, 10A, B and C).\textsuperscript{112} The quaternary

![Chemical structures of SiPc derivatives, 3e–3j.](image)

![Chemical structures of bis-ester SiPc derivatives, 5a–5h.](image)

Table 1  The performances and device characteristics of photovoltaic devices containing various SiPc and SiPc derivatives as additives, donor or acceptor materials

<table>
<thead>
<tr>
<th>Type of solar cell</th>
<th>Donor/acceptor pair</th>
<th>Additive dye(s)</th>
<th>Loading (wt%)</th>
<th>$J_{SC}$/mA cm$^{-2}$</th>
<th>$V_{OC}$/V</th>
<th>FF</th>
<th>PCE (%)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>—</td>
<td>—</td>
<td>8.48</td>
<td>0.57</td>
<td>0.56</td>
<td>2.74</td>
<td>30</td>
</tr>
<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>4b</td>
<td>10.3</td>
<td>0.57</td>
<td>0.69</td>
<td>4.1</td>
<td>112</td>
<td></td>
</tr>
<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>4b</td>
<td>9.84</td>
<td>0.57</td>
<td>0.59</td>
<td>3.29</td>
<td>30</td>
<td></td>
</tr>
<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>4c</td>
<td>10.0</td>
<td>0.36</td>
<td>0.59</td>
<td>3.30</td>
<td>40</td>
<td></td>
</tr>
<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>4d</td>
<td>10.1</td>
<td>0.57</td>
<td>0.60</td>
<td>3.40</td>
<td>40</td>
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<tr>
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<td>3e</td>
<td>9.49</td>
<td>0.59</td>
<td>0.54</td>
<td>3.02</td>
<td>30</td>
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<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
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<td>20.02</td>
<td>0.65</td>
<td>0.63</td>
<td>4.14</td>
<td>115</td>
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<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>26</td>
<td>8.0</td>
<td>11.4</td>
<td>0.62</td>
<td>4.46</td>
<td>116</td>
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<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>4b/27</td>
<td>4.8/1.5</td>
<td>10.9</td>
<td>0.57</td>
<td>0.69</td>
<td>4.3</td>
<td>112</td>
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<tr>
<td>BHJ</td>
<td>P3HT/PCBM</td>
<td>5b/28</td>
<td>7.5/5</td>
<td>10.80</td>
<td>0.65</td>
<td>0.55</td>
<td>3.89</td>
<td>117</td>
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<tr>
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<td>28</td>
<td>10</td>
<td>9.72</td>
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<td>0.58</td>
<td>3.68</td>
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<tr>
<td>DSSC</td>
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<td>4.5</td>
<td>118</td>
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<td>0.99</td>
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<td>0.78</td>
<td>40</td>
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<tr>
<td>BHJ</td>
<td>P3HT/4d</td>
<td>—</td>
<td>3.0</td>
<td>0.84</td>
<td>0.43</td>
<td>1.07</td>
<td>40</td>
<td></td>
</tr>
<tr>
<td>PHJ</td>
<td>1/C$_{60}$</td>
<td>—</td>
<td>0.47</td>
<td>0.24</td>
<td>0.32</td>
<td>0.04</td>
<td>36</td>
<td></td>
</tr>
<tr>
<td>PHJ</td>
<td>3f/C$_{60}$</td>
<td>—</td>
<td>1.4</td>
<td>0.76</td>
<td>0.31</td>
<td>0.34</td>
<td>36</td>
<td></td>
</tr>
<tr>
<td>PHJ</td>
<td>3g/C$_{60}$</td>
<td>—</td>
<td>4.6</td>
<td>0.87</td>
<td>0.44</td>
<td>1.8</td>
<td>119</td>
<td></td>
</tr>
<tr>
<td>PHJ</td>
<td>α-6 T/3h</td>
<td>—</td>
<td>3.5</td>
<td>0.66</td>
<td>0.43</td>
<td>1.0</td>
<td>119</td>
<td></td>
</tr>
<tr>
<td>BHJ</td>
<td>PTB7/5c</td>
<td>—</td>
<td>0.60</td>
<td>1.00</td>
<td>0.45</td>
<td>0.27</td>
<td>120</td>
<td></td>
</tr>
<tr>
<td>BHJ</td>
<td>PTB7/5d</td>
<td>—</td>
<td>6.18</td>
<td>1.03</td>
<td>0.42</td>
<td>2.67</td>
<td>120</td>
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</tr>
</tbody>
</table>

C$_{60}$ is fullerene. α-6 T is α-sexithiophene. PTB7 is an organic semiconducting polymer. All values are taken from the references as listed in the last column.
3.2.2 SiPc as acceptors and donors in photovoltaic solar cells. The promising traits of the SiPc dyes as additives prompted researchers to explore their donor and acceptor properties. Preliminary assessment of few SiPc variants (4b, 4c and 4d) as possible acceptors in binary blends P3HT/SiPc was carried out for the first time by Bender and co-workers (Fig. 7).

The studies led to the conclusion that replacement of PCBM by SiPc results in ~50% increase in Voc values, but a much lower PCE efficiencies of only ~1% when compared to the standard P3HT/PCBM donor acceptor pair (Table 1). In another report, the electron donating and accepting dual-nature functionalities of SiPcCl2, 1 and bis(pentafluorophenoxo)SiPc, 3f (Fig. 8) in un-optimized planar heterojunction (PHJ) solar cell devices were demonstrated (Fig. 8, Table 1). Compound 3f displayed superior donor and acceptor properties which improved the device performance as compared to 1. These striking device characteristics of 3f were partially attributed to the solid-state arrangement and improved π-shielding as evident from its X-ray crystal structure. On the contrary, such π–π interactions were completely missing in the crystals of 1. It was further noted that the frequency and position of the fluorine atoms in the axial phenoxy group governed the dual-characteristics of the SiPc derivatives. The crude structure-activity analysis revealed SiPc analogues, 3g and 3h as potential donors and acceptors, respectively for future device configuration (Fig. 8, Table 1).

Samuel and co-workers investigated symmetrical SiPcs with axial bis-esters of eicosanoic acid (5e) and 3,5-di-tert-butylbenzoic acid (5d) as NIR absorbing dyes and electron acceptors in binary BHJ organic solar cells (Fig. 9, Table 1). The best device characteristics were obtained on using PTB7, a dithiophene based polymer and 5d as the donor acceptor pair at a 1:1.5 weight ratio and were attributed to the increased photocurrent and strong visible absorption of 5d.

3.2.3 SiPc as crosslinking agent in photovoltaic solar cells. Lessard et al. reported a bis(6-azidohexanoate)SiPc dye, 5e bearing crosslinking groups. The incorporation of this dye in a P3HT/PCBM BHJ solar cell led to dual functions as a ternary additive and an effective crosslinker (Fig. 9). The baseline P3HT/PCBM binary device (without 5e) suffered significant loss in Jsc and PCE values on exposure to a thermal ageing of 150 °C for 23 h owing to the phase separation and crystallization of the PCBM. Loading of the dye, 5e in the device boosted the photocurrent generation in the near-infra red region implying a synergistic additive influence. Interestingly, the crosslinking ability of 5e via the axial azides, preserved the BHJ morphology and retained the overall performance (PCE) by 97% after thermal ageing (Fig. 12).

3.4 Applications of SiPc in OLEDs

SiPcs exhibit intense and narrow NIR luminescence spectra and thus find applications as emitters in organic light emitting diodes (OLEDs). Samuel and co-workers recently studied the optoelectronic properties and the electroluminescence of two SiPc derivatives, 5e and 5d (Fig. 9). Both derivatives displayed electroluminescence features in OLEDs similar to their
photoluminescence profiles with a maxima around 698–709 nm and a narrow full width at half maxima (FWHM) of 21–27 nm. It was observed that doping of the emitting layer of the device with 10% 5e resulted in better quality of OLEDs with fewer pinholes and reduced roughness as compared to 5d indicating that structure of the axial ligands influence device characteristics (Fig. 13).

Instead of using long chain carbons to reduce aggregation and confer solubility, Greenham and co-workers chose carbon-efficient and electron withdrawing pentafluorophenoxy moieties as the axial ligands.122 They deployed the SiPc derivative, 3f as dopant red emitters in both solution-processed and vapour-deposited OLEDs (Fig. 8). Compound 3f displayed a photoluminescence quantum yield value of 0.66 and a band gap of 1.77 eV. The OLED devices constructed with a polymeric host and 3f as a guest showed narrow electroluminescence spectra (FWHM of 20 nm) with a maximum external quantum efficiency (EQE) value of 2.5%. The vapour-deposited host: guest OLEDs incorporating 5 wt% doped 3f displayed similar emission profile with maximum around 715 nm, albeit, with a much lower EQE value of ~0.09%. Though the energy transfer was clearly confirmed in this device, low EQE values indicated that a proper choice of host is crucial in order to achieve a significantly higher dopant emission over the host emission.

3.5 Applications of SiPc in OTFTs

Like other nitrogen heterocycle-based compounds,123,124 certain SiPcs also possess electronic properties of n-type semiconductors.26 Additionally, the readily tunable axial ligands, high field-effect mobility, robust environmental stability and ease of processability make them ideal to perform as organic thin film transistors (OTFTs).

Lessard et al., first deployed SiPcs as n-type semiconductors in bottom-gate bottom-contact (BGBC) OTFT devices.125 SiPc bis-esters with 1-benzoic (5f), 1-naphthoic (5g) and 9-anthroic (5h) acid axial pendants were explored to establish a link between solid state arrangements of the semiconductors and the overall device performances (Fig. 9). The varying degree of π-bonding interactions of the Pc chromophores was expected to partly affect the 2D charge transfer pathways in the devices. The crystal structures revealed that 5f and 5g showed significant parallel stacking of the Pc rings. This probably led to higher electron field effect mobility (μe) and increased on-current (I_on) in devices with 5f and 5g. Although 5g revealed the closest π–π interactions, its device performance was lower than that of 5f. On the other hand, the bulky anthracene group hindered the overlap of the Pc rings and devices with 5h displayed lowest μe and I_on values. However, devices with 5h exhibited the lowest threshold voltage (V_T). This is possibly because the lowest unoccupied molecular orbital (LUMO) of 5h was energetically comparable to the work function of gold. Overall, 5f outclassed the other SiPcs in the series in terms of device performance. Further increment of device performance of 5f was achieved with octytrichlorosilane (OTS) dielectric modification and a substrate temperature of 200 °C maintained during deposition.

Motivated by these results, Lessard et al., investigated the behaviour of fluorophenoxy derivatives of SiPc as n-type transistors.126,127 It was noticed that the frequency and position of fluorine atoms affected electron mobility and the compounds 3f–5h exhibited some ambipolarity of charge transport (Fig. 8). In the series, devices built with 3f deposited on OTS-modified surface displayed superior performance. Taken together, these reports demonstrated the potential roles of axial substituents in governing the device performances and hinted towards a possibility of obtaining high performance SiPc based n-type semiconductors.

3.6 Applications in photocatalysis

The design of tetrapyrrrole complexes and investigation of their photocatalytic behaviour stemmed from the key roles played by naturally occurring tetrapyrrrole complexes (e.g. chlorophyll) in photosynthesis.128,129 In this context, SiPcs are recently
being explored for potential applications in solar energy conversion and homogeneous photocatalysis. The fulfilling criteria of SiPcs to function as photocatalysts include: (i) robust photostability, (ii) intense absorption features in the solar spectrum which are tunable at the molecular level and (iii) ability to initiate photoinduced electron transfer reactions.130

Knör and co-workers synthesized a cationic, water soluble SiPc, 3i and investigated its potential for construction of hydrogen evolving photocatalytic systems (Fig. 8, see above).131 Compound 3i reacted with aerial oxygen predominantly through a type II sensitization mechanism involving formation of singlet oxygen. Trace amounts of 1,4-benzoquinone and triethanolamine efficiently quenched the fluorescence signal of 3i following a Stern–Volmer kinetics of bimolecular quenching mechanism. Excess of triethanolamine led to rapid accumulation of permanent photoredox products of 3i. This confirmed the presence of a photoinduced electron transfer process possibly involving reduced Pc π-radical anion intermediates. Such radical species play important roles in mediating photochemical water splitting, thereby making SiPcs attractive candidates for construction of solar energy driven catalytic systems.

In another study, the catalytic behaviour of another SiPc derivative, 3j, was investigated using the 4-nitrophenol degradation probe reaction (Fig. 8, see above).132 Visible light induced degradation of 4-nitrophenol was observed with catalytic amounts of 3j and hydrogen peroxide as a source of oxygen. Hydroquinone and benzoquinone were determined as the major and minor photoproducts respectively. A substrate/oxygen/catalyst ratio of 2000/500/1 resulted in 88% conversion with a turnover number of 1758 and turnover frequency of 586 for 3j. Interestingly, photoconversion (37%) of 4-nitrophenol was observed without hydrogen peroxide, indicating the ability of 3j to act as a photocatalyst in absence of any oxygen source.

Recent reports exploited covalently functionalized SiPc-graphene based organic hybrid composites as improved solar catalysts.133,134 SiPc covalently functionalized with N-doped ultrasmall reduced graphene oxide (N-usRGO/SiPc) displayed strong absorption in the visible spectrum.133 The fluorescence quenching (90%) of SiPc and improved photocurrent responses for N-usRGO/SiPc demonstrated the efficient photoinduced electron transfer process from SiPc to N-usRGO through covalent bonds (Fig. 14A). These properties attributed to the SiPc and improved photocurrent responses for N-usRGO/SiPc demonstrated the efficient photoinduced electron transfer process from SiPc to N-usRGO through covalent bonds (Fig. 14A). These properties attributed to higher visible light induced catalytic activity of (N-usRGO/SiPc) in comparison to SiPc free N-usRGO. Platinum nanoparticles (Pt NP, 5 wt% Pt) deposited on N-heteroatoms of graphene acted as cocatalyst and further enhanced the catalytic performance of the composite (Fig. 14B). Pt NP decorated N-usRGO/SiPc construct produced a maximum of 4.5 μmol mg⁻¹ of hydrogen from water on 6 h of UV-vis irradiation. The proposed mechanism suggests sequential shuttling of electrons from light harvesting SiPc to graphene layers and consequently to Pt NP where water molecules accept electrons and evolve hydrogen (Fig. 14B). Finally, triethanolamine (TEA) donates electron to excited SiPc and brings it back to the ground state (Fig. 14B).

Lu et al. observed that addition of surfactants like cetyltrimethylammonium bromide (CTAB) prevented aggregation and boosted the photocatalytic behaviour of a SiPc(phenyl) covalently functionalized and Pt NP loaded graphene construct, SiPc(phenyl)₂G₂/Pt (Ref: IntJHydE2016).134 The integrated photocatalytic system, SiPc(phenyl)₂G₂/Pt/CTAB produced 18.2 μmol mg⁻¹ of hydrogen on 10 h or UV-visible irradiation. These studies together established the potential of SiPcs in construction of water splitting photocatalytic systems.

4. Conclusions and perspectives

Despite their long existence, SiPcs continue to attract unabated attention due to their emerging properties and applications. Like most other phthalocyanines, SiPcs have gained profound recognition as far-red and NIR photodynamic therapeutic and fluorescence imaging agents. Recent findings corroborated the photodissociative nature of the axial ligands and added a new
dimension to these old molecules. Novel mechanisms proposed to be involved in photouncaging and photoinmunotherapy have recommenced exploration of SiPcs in light-based therapeutic approaches, other than PDT. While the Si-O bonds exhibit additive and/or oxygen dependent photolability, the Si-C bonds can rapidly dissociate without any dependence on oxygen or redox reagents. Besides medicinal applications, SiPcs are rapidly acquiring a foothold in the emerging fields of photovoltaics, organic electronics and photocatalysis. However, thoughtful considerations of energy levels, bulk heterojunction morphology, solubility and crystallinity are essential while designing SiPc variants in order to observe enhanced efficiency and performances of the devices. Though few SiPc molecules seem propitious, further structure–activity studies are required to correlate the structural modifications with the device characteristics. Altogether, the promising results bode well for future explorations of SiPc in multifaceted research areas.

However, these outstanding applications rest heavily on the structural modifications of the axial and phthalocyanine ring positions of SiPcs. The synthetic challenges to construct SiPc variants have been the bottleneck of this field and have impeded many applications. Oftentimes, the syntheses of SiPcs employ harsh reagents and drastic conditions. Furthermore, the reactions suffer from low synthetic yields and produce multiple undesired products. The purification techniques are complicated and laborious. Thus, the current scenario highlights the need for convenient and reliable synthetic methodology using non-hazardous and affordable reagents. Future efforts to define elegant synthetic approaches will be of paramount importance for complete realization of potential of SiPcs in phototherapy and energy harvesting applications.

Conflicts of interest

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Notes and references

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