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# 1D alignment of Co(II) metalated porphyrin–naphthalimide based self-assembled nanowires for photocatalytic hydrogen evolution†

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The splitting of water into hydrogen and oxygen under visible light is an emerging phenomenon in green energy technology. Nevertheless, selecting an appropriate photocatalyst is rather significant to enhance hydrogen production on a large scale. In this context, organic photocatalysts have received considerable attention owing to their larger surface area, control in diffusion adsorption, nanostructures and electronic properties. Herein, we have developed five either free base or transition metalated porphyrin–naphthalimide based donor–acceptor systems (**PN1–PN5**) and studied their morphology, electronic properties and catalytic behaviour. Detailed studies suggest that the Co(II) substituent D–A system (**PN2**) displayed a well-aligned one-dimensional (1D) nanowire with high electrical conductivity promoting remarkable photocatalytic hydrogen production rate (18 mM g<sup>−1</sup> h<sup>−1</sup>) when compared to that of porphyrin-based derivatives reported until now. Thus, these results propose to investigate diverse metalated  $\pi$ -conjugated materials as photocatalysts for hydrogen production.

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## Introduction

Light-stimulated water splitting into hydrogen has received significant attention for the cleanest energy production and storage.<sup>1</sup> In this context, several inorganic semiconductors were utilized as photocatalysts for water splitting towards hydrogen evolution.<sup>2</sup> However, lack of control on the surface area of nanostructures and band gap alignments has resulted in finding alternative semiconductors. Hence, semiconductors with fine-tuning of their critical parameters such as crystallinity, band gap, stability of transient species and exciton migration are rather important. In this regard, organic semiconductors are considered alternative catalysts for H<sub>2</sub> evolution.<sup>1</sup> Organic semiconductors with low-lying HOMO levels are highly desirable to improve the catalytic process under visible light and also to overcome the kinetic barrier of the process. Thus, so far, very few examples are reported on

organic photocatalysts such as carbon nitrides, porous and linear conjugated polymers and organic frameworks.<sup>3–8</sup> However, exciton recombination, low surface area and poor charge separation direct us to develop small  $\pi$ -conjugated materials with high photocatalytic activity.<sup>9,10</sup> Viewing this, the porphyrin-based  $\pi$ -conjugated system has attracted much attention in organic electronics due to its high ionization potential, electron affinity, planarity and photo/thermal stability. The extended  $\pi$ -conjugated structure of porphyrin promotes self-assembled nanostructures *via* supramolecular interactions.<sup>11</sup> On the other hand, porphyrins have been employed as a photosensitizer in third-generation solar cells, however, their photocatalytic performance is not much explored. Herein, we have developed porphyrin comprised donor–acceptor systems in which porphyrin acts as an electron donor and naphthalimide as an electron acceptor. Naphthalimide (NMI) is a well-known electron acceptor having miscellaneous optoelectronic applications. Additionally, the NMI structure possesses rigidity and planarity, which facilitates self-assembly by C=O $\cdots\pi$  and  $\pi$ – $\pi$  interactions.<sup>12</sup> Indeed, porphyrin and NMI individually exhibit excellent well-defined nanostructures, and designing the combination of porphyrin–NMI in  $\pi$ -conjugated structure can serve as a donor–acceptor (D–A) system, which aid to improve the electronic properties through the self-assembly.

Thereby, we have designed five porphyrin–NMI based D–A derivatives (**PN1–PN5**) with a free base (**PN1**), and incorporated

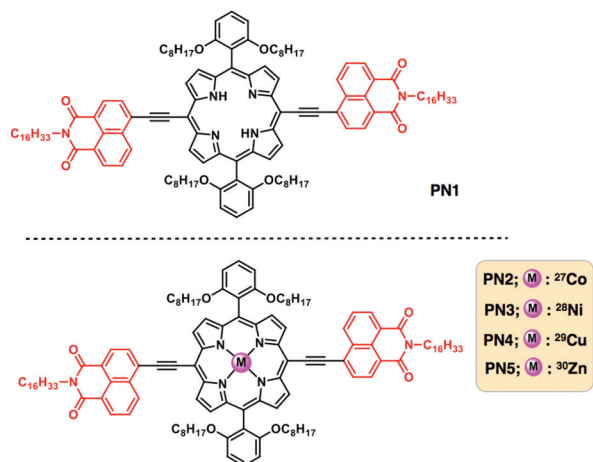
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**Fig. 1** Chemical structures of the free base and transition metalated porphyrin–naphthalimide series and represented **PN1** for the free base, **PN2–PN5** for Co(II), Ni(II), Cu(II) and Zn(II) metalated derivatives.

transition metals (**PN2–PN5**): cobalt (Co), nickel (Ni), copper (Cu) and zinc (Zn) (Fig. 1). As is known, the energy structure of porphyrin is non-planar and shows  $S_4$  symmetry, while transition metal-doped porphyrins exhibit planarity which results in the enhancement of symmetry to  $D_{4h}$ . On the other hand, transition metalated porphyrins depict high ionization potential, binding energy, electronic affinity and high chemical hardness. Among four transition metals, Co(II) metalated porphyrin exhibits high stability and poor reactivity due to low electron affinity, high chemical hardness and binding energy between the nitrogen complex with cobalt metal.<sup>13</sup> These molecules were synthesized using sonogashira coupling reactions, followed by metalation. Subsequently, these molecules were characterized by using spectroscopic techniques. These



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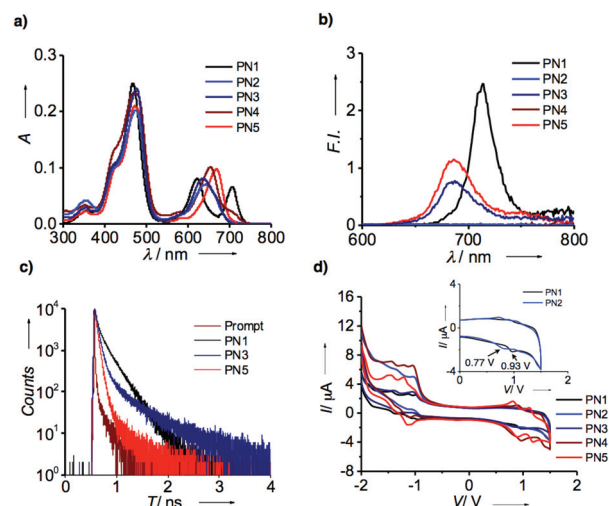
Seelam Prasanthkumar received his PhD degree in organic chemistry from CSIR-National Institute of Interdisciplinary Science and Technology (NIIST), Trivandrum, India in 2012. Afterwards, he joined as a post-doctoral fellow in the group of Prof. Takuzo Aida, RIKEN, Japan. Subsequently, he was selected for DST-Inspire faculty award and joined as a Inspire faculty fellow in CSIR-Indian Institute of Chemical Technology

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five derivatives showed broad range absorption from ultraviolet to the visible region and their redox nature facilitates plausible electron transfer from the donor to acceptor. Subsequently, these molecules form self-assembled one-dimensional nanostructures using supramolecular interactions resulting in high bulk conductivity, however, **PN2** is remarkable amongst them. Theoretical calculations also reveal that the **PN2** exhibits low lying HOMO and LUMO energy levels suggest that the desirable catalyst for hydrogen production. Consequently, **PN2** implies photocatalytic hydrogen production with a high rate promotes transition metalated linear  $\pi$ -conjugated D–A systems are highly enviable photocatalysts for future cleanest energy production and storage applications.

## Results and discussion

UV-visible absorption spectroscopic analysis of freebase and metalated porphyrin-based D–A systems were carried out in solution state at a concentration of  $1 \times 10^{-4}$  M. Fig. 2a represents the absorption spectra of **PN1–PN5** in chloroform showed prominent absorption bands at 400–500 nm and 570–750 nm, respectively. The typical absorption spectrum of porphyrin exhibits two significant bands at 400 nm and 550 nm, which correspond to strong and weak electronic transitions from  $S_0$  to  $S_2$  and  $S_0$  to  $S_1$ . These absorption bands indicate the Soret band or B-band and Q-bands, which arise due to  $\pi$ – $\pi^*$  transitions of the porphyrin moiety. Whereas in **PN** series (**PN1–PN5**), the spectral bands show bathochromic shift compared to the typical porphyrin core suggesting that the **PN** series consisting of porphyrin-linked naphthalimide facilitates



**Fig. 2** (a) Ultraviolet-visible absorption spectra of **PN** series (**PN1–PN5**) in  $\text{CHCl}_3$  at a concentration of  $1 \times 10^{-4}$  M. (b) Fluorescence spectra of **PN** Series at an excitation wavelength of 470 nm. (c) Time correlated single photon counting spectra of **PN1**, **PN3** and **PN5** using an excitation wavelength of 440 nm. (d) Cyclic voltammetry of **PN** series in  $\text{CHCl}_3$  at a scan rate of  $200 \text{ mV s}^{-1}$  (inset represents the oxidation potentials of **PN1** and **PN2** at 0–2 V).

extended  $\pi$ -conjugation, which promotes red shift absorption towards the visible region (Table S1†). However, transition metalated derivatives (**PN2**–**PN5**) exhibit a blue shift at the Q-band region compared to the free base (**PN1**), indicating that the vibronic states were stabilized upon metal binding in the porphyrin core. On the other hand, **PN2** and **PN3** showed a significant hypsochromic shift of  $\sim 20$ – $30$  nm than that of **PN4** and **PN5**, suggesting that Co, Ni metalated **PN** series has low lying ground state energy levels and higher stability relatively. Thus, the ground state spectra of the **PN** series showed a broad absorption from UV to visible range and also considerable spectral variations in the free base and metalated derivatives.

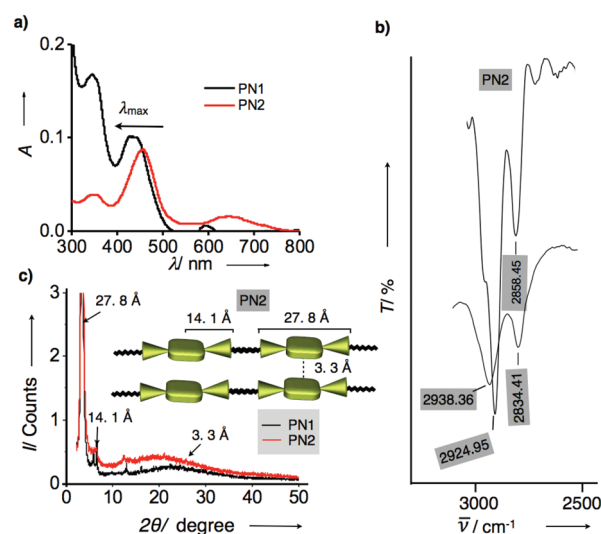
Subsequently, emission spectral data on the **PN** series in chloroform were recorded upon excitation using the wavelength of 470 nm with an optical density of 0.15 (Fig. 2b). Fluorescence spectra of **PN1** showed the emission maximum at 712 nm, while **PN3** and **PN5** at 685 nm with significant quenching of the emission intensity suggest the possibility of electron transfer in metalated derivatives. On the contrary, **PN2** and **PN4** did not show emission intensity due to the paramagnetic nature of Co and Cu substituent's influence on the loss of excited-state energy by passing through the intersystem crossing. Furthermore, lifetime analysis of **PN1**, **PN3** and **PN5** were performed to confirm the electron transfer mechanism using the time-correlated single-photon counting technique with an excitation wavelength of 440 nm. The resultant spectral data revealed that three derivatives displayed a biexponential decay profile with significant differences in lifetime values suggesting that the electron transfer is more favoured in metalated derivatives than the free base (Fig. 2c). As aforementioned, **PN2** and **PN4** did not show fluorescence thereby lifetime values were not observed. The ground and excited-state electronic properties confirmed that porphyrin and NMI appended **PN** series depict extended visible absorption and efficient electron transfer phenomenon in metalated derivatives.

Later, the redox-active nature of the **PN** series was analyzed by cyclic voltammetry (CV) at a scan rate of  $200 \text{ mV s}^{-1}$ . To examine their electrochemical properties, we measured each sample from the **PN** series in 0.1 M tetrabutylammonium hexafluorophosphate ( $\text{NBu}_4\text{PF}_6$ ) as a supporting electrolyte in chloroform followed by the three electrode system; saturated calomel, glassy carbon, platinum (Pt) wire as a reference, working and counter electrode, respectively. Fig. 2d represents the redox spectra of **PN1**–**PN5** showed quasi-reversible oxidation and reduction potentials (Table S1†). Freebase **PN1** displays oxidation potentials of 0.934 V, while **PN2** exhibits 0.77 V (inset Fig. 1b). Likewise, other derivatives **PN3**–**PN5** perform relatively similar to each other (Table S1†). Detailed CV analysis revealed that **PN2** showed lower oxidation potentials compared to that of free base and other metalated **PN** series indicating that Co(II) substituent plays a vital role to ease of oxidation among other metal substituents, which entails an efficient electron transfer from the donor to acceptor.

Having confirmed that the optical and electrochemical properties of metalated derivatives had prominent ground/excited

state electron properties and superior redox potentials prompts us to investigate their surface morphology. As mentioned above, porphyrins possess planarity and symmetry aid to form hierarchical self-assembled nanostructures. Whereas, in the **PN** series, tethering of porphyrin and NMI enhances conjugation leading to extended  $\pi$ -structure perhaps promoting long axial growth of well-defined nanostructures when compared to individual porphyrin. Thus, we have prepared aggregates of **PN** series by the methanol vapour diffusion approach mechanism. Subsequently, each of **PN** series were taken in 1 ml chloroform solution separately in a vial and placed in methanol for 24 h, resulting in green-coloured aggregates. These aliquots were analyzed using optical, X-ray diffraction and electron microscopic techniques. Initially, UV-visible absorption studies of aggregates were recorded in methanol at 25 °C. Fig. 3a represents absorption spectra of **PN1** and **PN2** showing the hypsochromic shift at the Soret band Q-band when compared to solution-state, indicating that these molecules exist as H-aggregates (Fig. 3a). Likewise, aggregates of **PN3**–**PN5** also performed in methanol and showed similar trends (Fig. S15†). Therefore, absorption spectra of the **PN** series suggest that these five molecules are arranged in head-to-head stacking, however, variations in a blue shift of each **PN** derivative might differ in molecular packing whilst undergoing self-assembly.

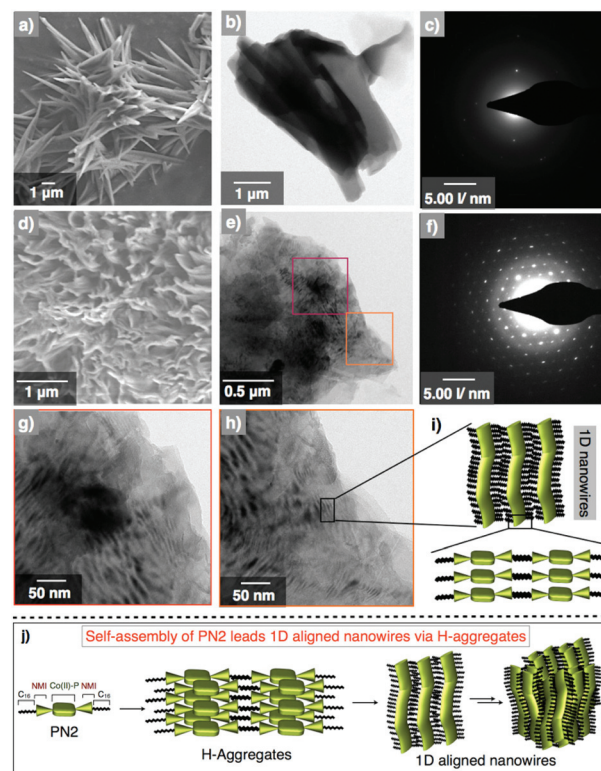
Furthermore, Fourier transform infrared spectroscopic analysis (FT-IR) of **PN** series were carried out in the solution and aggregate state using chloroform and methanol to ensure the stretching of long alkyl chains whilst aggregation (Fig. S15†). In chloroform, the  $\text{CH}_2$  stretching frequencies of **PN2** were



**Fig. 3** (a) UV-visible absorption spectra of **PN1** and **PN2** aggregates in methanol at 25 °C. (b) FT-IR spectra of **PN2** in solution and aggregate state recorded in chloroform and methanol. (c) Powder X-ray diffraction analysis of **PN1** (black line) and **PN2** (red line) aggregates and denoted  $d$ -spacing values for **PN2**. Inset shows the schematic illustration of molecular stacking in **PN2** aggregates and represented  $d$ -spacing values for each length of end-to-end, center to end and  $\pi$ - $\pi$  stacking.

observed at 2924.95 and 2858.45  $\text{cm}^{-1}$ , respectively. Whereas,  $\text{CH}_2$  stretching frequencies of aggregated samples have shown a significant shift to 2938.36 and 2834.41  $\text{cm}^{-1}$  as methanol solvent interference in  $\text{CH}_2$  stretching frequency results in the shift of asymmetric and symmetric methylene stretching frequencies. Thus, FT-IR analysis revealed that the presence of hexadecyl chains in **PN2** are stretched confirmation, which facilitated interdigitation with other molecules whilst under self-assembly (Fig. 3b and S15†).<sup>14</sup> Subsequently, the other four derivatives **PN1**, **PN3–PN5** were also studied, and their spectral characteristics resembled **PN2**. However, minor changes in the stretching frequencies indicate that the presence of distinct metals and freebase in the **PN** series directs variation in molecular packing, which defines divergent nanostructures from one to another. Furthermore, aggregates were analyzed using the powder X-ray diffraction method to confirm the molecular arrangement and their crystallinity whilst self-assembly at 25 °C. Fig. 3c represents that **PN1** and **PN2** exhibit sharp diffraction peaks at small and wide-angle regions signifying that molecules exhibit crystalline nature. **PN2** showed a broad peak observed at a small angle region, which corresponds to the  $d$ -spacing values of 3.3 Å, suggesting intermolecular  $\pi$ - $\pi$  stacking between molecules. However, other diffraction peaks with  $d$ -spacing values of 27.8 Å and 14.1 Å represent the end-to-end length of the NMI-porphyrin-NMI and metal centre to NMI reveals that the rigidity of the  $\pi$ -conjugated structure is extending over supramolecular nanostructures (Inset Fig. 3c). Along with that, hexadecyl chains also play a vital role in aggregation *via* van der Waal's interactions. On the other hand, **PN1**, **PN3–PN5** also showed similar diffraction peaks similar to **PN2** but considerable differences in  $d$ -spacing values suggest that stacking patterns differ from each other (Fig. S16†). Thus, the resultant UV-visible absorption, IR analysis and PXRD depict that the **PN** series undergoes H-aggregation leading to extended supramolecular nanostructures *via*  $\pi$ - $\pi$  stacking and van der Waals interactions. Nevertheless, the variation in molecular packing of each **PN** series is still unclear, thereby, microscopic visualization and their electron diffraction pattern are considered as a significant approach to studying the molecular stacking, alignment and supramolecular structures at the nanoscopic level.

Furthermore, we measured the surface morphology of aggregates of the **PN** series to check their self-assembled nanostructures and alignment by using electron microscopic and diffraction techniques. Scanning electron microscopy of **PN1** revealed nanorod-shaped structures with an average width of 0.5–1 micrometer and 3–4  $\mu\text{m}$  in length (Fig. 4a). Subsequently, transmission electron microscopy studies have been conducted by drop-casting the suspension of **PN1** in methanol at 25 °C. The TEM image of **PN1** displayed nanorod morphology with their dimensions similar to the SEM image (Fig. 4b). Eventually, we have performed electron diffraction analysis on the area same as in the TEM image and found that **PN1** did not show the precise molecular alignment in nanorod morphology (Fig. 4c). In contrast, the SEM image of **PN2**



**Fig. 4** (a, b) SEM and TEM images of an air-dried suspension of **PN1** at a concentration of  $1 \times 10^{-4}$  M at 25 °C. (c) The corresponding diffraction pattern of the **PN1** image. (d, e) SEM and TEM of an air-dried suspension of **PN2**. Purple- and orange-coloured boxes denote the area for high-resolution imaging. (f) Diffraction pattern recorded on the TEM images of **PN2**. (g, h) High-resolution TEM images of **PN2** from the selected area of image-e. (i) Proposed scheme of 1D nanowires from the molecular aggregates. (j) Schematic illustration of **PN2** undergoes self-assembly leading to 1D nanowires via H-aggregates.

depicts mesh wreath-type morphology wherein thin nanowires were aligned and joined together (Fig. 4d). In addition, small pores were observed in this morphology that showed a greater advantage when applied to photocatalytic hydrogen production. Interestingly, TEM images revealed well-aligned nanowires with an average width of 10 nm and length of 50–300 nm (Fig. 4e). Whereas, high-resolution TEM images show that these nanowires are unidirectionally aligned and connected to each other (Fig. 4g and h). Afterwards, diffraction analysis confirmed that the molecules undergo co-facial stacking, which was aligned in a specific direction, resulting in an ordered one-dimensional nanowires (Fig. 4f and i). Likewise, the morphology of three derivatives **PN3–PN5** was also studied and **PN3** showed nanorod formation with an average dimension of 0.5–1  $\mu\text{m}$  and several micrometers in length. In some places, these bundles of nanorods were wrapped together forming supramolecular nanoropes. Whereas, **PN4** exhibits nanowire morphology with several micrometers in length, which is in contrast to **PN2**. Subsequently, **PN5** was also studied and it showed nanorods shaped structures similar to **PN1** and **PN3** (Fig. S17†). Also, we performed the electron diffraction analysis

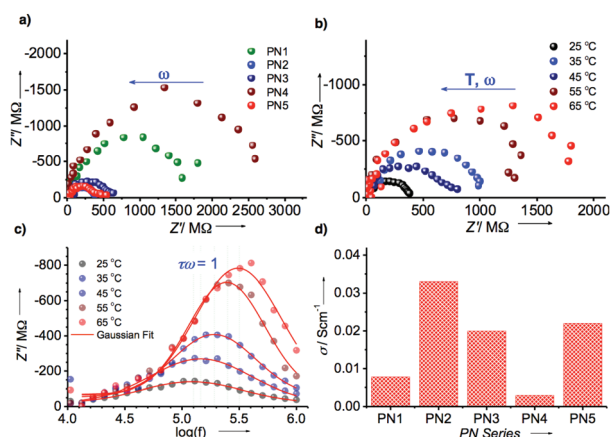
on **PN3–PN5**, and their corresponding images reveal a significant alignment of molecular arrangement in supramolecular nanostructures when compared to **PN1**, but relatively less than that of **PN2**. Thus, microscopic analysis suggests that the **PN2** ascertain well-aligned molecular arrangement leading to unidirectionality among 1D nanostructures (Fig. 4j). While **PN** series showed significant molecular alignment in 1D nanostructures, prompting us to perform their electronic characterization for organic electronic applications.

Hence, the electrical conductivity of the **PN** series was analyzed using electrochemical impedance studies (Fig. 5, S18–S22 and Table S2†). To assess the conductivity behavior of these assemblies, we coated the aggregates of each **PN** series on indium tin oxide glass plate suitably etched to provide  $1 \times 1$  cm cell dimensions and recorded the spectra. Fig. 5a represents Nyquist plots of **PN1–PN5** at 25 °C that revealed typical depressed semi-circular curves characteristic of contributions from both bulk resistance ( $R_b$ ) and capacitance ( $C_b$ ) when in a parallel combination. Amongst all, Co(II) metalated **PN2** showed the least bulk resistance ( $R_b$ ) of 400 M $\Omega$ , which corresponds to a specific conductivity of  $\sim 33$  mS cm $^{-1}$ . Whereas, other **PN** derivatives displayed slightly higher resistance in the range of 500–2000 M $\Omega$  that relates to specific conductivities in the range of 3–23 mS cm $^{-1}$ , respectively (Fig. 5d and Table S2†). Although metalated **PN** series have achieved remarkably higher bulk conductivity than **PN1**, Cu(II) substituent **PN4** yielded lower conductivity possibly to poor Ohmic contacts at the electrode interface with irregular and entangled nanowires. Subsequently, the temperature-dependent behavior of the **PN** series was recorded as a function of temperature from 25 °C to 65 °C that displayed a steady rise in the resistance of bulk (Fig. 5b and S18–S22†). This steady increase in the resistance against temperature is indicative of the predominant contribution of electronic conduction within the bulk (Fig. S18†). Furthermore, the bulk capacitance of **PN2** was esti-

mated from the significant parameters at the apogee of the semicircular arc where the condition  $\omega RC_{\text{bulk}} = 1$  is satisfied. From the radial peak frequency ( $\omega_{\text{peak}} = 2\pi f_{\text{peak}}$ ) obtained from the spectroscopic plots of imaginary impedance ( $Z''$ ) against  $\log(f)$  the bulk relaxation time ( $\tau_b$ ), of **PN2** and rest of the samples were estimated (Fig. 5c and Table S2†). Consequently, the calculated capacitance of **PN2** is 3.7 nF at 25 °C. Likewise, studies on **PN1**, **PN3–PN5** were also conducted and the bulk capacitance values were calculated with the variation of temperature from 25 °C to 65 °C (Fig. S19–S22 and Table S2†). Overall, these results confirmed that **PN2** exhibits the highest bulk electrical conductivity and capacitance values among the **PN** series.

As mentioned above, microscopic and impedance analysis of **PN2** displayed well-aligned 1D nanowires with high electrical conductivity among other transition metal substituted **PN** series, prompting us to utilize **PN2** as a catalyst for hydrogen evolution. Alongside, we also performed the density functional theory (DFT) calculations to check the appropriate energy levels of the **PN** series with the TiO $_2$  electrode, are a foremost criterion to be considered for the catalyst. Thereby, theoretical calculations of the **PN** series were studied using Gaussian 09 package with a functional basis set of the B3LYP/6-31G(d,p) level. The optimized structure of **PN1** shows the electron density distribution on the highest occupied molecular orbital (HOMO) on the porphyrin moiety and the lowest unoccupied molecular orbital (LUMO) on the porphyrin–NMI and the resultant values were 5.03 and 2.98 eV, respectively (Fig. 6). On the other hand, electron density allocation of HOMO–1 and LUMO+1 on porphyrin and naphthalimide exhibits unequivocal discrimination (Table S3†). The calculated band gap of **PN1** from the HOMO and LUMO yielded 2.05 eV. Subsequently, values for transition metalated **PN** series (**PN2–PN5**) were also calculated and the resultant HOMO and LUMO values were lower than **PN1** (Fig. 6 and Table S3†). Electrostatic potentials (ESP) of the **PN** series displayed similar electron density distribution in five derivatives (Table S3†). However, low lying HOMO levels and band gap variations suggest that the metalated derivatives were more suitable with TiO $_2$  nanoparticles than freebase (**PN1**). Amongst, Co(II) substituent **PN2** exhibits low lying HOMO and LUMO energy levels, hence **PN2** is a potential candidate to apply as a catalyst for hydrogen production.

Thus, photocatalytic hydrogen evolution experiments were performed using **PN2** as a catalyst with aq. triethanolamine (TEOA) solution (20 vol%) as a sacrificial agent and platinum as the cocatalyst for titanium oxide (TiO $_2$ ) electrode. Fig. 7a represents hydrogen yields of **PN2** recorded under visible light irradiations ( $\lambda = 420$  nm) up to 4 h at neutral, acidic and basic p $^H$  conditions. At neutral p $^H$ , hydrogen yields exhibit 15 mmol g $^{-1}$  until 4 h irradiation. Later, acidic and basic pH values were also recorded by controlled addition of concentrated HCl which results in higher H $_2$  yield in basic media when compared to acidic and neutral p $^H$ . Subsequently, the H $_2$  production rate was calculated and those values were remarkably higher than the previous porphyrin-based D–A systems



**Fig. 5** (a) Nyquist plot of **PN1–PN5** derivatives. (b) Temperature-dependent Nyquist plot of **PN2** with ranging from 25 °C–65 °C. (c) Temperature-dependent logarithmic frequency ( $\log(f)$ ) against imaginary impedance of **PN2**. (d) The bar diagram of the **PN** series vs. specific conductivity at 25 °C.

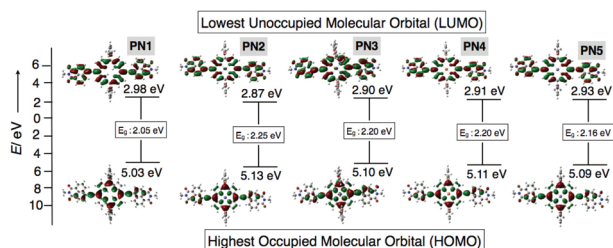


Fig. 6 Optimized structures of PN series (PN1–PN5) from DFT calculations. The corresponding HOMO/LUMO energy levels and band gap values.

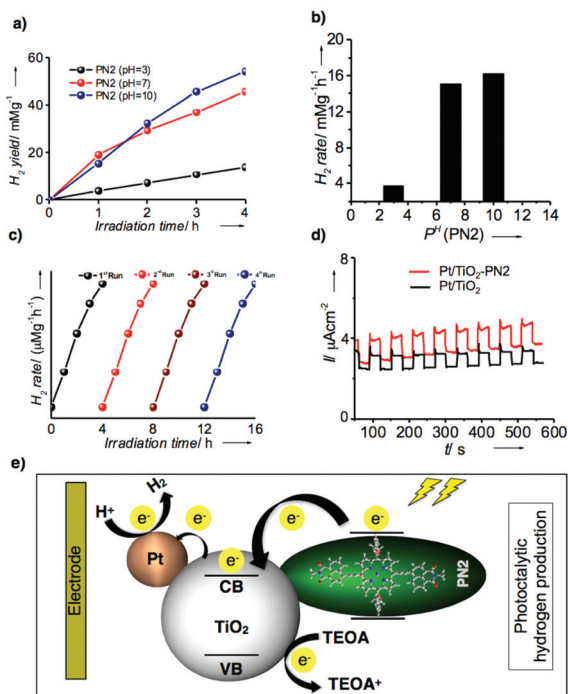


Fig. 7 (a) Photocatalytic activity of PN2 at different P<sup>H</sup> variations (acidic, neutral and basic conditions). (b) Bar diagram of H<sub>2</sub> rate against P<sup>H</sup> of PN2. Stability studies: (c) hydrogen production rate against the time variation from 0 h to 16 h. (d) Photocurrent response vs. time (seconds) between Pt/TiO<sub>2</sub>-PN2 and Pt/TiO<sub>2</sub>. (e) Schematic representation of possible electron transfer mechanism and hydrogen evolution under visible light.

reported until now.<sup>15</sup> The resultant H<sub>2</sub> rate of PN2 is 17.5 mmol g<sup>-1</sup> h<sup>-1</sup> (p<sup>H</sup> = 10), 15.7 mmol g<sup>-1</sup> h<sup>-1</sup> (p<sup>H</sup> = 7) and 4.0 mmol g<sup>-1</sup> h<sup>-1</sup> (p<sup>H</sup> = 3) (Fig. 7b). In addition, turnover number (TON) and TOF were calculated for PN2 to determine the number of reacting molecules in comparison with the active sites whilst irradiation time. The resultant values are 64 790.84 and 16 197.71, respectively. Nevertheless, the stability of the photocatalyst is rather significant for hydrogen production. Thereby, we conducted the stability studies of PN2 by extending the irradiation time from 4 h to 16 h revealed that the H<sub>2</sub> production rate remains constant still for a longer time (Fig. 7c). Furthermore, we recorded photocurrent against time

under the light on/off conditions suggesting that the photocurrent of Pt/TiO<sub>2</sub>-PN2 is higher than that of Pt/TiO<sub>2</sub> alone and stable until 550 s (Fig. 7d). Overall, Co(II) substituent PN2 showed a higher H<sub>2</sub> production rate and stability owing to the LUMO energy levels close to the conduction band of TiO<sub>2</sub>, which implies efficient electron transfer from the dye to the TiO<sub>2</sub> electrode (Fig. 7e). Therefore, a well-defined 1D molecular orientation of Co(II) substituted PN2 with high electrical conductivity leading to photocatalytic H<sub>2</sub> production facilitates enhancing the performance of organic electronic devices.

## Conclusions

In conclusion, we have demonstrated the five porphyrin-naphthelime (PN1–PN5) as donor–acceptor systems in which PN2 showed well-aligned molecular arrangements directing 1D nanowires, which exhibit high electrical conductivity and photocatalytic hydrogen production. The optical and electrochemical properties revealed that metalated derivatives displayed efficient intramolecular electron transfer from the donor to acceptor than free base. Microscopic data and diffraction analysis suggested that the precise molecular packing of Co(II) substituent PN2 exclusively formed aligned nanowires among other derivatives. Hence, it showed remarkable bulk electrical conductivity and accomplished high hydrogen production rate under visible light. Consequently, transition metal substituted macrocyclic-based D–A systems to be explored in detail for realistic challenges in clean and green technology.

## Author contributions

S. P., B. B. and L. G. designed the experiment, analysis of data and wrote the manuscript. B. B. and G. R. executed the synthesis, characterization, spectroscopic measurements of the PN series. N. C. and U. P. performed the photocatalytic hydrogen production analysis. P. B. conducted electrochemical impedance studies. V. K. measured the microscopic and electron diffraction analyses. All authors participated in the discussion of results and revision of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## Notes and references

- 1 S. Chen, T. Takata and K. Domen, *Nat. Rev. Mater.*, 2017, **2**, 17050; Y. Wang, H. Suzuki, J. Xie, O. Tomita, D. J. Martin, M. Higashi, D. Kong, R. Abe and J. Tang, *Chem. Rev.*, 2018, **118**, 5201; C. Zhao, Z. Chen, R. Shi, X. Yang and T. Zhang, *Adv. Mater.*, 2020, **32**, 1907296; Y. Bai, K. Hippalgaonkar and R. S. Sprick, *J. Mater. Chem. A*, 2021, **9**, 16222; F. Liu, M. Wang, X. Liu, B. Wang, C. Li, C. Liu, Z. Lin and F. Huang, *Nano Lett.*, 2021, **21**, 1643; K. Kondo, Y. Watanabe, J. Kuno, Y. Ishii, S. Kawasaki, M. Kato, G. Kalita, Y. Hattori, O. Mashkov, M. Sytnyk and W. Heiss, *Energy Technol.*, 2021, **9**, 2100123.
- 2 A. Kudo and Y. Miseki, *Chem. Soc. Rev.*, 2009, **38**, 253; M. G. Kibria, F. A. Chowdhury, S. Zhao, B. Alotaibi, M. L. Trudeau, H. Guo and Z. Mi, *Nat. Commun.*, 2015, **6**, 6797.
- 3 X. Wang, K. Maeda, A. Thomas, K. Takanabe, G. Xin, J. M. Carlsson, K. Domen and M. Antonietti, *Nat. Mater.*, 2009, **8**, 76; J. Liu, Y. Liu, N. Liu, Y. Han, X. Zhang, H. Huang, Y. Lifshitz, S.-T. Lee, J. Zhong and Z. Kang, *Science*, 2015, **347**, 970.
- 4 L. Li, Z. Cai, Q. Wu, W. Y. Lo, N. Zhang, L. X. Chen and L. Yu, *J. Am. Chem. Soc.*, 2016, **138**, 7681.
- 5 G. Zhang, Z.-A. Lan and X. Wang, *Angew. Chem., Int. Ed.*, 2016, **55**, 15712; J. Kosco, M. Sachs, R. Godin, M. Kirkus, L. Francas, M. Bidwell, M. Qureshi, D. Anjum, J. R. Durrant and I. McCulloch, *Adv. Energy Mater.*, 2018, **8**, 1802181; C. Shu, C. Han, X. Yang, C. Zhang, Y. Chen, S. Ren, F. Wang, F. Huang and J.-X. Jiang, *Adv. Mater.*, 2021, **33**, 2008498.
- 6 R. S. Sprick, J. X. Jiang, B. Bonillo, S. Ren, T. Ratvijitvech, P. Guiglion, M. A. Zwiijnenburg, D. J. Adams and A. I. Cooper, *J. Am. Chem. Soc.*, 2015, **137**, 3265; Y. S. Kochergin, D. Schwarz, A. Acharjya, A. Ichangi, R. Kulkarni, P. Eliášová, J. Vacek, J. Schmidt, A. Thomas and M. J. Bojdys, *Angew. Chem., Int. Ed.*, 2018, **57**, 14188; R. S. Sprick, C. M. Aitchison, E. Berardo, L. Turcani, L. Wilbraham, B. M. Alston, K. E. Jelfs, M. A. Zwiijnenburg and A. I. Cooper, *J. Mater. Chem. A*, 2018, **6**, 11994; Y. Bai, L. Wilbraham, B. J. Slater, M. A. Zwiijnenburg, R. S. Sprick and A. I. Cooper, *J. Am. Chem. Soc.*, 2019, **141**, 9063.
- 7 E. Jin, Z. Lan, Q. Jiang, K. Geng, G. Li, X. Wang and D. Jiang, *Chem*, 2019, **5**, 1632; P. Pachfule, A. Acharjya, J. Roeser, T. Langenhahn, M. Schwarze, R. Schomcker, A. Thomas and J. Schmidt, *J. Am. Chem. Soc.*, 2018, **140**, 1423; K. C. Ranjeesh, R. Illathvalappil, V. C. Wakchaure, G. Gouda, S. Kurungot and S. S. Babu, *ACS Appl. Energy Mater.*, 2018, **1**, 6442; K. C. Ranjeesh, L. George, A. Maibam, S. Krishnamurthy and S. S. Babu, *ChemCatChem*, 2021, **13**, 1717.
- 8 T. Banerjee, K. Gottschling, G. Savasci, C. Ochsenfeld and B. V. Lotsch, *ACS Energy Lett.*, 2018, **3**, 400; S. Ghosh, A. Nakada, M. A. Springer, T. Kawaguchi, K. Suzuki, H. Kaji, I. Baburin, A. Kuc, T. Heine, H. Suzuki, R. Abe and S. Seki, *J. Am. Chem. Soc.*, 2020, **142**, 9752.
- 9 Y. Zhong, J. F. Wang, R. F. Zhang, W. B. Wei, H. M. Wang, X. P. Lu, F. Bai, H. M. Wu, R. Haddad and H. Y. Fan, *Nano Lett.*, 2014, **14**, 7175; Y. Z. Chen, A. X. Li, Z. H. Huang, L. N. Wang and F. Y. Kang, *Nanomaterials*, 2016, **6**, 51; N. Zhang, L. Wang, H. M. Wang, R. H. Cao, J. F. Wang, F. Bai and H. Y. Fan, *Nano Lett.*, 2018, **18**, 560; Q. Li, N. N. Zhao and F. Bai, *MRS Bull.*, 2019, **44**, 172.
- 10 A. G. Mojarad and S. Zakavi, *ACS Appl. Mater. Interfaces*, 2020, **12**, 46190; G. B. Bodedla, J. Huang, W. Wong and X. Zhu, *ACS Appl. Nano Mater.*, 2020, **3**, 7040; F. Zeng, J. Liao, M. Ding and G. Ou, *Dyes Pigm.*, 2021, **192**, 109430; V. Nikolaou, G. Charalambidis and A. G. Coutsolelos, *Chem. Commun.*, 2021, **57**, 40554058; Y. Kobayashi, A. Muranaka, K. Kato, A. Saeki, T. Tanaka, M. Uchiyama, A. Osuka, T. Aida and T. Sakurai, *Chem. Commun.*, 2021, **57**, 120; E. Nikoloudakis, M. Pigiaki, M. N. Polychronaki, A. Margaritopoulou, G. Charalambidis, E. Serpetzoglou, A. Mitraki, P. A. Loukakos and A. G. Coutsolelos, *ACS Sustainable Chem. Eng.*, 2021, **9**, 7781.
- 11 M. Mrinalini, B. S. P. Achary, S. Ghosh, D. Koteswar, S. Prasanthkumar and L. Giribabu, *J. Phys. Chem. C*, 2018, **122**, 10255; M. Mrinalini, S. S. Pathak, B. S. Achary, L. S. Panchakarla and S. Prasanthkumar, *Chem. – Asian J.*, 2019, **14**, 537; M. Mrinalini, M. Naresh, S. Prasanthkumar and L. Giribabu, *J. Porphyrins Phthalocyanines*, 2021, **25**, 382; B. Bhavani, M. Mrinalini, J. V. S. Krishna, P. Basak, L. Giribabu and S. Prasanthkumar, *ACS Appl. Electron. Mater.*, 2021, **3**, 176.
- 12 D. L. Reger, J. J. Horger and M. D. Smith, *Chem. Commun.*, 2011, **47**, 2805; J. A. Kitchen, P. N. Martinho, G. G. Morgan and T. Gunnlaugsson, *Dalton Trans.*, 2014, **43**, 6468; S. Pagidi, N. K. Kalluvettukuzhy and P. Thilagar, *Organometallics*, 2018, **37**, 1900; N. Meher and P. K. Iyer, *Angew. Chem., Int. Ed.*, 2018, **57**, 8488; S. Sahin Un, S. Zehra Topal and Y. Zorlu and, *J. Lumin.*, 2017, **190**, 23; G. B. Bodedla, L. L. Li, Y. Y. Che, Y. J. Jiang, J. Huang, J. Z. Zhao and X. J. Zhu, *Chem. Commun.*, 2018, **54**, 11614; G. B. Bodedla, G. Tang, J. Zhao and X. A. Zhu, *Sustainable Energy Fuels*, 2020, **4**, 2675.
- 13 D. R. Roy, E. V. Shah and S. M. Roy, *Spectrochim. Acta, Part A*, 2018, **190**, 121; E. V. Shah, V. Kumar, B. K. Sharma, K. Rajput, V. P. Chaudhary and D. R. Roy, *J. Mol. Model.*, 2018, **24**, 239.
- 14 W. Jin, Y. Yamamoto, T. Fukushima, N. Ishii, J. Kim, K. Kato, M. Takata and T. Aida, *J. Am. Chem. Soc.*, 2008, **130**, 9434.
- 15 P. Y. Ho, M. F. Mark, Y. Wang, S. C. Yiu, W. H. Yu, C. L. Ho, D. W. McCamant, R. Eisenberg and S. Huang, *ChemSusChem*, 2018, **11**, 2517; Z. Chen, J. Wang, S. Zhang, Y. Zhang, J. Zhang, R. Li and T. Peng, *ACS Appl. Energy Mater.*, 2019, **2**, 5665; L. Y. Huang, J. F. Huang, Y. Lei, S. Qin and J. M. Liu, *Catalysts*, 2020, **10**, 656; N. Chanda, K. Devulapally, S. Gonuguntla, S. Bojja, U. Pal and L. Giribabu, *Mater. Adv.*, 2021, **2**, 4762; P. S. Gangadhar, S. S. Gonuguntla, S. Madanaboina, N. Islavath, U. Pal and L. Giribabu, *J. Photochem. Photobiol., A*, 2020, **392**, 112408.