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# Sensing properties of light-emitting single walled carbon nanotubes prepared *via* click chemistry of ylides bound to the nanotube surface†

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Pyridinium ylide functionalized single-walled carbon nanotubes (SWCNTs) generated from simple quaternary pyridinium salts covalently bound to the SWCNT surface undergo a 1, 3 dipolar cycloaddition with dimethyl acetylenedicarboxylate in a 'click' chemistry type fashion to yield indolizine modified light-emitting SWCNTs. Conversion of quaternary pyridinium salts into indolizines on the SWCNT surface was confirmed by XPS, fluorescence spectroscopy and optical microscopy. The resulting modified SWCNTs were found to emit blue light when excited at 330 nm. Fluorescence quenching experiments performed *via* nitroaromatic compounds displayed that the fluorescence of the modified SWCNTs was quenched by nitrophenol derivatives in both solution-state and solid-state, probably due to strong H-bonding interaction between host and guest molecules. The functionalized SWCNTs were further characterized using TGA-MS, FTIR and UV-vis-NIR spectroscopy.

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## Introduction

Chemical sensing has gained much attention due to increasing demand for monitoring environmental pollution, detecting warfare and security threats, developing diagnostics for medical care and providing efficient and sensitive tools for quality control in industrial applications.<sup>1</sup> Due to their interesting chemical and physical properties carbon nanotubes (CNTs) have been used for chemical sensor applications, each based on alterations of a particular feature of a functional material including conductivity for chemiresistors<sup>2,3</sup> and back gate field effect transistors<sup>3–6</sup> and the intensity of scattered light resulting from the adsorption of chemical species for photo chemical or optical sensors.<sup>7–11</sup> Substantial effort, therefore, has been devoted in fabricating SWCNT based sensors and the surface

functionalization of CNTs in order for utilizing them for sensing. Following the discovery of bright fluorescence in the 800 to 1600 nm wavelength range of the near infrared from one-dimensional direct band gap semiconducting tubes,<sup>12</sup> a SWCNT based near-infrared optical sensor for glucose detection was fabricated by assembling a monolayer of the enzyme on nanotube surface followed by ferricyanide functionalisation.<sup>13</sup> Pyrene-modified  $\beta$ -cyclodextrin-decorated SWCNT field-effect transistor device as a photosensor was used to probe a fluorescent adamantyl-modified Ru complex through a charge-transfer mechanism from pyrenecyclodextrin-SWCNT hybrids to adamantyl-modified Ru complex.<sup>14</sup> A chemiresistive sensor based on SWCNT and hexafluoroisopropanol functionalised polythiophene for the detection of chemical warfare agents including sarin gas and dimethyl methylphosphonate has been shown by a simple spin-casting technique,<sup>3</sup> where the sensory response was investigated measuring the conductance of the SWCNT deposition. Fabrication of fluorescent chemosensors were achieved by covalently quinoline modified CNTs for  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  ions<sup>15,16</sup> and CNTs bearing adsorbed DNA with 6-carboxyfluorescein groups for  $\text{Hg}^{2+}$ .<sup>17</sup>

Introducing surface groups also enhanced the processability of CNTs, particularly SWCNTs, in both aqueous and non-aqueous solvents by interrupting the strong van der Waals forces causing aggregation or bundling of the material. Covalent strategies to functionalize SWCNTs have included carbene,<sup>18</sup> radical,<sup>19,20</sup> cycloaddition,<sup>21–24</sup> organolithium addition<sup>25,26</sup> and oxidation<sup>27,28</sup> reactions and been recently reviewed.<sup>29</sup> Among radical reactions aryl diazonium salts addition<sup>19,30–34</sup> has widely been studied using diazonium salts directly<sup>30</sup> or generating them *in situ* by oxidation of the

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† Electronic supplementary information (ESI) available: Normalized (at 330 nm) UV-vis-NIR spectra, recorded in *N,N*-dimethylformamide, of purified SWCNTs and SWCNT-Py (Fig. S1). Schematic representation of the synthesis of 3-ethyl 1,2-dimethyl indolizine-1,2,3-tricarboxylate (free indolizine) (Scheme S1). Deconvoluted N 1s XPS spectrum of **3c** (Fig. S3). Fluorescent spectra of **1b**, P, 2NP, 3NP, 4NP, 4NT and 2\_4NT in  $\text{CH}_3\text{CN}$  (Fig. S4). Percentage decrease in fluorescence intensity of **1b** and FI upon the addition of P, 2NP, 3NP, 4NP, 4NT and 2\_4NT (Fig. S5). Fluorescence spectra of **1b** in the presence of 2, 4, 6, 8, 10, 15, 20, 25, 50, 75, 100, 200  $\mu\text{L}$  equivalent of P, 2NP, 3NP, 4NP and 2\_4NT (Fig. S6–S10). Fluorescence spectra of FI in the presence of 2, 4, 6, 8, 10, 15, 20, 25, 50, 75, 100, 200  $\mu\text{L}$  equivalent of 4NP (Fig. S11).  $^1\text{H}$  NMR spectra of 4NP, FI and possible FI:4NP complex formed (Fig. S12). FTIR spectra of FI, 4NP and possible FI:4NP complex formed (Fig. S13). See DOI: 10.1039/c5ra04330a

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corresponding aniline with inorganic<sup>31</sup> or organic<sup>32</sup> oxidants. In our research group, it was shown that pyridine diazonium salts underwent a radical addition reaction to SWCNTs to yield pyridine-functionalized SWCNTs which were able to act as cross-linkers and hydrogen bond to poly(acrylic acid) to form SWCNT hydrogels.<sup>35</sup> 'Click' chemistry, defined as a modular reaction bringing together two simple units to generate a more complex substance, is gaining momentum in materials science.<sup>21,36</sup> For CNTs 'click' chemistry is predominately based around the Huisgen cycloaddition of azides to alkynes and has been used to attach polystyrene,<sup>37</sup>  $\beta$ -cyclodextrin<sup>38</sup> and phthalocyanines<sup>39</sup> to the surface of nanotubes. Although other simple 1, 3-dipolar cycloaddition reactions also fit the criteria of 'click' chemistry such as the reaction described here.

Herein diazonium salt addition and 1, 3-dipolar cycloaddition reactions were utilized in a modular type fashion to introduce light emitting groups to the surface of SWCNTs in a 'click' chemistry type fashion at low temperatures. Fluorescent SWCNTs were tested as sensor to probe nitroaromatic compounds in both solution-state and solid-state.

## Experimental

### Material preparation

**1. SWCNTs.** Purified SWCNTs, produced by the HiPco method, were purchased from Unidym, USA, and further purified by heating in air at 400 °C and soaking in 6 M HCl overnight. The purified SWCNTs were isolated by filtration over a polycarbonate membrane (0.2  $\mu$ m, Whatman), and washed with copious amounts of high-purity water until pH neutral. The purified SWCNTs were annealed under vacuum ( $10^{-2}$  mbar) at 900 °C to remove residual oxygen containing functional groups and any adsorbed gases or solvents.

**2. SWCNT-pyridine (SWCNT-Py).** SWCNT-Py was prepared following the literature procedure.<sup>35</sup> In brief, pyridine diazonium salt generated was reacted with SWCNTs at 0 °C in *N,N*-dimethylformamide (DMF). The modified SWCNTs were filtered through a nylon membrane (0.2  $\mu$ m, Whatman), re-dispersed in 2 M HCl (100 mL), filtered and washed with copious amount of water until pH neutral. The resulting solid material was dispersed in 2 M NaOH (100 mL) and stirred overnight to ensure deprotonation of the pyridinium salt to pyridine. The functionalized SWCNTs were then isolated by filtration through a nylon membrane (0.2  $\mu$ m, Whatman) and washed with water until pH neutral and re-dispersed and filtered using THF (2  $\times$  30 mL), acetone (2  $\times$  30 mL) and ethanol (2  $\times$  30 mL), respectively and dried overnight at 80 °C to afford SWCNT-Py.

**3. SWCNT-pyridinium ester salt (1a).** To a dispersion of SWCNT-Py (5 mg) in dry DMF (10 mL), dispersed using an ultrasonic bath (Ultrawave U50, 30–40 kHz) for 5 min, was added ethyl-2-bromoacetate (5 mL, 7.53 g, 45.09 mmol) and the reaction mixture stirred at room temperature for 12 hours. The SWCNTs were then filtered through a nylon membrane (0.2  $\mu$ m, Whatman), re-dispersed in acetone (30 mL), filtered and washed with acetone (2  $\times$  30 mL) and dried overnight at 80 °C to afford SWCNT-pyridinium ester (1a).

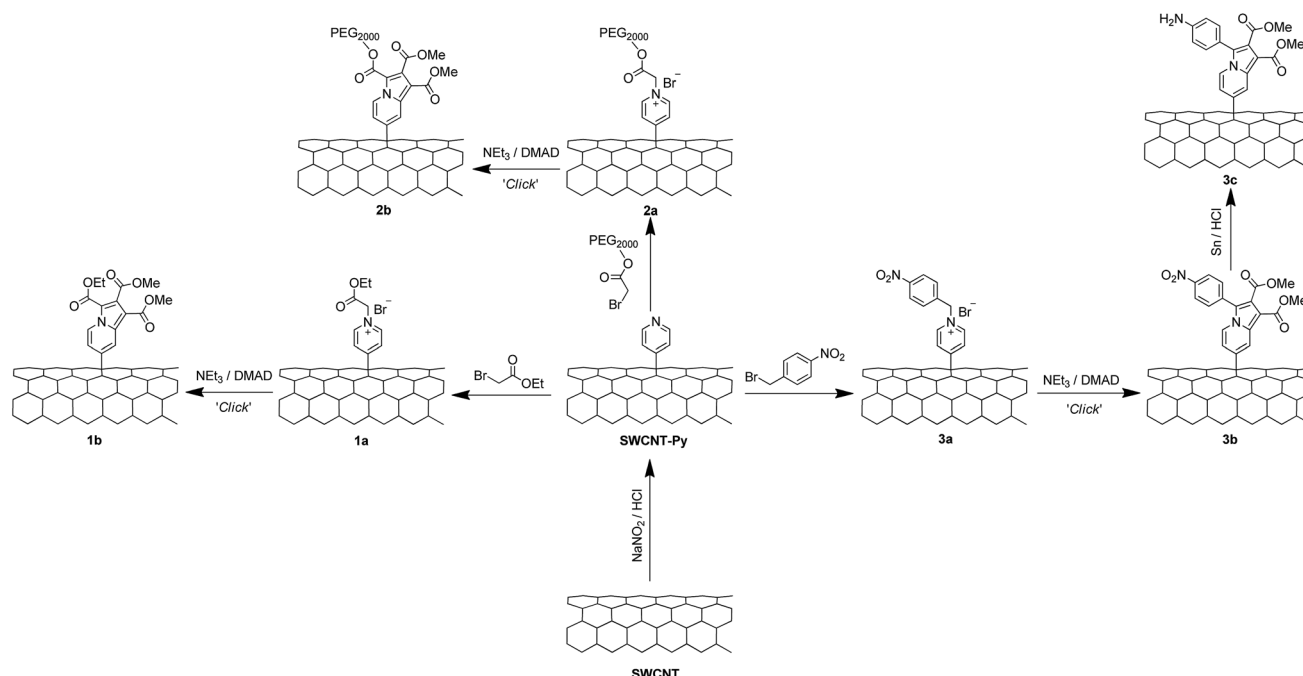
## Results and discussion

Herein, an elegant synthetic protocol to introduce light-emitting groups to the surface of SWCNTs *via* diazonium salt addition followed by 'click' chemistry is demonstrated using the reactivity of pyridine lone pair. The pyridinium ylide, rather than reacting with the CNT surface directly, is covalently bonded to the nanotube which then undergoes 1, 3-dipolar cycloaddition with dimethyl acetylenedicarboxylate (DMAD) to yield fluorescent indolizines, Scheme 1. In contrast to previously reported method,<sup>21,23</sup> combined diazonium salt-click chemistry approach described here offers high yielding indolizine formations at low temperatures, more control on reaction conditions, possible choice of dipoles/dipolarophiles with desired side groups (*e.g.* cyclodextrins,<sup>40</sup> activated alkynes or alkenes at low temperatures<sup>41</sup>) and the characterization of reaction intermediates. It may further provide a route to generate various quaternary pyridinium salts bearing alkyl, aryl or polymers (*e.g.* poly(ethylene glycol)) that is likely to enhance the solubility and processability of SWCNT material. However, multistep surface functionalization of SWCNTs resulted in low yields of isolated products that had noteworthy effect on the characterization of modified SWCNTs by concentration/thickness dependent techniques including FTIR and XPS.

In agreement with the previous literature findings,<sup>35</sup> electronic spectrum of SWCNT-Py in DMF showed suppressed van Hove singularities, indicative of covalent surface modification (see ESI Fig. S1†). A stable dispersion of SWCNT-Py had a concentration of  $\sim 160 \mu\text{g mL}^{-1}$ , calculated using the Beer's law and an extinction coefficient of  $30 \text{ mL mg}^{-1} \text{ cm}^{-1}$  and absorption values at  $\lambda = 700 \text{ nm}$ ,<sup>42</sup> which decreased dramatically to  $0.2 \mu\text{g mL}^{-1}$  upon the addition of ethyl-2-bromoacetate to form the Kröhnke salt (1a) which increased to  $40 \mu\text{g mL}^{-1}$  following 'click' reaction to form the indolizine functionalized SWCNTs (1b). Table 1 shows the solubility of the stable dispersion of Kröhnke salt modified SWCNTs (1a, 2a and 3a), indolizine functionalized SWCNTs (1b, 2b and 3b) and pyridine functionalized SWCNTs (SWCNT-Py) in both DMF and ethanol (EtOH). Solubility of SWCNT-Py in both DMF and EtOH decreased after the formation of 1a, 2a and 3a. However, solubility in DMF increased after the cycloaddition reaction to yield indolizines while the solubility in EtOH decreased, probably due to solute/solvent interactions.

The ATR-FTIR spectra of 1b–3b, Fig. 1, displayed the presence of the ester groups attached to the indolizine ring as a broad band ( $\nu_{\text{CO}}$ ) centred around  $1730 \text{ cm}^{-1}$  (along with multiple bands between  $1600\text{--}1300 \text{ cm}^{-1}$ , presumably from the formation of the indolizine).<sup>43</sup> The bands *ca.*  $2850\text{--}2990 \text{ cm}^{-1}$  in the spectrum of 3b that are up-shifted compared with 1b were attributed to the aromatic C–H stretching, and the band at  $1270 \text{ cm}^{-1}$  indicated the  $\text{--NO}_2$  symmetric stretching ( $\nu_{\text{NO}}$   $1270 \text{ cm}^{-1}$ ), confirming the presence of nitro groups.<sup>44,45</sup> The FTIR spectrum of 2b was noisy compared to others, probably due to the thickness of the film prepared.





Scheme 1 Multistep synthesis of quaternary pyridinium salt and indolizine modified SWCNTs (1a–3a, 1b–3b and 3c).

Table 1 Solubility data ( $\mu\text{g mL}^{-1}$ ) of 1a–3a, 1b–3b and SWCNT-Py in both DMF and EtOH

	SWCNT-Py	1a	2a	3a	1b	2b	3b
DMF	160	0.2	8.4	3.9	40.0	47.4	10.1
EtOH	72.4	34.0	28.5	3.1	16.4	19.3	1.26

TGA of **1b**, **2b** and **3b** showed a weight loss of ~38, 35 and 64% respectively, at 600 °C compared to *ca.* 7% for purified SWCNTs (Fig. 2). Weight losses of **1b**, **2b** and **3b** correspond to the presence of approximately 1 indolizine group for every 56 (1.77 N%), 76 (1.32 N%) and 142 (0.70 N%) nanotube carbon atoms, respectively, assuming the whole weight loss is due to removal of surface indolizines. Mass spectrometry results revealed that the indolizine groups were covalently attached to nanotube surface, showing common fragments at high temperatures for the indolizine modified SWCNTs (Fig. 3). The mass fragments detected by mass spectrometry demonstrated that **1b** fragmented into methyl ester

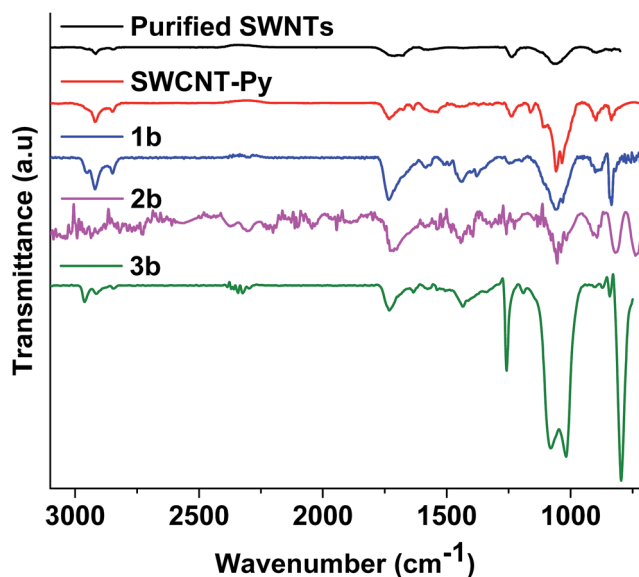


Fig. 1 Normalized and offset ATR-FTIR spectra of purified SWCNTs (black line), SWCNT-Py (red line), 1b (blue line), 2b (magenta line) and 3b (olive line). All spectra were recorded using SWCNT films. Curves are in top-down, respectively.

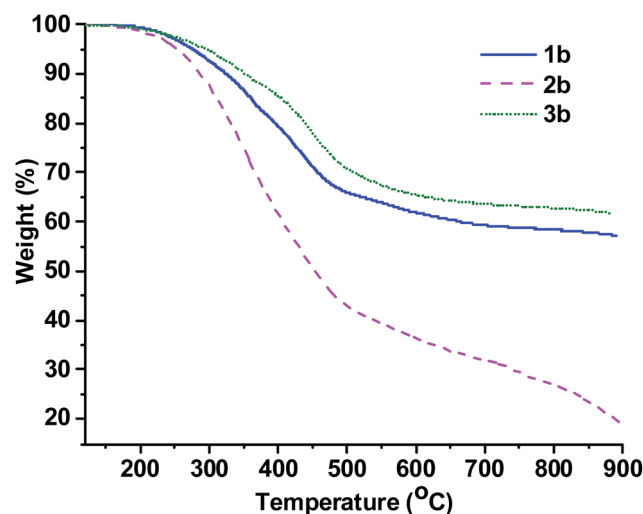


Fig. 2 TGA data of **1b** (blue line), **2b** (magenta dashed line) and **3b** (green dotted line). Data were recorded in flowing He (20 mL min<sup>-1</sup>) at a ramp rate of 10 °C min<sup>-1</sup> to 900 °C after being held at 120 °C for 30 min to remove any residual solvent.



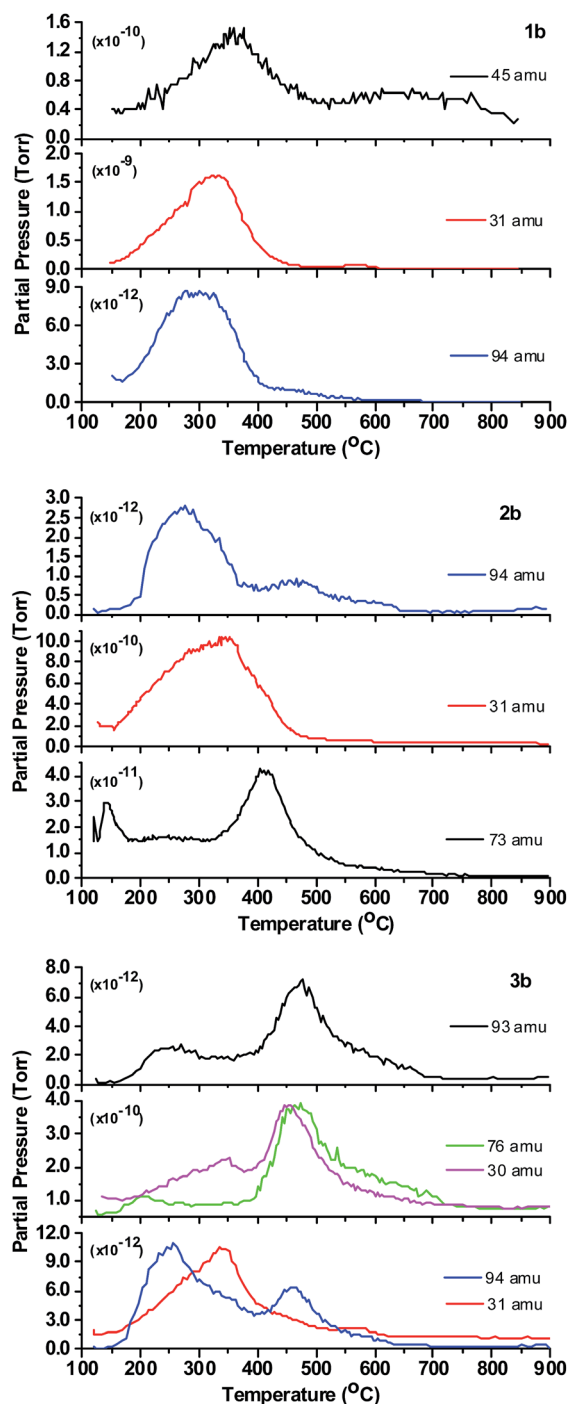


Fig. 3 Mass trace of ethoxy (45 amu), methoxy (31 amu) and methylpyridinium (94 amu) fragment given off during the heating of **1b**; mass trace of methylpyridinium (94 amu), methoxy (31 amu) and ethyl formyl (73 amu) fragment given off during the heating of **2b**; mass trace of phenoxy (93 amu), (green) phenyl (76 amu), (magenta) nitroso (30 amu), (red) methoxy (31 amu) and (blue) methylpyridinium (94 amu) fragment given off during the heating of **3b**. Data were recorded in flowing He (20 mL min<sup>-1</sup>) at a ramp rate of 10 °C min<sup>-1</sup> to 900 °C after being held at 120 °C for 30 min to remove any residual solvent.

group (OMe, 31 amu), ethyl ester group (OEt, 45 amu) and *N*-methylpyridinium (C<sub>6</sub>H<sub>8</sub>N, 94 amu) ca. 300–500 °C, although the parent ion of the indolizine was not observed.

Mass spectrometry data recorded for the poly(ethylene glycol) containing indolizine modified SWCNTs (**2b**) displayed the presence of methyl ester (OMe, 31 amu), *N*-methylpyridinium (C<sub>6</sub>H<sub>8</sub>N, 94 amu) and ethyl formyl group (OCOCH<sub>2</sub>CH<sub>3</sub>, 73 amu) as an indicative of the poly(ethylene glycol) grafted indolizine modified SWCNTs. Fragmented methyl ester group (OMe, 31 amu), *N*-methylpyridinium (C<sub>6</sub>H<sub>8</sub>N, 94 amu), phenyl group (C<sub>6</sub>H<sub>4</sub>, 76 amu), nitroso group (NO, 30 amu) and phenoxy group (C<sub>6</sub>H<sub>4</sub>O, 93 amu) confirmed the presence of a nitro phenyl group attached on **3b**.

In order to confirm the pyridinium salt formation and their conversion into indolizine *via* 'click' reaction, quaternary pyridinium salt modified (**1a** and **3a**) and indolizine modified (**1b** and **3b**) SWCNTs were further analyzed by X-ray photoelectron spectroscopy (XPS). Fig. 4 shows deconvoluted N 1s XPS spectra of (**1a**, **1b**, **3a** and **3b**). The deconvoluted N 1s XPS spectrum of **1a** showed three peaks that can be probably attributed to the quaternary nitrogen (N<sup>+</sup>) at ca. 402.3 eV,<sup>23,46</sup> pyridinic nitrogen at 398.9 eV (ref. 21, 23 and 35) and pyridine radical cation (N<sup>•+</sup>) at 400.2 eV.<sup>47,48</sup> After the 'click' reaction of **1a** with DMAD, the deconvoluted N 1s XPS spectrum of **1b** displayed two unchanged peaks positioned at ca. 399.9 and 398.9 eV and a new peak generated at 401.4 eV that probably correspond to indolizine ring nitrogen (–N<).<sup>49</sup> The deconvoluted N 1s XPS spectrum of **3a** showed signature of nitro group (–NO<sub>2</sub>),<sup>23</sup> pyridine diazo (–N=N),<sup>50</sup> quaternary nitrogen (N<sup>+</sup>)<sup>23</sup> and pyridine radical cation (N<sup>•+</sup>)<sup>47,48</sup> at ca. 406.2, 403.8, 402.4 and 400.1, respectively. Similar to **1b**, after the 'click' reaction, the deconvoluted N 1s XPS spectrum of **3b** displayed a new peak formation positioned at ca. 401.5 eV as an indication of indolizine formation. N 1s XPS spectra of both **1b** and **3b** showed no peak related to quaternary pyridinium nitrogen (N<sup>+</sup>), confirming that

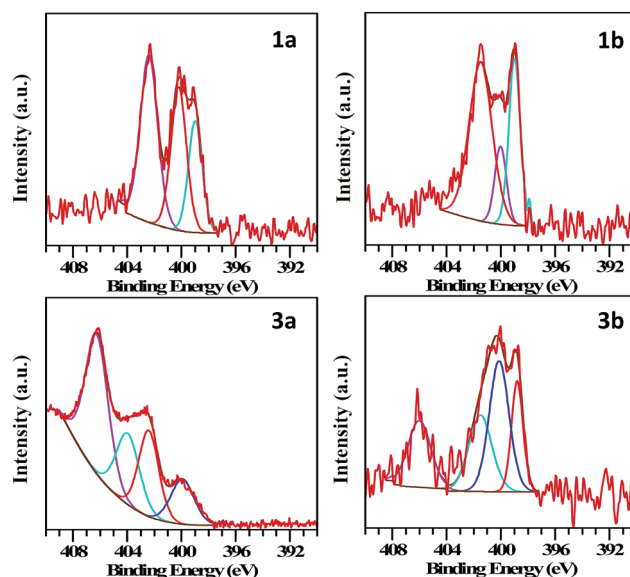


Fig. 4 Deconvoluted N 1s XPS spectra of the pyridinium salt modified SWCNTs (**1a** and **3a**) and indolizine modified SWCNTs (**1b** and **3b**). The data were fitted using a Tougaard background. Peak positions were constrained to ±0.2 eV and were fitted using a GL(30) lineshape. Peak FWHM was also constrained in the region 1–3.





pyridinium salts were successfully converted into pyridinium ylides, which subsequently reacted with DMAD to yield indolizines. Compared to the atomic per cent (at%) nitrogen (2.90%) found in SWCNT-Py, XPS elemental analysis results of **1b**, **2b** and **3b** verified the presence of  $2.22 \pm 0.33$ ,  $1.60 \pm 0.22$  and  $2.16 \pm 0.56$  atomic per cent (at%) nitrogen, respectively, suggesting nitrogen free new group addition (*e.g.* DMAD, PEG, benzene ring *etc.*) to SWCNT surface.

### Fluorescence properties of indolizine modified light-emitting SWCNTs

The presence of fluorescent groups, so-called indolizine heterocycles, on nanotube surface was also confirmed by fluorescence spectroscopy where the studies showed that the indolizine functionalized SWCNTs (**1b** and **2b**) emitted light upon excitation (Fig. 5). Excitation of **1b** and **2b** at 330 nm resulted in a blue emission at *ca.* 404 nm. To compare, free indolizine (FI), 3-ethyl 1,2-dimethyl indolizine-1,2,3-tricarboxylate was synthesised

following the literature<sup>45</sup> via the addition of ethyl-2-bromoacetate to pyridine to afford *N*-(ethoxycarbonylmethyl)-pyridinium bromide which undergoes the 1, 3-dipolar cycloaddition reaction with DMAD in the presence of a base ( $\text{NEt}_3$ ) (see Experimental section and ESI Scheme S1†). Compared to **1b** and **2b**, excitation of FI at 330 nm resulted in a blue shifted emission at *ca.* 382 nm, suggesting an electron transfer between electron rich indolizine and SWCNT upon the covalent attachment of indolizines to SWCNT surface. Control experiments mechanically mixing the FI with purified SWCNTs followed by washing with THF, acetone and ethanol showed no sign of fluorescence indicating that the fluorescence is not due to the physically adsorbed fluorescent indolizines (Fig. 5). In contrast to **1b** and **2b**, excitation of **3b** at 330 nm showed no sign of fluorescence (Fig. 5). It was no surprise since the structures like **3b** bearing strong electron withdrawing groups (*e.g.*  $-\text{NO}_2$  groups) are likely to be quenched.<sup>51</sup> Nitro groups of **3b** were reduced to amine ( $-\text{NH}_2$ ) moieties using  $\text{Sn}/\text{HCl}$ <sup>52</sup> to synthesize amino group modified SWCNTs (**3c**) (Scheme 1). As expected, excitation of **3c** at 330 nm emitted at 430 nm following the conversion (Fig. 5). XPS was used to confirm the reduction of nitro ( $\text{NO}_2$ ) groups into amines ( $\text{NH}_2$ ). N 1s XPS spectrum of **3c** showed three components positioned at 406.4, 401.0 and 399.2 eV, corresponding to nitro groups ( $-\text{NO}_2$ ),<sup>23</sup> the indolizine ring nitrogen ( $-\text{N}<$ )<sup>49</sup> and pyridinic nitrogen,<sup>35</sup> respectively (see ESI Fig. S2†). Decrease in the ratio of the relative peak intensity between the peak positioned at *ca.* 406.4 eV (nitro group nitrogen) and  $\sim 400$  eV (pyridinic nitrogen) ( $I_{406}/I_{400} = 0.57$ ) in N 1s XPS spectrum of **3b** (see Fig. 4) suggested that nitro groups were almost reduced when compared to the N 1s XPS spectrum of **3c** with  $I_{406}/I_{400}$  ratio of 0.08 (see ESI Fig. S2†).

Furthermore, the presence of a significant tail in the green region ( $\sim 500$  nm) of the emission spectra allowed us to optically image modified SWCNT materials (**1b**, **2b** and **3c**). Fig. 6 shows the white light optical image (left) and the epi-fluorescence image (right) of indolizine modified SWCNTs (**1b**, **2b** and **3c**). Fluorescence images were recorded using excitation light transmitted through a 420 to 480 nm excitation filter and fluorescence observed after passage through a 500 nm emission filter (Olympus CX-DMB-2 cube). The green spots can be clearly seen on the epi-fluorescence image (right) of modified SWCNTs (**1b**, **2b** and **3c**) as an indication of covalently attached light emitting indolizine groups on SWCNT surface, suggesting that the quantum efficiency of the emitted light is probably sufficient for practical applications (*e.g.* fluorescent imaging) (Fig. 6). However, we were unable to calculate the absolute quantum efficiencies of the indolizine modified SWCNTs (**1b**, **2b** and **3c**) due to the strong scattering and absorption of the SWCNTs. As comparison it was possible to determine the quantum yield of the FI heterocycle. The fluorescence quantum yield ( $\Phi_f$ ) of FI was determined from the corrected fluorescence spectra as 0.055, using anthracene dissolved in ethanol as a standard ( $\Phi_f^{\text{st}} = 0.27$  at  $25^\circ\text{C}$ ).<sup>51</sup>

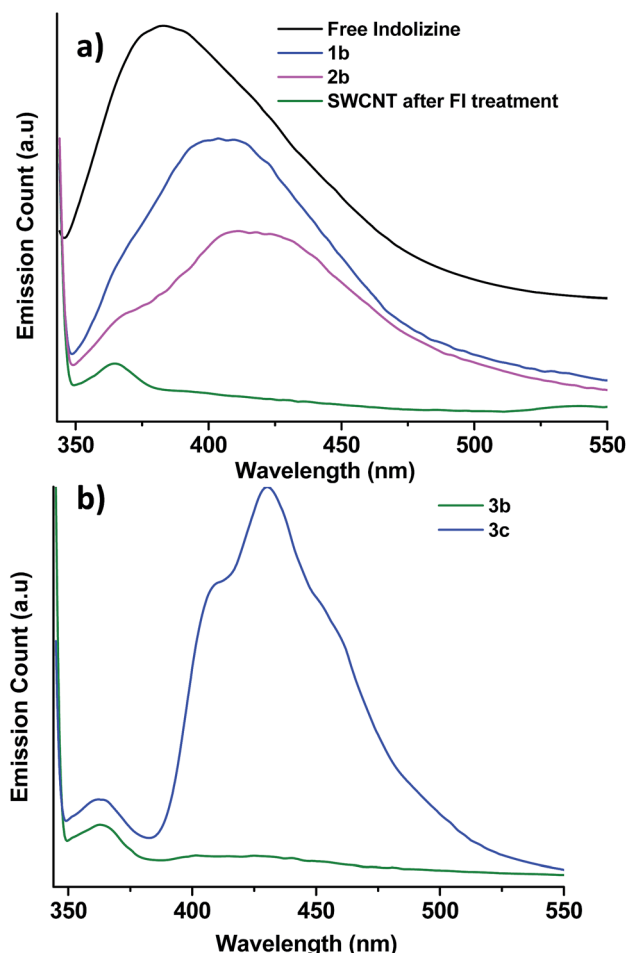


Fig. 5 Normalized and offset fluorescence spectra of (a) synthesized free indolizine (FI) structure (black line), **1b** (blue line), **2b** (magenta line), the control experiment of mechanically mixing purified SWCNTs with FI (olive line) recorded in  $\text{CH}_3\text{CN}$ ; (b) **3b** (olive line) and **3c** (blue line) recorded in  $\text{CH}_3\text{CN}$ . Curves in Fig. 6a and b are in top-down and down-top order, respectively.

### Solution state photochemical sensing properties of indolizine modified light-emitting SWCNTs

With the given fluorescent properties it is expected that indolizine modified SWCNTs may be used as molecular sensing



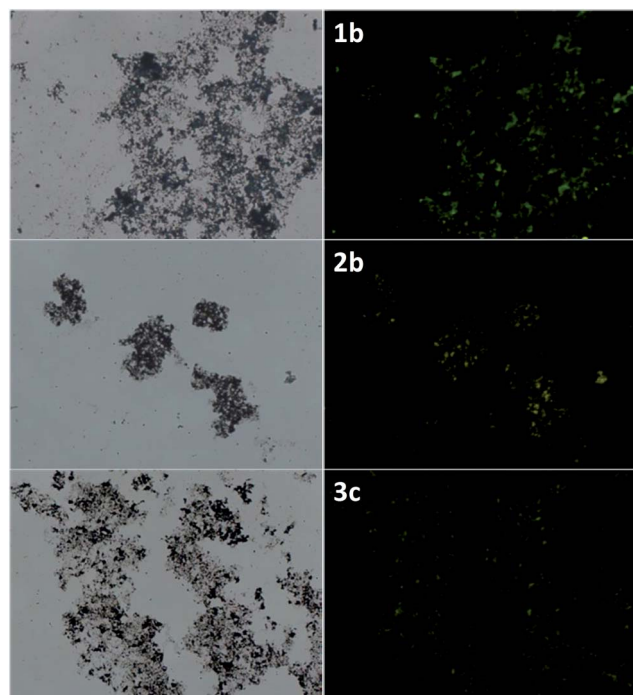


Fig. 6 Unmodified white light transmission image (left) and fluorescence image (right) of indolizine functionalized SWCNTs (**1b**, **2b** and **3c**) taken using an Olympus CX41 optical microscope, field of view  $718 \times 533 \mu\text{m}$ .

probes. Indolizine modified light-emitting SWCNTs (**1b**) was used as a *host molecule* to probe potentially dangerous nitroaromatics photochemically in solution-state. Phenol (P), 2-nitrophenol (2NP), 3-nitrophenol (3NP), 4-nitrophenol (4NP), 4-nitrotoluene (4NT) and 2,4-dinitrotoluene (2\_4NT) were selected as guest molecules. A literature method has been followed to evaluate the solution phase photochemical sensing properties of **1b**.<sup>53</sup> Shortly, a stock solution of **1b** ( $\sim 1.50 \mu\text{g mL}^{-1}$ ,  $\sim 1.25 \times 10^{-4} \text{ mol L}^{-1}$ ) and the guest molecules ( $1 \times 10^{-4} \text{ mol L}^{-1}$ ) were prepared in acetonitrile ( $\text{CH}_3\text{CN}$ ) for fluorescence spectral analysis. Each time a 3 mL solution of **1b** was filled in a quartz cell of 1 cm optical path length, and stock solutions of guest molecules were added into quartz cell portion wise (2, 2, 2, 2, 2, 5, 5, 5, 25, 25, 25, 100  $\mu\text{L}$ ) and mixed for 10 seconds. Measurements were performed immediately after mixing. An excitation wavelength of 330 nm and a temperature of  $25^\circ\text{C}$  were employed in all experiments. Emission spectra of the guest molecules in  $\text{CH}_3\text{CN}$  showed no emission upon excitation at 330 nm (see ESI Fig. S3†). Fluorescence response of **1b** (3 mL,  $\sim 1.25 \times 10^{-4} \text{ mol L}^{-1}$ ) to the guest molecules was initially tested by adding known amount of aromatic compound (P, 2NP, 3NP, 4NP, 4NT and 2\_4NT) (200  $\mu\text{L}$ ,  $1 \times 10^{-4} \text{ mol L}^{-1}$ ). Aromatic nitrophenol derivatives (2NP, 3NP and 4NP) caused a significant decrease in the fluorescence intensity of **1b** compared to P and nitrotoluene derivatives (4NT and 2\_4NT) (see ESI Fig. S4†). Fluorescence spectra of **1b** after the guest molecule addition suggested that fluorescent quenching efficiency (FQE) of 4NP was the highest among the guest molecules although it was not the least electron deficient one. FQE was observed in the order

of  $4\text{NP} > 2\text{NP} > 3\text{NP} > \text{P} > 4\text{NT} > 2,4\text{NT}$  (see ESI Fig. S4†). As expected, portion wise addition of 4NP (2, 2, 2, 2, 2, 5, 5, 5, 25, 25, 25, 100  $\mu\text{L}$ ) showed a significant, regular decrease trend in fluorescence intensity of **1b**, suggesting an intermolecular interaction between **1b** and 4NP (Fig. 7).

Detection limit of photochemical sensing was estimated as  $6.66 \times 10^{-8} \text{ mol L}^{-1}$ , considering the final concentration of 4NP after 2  $\mu\text{L}$  addition. However, the detection limit may be lower than the estimated value since some of 4NP molecules are likely to be physically adsorbed by SWCNT surface and reduce the concentration of free 4NP in analysed solution. Adsorption of 4NPs was previously shown where a SWCNT based device was developed assembling nanotubes to electrodes using AC dielectrophoresis technique which allows the real-time detection of adsorbed nitrophenols in aqueous solution.<sup>54</sup> The Stern–Volmer relationship allows us to investigate the kinetics of a photophysical intermolecular deactivation process and, luminescence in solution obeys the linear Stern–Volmer relationship,<sup>55</sup>  $F_0/F = 1 + k_q\tau_0[\text{Q}]$ , where  $F_0$  denotes the initial intensity of the fluorescent species without quencher,  $F$  represents the

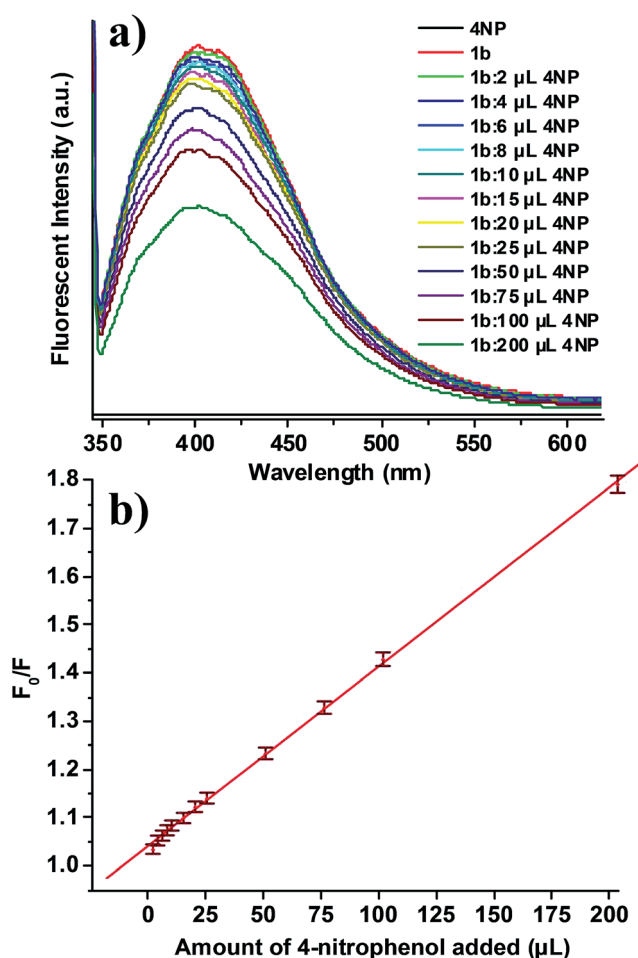


Fig. 7 (a) Change in the fluorescence spectra of **1b** ( $1.25 \times 10^{-4} \text{ M}$ ) excited at 330 nm after the addition of 2, 4, 6, 8, 10, 15, 20, 25, 50, 75, 100, 200  $\mu\text{L}$  equivalent of 4NP ( $1 \times 10^{-4} \text{ M}$ ) dissolved in  $\text{CH}_3\text{CN}$ ; (b) Stern–Volmer plot of the emission data provided by the change in fluorescence intensity after each 4NP addition.



final intensity of the fluorescent species with quencher,  $k_q$  is the quencher rate coefficient,  $\tau_0$  is the fluorescence lifetime of the fluorescent species without a quencher and  $[Q]$  is the concentration of the quencher. The Stern–Volmer plot constructed by the emission data obtained from the addition of 4NP to **1b** displayed a linear profile, probably indicating a single set of fluorophores, all equally accessible to the quencher (Fig. 8).<sup>56</sup> Similarly, fluorescent quenching of **1b** using other guest molecules (P, 2NP, 3NP, 4NT and 2,4NT) was also studied. Addition of 2NP and 3NP displayed similar decreasing trend in fluorescent intensity of **1b** while the change in the fluorescent intensity upon the addition of P, 4NT and 2,4NT is not significant (see ESI Fig. S5–S9†).

In order to further explore the intermolecular interactions between **1b** and 4NP, photochemical sensing properties of free indolizine (FI) structure, as a model, were also studied by fluorescence spectroscopy, FTIR and NMR. In agreement with **1b**, fluorescence spectral analysis of FI (3 mL,  $2.09 \times 10^{-7}$  M) in the presence of P, 2NP, 3NP, 4NP, 4NT and 2,4NT ( $200 \mu\text{L}$ ,  $1 \times 10^{-4}$  mol  $\text{L}^{-1}$ ) showed similar decreasing trend in the fluorescent intensity of FI, displaying a FQE order of  $4\text{NP} > 2\text{NP} > 3\text{NP} > \text{P} > 4\text{NT} > 2,4\text{NT}$  (see ESI Fig. S4†). Given the highest ability of 4NP to quench the fluorescence of FI this process was further investigated performing binding experiments. Similar to **1b**, FI was titrated with 4NP. As expected, similar trend in the fluorescence intensity of FI was observed after the addition of 4NP, confirming that the change in intensity is purely due to the intermolecular interactions between host and guest molecules (see ESI Fig. S10†). It further confirmed that SWCNTs were functionalized by light-emitting indolizine groups that are able to sense particularly nitrophenol derivatives.

To investigate the nature of intermolecular interactions between FI and 4NP, spectroscopic analysis of the FI, 4NP and possible FI:4NP complex were carried out by NMR and FTIR. Equimolar amounts of vacuum-dried FI (10.64 mg,  $3.28 \times 10^{-2}$  mmol) and 4NP (4.5 mg,  $3.28 \times 10^{-2}$  mmol) were dissolved in 1 mL of anhydrous freeze-dried  $\text{CDCl}_3$  for NMR spectroscopy

analysis. Dry experimental conditions ensured that there was no possible proton exchange with water. Fig. S11† shows the  $^1\text{H}$  NMR spectra of FI, 4NP and the mixture (FI:4NP). The  $^1\text{H}$  NMR spectrum of FI:4NP showed that the hydroxyl proton of 4NP ( $\text{H}_a$ ) at 5.74 ppm disappeared upon mixing with FI, suggesting a strong intermolecular interaction that can probably be attributed to H-bonding interaction/proton transfer between hydroxyl proton of 4NP and FI.<sup>43,57,58</sup> However, no significant signal shift was observed for the aromatic protons  $\text{H}_c$ ,  $\text{H}_e$ ,  $\text{H}_j$  and  $\text{H}_k$ , probably indicating weak or no  $\pi$ – $\pi$  interactions between FI and 4NP. Complementary FTIR spectroscopy further confirmed the possible H-bonding interaction between FI and 4NP. FTIR spectra of compounds were recorded by dropping the stock solutions of the compounds (FI and 4NP) and the mixture (FI:4NP) on vacuum dried KBr disk (see ESI Fig. S12†). FTIR spectrum of FI displayed three  $\nu_{\text{C=O}}$  stretching vibrations at 1740, 1710 and  $1695 \text{ cm}^{-1}$  due to the presence of ethoxy and methoxy ester carbonyl groups. In contrast, FTIR spectrum of the mixture (FI:4NP) showed a strong broad  $\nu_{\text{C=O}}$  stretching at  $1700 \text{ cm}^{-1}$ . Significant change in shape and position of the  $\nu_{\text{C=O}}$  stretching bands of FI after the addition of 4NP revealed a strong intermolecular interaction, probably H-bonding between ester carbonyl of FI and hydroxyl proton of 4NP.<sup>43,53,59</sup>

### Solid state photochemical sensing properties of indolizine modified light-emitting SWCNTs

Due to their exceptional physical and chemical properties, SWCNTs can be used as solid support materials for various applications including sensors.<sup>60,61</sup> Previously, it was shown that calculated photoluminescence quantum yield (PLQY) of fluorescent systems could be used as a tool to probe the sensing properties of fluorescent materials.<sup>62</sup> Similar to previous work, solid state photophysical properties of **1b** and **3c** were studied by exposing them to saturated 4NP vapour using a commercial spectrometer coupled with an integrating sphere.<sup>62</sup> The vapour pressure of 4NP at room temperature was previously reported as  $1.9 \times 10^{-5}$  mm Hg that corresponds to a vapour concentration of 19 ppb.<sup>63</sup> Fluorimetry experiments were performed on a quartz substrate coated with light-emitting SWCNTs. An excitation wavelength of 340 nm was selected for all measurements. PLQY of **1b** and **3c** was determined using the formalism outlined by de Mello where  $\Phi_{\text{PL}}$  is PLQY,  $E_i(\lambda)$  and  $E_0(\lambda)$  are, respectively, the integrated luminescence as a result of direct excitation of SWCNTs and secondary excitation,  $L_i(\lambda)$  and  $L_0(\lambda)$  are the integrated excitation when SWCNTs are directly excited and the excitation light first hits the sphere wall, respectively and,  $L_e(\lambda)$  is the integrated excitation profile for an empty sphere.<sup>64</sup>

$$\Phi_{\text{PL}} = \frac{E_i(\lambda) - (1 - A)E_0(\lambda)}{L_e(\lambda)A} \quad A = \frac{L_0(\lambda) - L_i(\lambda)}{L_0(\lambda)}$$

Using the equation, PLQY of **1b** and **3c** was calculated as 0.9 and 0.3%, respectively (Fig. 8). As expected the PLQY of the modified SWCNTs was less than the free indolizines (FI and  $\text{FI}_{\text{NH}_2}$  (ref. 43)) with calculated PLQY of 5.5 and 3.5%,

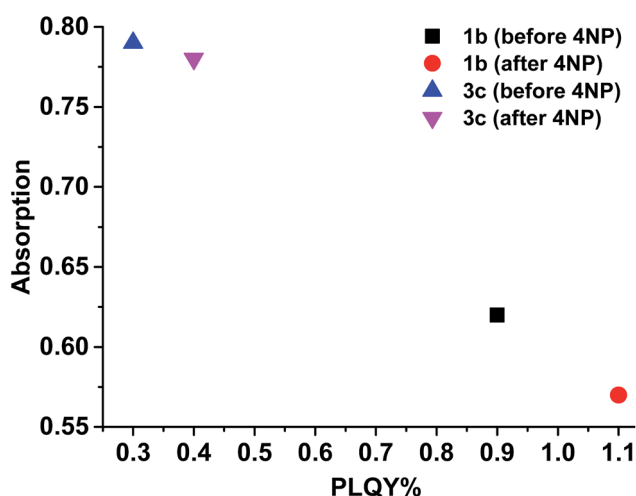


Fig. 8 Calculated photoluminescence quantum yield of **1b** and **3c** versus absorption of **1b** and **3c** before/after exposing them to 4NP vapour at room temperature.





respectively, probably due to the charge transfer from the electron rich indolizine system to the electron poor SWCNTs. These findings were in agreement with the blue-shift in emission spectra of **1b** and **3c** compared to pure spectra of FI and FI<sub>NH</sub>.<sup>43</sup> Similar decreasing trend in PLQY due to interaction between surface groups and SWCNT walls was previously reported in which the emission quantum yield of 2,4,6-triphenylpyrylium functionalized SWCNTs ( $\Phi = 1\%$ ) was lower than 2,4,6-triphenylpyrylium alone ( $\Phi = 33\%$ ).<sup>65</sup>

The change in absorption ( $A$  in the equation) of the corresponding indolizine modified SWCNTs (**1b** and **3c**) can be used as a measure of the photochemical response to 4NP vapour since the absorption of fluorescent material is expected to change upon the adsorption of 4NP to modified SWCNT surface.<sup>62</sup> Before exposing to 4NP vapour, the absorption values of **1b** and **3c** were determined as 0.62 and 0.79, respectively. The values decreased to 0.57 and 0.78 after exposing fluorescent SWCNTs to 4NP vapour for 10 seconds (Fig. 8). Furthermore, the calculated PLQY of **1b** and **3c** slightly increased to 1.1 and 0.4%, respectively following 4NP exposure (Fig. 8). The change in absorption ( $A$  in the equation) of the corresponding indolizine modified SWCNTs (**1b** and **3c**) together with the change in the calculated PLQYs revealed that nitrophenol derivatives can be photochemically probed by indolizine modified light-emitting SWCNTs in solid-state at room temperature on a solid substrate.

## Conclusions

A low temperature, highly controllable and efficient synthetic protocol to prepare indolizine modified light-emitting SWCNTs from pyridinium ylides generated on SWCNT surface was presented. Reactivity of the pyridine nitrogen lone pairs allowed the preparation of several quaternary pyridinium salt modified SWCNTs, which were subsequently used to synthesize pyridinium ylide functionalized SWCNTs and indolizine functionalized SWCNTs via 'click' chemistry. Shortly, the current study proposed a new perspective for the synthesis of carbon nanotube supported fluorescent molecules and their use in sensing of potentially dangerous nitrophenol derivatives. The indolizine modified light-emitting SWCNTs (**1b**, **2b** and **3c**) were shown to emit blue light (400–450 nm) upon excitation at ca. 330 nm. Fluorescent quenching of **1b** demonstrated that trace analysis of nitrophenol derivatives could be performed by indolizine modified light-emitting SWCNTs in both solution-state and solid-state. The fluorescent quenching was discussed work equally as well with **3c**. <sup>1</sup>H NMR and FTIR analysis of free indolizine (FI) structure suggested that the fluorescence of the modified materials quenched via probably an electron transfer from electron-rich indolizine to electron-deficient nitrophenolics through H-bonding interaction between ester carbonyls of indolizine and hydroxyls of nitrophenol. Optical microscopy images together with solid-state photophysical data revealed that indolizine modified fluorescent SWCNTs may potentially be used for imaging in cell transfection and drug delivery due to their high enough quantum efficiencies. Described methodology can be further expanded to introduce

various molecules including cyclodextrins and cucurbituril with fluorescent indolizine tags on SWCNT surface that may be used for protein sensing and crystallisation. Furthermore the preparation of fluorescent hierarchical networks of carbon nanomaterials (e.g. carbon nanotubes and graphene) are anticipated via the reaction of bisdipolarophiles (e.g. bisacetylenes) with pyridinium ylides generated on carbon materials. The principles outlined here are expected to be applicable to other low dimensional materials including graphene and related inorganic layered materials.

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