

# A near-infrared colorimetric fluorescent chemodosimeter for the detection of glutathione in living cells†

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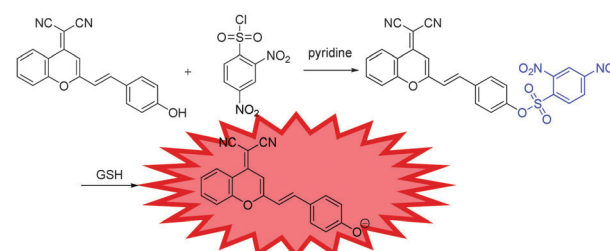
**A novel near-infrared (NIR) and colorimetric fluorescent molecular probe based on a dicyanomethylene-4*H*-pyran chromophore for the selective detection of glutathione in living cells has been developed. The fluorescence OFF–ON switch is triggered by cleavage of the 2,4-dinitrobenzenesulfonyl (DNBS) unit by the interaction with GSH.**

Biothiols such as glutathione (GSH), cysteine (Cys), and homocysteine (HCys) play a crucial role in maintaining appropriate redox homeostasis in biological systems.<sup>1,2</sup> GSH, the most abundant cellular thiol, is of great importance in cellular defense against toxins and free radicals.<sup>3</sup> Changes in the level of GSH concentration have been correlated with various diseases, such as AIDS, leukocyte loss, liver damage, cancer and neurodegenerative diseases.<sup>4</sup> Therefore, much effort has been devoted to the development of efficient methods for the detection of GSH, which is of significant interest in the fields of chemical, environmental and biological sciences.<sup>5,6</sup> Many fluorescent sensors for the detection of GSH have been developed based on various fluorophores, including BODIPY,<sup>7,8</sup> coumarin,<sup>9,10</sup> rhodamine,<sup>11,12</sup> squaraine,<sup>13</sup> fluorescein<sup>14,15</sup> and lanthanide based probes.<sup>16</sup> However, the chromophore of dicyanomethylene-4*H*-pyran has seldom been exploited in the detection of thiols.<sup>17,18</sup> Due to their emission located at the red or near infra-red (NIR) region, (DCM)-type derivatives are particularly suitable for application in biological samples, since they produce lower background fluorescence with less scattering, penetrate much deeper into tissues than UV and visible light and cause little damage to living cells.<sup>19,20</sup> The DCM fluorophore is more photostable than comparable NIR fluorophores such as squaraine and cyanine. Although our previously reported BODIPY fluorophore<sup>21</sup> is photostable it is much harder to synthesise than the DCM chromophore described here.

A number of sensing mechanisms have been employed for thiol detection, including Michael addition,<sup>22</sup> to –CHO attached fluorophores,<sup>23</sup> and deprotection of 2,4-dinitrobenzenesulfonyl (DNBS),<sup>24–26</sup> *etc.* Among them, the DNBS group is especially preferable as an efficient recognition unit for thiols due to its unique sensitivity and high reactivity toward thiolate anions with OFF–ON signalling. With these in mind, we demonstrate the development of a colorimetric and NIR fluorescence turn-on thiol probe **1** (Scheme 1) containing DCM as the fluorophore and DNBS as the fluorescence quencher and recognition moiety. Specifically, we attached a benzene unit to a dicyanopyran moiety, which extended the conjugated system and produced an emission band centred at about 690 nm, thus successfully making the wavelength fall in the desired NIR region.<sup>27</sup> Particularly, the developed probe **1** exhibits several meritorious features: (i) high sensitivity and selectivity to sulfhydryl-containing species; (ii) significant enhancement of NIR fluorescence upon removal of DNBS by GSH; (iii) excellent performance in live cell imaging.

Probe **1** is synthesized by stirring 2-(2-(4-hydroxystyryl)-4*H*-chromen-4-ylidene)malononitrile and 2,4-dinitrobenzene-1-sulfonyl chloride in the presence of pyridine at room temperature, to give probe **1** in 45% yield, which was fully characterized by <sup>1</sup>H NMR, <sup>13</sup>C NMR, and HRMS, as shown in the ESI† (Fig. S3 and S8). The GSH-promoted specific O–S cleavage of probe **1** liberates DCPO<sup>–</sup> with an intramolecular charge transfer (ICT) emission band falling in the desired NIR region at about 690 nm (Scheme 1).

Initially, UV-vis absorption of probe **1** was investigated. Probe **1** has an intense absorption at 414 nm (Fig. 1a) in a



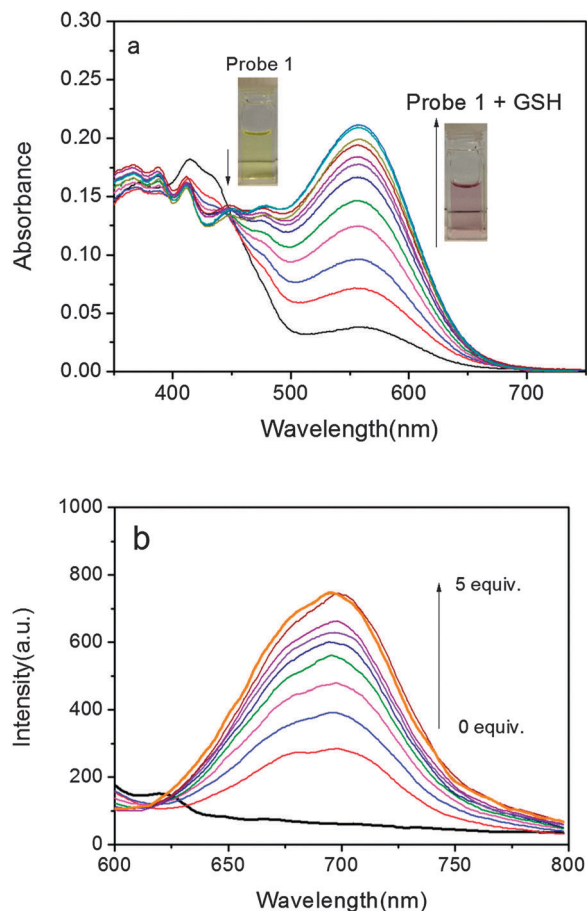
Scheme 1 Synthesis of probe **1** and GSH-promoted release of DCPO<sup>–</sup>.

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**Fig. 1** (a) Absorption spectra of probe **1** ( $1 \times 10^{-5}$  M) at different concentrations of GSH (0, 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 1.0, 2.0, 5.0 equiv.) in a mixture of DMSO–water (50 : 50, v/v) with a PBS buffer solution (10 mM, pH = 7.4). Insets: colour changes of probe **1** upon addition of GSH (5 equiv.). (b) Fluorescence spectra of probe **1** ( $1 \times 10^{-5}$  M) at different concentrations of GSH when excited at 560 nm.

DMSO–PBS buffer solution (pH = 7.4, 50/50, v/v) at 37 °C. Upon the addition of GSH, the colour of the solution turned from slight yellow to pink, which was clearly recognizable by the naked eyes. At the same time, a concomitant increase of a new absorption band at 560 nm was observed with an isosbestic point at 446 nm. As illustrated in Fig. 1a, due to the specific O–S cleavage, a distinct 146 nm red shift in absorbance was observed (Scheme 1). Since the phenolate group is a much stronger electron donor than the sulfonate group, the ICT efficiency should be significantly enhanced by the interaction of probe **1** with GSH and thus shifting the absorption to a longer wavelength.

Subsequently, the fluorescence spectra of probe **1** in the absence and presence of GSH were recorded. The probe alone exhibits almost no emission when excited at 560 nm and no distinct variations are observed over time, suggesting that probe **1** is stable and has no tendency to convert into DCPO<sup>−</sup> under the measurement conditions. In the presence of GSH, however, fluorescence at 690 nm was dramatically enhanced (Fig. 1b).<sup>‡</sup> The departure of the electron-withdrawing DNBS moiety *via* a GSH-induced O–S bond cleavage releases DCPO<sup>−</sup>, which possesses strong ICT and induces a turn-on NIR fluorescence response with a high off/on ratio. On the other

hand, the large Stokes shift of ~130 nm is also conducive to decrease the background fluorescence, enhancing the signal fidelity. Moreover, the response rate of **1** to GSH was tested by time-course fluorescence measurements. In the presence of GSH, the fluorescence intensity at 690 nm increases gradually and reaches a plateau after about 5 min (Fig. S1<sup>†</sup>), indicating the completion of the reaction.

To evaluate the selectivity of the developed probe **1** for sulfhydryl-containing species, a series of amino acids were examined. Not surprisingly, except for GSH, probe **1** shows similar response to other sulfhydryl-containing compounds such as cysteine, homocysteine, dithiothreitol, while no measurable fluorescence enhancement could be triggered by the treatment of other amino acids (Fig. 2). In fact, the plausible disturbance of DTT, Cys and HCys can be neglected due to their relatively low concentration in biological systems.<sup>28</sup> To gain insight into the sensing mechanism, mass spectral analysis was further performed. The reaction between probe **1** and GSH that occurred by the cleavage of O–S with the release of DCPO<sup>−</sup> was confirmed by the appearance of the peak at 311.0828 in the mass spectrum (Fig. S3<sup>†</sup>). Furthermore, the fluorescence intensity at 690 nm shows a linear relationship ( $R = 0.98$ ) with GSH concentrations from 1 to 10  $\mu$ M, and the detection limit was evaluated to be  $1.8 \times 10^{-8}$  M, indicating that probe **1** is particularly sensitive to the detection of GSH (Fig. S2<sup>†</sup>).

The effect of pH on the photophysical properties of DCPO and probe **1** was then investigated (Fig. S4<sup>†</sup> and S6<sup>†</sup>). As illustrated, we believe that the probe is quite stable in aqueous media up to pH 8, whereas DCPO displays a corresponding fluorescence enhancement in the pH range of 5.7–8.0, which clearly demonstrates that the fluorescence response of probe **1** in the physiological pH range is due to the presence of thiols and verifies the generation of DCPO<sup>−</sup> after treatment with GSH. Therefore, we decided to employ the probe for fluorescent imaging of cellular thiols. The  $pK_a$  of DCPO was calculated to be 8.70 (Fig. S5<sup>†</sup>).

The ability of probe **1** to detect GSH in HeLa cells was examined using confocal fluorescence microscopy (Fig. 3). HeLa cells were



**Fig. 2** Fluorescence spectra of probe **1** in the absence and presence of various amino acids and GSH, DTT, Cys, HCys in a mixture of DMSO–water (50 : 50, v/v) with a PBS buffer solution (10 mM, pH = 7.4). amino acids are arginine, threonine, serine, isoleucine, asparaginic acid, sarcosine, valine, glutamine, tryptophan, glutamic acid, proline, leucine, histidine, glycine, phenylalanine, alanine, asparagine, methionine.



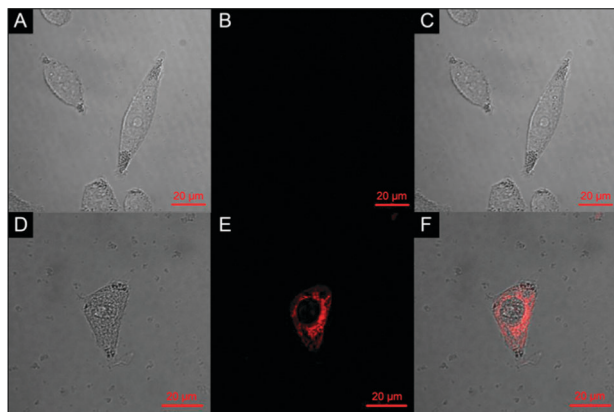


Fig. 3 Confocal fluorescence images in HeLa cells: (top) (A–C) cell incubated without probe **1**; (bottom) (D–F) cell incubated with probe **1** (20  $\mu$ M) for 0.5 h. Emission was collected at 660–740 nm upon excitation at 488 nm. Bright field (A and D), fluorescence (B and E) and overlap field (C and F).

cultured in RPMI 1640 supplemented with 10% FCS and then incubated with 10% fetal bovine serum followed by 20  $\mu$ M of probe **1** in PBS for 30 min. 2.5% DMSO was used in the cell culture process. Bright-field measurements confirmed that the cells after treatment with probe **1** were viable throughout the imaging experiments, indicating the superior biocompatibility of probe **1**. After the co-culture with probe **1**, remarkable intracellular NIR fluorescence could be detected. The overlay of fluorescence and bright-field images indicates that the fluorescence localizes mainly in the cytosol, indicative of the subcellular distribution of GSH and excellent membrane permeability of probe **1**. In contrast, cells that are not treated with probe **1** display nearly no fluorescence, verifying the fluorescence source of the **1**-treated cells (Fig. 3B). Moreover, the images also indicate that the probe has low toxicity since the cellular morphology is maintained. With this bioimaging experiment we have demonstrated the potential of probe **1** for the imaging of GSH in living cells.

In conclusion, we have designed and synthesized a NIR fluorescent chemodosimeter for GSH based on a DCM framework, showing a subcellular distribution of GSH and good cell membrane permeability. The fluorescence enhancement mechanism is based on the cleavage of DNBS from the fluorophore by GSH, which switches the dark excited state to an emissive excited state. Probe **1** displays a colour change from yellow to pink upon addition of GSH, and thus can serve as a “naked-eye” probe for GSH. Furthermore, we have also demonstrated that the probe can be used for the fluorescent imaging of cellular thiols. Since our DCM based system is very easy to prepare and has a long emission wavelength and high photo-stability, we believe that it will become the probe of choice for biological applications.

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## Notes and references

‡ When probe **1** is excited at the isosbestic point (446 nm, Fig. S7<sup>†</sup>) a fluorescence increase at 581 nm is observed which is ascribed to release of the DNBS quenching group. Conversely, when probe **1** is excited at 560 nm the large observed increase in fluorescence at 690 nm is due to the fluorescence of the DCPO<sup>−</sup> produced upon cleavage of DNBS from probe **1**.

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