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A novel highly specific and ultrasensitive fluorescent probe for monitoring hypochlorous acid and its application in live cells†

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Based on the important role of hypochlorous acid (HOCl) in the immune system and numerous physiological processes, the detection of intracellular basal HOCl is of significant interest. In this work, we present a simple thiocarbamate-protected fluorescein fluorescent probe, TCFL, for imaging basal HOCl in live cells. Surprisingly, probe TCFL could determine HOCl quantitatively in a large concentration range with a detection limit of 0.65 nM. In addition, probe TCFL showed excellent specificity for HOCl in the presence of other higher concentration analytes (1 mM). Moreover, probe TCFL exhibited a rapid response (within 3 s) to HOCl and thus could provide a tool for real-time monitoring of HOCl. Importantly, probe TCFL with outstanding response features could be applied for monitoring basal HOCl in live cells.

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1. Introduction

As an important reactive oxygen species (ROS), endogenous HOCl is generated from the peroxidation of Cl^- and H_2O_2 assisted by myeloperoxidase in biological systems.^{1,2} Hypochlorous acid at low concentration plays an important role in immune defense and cellular signal transduction in bio-systems.^{3–5} However, abnormal production of hypochlorous acid can lead to various health disorders because of the highly reactive characteristics of HOCl.^{6–8} Therefore, the development of a proper tool for tracking basal HOCl in live cells is essential.

Fluorescent probes have attracted much attention owing to their outstanding sensitivity and unique bioimaging applications in live cells.^{9–12} To date, although many fluorescent probes for HOCl have been explored, most of them cannot specifically detect intracellular basal HOCl because of their delayed response, and unsatisfactory sensitivity and selectivity.^{13–45} Additionally, most of the available probes possess poor water solubility, which hinders their bioimaging applications. Therefore, constructing water soluble, highly selective and sensitive fast-response fluorescent probes is of great importance to monitor intracellular basal HOCl levels.

Herein, we present a water soluble thiocarbamate-caged fluorescein fluorescent probe, TCFL, for detecting HOCl with high selectivity and sensitivity. The TCFL probe exhibited a fast response (within 3 s), ultrasensitivity (detection limit = 0.65 nM) and preeminent specificity for HOCl. Importantly, the bioimaging results indicated that the TCFL probe could achieve specific monitoring of basal HOCl levels in live cells.

2. Experimental section

2.1. Materials and instruments

All chemicals in this work were obtained commercially and used without further purification. High resolution mass spectra (HRMS) were recorded on a LC-MS 2010A (Shimadzu) instrument. ^1H and ^{13}C NMR data were acquired on a Bruker AV-400 NMR spectrometer. Absorption data were measured on a UV-3101PC spectrophotometer. Fluorescence data were obtained on a Horiba FluoroMax-4 spectrophotometer. Fluorescence imaging of HOCl in live RAW 264.7 macrophage cells was performed on a Leica TCS SP5 microscope.

2.2. Synthesis of probe TCFL

Dimethylthiocarbamoyl chloride (1.24 g, 10 mmol) was slowly added to fluorescein (332 mg, 1 mmol) in dry CH_2Cl_2 (15 mL) in batches. Subsequently, *N*-diisopropylethylamine (500 μL) was slowly added to the reaction mixture in batches. The resulting mixture was stirred at room temperature until the reaction was complete. After removing the solvents by decompression-evaporation, the residues were purified by silica column chromatography (CH_2Cl_2 as an eluent) to

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achieve pure fluorescent probe **TCFL** (436 mg, 86%). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm): 3.311 (s, 3H), 3.361 (s, 3H), 6.583–6.635 (m, 2H), 6.725 (d, $J = 1.6$ Hz, 1H), 6.794 (d, $J = 8.8$ Hz, 1H), 6.855 (d, $J = 8.4$ Hz, 1H), 7.161 (d, $J = 2.4$ Hz, 1H), 7.333 (d, $J = 3.6$ Hz, 1H), 7.746 (t, $J = 7.2$ Hz, 1H), 7.820 (t, $J = 8.0$ Hz, 1H), 8.036 (d, $J = 7.6$ Hz, 1H), 10.229 (s, 1H); $^{13}\text{C-NMR}$ (100 MHz, $\text{DMSO}-d_6$) δ (ppm): 39.11, 43.34, 82.57, 102.73, 109.60, 111.71, 113.54, 116.80, 119.70, 124.52, 125.29, 126.33, 128.99, 129.60, 130.78, 136.27, 151.43, 152.06, 152.77, 155.44, 160.18, 169.01, 186.05. HRMS (ESI): calcd for $\text{C}_{23}\text{H}_{18}\text{NO}_5\text{S}$ $[\text{M} + \text{H}]^+$ 420.0900; found, 420.0906.

3. Results and discussion

3.1. Design and synthesis of the fluorescent probe **TCFL**

Recently, Falck *et al.* demonstrated that a dimethylthiocarbamate (DMTC) group is a desirable protecting group of hydroxyl moieties, and its cleavage can be realized by means of the oxidation of a sulfide.⁴⁶ Thus, we reasonably envisioned that the DMTC moiety might be a preferred recognition receptor of HOCl with strong oxidation capabilities. Fluorescein, as a commercially available dye, has been successfully used in the development of different fluorescent probes for monitoring various bioactive molecules in biosystems owing to its excellent photophysical properties.^{47–49} Additionally, modification of the hydroxyl moiety results in a spironolactone structure with suppressed fluorescence.^{50–52} Therefore, we deemed that DMTC-caged fluorescein with reduced fluorescence could once again exhibit enhanced fluorescence by means of the removal of DMTC, induced by the oxidation of HOCl. Bearing the above-mentioned considerations in mind, we synthesized DMTC-caged fluorescein as highly selective fluorescent probe **TCFL** for detecting HOCl with excellent sensitivity (Scheme 1).

3.2. The determination of HOCl by probe **TCFL**

Because probe **TCFL** possesses ideal water solubility, the fluorescence spectra of probe **TCFL** for detection of HOCl were evaluated in a buffered aqueous solution (pH 7.4, PBS 5 mM). As shown in Fig. 1, the free probe **TCFL** exhibited faint fluorescence. Surprisingly, an ultrafast (within 3 s) and enormous fluorescence enhancement at 515 nm was observed in the presence of HOCl. This feature suggested that probe **TCFL** has potential for real-time and ultrasensitive detection of HOCl in the environment and biological systems.

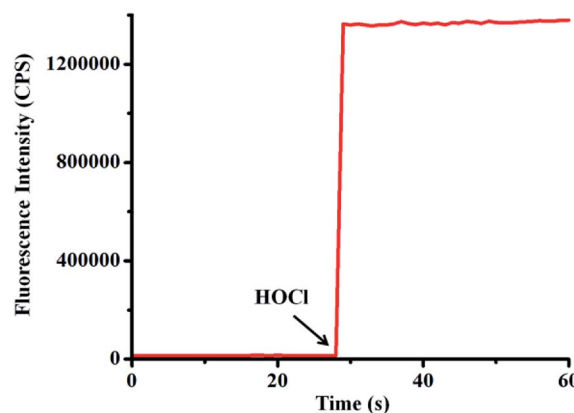
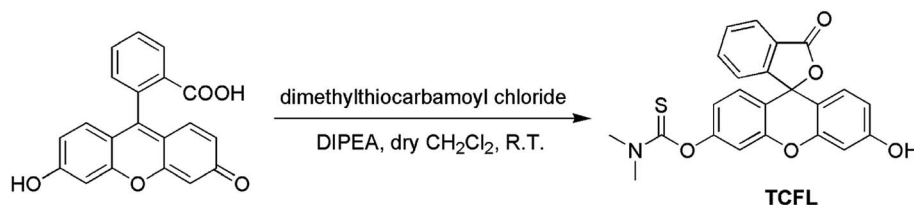


Fig. 1 Time-course of probe **TCFL** (5 μM) for monitoring HOCl (8 μM) in PBS solution (5 mM, pH = 7.4) at 25 $^\circ\text{C}$. $\lambda_{\text{ex}} = 470$ nm, $\lambda_{\text{em}} = 515$ nm. Slit widths: $W_{\text{ex}} = W_{\text{em}} = 1$ nm.

Additionally, the fluorescence intensity at 515 nm was enhanced gradually with an increasing amount of HOCl (Fig. 2a). The fluorescence intensity at 515 nm showed a satisfactory linear relationship with the concentration of HOCl ranging from 1–8 μM (linear equation: $y = 200\,470 \times [\text{HOCl}] (\mu\text{M}) - 91\,102$, $R^2 = 0.9988$) (Fig. 2b). Moreover, to examine the feasibility of using probe **TCFL** to measure extremely low levels of HOCl, the titration of HOCl at ultralow concentrations was carefully carried out. Satisfactorily, a good linear relationship between the fluorescence intensity at 515 nm and the HOCl concentration ranging from 0–30 nM was successfully obtained, which indicated that probe **TCFL** could accurately detect HOCl at the nanomolar level (Fig. 2c). Furthermore, the detection limit (DL) was calculated to be 0.65 nM ($3\sigma/k$). Consequently, we envisioned that **TCFL** would be capable of tracking the basal HOCl levels in live cells without exogenous stimuli.

Next, the absorption spectra of probe **TCFL** in the presence of HOCl were recorded. The absorption spectra at around 490 nm generally increased with the gradual addition of HOCl (0–8 μM) (Fig. 3a). Additionally, the absorbance of probe **TCFL** at 490 nm was linearly proportional ($R^2 = 0.9971$) to the concentration of HOCl ranging from 0–8 μM (Fig. 3b). These data indicated that probe **TCFL** could accurately detect HOCl with satisfactory sensitivity using the absorption spectrum method.



Scheme 1 The synthesis of the fluorescent probe **TCFL**.



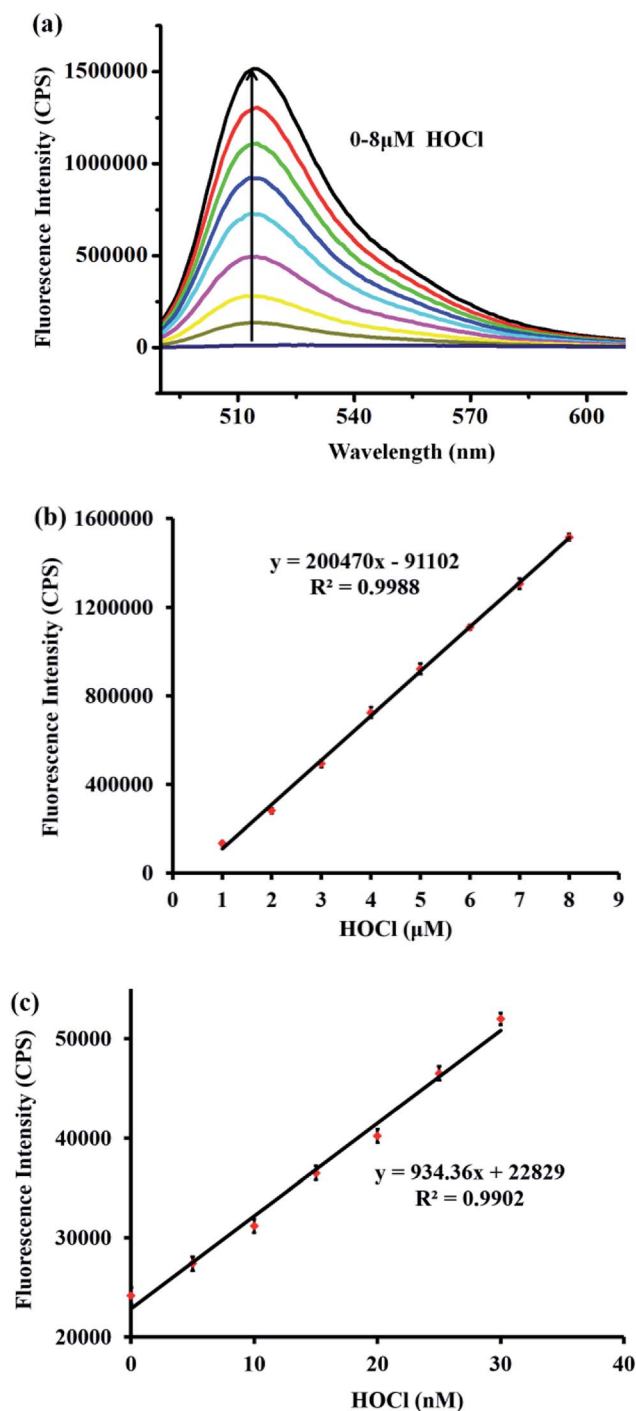


Fig. 2 (a) The fluorescence spectra of probe TCFL (5 μM) with increasing concentration of HOCl (0–8 μM) in PBS solution (5 mM, pH = 7.4). (b) The fluorescence intensity at 515 nm of probe TCFL (5 μM) as a function of HOCl concentration (1–8 μM). $\lambda_{\text{ex}} = 470$ nm. Slit widths: $W_{\text{ex}} = W_{\text{em}} = 1$ nm. (c) The fluorescence intensity at 515 nm of probe TCFL (1 μM) as a function of HOCl concentration (0–30 nM). $\lambda_{\text{ex}} = 470$ nm. Slit widths: $W_{\text{ex}} = 1$ nm, $W_{\text{em}} = 2$ nm. Each spectrum was recorded 1 min after HOCl addition at 25 $^{\circ}\text{C}$. Error bar = RSD ($n = 3$).

3.3. Specificity of probe TCFL for detecting HOCl

Next, we studied the specificity of probe TCFL for measuring HOCl. As expected, compared with HOCl, various other substances including K^+ , Ca^{2+} , Mg^{2+} , Na^+ , Zn^{2+} , Al^{3+} , NO_3^- ,

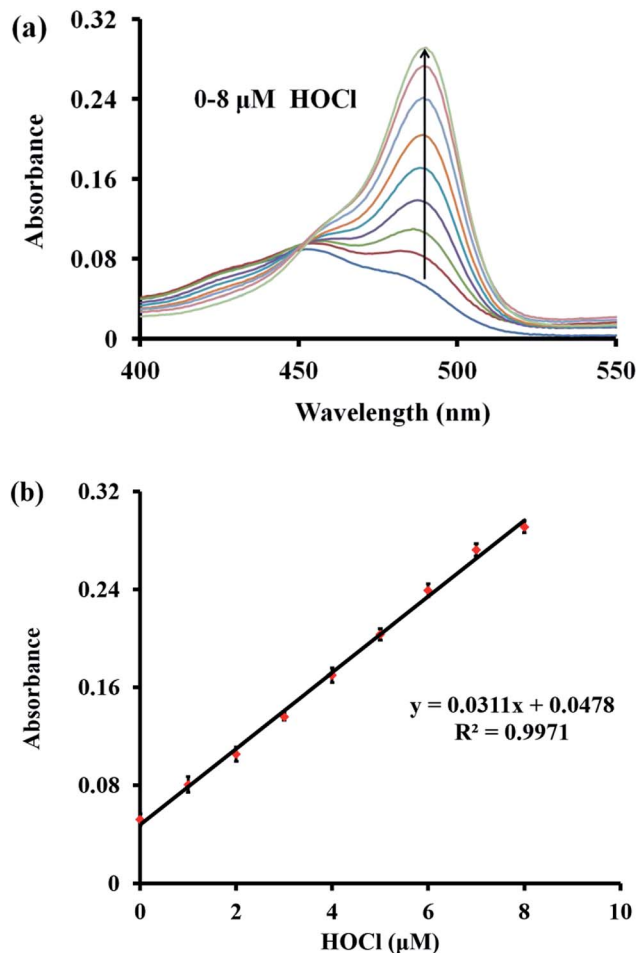


Fig. 3 (a) Absorption responses of probe TCFL (5 μM) in the presence of different concentrations of HOCl (0–8 μM) in PBS solution (5 mM, pH = 7.4). (b) The absorbance of TCFL at 490 nm as a function of HOCl concentration (0–8 μM). Each spectrum was recorded 1 min after HOCl addition at 25 $^{\circ}\text{C}$. Error bar = RSD ($n = 3$).

NO_2^- , Cu^{2+} , Fe^{3+} , Fe^{2+} , cysteine (Cys), H_2O_2 , *tert*-butylhydroperoxide (TBHP), hydroxyl radical ($\cdot\text{OH}$), *tert*-butoxy radical ($\cdot\text{O}^t\text{Bu}$), NO, $^1\text{O}_2$, O_2^- , glutathione (GSH), and ONOO^- at a higher concentration (1 mM) resulted in no significant change in fluorescence intensity at 515 nm, indicating that probe TCFL displays the excellent specificity toward HOCl (Fig. 4). The preeminent selectivity of probe TCFL for monitoring HOCl can be attributed to the distinctive DMTC recognition receptor.

3.4. Sensing mechanism of probe TCFL for detecting HOCl

To uncover the recognition mechanism of probe TCFL with HOCl, a reaction mixture of probe TCFL and HOCl was carefully analyzed by HRMS. When HOCl was added, the mass peak of probe TCFL (m/z 420.0906 [$\text{M} + \text{H}]^+$) disappeared, and a new mass peak at m/z 333.0756 was discovered, which indicated the production of fluorescein (calcd m/z 333.0757 [$\text{M} + \text{H}]^+$). Based on the published results,^{45,46} a plausible reaction mechanism of probe TCFL for detecting HOCl is presented in Scheme 2.



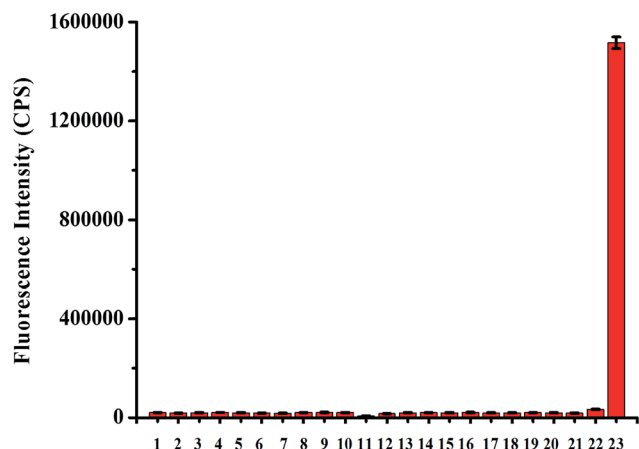


Fig. 4 The fluorescence response of probe TCFL (5 μ M) in the presence of various analytes (1 mM except for 8 μ M HOCl) in PBS solution (5 mM, pH = 7.4). 1. Blank; 2. K^+ ; 3. Ca^{2+} ; 4. Mg^{2+} ; 5. Na^+ ; 6. Zn^{2+} ; 7. Al^{3+} ; 8. NO_3^- ; 9. NO_2^- ; 10. Cu^{2+} ; 11. Fe^{3+} ; 12. Fe^{2+} ; 13. Cys; 14. H_2O_2 ; 15. TBHP; 16. $\cdot OH$; 17. $\cdot O^tBu$; 18. NO; 19. 1O_2 ; 20. O_2^- ; 21. GSH; 22. $ONOO^-$; 23. HOCl. The bars represent the fluorescence intensities at 515 nm and each spectrum was acquired 1 min after the addition of each analyte at 25 $^{\circ}C$. Error bar = RSD ($n = 3$).

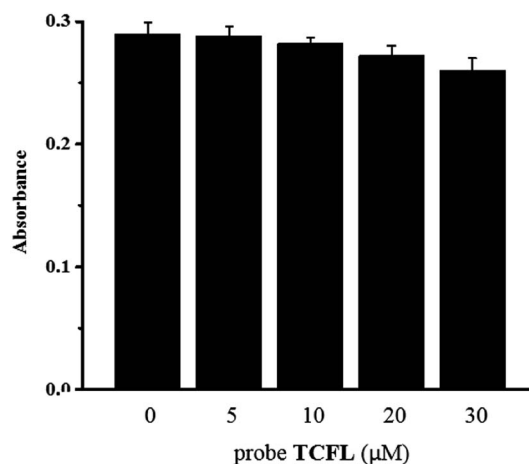
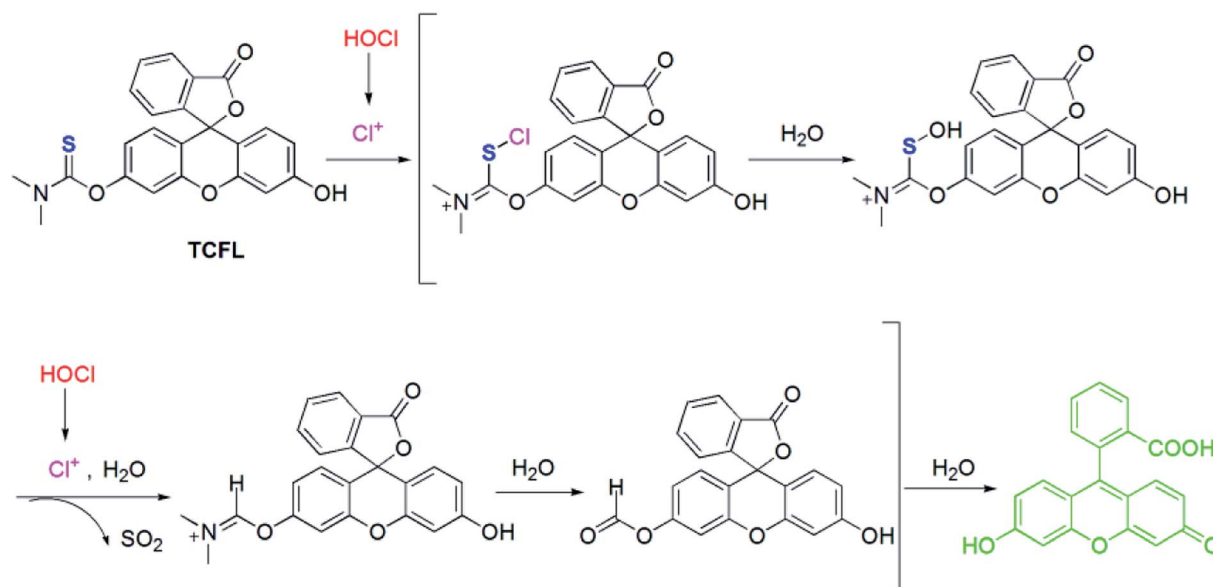


Fig. 5 Cytotoxicity tests of probe TCFL at different concentrations in RAW 264.7 macrophage cells.

3.5. Bioimaging applications of probe TCFL

Firstly, according to 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) tests, the cytotoxicity of probe TCFL was studied in live RAW 264.7 macrophage cells. As illustrated in Fig. 5, probe TCFL exhibits low cytotoxicity. Then, probe TCFL was applied to monitor HOCl in live RAW 264.7 macrophage cells. After incubation with the TCFL probe (10 μ M) for 30 min, the cells showed a medium green fluorescence (Fig. 6a and b), indicating that probe

TCFL was sensitive enough as a tool to monitor the basal HOCl levels in live cells without stimulation. The application of probe TCFL for the detection of exogenous HOCl was further performed (Fig. 6c and d). Additionally, cells pre-treated with PMA (a ROS stimulant) exhibited a relatively brighter fluorescence (Fig. 6e and f). Then, the hypochlorous acid in the cells was scavenged by NAC, and the cells displayed a dramatic reduction in fluorescence (Fig. 6g and h). These data demonstrated that probe TCFL could monitor the changes in endogenous/exogenous HOCl levels in live cells (Fig. 6i), which is very meaningful to unravel the physiological and pathological functions of HOCl. These results fully confirmed that our proposed probe TCFL can act as a robust tool for imaging basal HOCl and the fluctuations of endogenous/exogenous HOCl levels in live cells.



Scheme 2 The response mechanism of the fluorescent probe TCFL for monitoring HOCl.



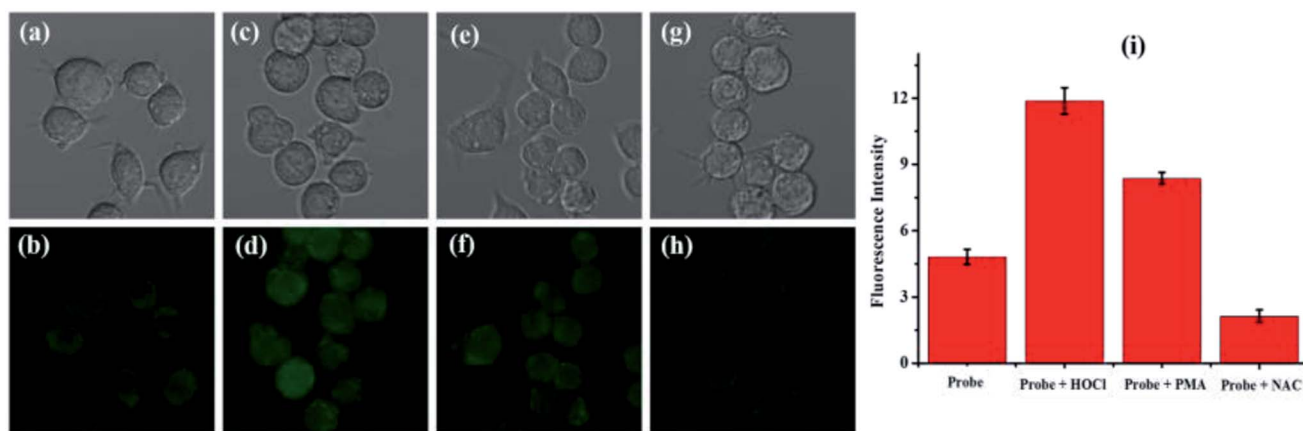


Fig. 6 Fluorescence imaging of HOCl in live RAW 264.7 macrophage cells: (a and b) the cells loaded with probe TCFL (10 μ M) for 30 min; (c and d) the cells incubated with probe TCFL (10 μ M) for 30 min followed by the addition of HOCl (20 μ M) for 10 min; (e and f) the cells loaded with probe TCFL (10 μ M) for another 30 min after preincubation with 2.0 μ g mL⁻¹ PMA for 60 min; (g and h) the cells loaded with probe TCFL (10 μ M) for another 30 min after preincubation with 500 μ M NAC for 60 min. (i) The bars represent the fluorescence intensities of the corresponding cells. The provided images of live RAW 264.7 macrophages are representative ones ($n = 10$ fields of cells).

4. Conclusions

In summary, an efficient DMTC-based specific fluorescent probe, TCFL, with excellent water solubility and ultra-sensitivity for monitoring HOCl in live cells has been designed. Probe TCFL can realize quantitative detection of HOCl from 0–30 nM. Moreover, probe TCFL exhibits noteworthy selectivity for HOCl over other biologically relevant species, which was attributed to the contribution of the unique DMTC recognition receptor. In particular, fluorescence imaging demonstrated that probe TCFL could be effectively applied in imaging basal HOCl levels in live cells. In brief, this highly specific and ultrasensitive fluorescent probe will provide a remarkable tool to track the HOCl levels in aqueous environments and biological systems.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

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