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A selective and sensitive near-infrared fluorescent probe for real-time detection of Cu(I)†

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The disruption of copper homeostasis (Cu^+/Cu^{2+}) may cause neurodegenerative disorders. Thus, the need for understanding the role of Cu^+ in physiological and pathological processes prompted the development of improved methods of Cu^+ analysis. Herein, a new near-infrared (NIR) fluorescent turn-on probe (**NPCu**) for the detection of Cu^+ was developed based on a Cu^+-mediated benzylic ether bond cleavage mechanism. The probe showed high selectivity and sensitivity toward Cu^+, and was successfully applied for bioimaging of Cu^+ in living cells.

Introduction

As an essential trace transition metal element, found in both the oxidized Cu^{2+} and reduced Cu^+ states in living organisms, copper is considered a vital redox-active cofactor for various cytosolic, mitochondrial and vesicular oxygen-processing enzymes, including cytochrome-*c*-oxidase,[1,2] copper/zinc superoxide dismutase[3] and metallothionein.[4] The disproportionation of Cu^+ in cells could produce reactive oxygen species (ROS), leading to oxidative damage of proteins, nucleic acids and lipids.[5–7] There is a growing body of evidence to suggest that the imbalance of Cu^+ may cause neurodegenerative disorders, such as prion, Parkinson's, Alzheimer's, Menkes, and Wilson's diseases and amyotrophic lateral sclerosis.[8–11] In addition, copper was also demonstrated to play a critical role in other diseases including urinary tract infection[12] and infertility.[13] It is of significance, therefore, to establish an effective and reliable strategy to monitor Cu^+ in complex environmental settings and biological systems. The involvement of Cu^+ in physiological and pathological processes promoted the development of methods in Cu^+ analysis. There are existing techniques such as electrochemistry,[14–17] chromatography,[18–21] pulse polarography,[22] voltammetry[23] and atomic absorption spectrophotometry (AAS),[24,25] but these generally require sophisticated procedures and expensive instrumentation. More importantly, these methods cannot provide the real-time visualization of labile Cu^+ *in situ*. Optical imaging techniques are non-invasive, highly sensitive, easily handled, and suitable for detecting analytes in biological system. In the past decade, a number of fluorescent probes have been designed for copper detection.[26–30] Among this collection, NIR fluorescent probes stand out due to their unique properties of high penetration through tissues,[31–34] low auto-fluorescence and less photodamage.[35,36] T. Govindarajua *et al.* reported on the development of a NIR fluorescent probe with a tripicolylamine (N_4) functionality for the detection of intracellular Cu^+.[37] B. R. Cho *et al.* and W. Wan *et al.* developed probes with the recognition group being bis(2-((2-(ethylthio)ethyl)-thio)ethyl) amine (BETA) containing electron rich S atoms.[38,39] Although these probes offer a promising strategy for intravital non-invasive quantitative imaging, they cannot completely discriminate Cu^+ from other interfering cations, such as Cu^{2+}, Co^{2+} or Hg^{2+}. Developing a highly selective tool to detect Cu^+, especially to visualize dynamic process of Cu^+ in living system, is a challenging endeavour.

Herein, we developed a new near-infrared (NIR) fluorescent turn-on probe (**NPCu**) for the detection of Cu^+, based on the mechanism of a Cu(I)-mediated benzylic ether bond cleavage. Our results demonstrate that the **NPCu** probe can effectively distinguish Cu^+ from most interfering cations *in vitro* and *in vivo*.

Experimental

Materials and instruments

All reagents were purchased from commercial suppliers and used without further purification (Adamas-beta® or Lab Network), unless otherwise noted. All chemicals and solvents were used without purification unless otherwise noted. Column chromatography was carried out on silica gel (200–300 mesh)

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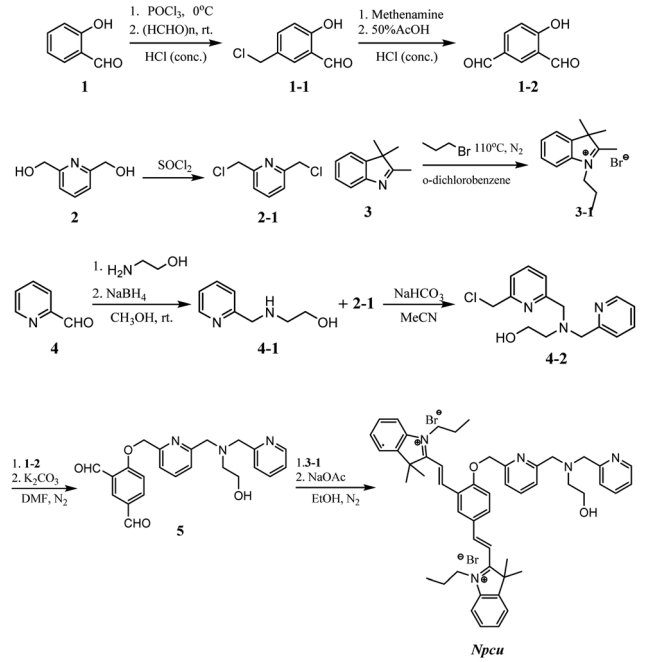

Scheme 1 Synthesis procedure of probe **NPCu**.

using an eluent of ethyl acetate and cyclohexane, dichloromethane/methanol, or ethyl acetate/methanol. TLC analyses were conducted on silica gel plates and visualized using UV (ZF-2 UV254). Mass spectrometry analyses were performed on an API 4000 (ESI-HRMS). NMR spectra were recorded at ^{1}H (400 MHz or 600 MHz) and ^{13}C (100 MHz) on a Bruker instrument. Chemical shifts (δ values) and coupling constants (J values) are given in ppm and hertz respectively. Melting points were determined on a Mel-Temp apparatus and were not corrected. UV/vis spectra were recorded on a UV-1800 (240V) spectrophotometer and fluorescence spectra were recorded on a FL3C-iHR320 spectrofluorometer (HORIBA Instrument Inc.) Absorption and emission were measured using a 1 cm path length quartz cells. Fluorescence imaging was carried out with Nikon Ti-E laser scanning microscope that excitation at 560 nm under a 40× objective.Water used for this study was purified with a Milli-Q filtration system (Scheme 1).

Synthesis of the NPCu probe

To a mixture of compound **5** (297.5 mg, 0.7 mmol) and **3-1** (517.7 mg, 1.8 mmol) in 4 mL of anhydrous ethanol was added sodium acetate (119.6 mg, 1.46 mmol) in 2 mL of ethanol under a nitrogen atmosphere. The reaction mixture was refluxed at 80 °C for 1 h under N_2 protection. When completed, the solvent was removed under reduced pressure and the residue was purified by silica gel column chromatography (MeOH/DCM from 0 to 10%) to yield a deep yellow solid, **NPCu** (150.0 mg, 21.9% yield). ^{1}H NMR (600 MHz, DMSO-d_6) δ: 9.41 (s, 1H), 8.62 (dd, J = 16.4, 8.5 Hz, 2H), 8.59–8.54 (m, 1H), 8.46 (d, J = 5.2 Hz, 1H), 8.15 (d, J = 16.4 Hz, 1H), 8.04–7.97 (m, 2H), 7.93 (dd, J = 9.2, 5.6 Hz, 3H), 7.90–7.86 (m, 1H), 7.78–7.72 (m, 1H), 7.70–7.62 (m, 5H), 7.62 (d, J = 8.8 Hz, 1H), 7.56 (d, J = 7.6 Hz, 2H), 7.23 (t, J = 5.6 Hz, 1H), 5.55 (s, 2H), 4.83 (t, J = 7.2 Hz, 2H), 4.75 (t, J = 7.2 Hz, 2H), 4.58 (s, 1H), 3.87 (s, 2H), 3.81 (s, 2H), 3.54 (d, J = 6.0 Hz, 2H), 2.61 (t, J = 6.4 Hz, 2H), 1.95–1.82 (m, 10H), 1.75 (s, 6H), 1.05 (t, J = 7.2 Hz, 3H), 0.97 (t, J = 7.2 Hz, 3H).^{13}C NMR (100 MHz, DMSO-d_6) δ: 182.51, 182.36, 162.29, 160.47, 159.96, 154.79, 153.02, 149.17, 147.45, 144.40, 144.22, 141.29, 141.27, 138.12, 137.84, 136.99, 134.02, 130.24, 129.93, 129.75, 129.62, 128.51, 124.09, 123.61, 123.58, 123.06, 122.68, 122.58, 121.18, 116.17, 115.87, 115.08, 114.99, 112.56, 72.39, 63.25, 60.64, 59.59, 56.62, 52.77, 52.75, 48.80, 48.34, 26.56, 26.37, 22.41, 22.36, 11.26, 11.23. HRMS (ESI) m/z: calcd for $[C_{51}H_{59}N_5O]^{2+}$ $[M]^{2+}$: 386.7335; found 386.6984.

Measurement of absorption and fluorescence

For UV/vis and fluorescence titrations stock solution of **NPCu** probe were prepared (c = 10 mM) in 25 mM PBS buffer (pH = 7.2). The solutions of cations were freshly prepared in PBS buffer solution. Working solutions of the probe and metal ions were prepared from the stock solutions. Excitation and emission spectra were carried out at 560 nm and 710 nm for **NPCu** with 4 nm slit widths. All buffers for pH titration were prepared in deionized water and adjusted to a suitable pH with HCl or NaOH aqueous solutions.

Selectivity of probes to Cu(I)

Each metal cation was diluted with PBS buffer solution to a 1 mM working solution for use, and mixed with 2 mL of **NPCu** solution (100 μM) in equal volume; in experiment 2, each metal cation was mixed with Cu(I), and each was configured. A mixed solution of 1 mM ion concentration was mixed with 2 mL of **NPCu** solution (50 μM) in equal volume. The samples from both experiments were incubated for 1 hour in a dark shaker under a 37 °C constant temperature shaker to determine the fluorescence intensity of each ion solution at an excitation wavelength of 560 nm and an emission wavelength of 710 nm. The molar absorption coefficients of probe **NPCu** without Cu^+ was calculated as $\varepsilon = 1.76 \times 10^4$ M^{-1} cm^{-1} while with Cu^+ was 0.76×10^4 M^{-1} cm^{-1}. **NPCu** after Cu^+ treatment exhibited a high quantum yield (Φ = 0.39). The quantum yields of **NPCu** was 0.024 in aqueous medium when excited at the λ_{max} (560 nm, $\varepsilon = 0.76 \times 10^4$ M^{-1} cm^{-1}).

Cell cytotoxicity and imaging

Human alveolar epithelial A549 cells were grown in RPMI-1640 medium, containing 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin, at 37 °C in a humidified incubator with an atmosphere of 5% CO_2/95% air. One day before the experiment, a 6-well culture plate was inoculated with logarithmic growth phase cells. The cells in the experimental group were incubated with Cu^+ (50 μM) for 6 h. In the control group, only medium was added. After that, cells were incubated with **NPCu** (10 μm) at 37 °C for 1 h. Finally, an inverted microscope was used to observe the cells with DIC and Cy-3 fluorescence filters.

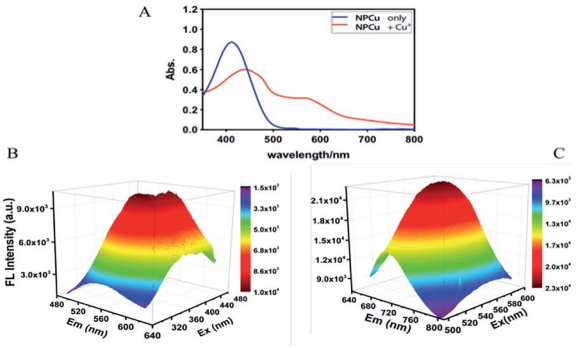

Fig. 1 UV-vis absorption spectra (A) and 3D fluorescence spectra (B and C) of **NPCu** (50 μM) incubated without or with Cu^+ (500 μM) in 25 mM PBS buffer (pH 7.2) containing 2 mM GSH.

Results and discussion

Sensitivity research

We first evaluated its spectral properties and determined its responsiveness towards Cu(I). As shown in Fig. 1, the absorbance, extraction and emission spectra of **NPCu** peaked at 415 nm, 430 nm, and 560 nm respectively in 25 mM PBS (pH 7.2) containing 2 mM glutathione (GSH). The addition of Cu^+ (10 eq.) induced significant signal changes in the optical properties of **NPCu** solutions. As shown in Fig. 1A, a marked redshift (~150 nm) was noted in absorption. Meanwhile, in the emission spectra, a turn-on response from 560 nm to 710 nm was noted upon introduction of Cu^+ to the probe. After the incubation with Cu^+, large Stokes shifts (>150 nm) both in the excitation and emission spectra were observed (Fig. 1B and C). This is considered a highly desirable feature as it increases the signal-to-noise ratio.

To investigate whether **NPCu** had the sensitivity to measure small fluctuations in Cu^+ level in aqueous solutions, the probe (50 μM) was exposed to a range of Cu^+ concentrations. The fluorescence intensity at 710 nm was plotted and exhibited good linear correlation ($R^2 = 0.995$) with Cu(I) levels from 0 to 1000 μM. A typical calibration curve is shown in Fig. 2 (inset). The detection limit for Cu^+ was 9.1×10^{-5} M (S/N = 3). It was shown that the fluorescence intensity of the probe progressively increased with increasing cocentration of Cu^+.

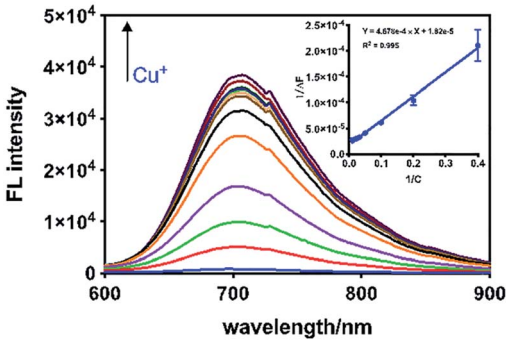

Fig. 2 Titration of **NPCu** (50 μM) in pH 7.2 buffer with different concentrations of Cu^+ at 0–1000 μM (λ_{ex} = 560 nm). Inset: linear correlation between emission intensity and Cu^+ concentration from 0 to 1 mM.

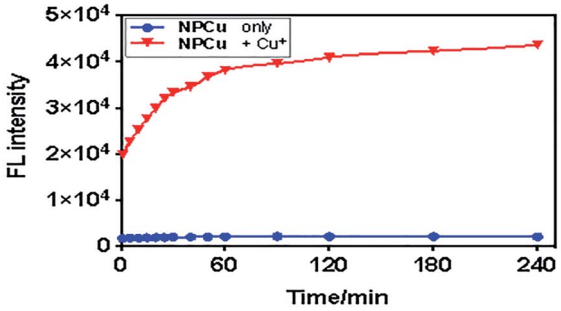

Fig. 3 Change of fluorescent intensity: **NPCu** (50 μM) reacted with Cu^+ (1000 μM) at various time points. $\lambda_{ex}/\lambda_{em}$ = 560/710 nm, 25 mM PBS (pH 7.2) containing 2 mM GSH.

Fluorescence intensity changes of NPCu with time

To 2 mL of **NPCu** solution (50 μM) was added the same volume of Cu^+ (10 eq.) in 25 mM PBS buffer (pH 7.2) containing 2 mM GSH and incubated at 37 °C for different lengths of time. The fluorescence intensity was measured at 1, 5, 10, 15, 20, 25, 30, 40, 50, 60, 90, 120, 180, and 240 min (Fig. 1). The correlation between **NPCu** responsiveness and different concentrations of Cu^+ was determined in 25 mM PBS (pH 7.2) containing 2 mM GSH. To the EP tubes (1.5 mL), each containing 500 μL of Cu^+ solution at various concentrations from 0 to 1 mM, was added the same volume of **NPCu** (250 μM) and incubated at 37 °C for 1 hour. It was found that the colour of the reaction mixture deepened with increasing Cu^+ concentration (Fig. S1†) (Fig. 3).

Selectivity research

NPCu can be conveniently used as a 'switch-on' probe for the detection of Cu^+ without interference from pH-related effects in physiological environments. As shown in Fig. 4, the specificity

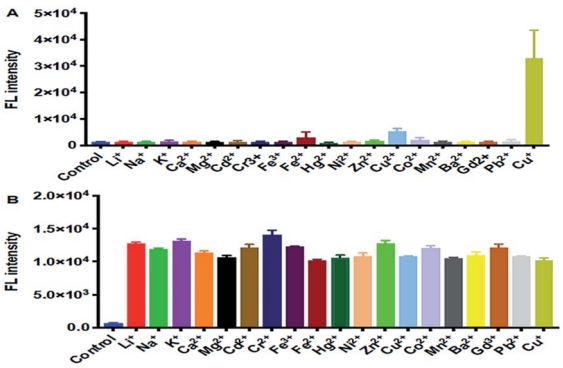

Fig. 4 Competition tests. Fluorescence responses of 50 μM **NPCu** to various metal ions (10 eq.). (A) Addition of competing metal ions to the solution of the receptor. (B) Addition of equal amounts of Cu^+ to the solution containing the other metal. Excitation was at 560 nm. Spectra were acquired in 25 mM PBS (pH 7.2) + 2 mM GSH.

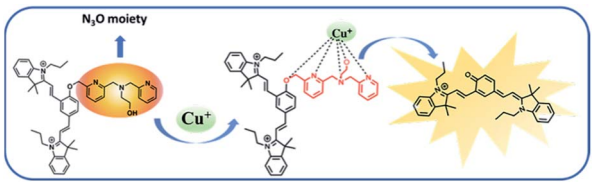

Fig. 5 The chemical structure of **NPCu** and its Cu^{+} sensing mechanism.

of **NPCu** ion detection was tested with various metal ions including Cu^{2+}, Co^{2+}, Hg^{2+}, and Fe^{3+}. **NPCu** exhibited a remarkable 30-fold enhancement in the NIR region ($\lambda_{ex} = 560$ nm) upon addition of Cu^{+} (10 eq.) after 2 h, whereas the responses toward other ions were negligible. The **NPCu** exhibited high selectivity for Cu^{+} over other biologically relevant interfering metals, including redox-active copper and cobalt transition metals.

Fluorimetric pH titration of NPCu

We further assessed the probe's capability of detecting Cu^{+} at different pH's.The pH was adjusted between 4.0 and 10.0 by adding 1 M HCl, and the fluorescence titration was performed in a quartz colorimetric dish with a path length of 1 cm (constant temperature set to 25 °C). The absorption spectra (Fig. S2†) were collected from 200–410 nm and the fluorescence intensity was measured at 710 nm with excitation at 560 nm. In the absence of Cu^{+}, there was almost no change fluorescence intensity in the pH range of 4–10, which suggested this probe has considerable stability. Under neutral to mildly alkaline conditions, **NPCu** can maintain high responsiveness to Cu^{+}.

Sensing mechanisms

According to the structural characteristics of **NPCu** and literature reports,[40,41] we speculate a new approach for the detection of Cu^{+} in live cells through the development of a NIR probe (**NPCu**), which is comprised of two discrete elements: a NIR core (Fig. 5, black), a Cu^{+} reactive moiety (Fig. 5, red). **NPCu** with a tetradentate ligand N_3O moiety, which is first employed as a highly selective trigger for the detection of Cu^{+}. Upon binding to Cu^{+}, the benzylic ether bond (C–O) of **NPCu** was cleaved, and cyanine-quinone dye was released from **NPCu**, leading to a robust NIR fluorescence enhancement. This mechanism was confirmed by mass spectrometry (Fig. S4†).

We also used orbital theory to better verify our hypothesis. LUMO and HOMO levels of **NPCu** in the absence or presence of Cu^{+} all have obvious delocalization effect on the entire molecular skeleton. As shown in Fig. 6, HOMO and LUMO energies of **NPCu** were calculated to be −3.55 eV and −1.91 eV while the energies of **NPCu + Cu^{+}** were −3.37 eV and −1.02 eV. The energy gaps (LUMO–HOMO) of **NPCu** and **NPCu + Cu^{+}** are 2.45 eV and 1.64 eV, respectively. It is worth noting that after reacting with Cu^{+}, the UV absorption peak of **NPCu** was observed to have red-shifted. Theoretical predictions were consistent with the spectroscopy experimental data, indicating that the proposed reaction mechanisms were reasonable.

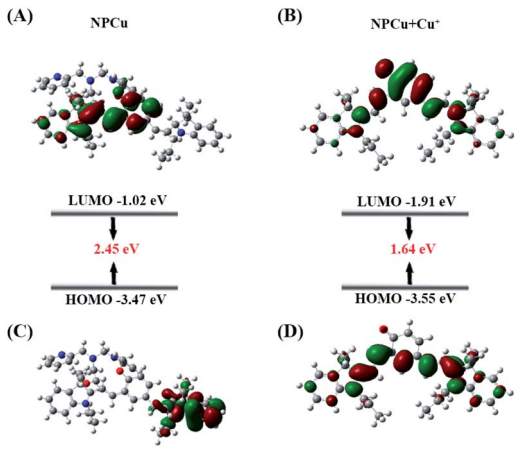

Fig. 6 The HOMO and LUMO of **NPCu** and **NPCu + Cu^{+}**. Red, blue, grey and white balls represent O, N, C, and H atoms, respectively.

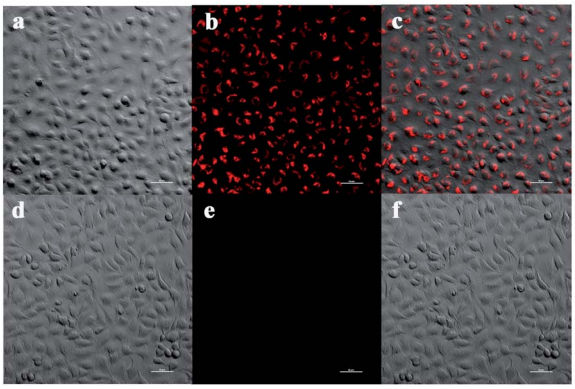

Fig. 7 Confocal microscopy images of A549 cells incubated with 10 μM **NPCu** for 1 h at 37 °C. Upper: incubated with 50 μM Cu^{+} prior to **NPCu** staining. Lower: incubated with vehicle prior to **NPCu** staining. (a and d) Bright field images; (b and e) fluorescence images; (c and f): overlay of fluorescence and bright field images. Scale bar = 50 μm.

Cytotoxicity and cell imaging

The cytotoxicity of **NPCu** was evaluated using the methylthiazolyldiphenyltetrazolium (MTT) viability assay. As indicated in Fig. S4,† A549 cells incubated with **NPCu** showed little decrease in cell viability. Viability was >70% at 30 μM **NPCu**, which indicates that **NPCu** is suitable for bioimaging applications at 10 μM. Next, we tested the utility of this probe for detecting Cu^{+} in cultured A549 cells by confocal microscopy. The results clearly demonstrate that **NPCu** is permeable to membrane, and can react with intracellular Cu^{+} to release near infrared fluorescent Cy-quinone dye in living cells (Fig. 7).

Conclusions

In summary, we have developed a new NIR fluorescent turn-on Cu^{+} probe based on the cyanine-quinone fluorophore and novel Cu^{+} receptor N_3O-ol skeleton. The probe **NPCu** exhibited high

sensitivity, good selectivity, and considerable stability at physiological pH. More importantly, **NPCu** showed excellent biocompatibility and high permeability for penetrating cell membrane and tracking intracellular Cu^+. **NPCu** shows promising properties for non-invasive imaging of Cu^+ in living cells and provides a useful tool to better understand the contribution of Cu^+ in living systems.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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