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A biomimetic fluorescent chemosensor for highly sensitive zinc(II) detection and its application for cell imaging[†]

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To fabricate a novel biomimetic fluorescent chemosensor, PSaAEMA-*co*-PMPC was synthesized *via* atom transfer radical polymerization, and this copolymer could be used for the detection of zinc(II) and cell imaging. A series tests with various metal ions verified the specific fluorescence response behavior. This novel biomimetic fluorescent chemosensor exhibits excellent selectivity for Zn²⁺ ions over a wide range of tested metal ions in an aqueous solution. Moreover, cytotoxicity and bio-imaging tests were conducted to study the potential bio-application of the chemosensor. Owing to the biomimetic portion (phosphorylcholine), this copolymer possesses outstanding biocompatibility and could clearly image cells. The results indicated that PSaAEMA-*co*-PMPC has great potential for application in zinc(III) detection and cell imaging.

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Introduction

Zinc, the second most abundant transition metal ion in the human body, is often present in a tightly bound form in proteins. It plays several significant roles in various fundamental biological processes. The disruption of Zn(II) homeostasis could lead to Parkinson's disease, Alzheimer's disease, immune dysfunction, and infantile diarrhea.¹⁻⁵ As a consequence, the development of probes to effectively quantify and visualize Zn(II) concentration in biological systems is key to understanding the biological processes of Zn(II) *in vivo*.

Recently, there have been many studies on fluorescence probing methods and fluorescent chemosensors, such as the AIE probe technique,⁶⁻¹³ and fluorescent inorganic nanoparticles.¹⁴⁻¹⁸ Currently, there is great interest in the development of new methods with high sensitivity and the exploration of the role of $Zn(\pi)$ in medicine and biology as well as the environment. Among the various types of methods, fluorescent chemosensors have attracted much attention due to their convenient use and high sensitivity, and could be employed to clarify the various biological functions of targeted metal cations in living cells.^{19–25} A few $Zn(\pi)$ fluorescent chemosensors have been fabricated for physiological applications and some are now commercially available. Salicylaldimine Schiff bases, one of the common structures, have aroused great interest for fluorescent chemosensors due to their facile synthesis and high binding affinity towards many metal ions. Salicylaldimine Schiff bases have poor fluorescence, but can display strong fluorescence once they coordinate with special metal ions such as zinc ions via the phenol oxygen atom and the imine nitrogen atom, which results in a rigid structure.26 Until now, a large amount of small molecule fluorescent metal ion probes have been developed based on salicylaldimine Schiff bases in past decades.26-37 However, for the majority of them, their water solubility is very limited since these chemosensors are hydrophobic, which induces aggregation between them. Thus, detection could not be performed in pure water, and is often performed in mixed aqueous solutions with a certain portion of organic solvent. Furthermore, the accuracy and sensitivity of small molecule detection are not satisfactory. To this end, several elegant strategies have been developed to overcome the flaws of small molecule probes, for example conjugating or loading them into inorganic nanoparticles, micelles, polymers, and so forth.³⁸⁻⁴³ Of all of the above, polymer-based fluorescent chemosensors have several advantages over the others, such as water solubility, exceptional processability, and signal amplification. Hence, efforts have been dedicated to the fabrication of polymeric chemosensors, which contain salicylaldimine Schiff bases. Theato et al. prepared thermo- and light-responsive copolymers by grafting salicylaldehyde to amine pendantfunctionalized polymers.44 Cai et al. synthesized block copolymers and constructed a variety of coordinated polymeric aggregates with different metal ions.45-47 Zhang et al. constructed well-defined polymeric salicylaldimine Schiff bases via

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RAFT alternating copolymerization and they further confirmed the superb accuracy and sensitivity of copolymers compared with small molecule detection.⁴⁰ However, almost all of the reported studies on polymer-based fluorescent chemosensors primarily focus on easy fabrication processes, high solubility, accuracy and sensitivity, but cytocompatibility has not been taken seriously.

To be used safely, it is absolutely necessary for probes to have excellent biocompatibility properties. Phosphorylcholine (PC) groups, a major headgroup in phospholipids, which are the main constituents of plasma membranes, have been often conjugated into different polymers to mimic membrane structure and have made quite good progress.48 2-Methacryloyloxyethyl phosphorylcholine (MPC) is a methacrylate monomer with a phosphorylcholine group, which has been extensively synthesized to mimic cell membranes and is considered as a biocompatible material. On account of their simple molecular design and synthesis process, excellent biocompatibility, water solubility, and transparency in visible light, MPC polymers are considered promising in materials science and biomedical applications. In our previous studies,49,50 these polymers with PC shells exhibited great biostability and biocompatibility, which supported the polymers' normal functions in complex environments. Additionally, MPC polymers were also developed by Ji's group as a biocompatible imaging agent.51-54 It turned out that these biomimetic polymers were suitable for biological materials and imaging agents. Thus, the combination of salicylaldimine Schiff bases with MPC polymers would utilize well the advantages of both of their merits.

Herein, a biomimetic fluorescent chemosensor was designed and used for the delicate detection of zinc(π). To synthesize this copolymer, a three-step process was required. Firstly, ATRP polymerization was conducted with *N*-Boc-aminoethyl methacrylate and MPC using ethyl α -bromoisobutyrate as the initiator to produce PNBAEMA-*co*-PMPC. Then, PNBAEMA-*co*-PMPC was deprotected to obtain PAEMA-*co*-PMPC. Lastly, salicylaldehyde and PAEMA-*co*-PMPC were connected by Schiff base bonds (PSaAEMA-*co*-PMPC). The polymer chemosensor could efficiently bind Zn²⁺ ions, which specifically led to a change in fluorescence intensity. Furthermore, the cytotoxicity and cell uptake behaviors of PSaAEMA-*co*-PMPC were investigated to evaluate the biological performance of this material (Scheme 1).

backbone PC chains hydrophobic chains $e^{-\alpha}$

Experimental

Materials

Ethanolamine, *N*-Boc-aminoethyl, methacryloyl chloride and ethyl α -bromoisobutyrate were purchased from Aladdin. MPC was synthesized according to our previous studies. Chloride salts, MgSO₄, and nitrate salts were used for all of the sensing experiments. Water was prepared by our own filtration system. The human umbilical vein endothelial cell line (HUVEC) and HeLa cells were obtained from the Cell Bank of the Chinese Science Academy, Shanghai, China. All of the reagents obtained from commercial sources were used as received without further purification.

Instrumentations

Absorption spectra were recorded on an Analytic-jena Specord S600 (Germany) spectrophotometer in absorbance mode using quartz cuvettes of 1 cm path length. ¹H NMR measurements were carried out on a Bruker AV400 spectrometer. A Perkin Elmer LS-55 spectrofluorometer was used to measure the fluorescence spectra. Fluorescent pictures were taken on a ZEISS Axio Scope Fluorescence Microscope.

Synthesis of N-Boc-ethanolamine

Ethanolamine (1.10 g, 18.3 mmol) was added into 200 mL methanol and a solution of di-*tert*-butyl dicarbonate (0.01 mol) was added dropwise over a period of about 2 h at 0 °C. After stirring for 24 h, the solution was concentrated under reduced pressure. Then, the crude produce was extracted with methylene chloride three times. The extract was dried with anhydrous Na₂SO₄ and concentrated under vacuum. Yield: 80%. ¹H NMR (500 MHz, CDCl₃): d 1.42 (9H, $-C(CH_3)_3$), 3.15 (2H, HOCH₂- CH_2 CO–), 3.67 (2H, HOCH₂CH₂CO–).

Synthesis of N-Boc-aminoethyl methacrylate

N-Boc-aminoethyl methacrylate was synthesized by a condensation reaction between acyl chloride and hydroxyl groups. *N*-Boc-aminoethyl (20.2 g, 0.1 mol), triethylamine (10.1 g 0.1 mol) and 300 mL of methylene chloride were added into a flask at room temperature, and a methylene chloride solution (15 mL) of methacryloyl chloride (8.32 g, 0.08 mol) was added dropwise. After 12 h stirring, deionized water was added and extracted with DCM three times. The extract was dried with anhydrous Na₂SO₄ and concentrated under vacuum. Yield: 79%. ¹H NMR (500 MHz, CDCl₃): d 1.42 (9H, $-C(CH_3)_3$), 1,91 (3H, $CH_2 = CH(CH_3)-$) 3.15 (2H, HOCH₂CH₂CO-), 4.36 (2H, $-OCH_2CH_2CO-$), 5.56, 6.12 (2H, $CH_2 = CH(CH_3)-$).

Synthesis of P(*N*-Boc-aminoethyl methacrylate)-*co*-P(2methacryloyloxyethyl phosphorylcholine) (PNBAEMA-*co*-PMPC)

PNBAEMA-*co*-PMPC was synthesized by the ATRP polymerization of *N*-Boc-aminoethyl methacrylate and MPC using ethyl α bromoisobutyrate as the initiator. In a typical reaction, *N*-Bocaminoethyl methacrylate (4 g, 25 mmol), MPC (4 g, 25 mmol),

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ethyl α -bromoisobutyrate (124 mg, 0.625 mmol), CuBr (89 mg, 0.625 mmol), PMDETA (216 mg, 1.25 mmol) and 10 mL DMSO/ MeOH (1 : 1) were added into a 25 mL flask under a nitrogen atmosphere. The reaction mixture was degassed by three freezepump-thaw cycles and stirred at 30 °C for 12 h. Then, the reaction mixture was exposed to air and the solution was diluted with MeOH and passed through silica gel column chromatography to remove the catalyst. The filtrate was concentrated and precipitated into ethyl ether and the precipitate was dried under vacuum. Yield: 65%.

Synthesis of P(aminoethyl methacrylate)-*co*-P(2methacryloyloxyethyl phosphorylcholine)trifluoroacetate salt (PAEMA-*co*-PMPC)

0.15 g PNBAEMA-*co*-PMPC was dissolved in 3 mL methanol and 3 mL DMSO in a 25 mL flask. Then, 6 mL TFA was added and stirred at room temperature for 5 h. The reaction solution was concentrated under reduced pressure. Finally, the crude product was dialyzed against distilled water (MWCO 3500) for 2 days and lyophilized. Yield: 70%.

Synthesis of P(2-salicylaldehyde-aminoethyl ethanolamine methacrylate)-*co*-P(2-methacryloyloxyethyl phosphorylcholine) (PSaAEMA-*co*-PMPC)

PAEMA-*co*-PMPC (508 mg, 2 mmol) and anhydrous TEA (0.55 mL, 4 mmol) were dissolved in 10 mL anhydrous methanol, and then salicylaldehyde (0.23 mL, 2.2 mmol) was added. The reaction mixture was stirred at room temperature for 12 h, and the solution was then concentrated under reduced pressure. The crude product was precipitated into ethyl ether. After drying for two days in a vacuum, a yellow powder was obtained. Yield: 70%.

Absorption and fluorescence measurements

The copolymers (PSaAEMA-*co*-PMPC) were prepared and dissolved in PBS (20 mM; pH = 7.4) before use. Cationic solutions were prepared by dissolving metal salts in deionized water with a concentration of 0.1 M and were diluted as required before use. A solution of the test sample (4.0 mL, 20 μ M for UV-Vis and 40 μ M for fluorescence) was prepared, and UV-Vis and fluorescence spectra were recorded at room temperature. The total volume of each ionic solution introduced to the test solution was no more than 25 μ L, and the changes in fluorescence intensity were recorded after the solution was mixed for 10 min ($\lambda_{ex} = 368$ nm).

Cell cytotoxicity assay

The cytotoxicity of PSaAEMA-*co*-PMPC was evaluated by an *in vitro* MTT assay. HUVEC cells and HeLa cells were seeded in 96-well plates with 6×10^3 cells per well. After incubation for 24 h, the culture medium was replaced by fresh medium with PSaAEMA-*co*-PMPC at different concentrations (10 to 1000 µg mL⁻¹). After the cells were incubated for 48 h, the cells were subjected to an MTT assay.

Cell imaging

HeLa cells were used for cell imaging in this study. HeLa cells were seeded in 24-well plates with 2 \times 10⁴ cells per well. After incubation for 24 h, the culture medium was replaced by fresh medium with PSaAEMA-co-PMPC (20 μ M). After incubation for a further 1 h, the cells were washed with cold PBS three times. Then the cells were further treated with ZnCl₂ (25 mM) plus sodium pyrithione (5 mM). After this, the cells were washed with cold PBS three times and fixed with 4% formaldehyde for 15 min. Finally, the cells were observed using a fluorescence microscope.

Results and discussion

Synthesis of Zn²⁺ probe

The synthetic route for the preparation of the PSaAEMA-co-PMPC polymer is illustrated in Scheme 2. This copolymer was prepared in three steps. First, N-Boc-aminoethyl methacrylate was synthesized by N(Boc) protected ethanolamine and methacryloylchloride, and the product was sequentially polymerized with MPC via ATRP using EBIB as the initiator. Then, deprotection of the N-Boc of the copolymer using TFA exposed a free amine group which could react with salicylaldehyde. Finally, salicylaldehyde was grafted to the copolymer through Schiff base bonds. Fig. S1 and S2[†] show the ¹H NMR spectra of *N*(Boc) protected ethanolamine and N-Boc-aminoethyl methacrylate, respectively. The results indicated that the monomer was prepared successfully. Moreover, the ¹³C NMR spectrum (Fig. S3[†]) further confirmed the correctness of the structure. Fig. 1 shows the ¹H NMR spectrum of the polymer probe. The signal at $\delta = 8.5$ ppm (*j*) was assigned to the characteristic peaks of the Schiff bases, and the signal at $\delta = 3.6$ ppm (b) was assigned to the methylene protons of the choline units. The ratio of *j* and *b* is roughly equal to 1 : 2, which suggested that the two units were equivalent in PSaAEMA-co-PMPC. As the Mn content of PMPC-co-PMEMA could not be determined by GPC, the Mn content was estimated from the ¹H NMR spectrum by comparing the proportion of the signals at $\delta = 3.40$ (EBiB) and



Scheme 2 Detailed synthetic route of PSaAEMA-co-PMPC.



¹H NMR spectrum of PSaAEMA-co-PMPC in CD3OD/DMSO-Fig. 1

 $\delta = 3.6$ ppm (b); the Mn content of PMPC-b-PMEMA was about 6494 g mol^{-1} .

It is known that salicylaldimine Schiff bases can coordinate with various metal ions, especially zinc complexes, which often leads to strong fluorescence emission. Therefore, we designed a monomer that possessed an amine group to form a Schiff base with salicylaldehyde and speculated that this part of the copolymer would show no fluorescence prior to the addition of zinc ions, but once the zinc ions were added, the copolymer would emit intense fluorescence. To verify our presumption, we examined the UV-Vis spectra of the copolymer with a gradual addition of Zn²⁺. Fig. 2 shows the absorption changes of the



Fig. 2 Absorption spectra of 10 mol L^{-1} copolymer with the addition of 0-15 mol L^{-1} Zn²⁺. Inset: The absorbance ratio (A₃₈₃/A₃₄₆) as a function of Zn²⁺ concentration.

copolymer upon a gradual addition of 0-1.0 equiv. Zn²⁺. It can be seen that with the addition of Zn²⁺ into the copolymer solution, the absorption bands at 254 and 316 nm decreased gradually, and new absorption bands at 363 nm emerged simultaneously. Three isosbestic points could be clearly seen (Fig. 2). The absorption bands at 316 nm gradually decrease which was attributed to the n- π^* transition of C=N. The above results indicated a conversion of the Schiff bases to Schiff base-Zn²⁺ complexes. No further changes were observed in the absorbance spectra with up to 1 equiv. Zn²⁺ addition, revealing that the binding ratio between Zn²⁺ and the copolymer should be 1 : 1.

To test the sensing ability of PSaAEMA-co-PMPC for Zn²⁺ in aqueous solutions, fluorescence titration of this copolymer with a gradual addition of Zn²⁺ was investigated and the results are shown in Fig. 3. The addition of Zn²⁺ leads to an increase in 450 nm emission intensity, which is ascribed to the formation of zinc complexes, which inhibit C=N isomerization and cause a chelation-enhanced fluorescence effect. In the fluorescence spectra under the same test conditions, more than an 800-fold fluorescence enhancement at around 450 nm occurred upon the addition of Zn^{2+} (0.1 equiv., Fig. 3). This increase is similar to that with the addition of 1 equiv. of Zn^{2+} . As shown in the inset, a linear relationship is observed between the intensity and the amount of Zn^{2+} in the range of 0–0.7 equiv., which corresponds to 0-14 mM Zn²⁺ (linearly dependent coefficient: $R^2 = 0.997$). The linear fitting of the titration curve further confirmed that the binding of the copolymer and Zn²⁺ was most probably a 1:1 metal-to-ligand ratio. This indicates that the copolymer has potential for application in the quantitative determination of Zn²⁺ in aqueous solutions.

Then, the PSaAEMA-co-PMPC solution was treated with various metal ions to verify the fluorescence response behavior. The measurements were conducted with excitation at 368 nm to examine the ion selective ability of the copolymer in aqueous solutions. As shown in Fig. 4A, the addition of Zn²⁺ causes an



Fig. 3 Fluorescence spectra ($\lambda_{ex} = 368$ nm) of (PSaAEMA-co-PMPC) (20 μ M) with different concentrations of Zn²⁺; inset: the fluorescence intensity at 450 nm as a function of Zn²⁺ concentration. Excitation was at 368 nm

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Fig. 4 (A) Fluorescence spectra of the fluorescence intensity of (PSaAEMA-*co*-PMPC) (20 μ M) upon the addition of 1.0 equiv. of various metal ions. (B) Corresponding fluorescence intensity of (PSaAEMA-*co*-PMPC) with different metal ions in the absence and presence of Zn²⁺. ($\lambda_{ex} = 368$ nm).

obvious 5000-fold emission enhancement at around 450 nm. However, the emission intensities at around 450 nm were barely enhanced with the addition of Na⁺, K⁺, Mg²⁺, Ca²⁺, Ba²⁺, Pb²⁺, Cu^{2+} , Ag^{2+} and Fe^{3+} . Therefore, the copolymer exhibits an excellent selectivity toward Zn²⁺ binding. Fig. 4B shows the fluorescence response of PSaAEMA-co-PMPC with different metal ions in the absence and presence of Zn²⁺. Most of the representative ions like Na^+ , K^+ , Ba^{2+} , Pb^{2+} and Ag^{2+} cause a slight interference in Zn²⁺ detection. The addition of Ca²⁺ causes a small increase in the emission intensity, whereas Mg²⁺ and Fe³⁺ partially quench the fluorescence, which is due to their competitive coordination to Zn2+, as verified by the previous reports. Besides, the presence of Cu2+ induces absolute quenching, since Cu²⁺ has a higher binding constant with the salicylaldimine Schiff bases as well as a heavy atom guenching effect.

Living cell imaging

Before PSaAEMA-*co*-PMPC can be used as a bio-imaging probe, it is necessary to evaluate its cytotoxicity. For this purpose, cultured HeLa cells and HUVEC cells were incubated with



Fig. 5 Cell cytotoxicity of HeLa and HUVEC cells incubated with various concentrations of PSaAEMA-*co*-PMPC.



Fig. 6 Fluorescence images of live HeLa cells. From left to right are bright field images, fluorescence images and overlay images. (A) HeLa cells were incubated with PSaAEMA-*co*-PMPC for 30 min and washed with PBS; (B) HeLa cells were incubated with PSaAEMA-*co*-PMPC, and then with Zn²⁺ (10 μ mol L⁻¹) and pyrithione (10 μ mol L⁻¹) for 20 min and washed with PBS.

different concentrations of copolymer solution (from 10 to $1000 \ \mu g \ mL^{-1}$) for 48 h. Subsequently, an MTT assay was carried out. Fig. 5 demonstrated that, even when treated with a certain high concentration of this copolymer (1000 $\ \mu g \ mL^{-1}$), 80% of cells were still alive. This result indicates that PSaAEMA-*co*-PMPC has a low cytotoxicity to cells and is suitable for intracellular imaging.

Then, the application of PSaAEMA-*co*-PMPC for the fluorescence imaging of Zn^{2+} in live HeLa cells was investigated. As observed by laser scanning confocal microscopy, HeLa cells incubated with 20 µg mL⁻¹ PSaAEMA-*co*-PMPC for 20 min at 37 °C showed no intracellular fluorescence. Then, the cells were cultured with 10 µmol L⁻¹ Zn²⁺ and 10 µmol L⁻¹ pyrithione, which is a zinc selective ionophore, for 20 min at 37 °C and then washed with PBS three times. A significant fluorescence increase from the intracellular area was shown (Fig. 6), and we could clearly observe the entire cellular outline. These results suggested that PSaAEMA-*co*-PMPC could be used as an effective chemosensor for Zn²⁺ imaging in living cells.

Conclusion

In summary, we have developed a novel salicylaldehyde based chemosensor PSaAEMA-*co*-PMPC, which displays strong fluorescence with high selectivity and sensitivity for Zn^{2+} compared to other competing metal ions. The sensor gives a fluorescence response in an aqueous solution upon binding with zinc(n) in a 1 : 1 ratio. Moreover, due to the structure of PC, the cytocompatibility of the copolymer was excellent, which is perfect for biological applications. Hence, the biological application of PSaAEMA-*co*-PMPC was evaluated for the detection of Zn^{2+} in living cells. The results showed that it had low cytotoxicity and excellent cell fluorescence illumination and suggested that PSaAEMA-*co*-PMPC was cell-permeable and capable of the intracellular detection of Zn^{2+} .

Conflicts of interest

There are no conflicts to declare.

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