Investigation of a halloysite-based fluorescence probe with a highly selective and sensitive “turn-on” response upon hydrogen peroxide†

Jingwei Dong, Zhihang Zhao, Rui Liu, Hailei Zhang, Yonggang Wu and Xinwu Ba*

Inorganic halloysite nanotubes (HNTs) were modified with an organic fluorescein derivative (PA) to prepare HNTs-based hybrid fluorescence probe (HNTs-PA). Thermogravimetric analysis (TGA), Fourier transform infrared spectroscopy (FTIR), solid-state 13C NMR spectroscopy, X-ray photoelectron spectroscopy (XPS), energy dispersive spectrum (EDS) and transmission electron microscopy (TEM) confirmed that PA was successfully grafted onto the lumen of HNTs to obtain HNTs-PA with a grafting degree of 6.0%. The established B–C bond endows a selective fluorescence response toward hydrogen peroxide (H2O2) to HNTs-PA. The new hybrid probe exhibited a highly specific “turn-on” fluorescence response to H2O2 over other reactive oxygen species (ROS) and common ions owing to their chemoselective boronate-to-phenol switch. The “turn-on” response could even be tracked when the additional amount of H2O2 was limited to 1/10 mol. Moreover, human lung cancer cells (A549 cells) were successfully stained and the staining intensity enhanced as time prolongs, which can be due to the overexpressed H2O2 of cancer cells. Thus, the as-prepared organic–inorganic hybrid fluorescence probe can have a broad range of applications for identification and diagnosis.

Introduction

Reactive oxygen species (ROS) often play important roles in cell signaling and are also regarded as biomarkers for some pathological changes in human beings.1,2 H2O2, one of the representative ROS, is often involved in inflammation, hemorrhage, tumor proliferation and so on.3–6 The accumulation of H2O2 can also destruct various cellular oxidants. Hence, it is important to trace the H2O2 at low levels in order to prevent and diagnose the abovementioned diseases, which presents the need for developing H2O2-sensitive fluorescent probes (Schemes 1 and 2).

Recent studies showed that the H2O2-triggered deprotection of arylboronates to phenols provided a reaction-based approach to the specific detection of H2O2, which results in the exploration of arylboronate-based fluorescent probes for H2O2.3,7 Fluorescein and its derivatives have attracted much attention primarily due to their high quantum yields, desirable photostability, and long emission wavelengths. Chang et al. reported a series of arylboronate-containing fluorescein derivatives. The H2O2-triggered deprotection of arylboronates to phenols can cause a dramatic increase in fluorescent intensity, where the “turn on” effect makes the synthesized fluorescein derivatives potential H2O2-sensitive fluorescent probes.3,7 However, the...
undesirable solubility in aqueous system restricts their applicability to a broader range of applications.

Halloysite nanotubes (HNTs) are a type of natural tubular nanoparticles with a broad range of length to diameter ratios.9,10 The exterior and interior surfaces of HNTs are composed of siloxane (Si–O–Si) groups and gibbsite-like arrays of aluminol (Al–OH) groups, respectively, which results in a negatively charged outer surface and a positively charged inner lumen.11,12 The peculiar composition allows selective functionalization at the inner or outer side, leading to the formation of a composite with a hierarchical nanostructure.13,14 Their attractive properties, including non-toxicity, harmlessness, high mechanical strength, and good biological adaptability,15–20 make HNTs find a variety of applications in fields such as electronics, catalysis, separation, and functional materials.21–25 Environment friendliness and biocompatible nature enable halloysite nanotubes (HNTs) to be a promising nanomaterial for developing organic/inorganic composites in the biomedicine field.16,26–32 The surface modification of HNTs would create new types of smart functional materials.33–40 The selective modification methods have been extensively summarized in a latest review by our group.41

In our previous studies, pyrenylboronic acid (PBA)-modified HNTs were prepared and the established Al–O–B linkage provided a “turn off” fluorescence response for H2O2.42 In addition, a H2O2-responsive drug delivery system was also constructed using the established Al–O–B linkage.43 In this study, the fluorescein derivative bearing arylboronic acid was conjugated to the inner surface of HNTs using the previously reported method. Moreover, the established Al–O–B linkage can endow a “turn on” fluorescence response for H2O2 toward the fluorescein-modified HNTs. The good water-dispensibility and biocompatibility properties of fluorescein-modified HNTs make them more suitable for imaging H2O2 signaling in living cells.

Experimental

Materials

HNTs were obtained from Guang Zhou Shinshi Metallurgy and Chemical Company Ltd (Guangzhou, China). Fluorescein was purchased from J&K Chemical Technology. Dimethyl sulfoxide (DMSO) was dried and distilled from CaH2 under vacuum before use. Distilled water was used throughout the study. The reactions were monitored by thin layer chromatography (TLC) with silica gel 60 F254 (Merck 0.2 mm). Column chromatography was carried out on silica gel (200–300 mesh). Peroxyfluoroboronic acid (PF1) was synthesized according to the literature.9

Purification of HNTs

HNTs were purified according to the method described in our previous study.44

Synthesis of boronic acid-contained fluorescein derivative (PA)

PF1 (940 mg, 1.7 mmol) and diethanolamine (400 mg, 3.8 mmol) were dissolved into an ether solution and then stirred at room temperature. After 50 min, the precipitate was filtered, washed with ether, and dried in vacuum to afford a brown powder. Then, the obtained solid was treated with 0.1 M HCl. After ultrasonication for about 30 min at room temperature, a yellow precipitate was obtained, which was centrifuged and washed with 0.1 M HCl to afford the product PA (435 mg, 66%) (Scheme S1†). 1H NMR ((CD3)2SO, 600 MHz, ppm): δ 8.34 (s, 1H), 8.05 (t, J = 8.0 Hz, 1H), 7.78 (s, 2H), 7.76 (d, J = 7.4 Hz, 1H), 7.75 (d, J = 7.4 Hz, 1H), 7.50 (t, J = 8.0 Hz, 2H), 7.28 (t, J = 7.6 Hz, 1H), 6.80 (d, J = 7.8 Hz, 2H). 13C NMR ((CD3)2SO, 150 MHz, ppm): δ 168.7, 152.9, 149.9, 135.8, 129.3, 126.6, 125.1, 124.9, 123.9, 122.4, 119.9, 81.5. FTIR (KBr): 3622, 3525, 3025, 2921, 1410 cm−1. Elemental analysis: calcd (%) for C20H14B2O7: C 61.92, H 3.64; found: C 61.81, H 3.69.

Preparation of HNTs-PA

A mixture of purified HNTs (120 mg) and PA (40 mg) in anhydrous DMSO was carefully degassed. The system was heated at 80 °C for 12 h under stirring in nitrogen. The mixture was cooled to room temperature and washed by diethyl ether. The residue was collected by centrifugation. After vacuum-drying, the HNTs-PA was obtained as a faint grey solid.

Generation of reactive oxygen species (ROS) and common ions

Nitric oxide (NO•) was generated from sodium nitroferricyanide (Na2Fe(CN)6)43 H2O2, and ClO− were delivered from 30% H2O2, and 5% NaClO aqueous solutions, respectively. Metal ions (Cu2+, Fe3+, Fe2+, Mg2+, Na+, Zn2+, K+, Ca2+) and anions (SO42−, H2PO4−, SO32−, NO3−, CO32−, Cl−) were derived from CuSO4, FeCl3, FeCl2, MgSO4, NaCl, ZnCl2, KCl, CaCl2, Na2SO4, NaHPO4, Na2SO3, NaNO3, Na2CO3, NaCl, respectively.

Cell cultures

Human lung cancer cells (A549 cells) and Hela cells were cultured in Dulbecco’s modified Eagles medium (DMEM, Beijing North T2-Biotech Develop, Co. Ltd.) with 10% fetal bovine serum and 100 U mL−1 of 1% antibiotics (penicillin/streptomycin). A549 cells were placed on a confocal dish and the number of cells on each confocal dish was about 5.0 × 103. Then the cells were maintained at 37 °C for 12 h in a 100% humidified atmosphere containing 5% CO2 for further use.

MTT assay

A549 cells or Hela cells were seeded in a 96-well plate at a density of 4 × 103 per well. The cells were provided by the Chinese Academy of Medical Sciences, Peking Union Medical College. Accurately weighed HNTs-PA probes were dissolved in phosphate buffered saline (PBS) to achieve mother liquors. Following this, the cells were further incubated in media containing various concentrations diluted from the mother liquors. The plates were maintained at 37 °C in a 5% CO2/95% air incubator for 24 h. After 24 h, the medium was removed and washed with PBS. MTT solution (0.5 mg mL−1, 100 μL) was then added to each well. After 4 h, the remaining MTT solution was removed and 100 μL of DMSO was added to each well to dissolve...
the formazan crystals. Absorbance was set at 520 nm by a Triturus microplate reader. Wells containing no samples were used as blank control. The cell viability (%) was calculated as the ratio of the number of surviving cells in the test samples to the control groups.

**Cell imaging**

Cells were grown in Minimum Essential Medium Eagle (MEM medium, Sigma) in a watch glass bottom poly-o-lysine coated Petri-dish for at least 24 h to enable adherence to the bottom. Accurately weighed amounts of HNTs-PA suspension or Mito-tracker Red were added to the medium at 37 °C in 5% CO2. The cells were maintained in an incubator for another 30 min to achieve a sample loaded MEM medium. Then, the cells were washed three times with DPBS buffer to remove the samples that were not taken up into the cells. Finally, DMEM was added. A Hitachi F-4500 spectrofluorometer (Tokyo, Japan) equipped with a xenon lamp was used to obtain the fluorescence spectra. All experimental parameters (the laser intensity, exposure time, objective lens) were stationary when the different fluorescence images were captured. The excitation wavelength was fixed at 488 nm and fluorescent signals were collected from 500 to 600 nm.

**Characterizations**

Microwave reactions were performed using a Discover (PE-CEM) reactor. UV-visible absorption spectra were obtained on a Shimadzu UV-visible spectrometer model UV-2550. Fluorescence spectra were recorded on a Shimadzu RF-5301PC. TGA was performed on Perkin-Elmer Pyris 6 under a nitrogen atmosphere. UV-visible absorption spectra were recorded on a Shimadzu UV-visible spectrometer model UV-2550. Fluorescence spectra were obtained on a Shimadzu RF-5301PC. TGA was performed on Perkin-Elmer Pyris 6 under a nitrogen atmosphere. UV-visible absorption spectra were recorded on a Shimadzu UV-visible spectrometer model UV-2550. Fluorescence spectra were obtained on a Shimadzu RF-5301PC. TGA was performed on Perkin-Elmer Pyris 6 under a nitrogen atmosphere. UV-visible absorption spectra were recorded on a Shimadzu UV-visible spectrometer model UV-2550. Fluorescence spectra were obtained on a Shimadzu RF-5301PC. TGA was performed on Perkin-Elmer Pyris 6 under a nitrogen atmosphere. UV-visible absorption spectra were recorded on a Shimadzu UV-visible spectrometer model UV-2550. Fluorescence spectra were obtained on a Shimadzu RF-5301PC. TGA was performed on Perkin-Elmer Pyris 6 under a nitrogen atmosphere.

**Results and discussion**

Previous studies have demonstrated that arylboronic acid can react with the hydroxyls spread on the surface of HNTs. In this study, a fluorescein derivative bearing an arylboronic acid group was synthesized and then bound to the interior surface of HNTs according to the above-mentioned method. The obtained nanocomposite (HNTs-PA) showed desirable dispersibility in aqueous solution. The differences between HNTs-PA and HNTs were revealed by FTIR, XPS and solid-state 13C NMR.

Fig. 1 shows the solid-state 13C NMR spectra of HNTs and HNTs-PA. HNTs show no visible resonance in the region from 0 to 200 ppm, implying the absence of carbon units in HNTs. Moreover, remarkable resonances ranging from 180 to 40 ppm can be clearly observed in the solid-state 13C NMR spectrum of HNTs-PA. The peaks at 170.7 and 83.8 are attributed to the carbon in C=O and the quaternary carbon in the PA units. The resources from 157.1 to 116.9 can be assigned to the aromatic carbons in the PA units. The remarkable difference in the 13C NMR spectra between HNTs and HNTs-PA indicated the presence of fluorescein groups in HNTs-PA.

The FTIR spectra of HNTs and HNTs-PA are shown in Fig. 2. In the spectrum of HNTs, two distinct peaks located at 3696 and 3622 cm⁻¹ are attributed to the inner hydroxyl groups lying between the tetrahedral and octahedral sheets and the surface hydroxyl groups in the lumen of the nanotubes, respectively. The band at 1034 cm⁻¹ is assigned to the stretching vibration of Si–O–Si. The FTIR spectrum matches well with the composition of HNTs.

As for HNTs-PA, the distinct band corresponding to the surface hydroxyl groups in the lumen at 3696 cm⁻¹ displays a significant decrease, suggesting the consumption of surface hydroxyl groups in lumen during the modification. Simultaneously, some new peaks at 3525, 3025, 2921, 1765 and 1410 cm⁻¹ for each sample on a Varian-640 spectrophotometer. The spectrum for each sample was obtained by averaging 32 scans over the selected wave number range. 13C solid-state NMR spectra were recorded on a Bruker Advance III spectrometer. X-ray photoelectron spectroscopy (XPS) was carried out on a Thermo Scientific ESCALab 250Xi using 200 W monochromated Al Kα radiation. The 500 μm X-ray spot was used for XPS analysis. The base pressure in the analysis chamber was about 3 × 10⁻¹⁰ mbar. Typically, the hydrocarbon C 1s line at 284.8 eV from adventitious carbon was used for energy referencing. The energy dispersive spectrum (EDS) was determined by a Hitachi S-4800 scanning electron microscope.

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The quantity of the fluorescein derivative grafted to the nanotube was calculated to be 6.0 wt% by thermogravimetric analysis (TGA, Fig. 3). The XPS spectra are presented in the ESI.† Original HNTs and HNTs-PA all show the presence of aluminium (Al 2s and Al 2p) and silicon (Si 2p), in accordance
with the composition of alumino-silicate clay. The newly emerged weak peak of B 1s at 191.5 eV supports the successful modifications, where the atomic percentage of boron on the surface of HNTs-PA is calculated to be 0.6%, which is much less than the grafting degree calculated by TGA results. Generally, the wall thickness of HNTs exceeds the penetration depth of XPS (10 nm). Therefore, it is difficult to disclose the chemical composition of the inner lumen surface. The weak signal of B 1s in the XPS spectrum of HNTs-PA can be attributed to the fractions grafted on the opening of the tubes. The XPS results indicate that a vast majority of the fluorescein units were grafted on the HNTs lumen, which matches well with our previous findings and the FTIR results. The chemical composition of HNTs-PA was also investigated by EDS (shown in ESI†). The atomic% of O, Al and Si were calculated as 38.17%, 22.37% and 24.12%, respectively. The results of the presence of O, Al and Si are in accordance with the composition of alumino-silicate clay. The presence of carbon units (atomic% = 11.25%) revealed that the fluorescein groups were immobilized onto the HNTs, which matched well with the solid-state 13C NMR results.

The TEM images of original HNTs and HNTs-PA are shown in Fig. 4. Fig. 4A shows that the original HNTs possess a cylindrical shape and contain a transparent central area that runs longitudinally along the cylinder. The nanotubular particles are hollow and open ended, with a length of 0.5–1.0 μm, an inner diameter of 20–30 nm and an outer diameter of 40–60 nm, which can be taken or phagocytized in cell cultures.

After modification, the tubes retain their smooth outermost surface. Moreover, the lumen is almost completely filled, suggesting that the fluorescein derivative was successfully grafted onto alumina at the tube lumen and did not bind the outer siloxane surface of the tube. The overall morphologies of these nanotubes were similar to those of the original nanotube, indicating that the nanotubes underwent slight damage during the grafting reaction.

The established B–C link in the modified HNTs can be degraded into B–OH and Ar–OH in the presence of H2O2 as well as the B–C link in the fluorescein derivative. The quantum yield (\(\Phi_f\)) of HNTs-PA was calculated as ca. 0.01, with quinine sulfate (\(\Phi_f = 0.55\) in 0.05 M H2SO4) as the reference standard. Moreover, the degradation product, fluorescein, possesses a significantly higher fluorescence quantum yield (\(\Phi_f = 0.94\) in 0.1 M NaOH) than modified HNTs and fluorescein derivative bearing arylboronic acid groups. Therefore, the degradation from HNTs-PA to fluorescein can dramatically increase the...
fluorescence intensity, which can be called a “turn-on” response. The “turn-on” response in fluorescence intensity toward H₂O₂ can endow a H₂O₂-detecting ability of modified HNTs.

Fig. 5 shows the fluorescence responses of HNTs-PA to H₂O₂. When 5 × 10⁻⁶ mol H₂O₂ was added, the HNTs-PA aqueous solution displayed a drastic increase in fluorescence intensity in 30 min. A 50-fold difference in fluorescence intensity can even be detected when the amount of added H₂O₂ decreased to 1 × 10⁻⁷ mol (shown in ESI†), where the concentration of H₂O₂ is 5 × 10⁻⁵ M. Previously, we developed a HNTs-based H₂O₂-sensitive fluorescence probe, for which the fluorescence response could not be tracked when the concentration of H₂O₂ decreased to 5 × 10⁻⁵ M. Therefore, the HNTs-PA shows a prominent advantage as compared to the previous probe.⁴² Owing to their chemoselective boronate switch, HNTs-PA probes retain highly specific responses to H₂O₂. Other groups did not show the “turn-on” fluorescence when treated with other ROS agents, including NO⁻ and ClO⁻ (shown in the ESI†).

The fluorescence intensity of HNTs-PA was found to be almost unaffected by the addition of metal ions (Cu²⁺, Fe³⁺, Fe²⁺, Mg²⁺, Na⁺, Zn²⁺, K⁺, Ca²⁺) and anions (SO₄²⁻, H₂PO₄⁻, S₂O₃²⁻, NO₃⁻, CO₃²⁻, Cl⁻). Such favorable performances make the modified HNTs a desirable H₂O₂-responsive fluorescence probe in vivo.

To evaluate the cytotoxicity of the modified HNTs, the growth inhibitory activity against A549 cells was calculated based on a MTT assay. The concentration expressed in the Fig. 6 represents the concentration of HNTs-PA in aqueous suspension. The results indicate that HNTs-PA shows low cytotoxicity (over 80% cell viability) within 24 h of incubation time, in which the concentration of HNTs-PA ranges from 0.05 to 1.0 mg mL⁻¹, which makes the modified HNTs a desirable biological materials at cellular levels. In addition, the cytotoxicity of HNTs-PA was also evaluated using Hela cells, where the IC₅₀ was calculated to be 1.11 mg mL⁻¹.

It is well-known that the overexpression of H₂O₂ usually occurs around the tumor cells. We next assessed the ability of HNTs-PA to stain within human lung cancer cells (A549 cells). A549 cells were incubated with 0.5 mg mL⁻¹ HNTs-PA for 1, 2, 4, 8 and 12 h at 37 °C (shown in Fig. 7). The excitation wavelength was fixed at 488 nm and fluorescent signals were collected from 500 to 600 nm. A reference test was also made by incubating A549 cells with Mitotracker Red, which is a commercially available fluorescent dye for cell imaging. The images indicated that the cells can be stained under the endogenous levels of H₂O₂.
H₂O₂ derived from the cancer cells, where the fluorescence intensity increased as time extends. Moreover, the fluorescence intensity of the cells treated by Mitotracker Red remained almost unchanged after 4 h. The difference can be due to the fact that the fluorescein unit gradually released from the lumen as the B-C bond broke into B-OH and Ar-OH in the presence of H₂O₂, which corresponds to the “turn on” response of HNTs-PA on H₂O₂. We also assessed the ability of HNTs-PA to stain within Hela cells. The results indicated that the cells can also be stained and the fluorescence intensity increased as time extends, which can also be due to the endogenous levels of H₂O₂. The H₂O₂ concentration in mammals commonly ranges from 1 nM to 0.5 μM and is usually overexpressed in cancer cells and tissues. Sometimes the H₂O₂ concentration in cancer cells can exceed 50 μM. The H₂O₂ in A549 cells and Hela cells should also be in this range, implying the HNTs-PA is capable of detecting the endogenic H₂O₂ at this level.

Conclusions

In summary, we have described the design and synthesis of a fluorescent probe HNTs-PA, which can be used for the detection of H₂O₂. The nanocomposite showed regular morphology and desirable dispersibility in aqueous solution. The established B-C linkage provides a highly specific H₂O₂ sensitivity to HNTs-PA; as a result, the probe exhibits a highly specific “turn-on” fluorescence response for hyperoxide. Moreover, HNTs-PA was found to be capable to monitor H₂O₂ derived from A549 cells under physiological conditions. The prepared HNTs-based fluorescent probe can open new applications in the detection of cancer cells and thus serve as a practical tool for cancer diagnosis.

Conflicts of interest

The authors declare that they have no conflict of interest.

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