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# An artificial protein-probe hybrid as a responsive probe for ratiometric detection and imaging of hydrogen peroxide in cells†

Xing Zhang,<sup>ab</sup> Youxin Fu, <sup>bb</sup> Guangren Qian, <sup>b\*</sup> Run Zhang <sup>bb</sup> and Zhi Ping Xu <sup>b\*</sup>

An artificial protein-probe hybrid, Cm-Np-B@BSA, was prepared via host-guest interactions between hydrogen peroxide ( $H_2O_2$ )-responsive Cm-Np-B molecule and bovine serum albumin (BSA). The Cm-Np-B@BSA probe exhibited high sensitivity and selectivity towards  $H_2O_2$  under physiological conditions and had excellent biocompatibility, allowing for sensitive ratiometric detection and imaging of endogenous  $H_2O_2$  in live cells.

Fluorescent protein has emerged as an attractive platform in designing responsive probes for real-time detection and monitoring of specific biomarkers in biological systems.<sup>1–3</sup> In the past few decades, many efforts have been made to produce genetically encoded fluorescent protein probes to respond to biomarkers and investigate their applications in biology and biomedicine.<sup>4–7</sup> Despite the advantages of fluorescent protein probes in biosensing and bioimaging (such as excellent biocompatibility), only a limited number of responsive fluorescent protein probes have been reported.<sup>4,8,9</sup> This is mainly because the tedious procedure for fluorescent protein production restricts their expression on a large scale.

In order to overcome this limitation, several artificial protein-probe hybrids were made by integrating a fluorophore (e.g. organic fluorescent dye, luminescent metal complex, or quantum dot) with a suitable protein via supermolecular and host-guest interactions, or chemical conjugations. This kind of hybrid provides an alternative scaffold for designing responsive fluorescent protein-probe hybrids for biological applications. Incorporating various fluorophores, especially fluorescent organic dyes, into proteins to yield artificial protein-probe hybrids enhances the biocompatibility and bioavailability of fluorescent dyes. More importantly, the artificial hybrid enables biomarker detection in aqueous solutions and physiological

conditions without fluorescence quenching caused by dye molecule aggregation.  $^{14,15}$  For example, coumarin was loaded into the cavities of bovine serum albumin (BSA),  $^{16-18}$  and emitted characteristic fluorescence that was quenched in its aggregation state in water. With interactions between the fluorophore and protein, several fluorescent dyes, including coumarin, have been reported for protein detection in aqueous solutions.  $^{12,13,19-21}$  In this work, we expanded this idea and developed an artificial protein-probe hybrid specific for hydrogen peroxide ( $H_2O_2$ ) detection, and also highlighted the strategy of developing artificial protein-probe hybrids beyond the aforementioned protein detection.

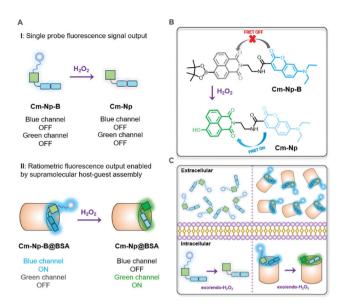
 $\rm H_2O_2$ , produced during mitochondrial respiration, is one of the most important reactive oxygen species (ROS) in biological systems.  $^{22-28}$  Endogenous  $\rm H_2O_2$  is the most stable ROS, having the lowest reactivity and the highest intracellular concentration ( $\sim 10^{-8}$ – $10^{-4}$  M).  $^{29}$  In living organisms, this biomolecule serves as an essential biomarker in regulating a wide variety of physiological events, such as activation of diverse signalling pathways to stimulate cell proliferation, differentiation, migration, and apoptosis.  $^{29-32}$  The role of  $\rm H_2O_2$  in biological systems is concentration-dependent.  $^{33}$  It is, therefore, highly demanded to develop suitable responsive fluorescence probes for  $\rm H_2O_2$  detection in live cells for a better understanding of  $\rm H_2O_2$ -mediated biofunctions.  $^{34}$ 

Different from previously reported molecule probes for  $H_2O_2$  detection, this communication presents a biocompatible responsive protein-probe hybrid for ratiometric detection of  $H_2O_2$  in aqueous solutions and living systems through a simple one-step self-assembly of organic dye **Cm-Np-B** with BSA. As shown in Scheme 1, the dye (**Cm-Np-B**) was designed based on the elegant modulation of the intramolecular charge transfer (ICT) and the Förster resonance energy transfer (FRET). The emission of energy transfer (ET) donor, *i.e.* coumarin (**Cm**, blue channel) can be activated after loading into the cavity of BSA through host–guest interactions in PBS buffer. Within the BSA system (**Cm-Np-B@BSA**), the emission of the ET acceptor, *i.e.* naphthalimide (**Np**, green channel), is quenched due to the inability of

<sup>&</sup>lt;sup>a</sup> School of Environmental Science and Chemical Engineering, Shanghai University, Shanghai 200444, China. E-mail: grqian@shu.edu.cn

b Australian Institute for Bioengineering and Nanotechnology, The University of Queensland, St Lucia, QLD 4072, Australia. E-mail: gordonxu@uq.edu.au

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Scheme 1 Design of an artificial protein-probe hybrid (Cm-Np-B@BSA) for responsive H<sub>2</sub>O<sub>2</sub> detection. (A) ON-OFF non-response of molecular probe (i) and response of BSA-probe (ii); (B) ON-OFF responsive mechanism; (C) cellular responses for H<sub>2</sub>O<sub>2</sub> detection.

the ICT process. Through an H2O2-triggered reaction to cleave boronate group,  $^{38}$  the "donor- $\pi$ -acceptor (D- $\pi$ -A)" hydroxyl-Np derivative (Cm-Np@BSA) is generated and thus the emission of Np is switched "ON". Moreover, the formation of Cm-Np@BSA also enables FRET from Cm to Np, allowing ratiometric fluorescence detection and bioimaging of H<sub>2</sub>O<sub>2</sub> with a blue-to-green fluorescence colour change.

The fluorescent dye, Cm-Np-B, was synthesized in a fourstep procedure with a reasonable yield (Scheme S1, ESI†), and the chemical structures of Cm-Np-B and the intermediates were identified with NMR and ESI-MS (Fig. S1-S5, ESI†). Then the protein-probe hybrid (Cm-Np-B@BSA) was made through simply mixing the probe and BSA in PBS buffer (Page S6, ESI†).

The blue channel activation of Cm-Np-B@BSA was first evaluated by UV-vis absorption and fluorescence spectra. As shown in Fig. 1A, Cm-Np-B was essentially non-fluorescent in both the blue and green channels in PBS buffer (pH 7.4, black dashed curve), largely attributed to the aggregation-caused

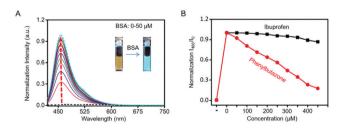


Fig. 1 Formation of Cm-Np-B@BSA and its fluorescence. (A) fluorescence spectra of Cm-Np-B (5  $\mu$ M) at different concentrations of BSA (0-50  $\mu$ M); (B) fluorescence intensity changes in Cm-Np-B@BSA (5/50  $\mu$ M, means 5  $\mu$ M probe with 50  $\mu$ M protein) with the concentration of ibuprofen (black solid line) and phenylbutazone (red solid line) increasing from 0 to 450  $\mu$ M in PBS buffer (0.5% DMSO (v/v), pH 7.4; excitation wavelength: 410 nm).

quenching (ACQ).<sup>39</sup> In sharp contrast, the fluorescence intensity was gradually increased upon mixing with more BSA (from 0 to 50 μM) and maximum enhancement of emission intensity (>145-fold) at 460 nm was achieved at 50 µM of BSA. Simultaneously, a new peak in the UV-vis absorption spectrum at 280 nm was observed, which is attributed to the characteristic absorbance of BSA (Fig. S6, ESI†). Similarly, Cm-Np-B is also responsive to human serum albumin (HSA). As shown in Fig. S7 (ESI†), the fluorescence intensity gradually increased upon mixing with more HSA (from 0 to 50 μM), presumably due to the similar 3D structure, chemical composition, and biological functions of these two albumins.

Dynamic light scattering (DLS) analysis showed a large size of molecule probe Cm-Np-B (600-700 nm) in PBS buffer (Fig. S8, ESI†), due to aggregation of Cm-Np-B molecules in PBS. Upon assembling Cm-Np-B with BSA, the hydrodynamic diameter of the Cm-Np-B@BSA hybrids was 17.8  $\pm$  1.3 nm, slightly larger than that of BSA (13.0  $\pm$  0.3 nm). The size increase and disappearance of Cm-Np-B aggregates together confirm formation of the guest-host complexes (Cm-Np-B@BSA) via binding Cm-Np-B into the hydrophobic pocket of BSA with a slightly big volume.

As shown in Fig. 1A, the emission intensity of Cm-Np-B dye was BSA-concentration dependent. Fluorescence Job's plot analysis showed a maximum intensity at the molecular fraction of 0.5 (Fig. S9, ESI†), indicating the 1:1 binding stoichiometry between Cm-Np-B and BSA. Cm-Np-B is thus located only at one binding site of BSA. Previous reports have shown that most small hydrophobic molecules are wrapped into the hydrophobic pockets of BSA, i.e. site I and site II.40-42 Phenylbutazone and ibuprofen are known to bind site I and site II of BSA, respectively. In order to determine the binding site, a competitive assay was carried out to displace Cm-Np-B from BSA using these two albumin-binding molecules. As shown in Fig. 1B, the intensity of blue emission at 460 nm was decreased upon adding phenylbutazone, while adding ibuprofen caused a negligible change in the fluorescence intensity. The selective replacement of Cm-Np-B by phenylbutazone suggests that the preferred binding site of Cm-Np-B is site I of BSA. In addition, the blue channel emission of the probe largely decreased upon adding trypsin, suggesting the release of Cm-Np-B into PBS buffer and subsequent aggregation after BSA degradation (Fig. S10, ESI†).

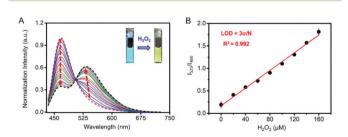
To test the fluorescence response of Cm-Np-B@BSA to H<sub>2</sub>O<sub>2</sub>, Cm-Np-B@BSA in PBS buffer was treated with H<sub>2</sub>O<sub>2</sub> at different concentrations and the changes in UV-vis absorbance and fluorescence spectra were recorded. As shown in Fig. S11A (ESI†), the absorbance at  $\lambda_{abs}$  = 350 nm was gradually decreased and the maximum absorbance band at  $\lambda_{abs}$  = 430 nm was gradually intensified upon the addition of H<sub>2</sub>O<sub>2</sub> (0-200 µM). The change of UV-vis absorption spectra is ascribed to the ICT feature in Cm-Np dye due to the transformation of Cm-Np-B@BSA to Cm-Np@BSA (Scheme 1). Clearly, the UV-vis absorption spectral change in Cm-Np-B@BSA towards H2O2 was more sensitive than that of Cm-Np-B (Fig. S11B, ESI†), suggesting the higher sensitivity of the artificially devised fluorescent protein-probe hybrid.

As shown in Fig. 2A, Cm-Np-B@BSA exhibited a blue emission band from the coumarin donor ( $\lambda_{ex} = 410 \text{ nm}$ ,  $\lambda_{em} = 460 \text{ nm}$ ) in the absence of  $H_2O_2$ . Upon the addition of  $H_2O_2$  to test solution, the green emission band from the **Np** acceptor at  $\lambda_{\rm em}=525$  nm was gradually intensified, accompanied by blue emission weakening. The green emission enhancement is contributed by the ICT-dominated fluorescence of **Cm-Np** and the FRET process from **Cm** to **Np**, while the blue emission decrease is ascribed to the FRET process. Similar to the unchanged UV-vis absorption spectrum with  $H_2O_2$ , **Cm-Np-B** showed no fluorescence response to  $H_2O_2$  (Fig. S12, ESI†), corroborating the superior capability of the artificial protein-probe hybrid in  $H_2O_2$  detection. Besides these, the time-dependent fluorescence response of probeprotein hybrid **Cm-Np-B@BSA** and **Cm-Np-B** toward  $H_2O_2$  in PBS buffer was also measured (Fig. S13, ESI†), and **Cm-Np-B@BSA** showed a faster response than **Cm-Np-B**.

The blue-to-green fluorescence emission change in **Cm-Np-B@BSA** enables ratiometric fluorescence detection of  $H_2O_2$  in PBS buffer. As shown in Fig. 2B, the dose-dependent fluorescence response  $(F_{525}/F_{460})$  of **Cm-Np-B@BSA** exhibited a good linearity against the concentration of  $H_2O_2$  in the range of 0–160  $\mu$ M. The detection limit  $(3\sigma/k)$  was calculated to be 1  $\mu$ M. The low detection limit and the broad detection range indicate the high sensitivity of artificial protein-probe hybrid **Cm-Np-B@BSA** for  $H_2O_2$  detection.

The selectivity of **Cm-Np-B@BSA** towards  $H_2O_2$  over other interfering species, including some other ROS, anions, and biomolecules, was also confirmed. **Cm-Np-B@BSA** in PBS buffer was treated with 200  $\mu$ M of  $H_2O_2$  and 500  $\mu$ M of other species, respectively. After 10 min incubation, remarkable changes of the fluorescence of **Cm-Np-B@BSA** were noticed in the presence of  $H_2O_2$ , but not in the presence of other species ( $F_{525/460}$ , Fig. S14, ESI†). Furthermore, the competitive selectivity of **Cm-Np-B@BSA** to  $H_2O_2$  over other species was also tested. As shown in Fig. S15 (ESI†), the fluorescence ( $F_{525/460}$ ) was increased exclusively to  $H_2O_2$  and the enhancement of fluorescence intensity was not affected in the presence of all competitive species. These data indicate the high selectivity of protein-probe hybrid **Cm-Np-B@BSA** for a ratiometric fluorescence response to  $H_2O_2$  in PBS buffer.

Then, the effect of pH on the ratiometric fluorescence response of Cm-Np-B@BSA to  $H_2O_2$  was examined in PBS buffer with pH 5-8. As shown in Fig. S16E (ESI $\dagger$ ), no obvious difference in the emission spectra was observed in solutions

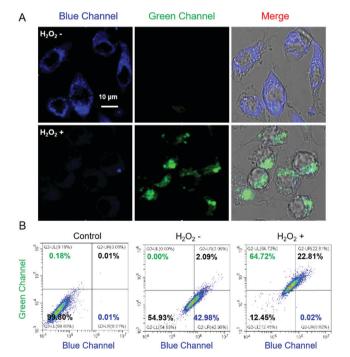


**Fig. 2** Sensitivity and selectivity of **Cm-Np-B@BSA** (5/50 μM) for  $H_2O_2$  detection. (A) Fluorescence spectral response of **Cm-Np-B@BSA** (5/50 μM) to  $H_2O_2$  at the concentration of 0–200 μM; (B) linear detection range and limit of detection (LOD) of **Cm-Np-B@BSA** (5/50 μM). All measurements were carried out in PBS buffer, pH 7.4, excitation wavelength: 410 nm.

with pH 6–8, suggesting that the Cm–Np-B@BSA probe is able to detect  $H_2O_2$  in an environment with pH 6–8, while the peak at 525 nm in pH 5 solution was relatively weaker, probably due to the acidity slowing down the  $H_2O_2$  detection reaction.

Encouraged by the excellent sensitivity and selectivity of the Cm–Np-B@BSA response to  $\rm H_2O_2$ , we further demonstrated the application of this artificial protein-probe hybrid for imaging  $\rm H_2O_2$  in living cells. MTT assay showed that Cm–Np-B@BSA had low cytotoxicity towards RAW 264.7 macrophage cells (Fig. S17, ESI†). For example, the cell viability remained over 85% after incubation of RAW 264.7 macrophage cells with Cm–Np-B@BSA (60  $\mu$ M/50  $\mu$ M) for 24 h. For safety, 5  $\mu$ M/50  $\mu$ M of Cm–Np-B@BSA was used in most subsequent experiments.

The capability of **Cm-Np-B@BSA** in visualizing intracellular  $H_2O_2$  in RAW 264.7 cells was then examined. As shown in Fig. 3A, clear blue fluorescence was noticed after incubating RAW 264.7 cells with **Cm-Np-B@BSA** for 2 h, which was remarkably increased in comparison with the control group (RAW 264.7 macrophage cells only) (Fig. S18, ESI†). The flow cytometric data showed a similar change. Both observations confirm the effective cell internalization of **Cm-Np-B@BSA** (Fig. S19, ESI†). Interestingly, supplying **Cm-Np-B@BSA**-internalized cells with  $H_2O_2$  (200  $\mu$ M), the intracellular blue channel fluorescence was largely quenched, while intense green channel fluorescence emerged (Fig. 3A). This remarkable change was also confirmed by flow cytometric analysis (Fig. 3B), in agreement with spectrometric



**Fig. 3** Fluorescence imaging and flow cytometric data of Raw 264.7 cells using **Cm–Np-B@BSA** (5/50 μM) as the probe. (A) Fluorescence imaging of living cells with ( $H_2O_2+$ ) or without ( $H_2O_2-$ ) exogenous  $H_2O_2$  (200 μM); (B) flow cytometric data of cells with ( $H_2O_2+$ ) or without ( $H_2O_2-$ ) exogenous  $H_2O_2$  (200 μM).  $\lambda_{ex}=405$  nm,  $\lambda_{em}=460-480$  and 520–560 nm for blue and green channels, respectively. The images were recorded by confocal laser-scanning microscopy (scale bar = 10 μm).

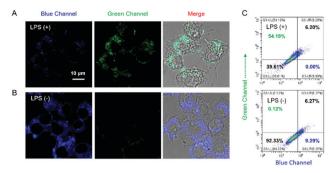


Fig. 4 Fluorescence imaging of Raw 264.7 cells pretreated with LPS using Cm-Np-B@BSA (5/50 µM) as the probe. (A) Fluorescence imaging of living cells pretreated with 80 ng  $mL^{-1}$  of LPS; (B) control group of cells; (C) flow cytometric data of Raw 264.7 cells.  $\lambda_{ex}$  = 405 nm,  $\lambda_{em}$  = 460–480 and 520-560 nm for blue and green channels, respectively. The images were visualized by confocal laser-scanning microscopy (scale bar =  $10 \mu m$ ).

analysis (Fig. 2A) and in sharp contrast to that using Cm-Np-B for the same fluorescence imaging and flow cytometric analysis, where only small alternation in the fluorescence blue and green channels was observed (Fig. S20, ESI†). Thus the sharp contrast corroborates the high sensitivity of the fluorescent protein-probe hybrid (Cm-Np-B@BSA) for intracellular H<sub>2</sub>O<sub>2</sub> detection.

The capability of Cm-Np-B@BSA for ratiometric fluorescence visualization of intracellular H<sub>2</sub>O<sub>2</sub> generation was further evaluated in LPS-stimulated RAW 264.7 macrophage cells. As shown in Fig. 4, obvious fluorescence emissions in both the blue and green channels were observed, suggesting that Cm-Np-B@BSA is able to detect endogenous H<sub>2</sub>O<sub>2</sub> in live cells.

In summary, we developed an artificial protein-probe hybrid, Cm-Np-B@BSA, for ratiometric fluorescence detection and imaging of H<sub>2</sub>O<sub>2</sub> in physiological buffers and live cells. Cm-Np-B@BSA was devised by assembling H2O2-responsive Cm-Np-B molecules with BSA via host-guest interactions. The strong binding of Cm-Np-B with BSA occurred at the preferred site I of BSA. In this complex, blue channel emission ( $\lambda_{em}$  = 460 nm) was activated (>145-fold enhancement) while the green channel ( $\lambda_{em} = 525$  nm) remained weakly fluorescent. Cleavage of a borate group of Cm-Np-B@BSA by the H2O2-initialized reaction generated a new product Cm-Np@BSA, and enabled green channel emission observable as the result of FRET from Cm to Np. This change allowed Cm-Np-B@BSA to be used in ratiometric fluorescence detection of  $H_2O_2$  in aqueous solution. This Cm-Np-B@BSA exhibited high sensitivity and selectivity for H<sub>2</sub>O<sub>2</sub> detection with good biocompatibility, and demonstrated ratiometric fluorescence detection of H<sub>2</sub>O<sub>2</sub> in live macrophage cells. Thus, the success in developing Cm-Np-B@BSA not only provides a useful tool to monitor H<sub>2</sub>O<sub>2</sub> generation in live cells, but also offers a feasible way to design new artificial proteinprobe hybrids for biomolecule detection and imaging.

#### Conflicts of interest

There are no conflicts to declare.

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