

Cite this: *Chem. Sci.*, 2017, 8, 4833

Target-triggered cascade assembly of a catalytic network as an artificial enzyme for highly efficient sensing†

Lei Zhang,‡ Fengjiao Ma,‡ Jianping Lei, * Jintong Liu and Huangxian Ju*

Determining the catalytic activity of artificial enzymes is an ongoing challenge. In this work, we design a porphyrin-based enzymatic network through the target-triggered cascade assembly of catalytic nanoparticles. The nanoparticles are synthesized *via* the covalent binding of hemin to amino-coated gold nanoparticles and then the axial coordination of the Fe center with a dual-functional imidazole or pyridine derivative. The network, which is specifically formed by coordination polymerization triggered by Hg²⁺ as the target, shows high catalytic activity due to the triple amplification of enzymatic activity during the cascade assembly. The catalytic dynamics are comparable to those of natural horseradish peroxidase. The catalytic characteristics can be ultrasensitively regulated by the target, leading to a selective methodology for the analysis of sub-attomolar Hg²⁺. It has also been used for "signal-on" imaging of reactive oxygen species in living cells. This work provides a new avenue for the design of enzyme mimics, and a powerful biocatalyst with signal switching for the development of biosensing protocols.

Received 31st March 2017

Accepted 25th April 2017

DOI: 10.1039/c7sc01453h

rsc.li/chemical-science

Introduction

In nature, enzymes often participate in self-assembly processes to form "soft" nanostructures, such as spheres and tubes, demonstrating their catalytic functions. Many synthetic catalysts that behave as enzymes can mimic the functions of natural systems.^{1–4} However, the performance of some artificial enzymes is very inferior to that of corresponding natural enzymes, possibly as a result of the aggregation of metal macrocycles, such as porphyrins and phthalocyanines, to form inactive dimers that are arranged in a disordered manner.⁵ To improve the catalytic efficiency, much effort has been devoted to the self-assembly of porphyrins into ever-larger structures involving porphyrinic dendrimers, nanocrystals, and square nanosheets,^{6–9} and the incorporation of porphyrin-containing hemin in a guanine quadruplex to dissociate the aggregate.^{10,11} The assembly of porphyrins on nanomaterials has been demonstrated to be a simple and efficient approach for achieving high catalytic activity, specific recognition and even signal transduction,^{12,13} and it was used to produce multifunctional nanoscale systems for artificial photosynthesis, catalysis, and biosensing.^{14–23} Porphyrinic metal-organic frameworks as

crystalline molecular materials have shown unique enzyme-like activities.^{24–26} Inspired by the promising applications of porphyrin assemblies as artificial enzymes, this work designs a target-triggered cascade assembly mechanism for the formation of a porphyrin-based enzymatic network, which leads to an ultrasensitive signal switch for target-related bioanalysis. The assembly of porphyrins on nanomaterial surfaces is generally performed through chemisorption, π - π stacking interactions, covalent binding and axial ligation.^{27–31} Owing to the lack of functional groups on biocompatible gold nanoparticles (AuNPs), which are extensively used in biomedicine, it is difficult to immobilize porphyrinic catalysts on AuNPs. Thus, this work describes the preparation of amino group modified AuNPs (N-AuNPs) using a rationally designed ligand, pamidronic acid disodium salt (PADS), for effective chemical combination with metalloporphyrins (hemin) *via* an amide reaction (Fig. 1A). The **Au-Hem** conjugates then assemble with two dual-functional small molecules: 5-(1*H*-imidazol-4-ylmethylene)-pyrimidine-2,4,6-trione (**MPT**) *via* the model shown in Fig. 1B, and 5-(pyridin-4-ylmethylene)-pyrimidine-2,4,6-trione (**PMPT**) through axial coordination. In the presence of Hg²⁺ as a target, the coordination polymerization of functionalized AuNPs (**Au-Hem-MPT** or **Au-Hem-PMPT**) *via* N-Hg^{II}-N bond connectivities produces enzyme-active networks (**Au-Hem-Net** and **Au-Hem-pNet**). Both of the networks show catalytic activities that are much higher than that of free hemin due to the triple amplification of enzymatic activity during the cascade assembly (Fig. 1C). The ordered network provides catalytic dynamics that are comparable to those of the natural enzyme horseradish

State Key Laboratory of Analytical Chemistry for Life Science, School of Chemistry and Chemical Engineering, Nanjing University, Nanjing 210023, P. R. China. E-mail: jpl@nju.edu.cn; hxju@nju.edu.cn; Tel: +86 25 89681922

† Electronic supplementary information (ESI) available: Experimental details, and supporting figures and tables. See DOI: 10.1039/c7sc01453h

‡ These authors contributed equally to this work.



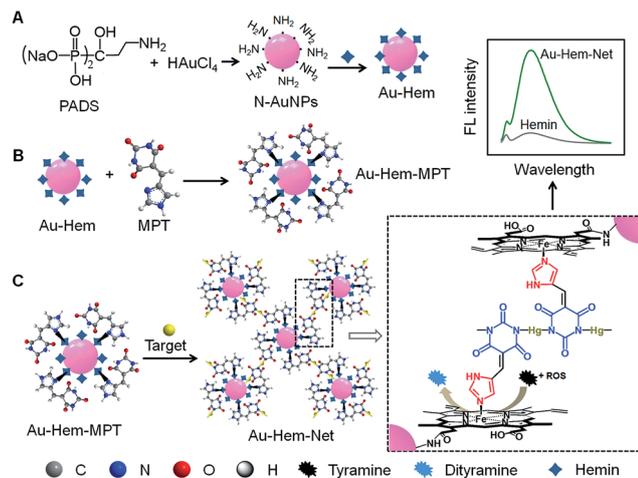


Fig. 1 Schematic representation of the cascade assembly of high catalytic networks. (A) The synthesis of N-AuNPs using PADS as both a reducing agent and a protection ligand, and Au-Hem conjugates *via* an amide reaction. (B) The assembly of MPT on Au-Hem through axial interactions. (C) The preparation of an enzymatic network *via* Hg²⁺-triggered polymerization and the catalytic oxidation of tyramine by the network in the presence of ROS.

peroxidase (HRP). Moreover, the catalytic characteristics can be flexibly regulated by trace amounts of Hg²⁺, a highly toxic metal ion that damages the central nervous and endocrine systems.^{32,33} This work provides a powerful protocol for the ultrasensitive sensing of sub-attomolar Hg²⁺. More importantly, the network showed low cytotoxicity and it can be conveniently used for the sensitive fluorescence imaging of intracellular enzyme-related species, such as reactive oxygen species (ROS). Thus, the cascade synthesis of target-controllable catalytic systems provides a new concept for the design and application of enzyme mimics.

Results and discussion

Construction and characterization of enzyme-active networks

The N-AuNPs were synthesized *via* the reaction of HAuCl₄ with PADS as both the reducing reagent and the protection ligand in a boiling aqueous solution. At different mass ratios of PADS:HAuCl₄, the obtained N-AuNPs showed different colours related to size and dispersion (Fig. 2A). The sharp absorption peak in their UV-*vis* spectra (Fig. 2B) and dynamic light scattering measurements (Table S1†) indicated a relatively narrow size distribution at ratios from 3.2 : 1 to 11.0 : 1. Transmission electron micrographs (TEMs) and zeta-potential analysis further demonstrated that the best dispersion of N-AuNPs was at a ratio of 5.0 : 1 (Fig. S1 and Table S1†), which showed the largest zeta potential, a diameter of around 17 nm, and the characteristic surface plasmon band centered at 525 nm.

The formation mechanism of N-AuNPs was studied using 1-hydroxyethane-1,1-diphosphonic acid (HEDP), which does not contain an amino group at the tertiary carbon, and 3-amino-propylphosphonic acid (APPA), which does not contain a hydroxyl group at the carbon adjacent to the phosphonic

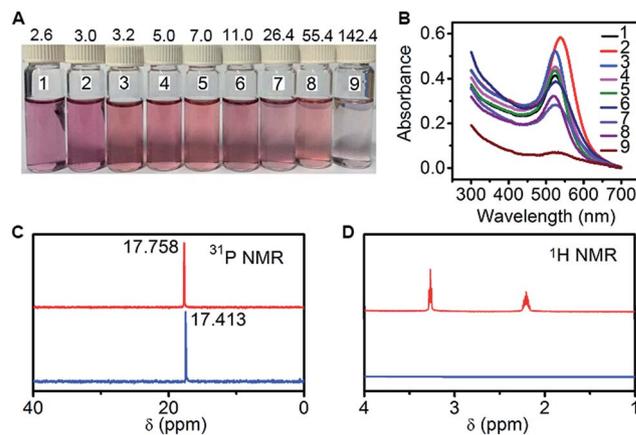


Fig. 2 (A) Photograph and (B) UV-*vis* spectra of AuNPs prepared with PADS and HAuCl₄ at mass ratios of 2.6, 3.0, 3.2, 5.0, 7.0, 11.0, 26.4, 55.4 and 142.4 from 1 to 9. (C) ³¹P NMR and (D) ¹H NMR spectra of PADS (red) and the reaction mixture at a mass ratio of 5.0 (blue).

group, as controls. An obvious colour change occurred in the mixtures of HAuCl₄ with PADS and HEDP, while no reaction and no plasmon absorption of the AuNPs were observed in the mixture of HAuCl₄ with APPA (Fig. S2†), demonstrating that the hydroxyl group at the tertiary carbon was involved in the oxidation reaction of PADS or HEDP by HAuCl₄.^{34,35} The oxidation reaction led to a shift in the ³¹P NMR peak to a higher magnetic field, accompanied by the signal disappearance of two multiple peaks that were attributed to two methylene groups in the ¹H NMR spectrum (Fig. 2C and D), helping to identify the formation of a C=O bond *via* the cleavage of another group from the carbon, as shown in the electrospray ionization mass spectra (Fig. S3†). The N-AuNPs prepared with PADS remained as a deep red suspension for several months and exhibited better stability and dispersion (Fig. S2†) due to the presence of an excess amount of PADS on their surfaces. The presence of excess PADS on the nanoparticles was supported by the characteristic vibration frequencies of primary amines at 3228.9 and 3406.1 cm⁻¹ for N-AuNPs in the infrared absorption spectrum (Fig. 3A). More importantly, the adsorbed PADS not only protected the AuNPs against aggregation, but also provided the -NH₂ group for covalent binding with hemin, which was characterized by the infrared absorption of the amide bond at 1698.4 and 1298.6 cm⁻¹ (Fig. 3A).

The formed Au-Hem showed a single Soret band at around 400 nm (Fig. 3B). The disappearance of the dimeric absorption of hemin at 385 nm and the presence of monomeric absorption at 400 nm and around 350 nm (Fig. S4A†) suggested that the hemin on the N-AuNPs was in its monomeric state, the preferred form for obtaining a high catalytic ability.³⁶ The number of hemin complexes on each Au nanoparticle was calculated to be 73 from the Soret band intensity of hemin using UV-*vis* spectroscopy (see the ESI Experimental section and Fig. S4B†), indicating an acceptable capacity of the enzyme mimic system for amplifying the catalytic activity.

The Au-Hem was further modified with MPT or PMPT *via* the axial coordination of their imidazole or pyridine group with the



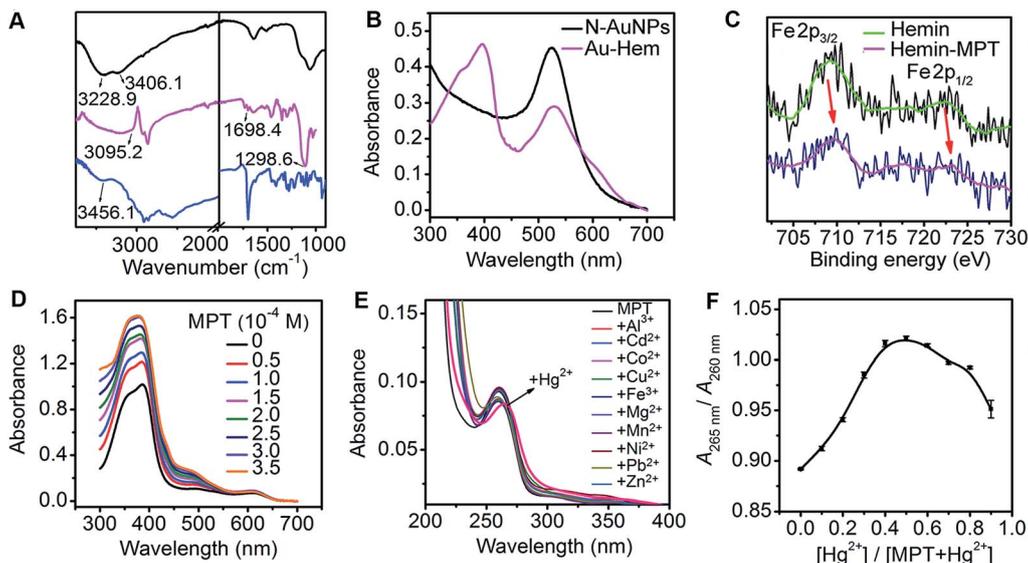


Fig. 3 (A) The Fourier transform infrared spectra of N-AuNPs (black), hemin (blue) and Au-Hem (magenta). (B) The UV-vis spectra of N-AuNPs and Au-Hem (1 μM equivalent hemin) solutions. (C) The Fe 2p XPS spectra of hemin and the hemin-MPT complex. (D) The change in the UV-vis spectrum upon the titration of MPT to hemin (20 μM) in pH 7.4 Tris-HCl at 36 $^{\circ}\text{C}$. (E) The UV-vis spectra of MPT (10 μM) in the presence of different metal ions (100 μM). (F) A Job's plot for the binding of Hg^{2+} to MPT. The total concentration of MPT and Hg^{2+} was kept constant at 100 μM in pH 7.4 Tris-HCl.

Fe center of hemin. Both **MPT** and **PMPT** were synthesized to contain a pyrimidinetrione moiety for the preparation of the enzyme-active network by coordination polymerization. The **MPT** and **PMPT** structures were firstly characterized using MS and NMR information (see the ESI Experimental section, and Fig. S5 and S6 \dagger). The Fe 2p X-ray photoelectron spectra (XPS) (Fig. 3C and S7A \dagger) demonstrate that the axial coordination interaction of hemin with both **MPT** and **PMPT** increased the Fe 2p $_{1/2}$ binding energy due to a change in the electron density on the Fe atom.²⁷ The coordination numbers and the binding constants (n and K) were determined *via* the titration of the hemin solution with **MPT** and **PMPT**,³⁷ which led to a slight red shift of the Soret band of hemin at 385 nm (Fig. 3D and S7B \dagger), further suggesting the formation of an axial ligation to the Fe center of hemin.³⁸ As expected, a six-coordinated structure of hemin with two **MPT** complexes at both axial sides could be formed. However, it occurred only at relatively low temperatures, and showed a much higher binding constant than the five-coordinated structure formed at relatively high temperatures due to the exothermic axial coordination reaction (Fig. S8 \dagger). Interestingly, only one **PMPT** complex could axially coordinate to the Fe center to form the five-coordinated hemin structure due to the weaker coordination ability of the pyridine group compared to that of the imidazole group, which led to relatively lower binding constants than those of the hemin-**MPT** complex (Table S2 \dagger). The $\Delta_r H$ and $\Delta_r S$ values for the formation of the hemin-**PMPT** complex were calculated to be $-156.30 \text{ kJ mol}^{-1}$ and $-432.33 \text{ J mol}^{-1} \text{ K}^{-1}$, respectively. Considering the fact that the dual axial coordination blocked the access of substrate to the Fe sites for catalytic oxidation,³⁹ the **Au-Hem-MPT** was synthesized at 37 $^{\circ}\text{C}$, the physiological temperature, for ensuring effective enzymatic catalytic activity.

The interaction between the imide site of thymine and Hg^{2+} to form a “N-Hg^(II)-N” structure is well known and is used for bioanalysis.^{40,41} The designed pyrimidinetrione moiety in both **MPT** and **PMPT** contained two imide sites. Using **MPT** as an example, the absorption peak of **MPT** was red-shifted from 260 nm to 265 nm upon the addition of Hg^{2+} in the **MPT** solution, while other ions did not change the absorption peak (Fig. 3E), demonstrating the specific recognition of the two imide sites of **MPT** to Hg^{2+} to form a “N-Hg^(II)-N” like binding motif.⁴² Using the absorbance ratio of $A_{265 \text{ nm}} / A_{260 \text{ nm}}$ in the UV-vis spectra, the Job's plot analysis revealed a 1 : 1 binding stoichiometry of the coordination reaction (Fig. 3F). Thus the structure of the **MPT-Hg**²⁺ coordination polymer is represented in the inset of Fig. S9A \dagger . The matrix-assisted laser desorption/ionization time-of-flight mass spectrum of the mixture of **MPT** and Hg^{2+} further verified the formation of the 1 : 1 **MPT-Hg**²⁺ coordination polymer (Fig. S9A \dagger), in which the mass peaks at m/z 406.6, 612.6, 655.9 and 847.6 could be attributed to the characteristic fragments **MPT** + Hg-1H, **2MPT** + Hg-2H, **2MPT** + 2Hg-2 imidazole groups-3H and **2MPT** + 2Hg + Cl-3H, respectively.

The 1 : 1 pyrimidinetrione- Hg^{2+} coordination reaction induced the formation of **Au-Hem-Net** and **Au-Hem-pNet**, as shown in Fig. 1C. After the Hg^{2+} -triggered polymerization of **Au-Hem-MPT**, both the dynamic light scattering measurements and the transmission electron micrographs showed a greatly increased nanoparticle size from ~ 30 to ~ 130 nm (Fig. 4A and B). The presence of two clusters of MS peaks at around m/z 1258.6 and 1465.0 also supported the formation of **Au-Hem-Net** (Fig. S9B \dagger). Here the characteristic peaks at m/z 652.0 and 859.4 corresponded to hemin and hemin-**MPT**, and the peaks at around m/z 1258.6 and 1465.0 could be attributed to hemin-**MPT** + 2Hg-2H and hemin-**MPT** + 2Hg + **MPT**-3H, respectively.



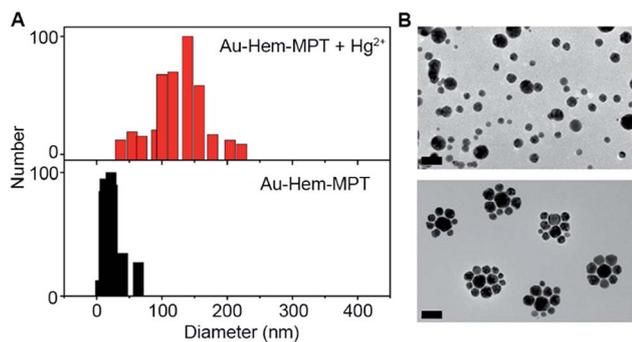


Fig. 4 (A) DLS assays of Au-Hem-MPT (0.1 μM equivalent hemin) before and after the addition of Hg^{2+} (0.1 μM). (B) The TEM images of Au-Hem-MPT (0.1 μM equivalent hemin) in the absence (top) and presence (bottom) of Hg^{2+} (0.1 μM). The scale bars represent 50 nm.

Catalytic performance

To evaluate the catalytic performance of the enzyme mimics proposed in this work, a fluorescence assay of the tyramine oxidation reaction was performed, in which the nonfluorescent tyramine (0.14 mM) was oxidized by hydrogen peroxide (0.4 mM) to give fluorescent dityramine,⁴³ thus the fluorescent signal represented the catalytic activity of these enzymes as catalysts. The amplified catalytic activity was demonstrated by the strong fluorescent signal of the oxidation product of tyramine by H_2O_2 in the presence of Au-Hem, when compared to the N-AuNPs and free hemin in aqueous solution (Fig. 5A).

On increasing the MPT concentration in Au-Hem, the catalytic activity was slightly enhanced (Fig. 5B), suggesting that the axial coordination of MPT to the Fe center of hemin changed its electronic and geometric structure, which increased the rate of the oxidation–reduction reaction.²⁷ However, the catalytic activity greatly increased upon the addition of MPT in the mixture of Au-Hem and Hg^{2+} , and reached a maximum at a hemin/MPT concentration ratio of 1 : 1. At this concentration ratio, the formed mimic produced a 2.6, 3.1 and 14.9 times higher fluorescence intensity than that of Au-Hem-MPT, Au-Hem and hemin, respectively, (Fig. 5A), suggesting an improved peroxidase activity in each assembly step. As a control, different concentrations of Hg^{2+} were added in the Au-Hem solution, and the catalytic oxidation product did not show any fluorescence change (Fig. 5C). More interestingly, this method showed high selectivity. Other metal ions did not produce an obvious fluorescence change (Fig. 5D). Thus, high peroxidase activity resulted from the Hg^{2+} -induced coordination polymerization of Au-Hem-MPT. That is, Hg^{2+} -triggered nanoparticle aggregation results in ordered multiscale organization and the strong interparticle coupling of surface plasmons between neighboring nanoparticles.^{44,45} Thus, the plasmon enhancement effect accelerates the electronic transfer between H_2O_2 and Fe^{III} hemin with the production of an initial intermediate (a ferrylporphyrin radical cation),⁴⁶ leading to the improved catalytic performance of the Hg^{2+} triggered network. In addition, the networks could be formed by the different sized N-AuNPs with average diameters of 26.1–6.4 nm, which was deduced from the

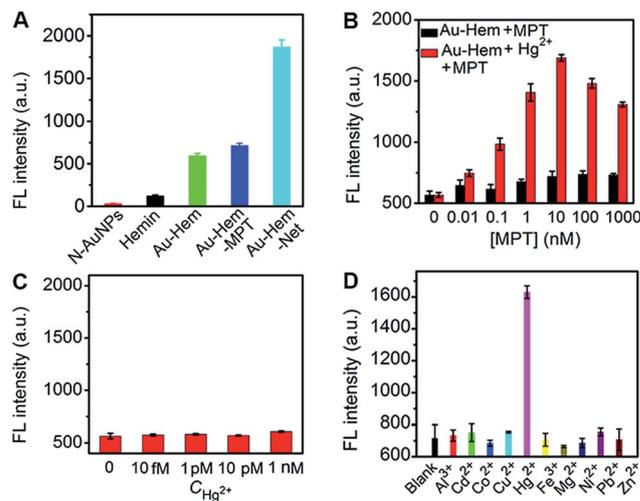


Fig. 5 (A) Fluorescence intensities of the oxidation product of tyramine (0.14 mM) catalyzed by N-AuNPs (1.0 nM), hemin, Au-Hem, Au-Hem-MPT and Au-Hem-Net (10 nM equivalent hemin) in the presence of H_2O_2 (0.4 mM) at 37 $^\circ\text{C}$. (B) The fluorescence responses of the oxidation product of tyramine (0.14 mM) by H_2O_2 (0.4 mM), which is catalyzed by the adduct of Au-Hem (10 nM equivalent hemin) and MPT at marked concentrations in the absence (black) and presence (red) of Hg^{2+} (1.0 nM) at 37 $^\circ\text{C}$. (C) The fluorescence response of the oxidation product of tyramine (0.14 mM) by 0.4 mM H_2O_2 for 15 min at 37 $^\circ\text{C}$ in the presence of an incubated mixture of Au-Hem (10 nM equivalent hemin) and different concentrations of Hg^{2+} . (D) The fluorescence intensities of the oxidation product of tyramine (0.14 mM) by H_2O_2 (0.4 mM), which is catalyzed by Au-Hem-MPT (10 nM equivalent hemin) in the presence of marked metal ions at 37 $^\circ\text{C}$. The concentration of Hg^{2+} is 1.0 nM, and the concentrations of the other metal ions are 10 nM.

increased fluorescence intensity after the addition of Hg^{2+} , among which small-sized particles had less of a tendency to form the network (Fig. S10[†]).

The Hg^{2+} -regulated catalytic characteristics of Au-Hem-MPT could be applied to the sensitive detection of Hg^{2+} using the catalytic fluorescence signal. Using the optimized incubation (9 min) and catalytic reaction (15 min) time (Fig. 6A and B), a remarkable fluorescence increase was observed upon addition of Hg^{2+} in the mixture of Au-Hem-MPT, tyramine and H_2O_2 at 37 $^\circ\text{C}$ (Fig. 6C). The plot of fluorescence intensity vs. the logarithm of Hg^{2+} concentration showed good linearity in the range 1.0 aM to 10 pM with a correlation coefficient of 0.992 (Fig. 6D). It is worth noting that the detection limit at 3σ was 0.30 aM, which was 7 orders of magnitude lower than that of conventional analytical methods,^{47–50} and also the most sensitive method available for detection of Hg^{2+} (Table S3[†]). Since Hg^{2+} is very difficult to biodegrade and can accumulate in organisms to cause various human diseases,⁵¹ the coordination polymerization-amplified catalytic reaction is of great importance in monitoring Hg^{2+} at ultralow abundances and the slow accumulation of Hg^{2+} in organisms.

The Hg^{2+} -triggered formation of Au-Hem-pNet could also be used for Hg^{2+} detection with the same detection system, and showed a detectable concentration range of 1.0 aM to 100 pM with a detection limit of 0.78 aM (Fig. S11A and B[†]). Similarly,



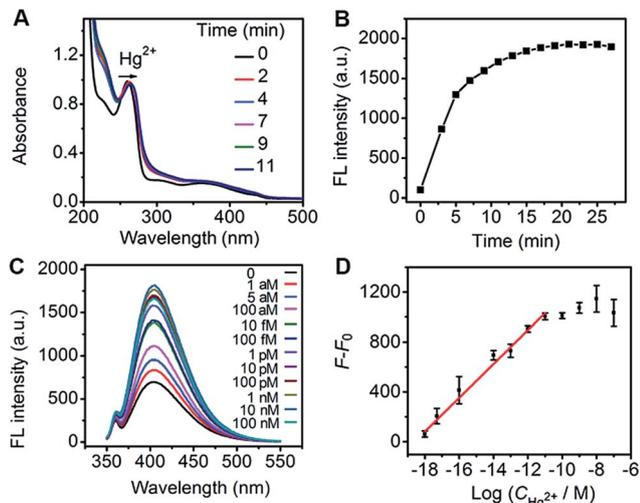


Fig. 6 (A) Time-dependent UV-vis spectra of MPT (100 μM) in the presence of Hg^{2+} (100 μM). (B) The time-dependent fluorescence intensity of the oxidation product of tyramine (0.14 mM) catalyzed by Au-Hem-Net (10 nM equivalent hemin) in the presence of 0.4 mM H_2O_2 at 37 $^\circ\text{C}$. (C) The fluorescence spectra of the oxidation product of tyramine (0.14 mM) by H_2O_2 (0.4 mM), which is catalyzed by Au-Hem-MPT (10 nM equivalent hemin) in the presence of Hg^{2+} at marked concentrations at 37 $^\circ\text{C}$. (D) A plot of $F - F_0$ vs. the logarithmic value of Hg^{2+} concentration. F_0 and F are the fluorescence intensity in the absence and presence of Hg^{2+} , respectively.

the formed highly catalytic network could be applied to the detection of H_2O_2 . Using Au-Hem-Net as an example, the mixture of Au-Hem-Net and tyramine showed linearly increasing fluorescence intensity with increasing H_2O_2 concentration in the range 4.4 to 100 μM (Fig. S11C and D†).

To evaluate the catalytic dynamics of the designed network, the oxidation product of pyrogallol by H_2O_2 was detected using UV-vis spectroscopy. At pyrogallol concentrations that were much higher than those of the enzyme mimics, three enzyme mimics, Au-Hem, Au-Hem-PMPT and Au-Hem-pNet, all showed a linear Lineweaver-Burk plot at the initial oxidation stage (Fig. 7), which led to two important kinetic parameters, K_m (the Michaelis constant) and k_{cat} (the catalytic kinetic constant), as listed in Table 1. The K_m value of Au-Hem was a surprisingly low value of 0.40 mM, which was less than that of HRP (0.81 mM), indicating a good affinity of pyrogallol to the Au-Hem conjugate. Meanwhile, Au-Hem-PMPT and Au-Hem-pNet showed comparable affinities to HRP. The k_{cat} values of Au-Hem (419.6 min^{-1}), Au-Hem-PMPT (714.3 min^{-1}) and Au-Hem-pNet (1054.5 min^{-1}) were 2–3 orders of magnitude higher than that of free hemin (2.4 min^{-1}) and higher than those of other carrier-supported hemin composites.^{20,21,52,53} Moreover, the kinetic constant of Au-Hem-pNet was comparable to that of natural HRP (1750 min^{-1}), highlighting the excellent catalytic activity of the enzyme-active network, which was greatly facilitated by the substrate being close to the active site of the enzyme and cooperative binding.⁵⁴

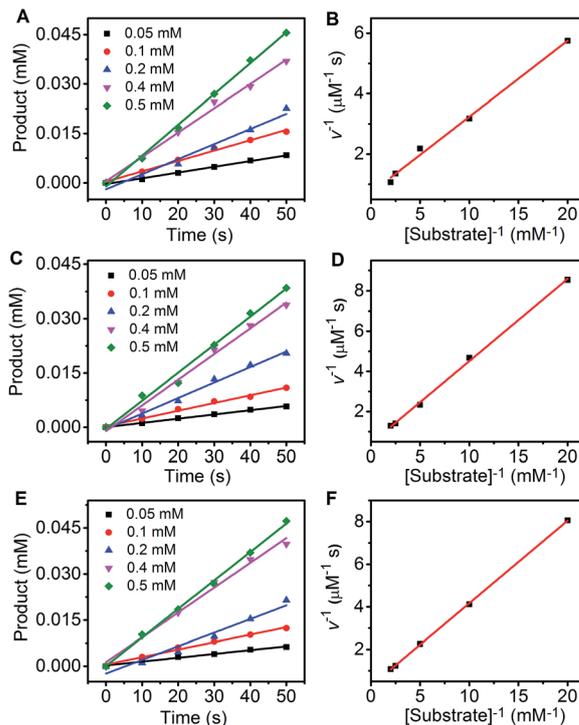


Fig. 7 The fitting curve of the initial pyrogallol oxidation profile at marked pyrogallol concentrations (A, C and E) and a Lineweaver-Burk plot of the pyrogallol oxidation (B, D and F) catalyzed by the Au-Hem conjugate (A and B, 0.2 μM hemin equivalent), Au-Hem-PMPT (C and D, 0.2 μM hemin equivalent) and Au-Hem-pNet (E and F, 0.2 μM hemin equivalent) in the presence of 0.4 mM H_2O_2 .

Fluorescence imaging of intracellular reactive oxygen species

In view of the high catalytic activity towards the substrate oxidation by different ROS (Fig. 8A), the cascade-assembled network, Au-Hem-Net, was used for the fluorescence imaging of ROS in living cells with *o*-phenylenediamine as the substrate. The cytotoxicity of Au-Hem-Net along with the substrate was examined using MTT assay (a colorimetric assay for assessing cell metabolic activity, see the ESI materials and reagents, Experimental section†) by incubating HeLa cells with the

Table 1 Kinetic parameters for the pyrogallol oxidation catalyzed by enzyme mimics at room temperature

Mimics	K_m (mM)	k_{cat} (min^{-1})	Ref.
Au-Hem	0.40	419.6	This work
Au-Hem-PMPT	0.97	714.3	This work
Au-Hem-pNet	1.36	1054.5	This work
Hemin	87.1	2.4	52
Hemin-graphene	1.22	246	21
Hemin-hydrogel	—	19.9	20
Hemin-polymer	—	7.6	20
FeTMPyP ^a	—	83	53
FeTMPyP-graphene	0.96	545	21
FeTMPyP-antibody	8.6	680	53
HRP	0.81	1750	53

^a FeTMPyP: tetramethylpyridylporphyrin iron.



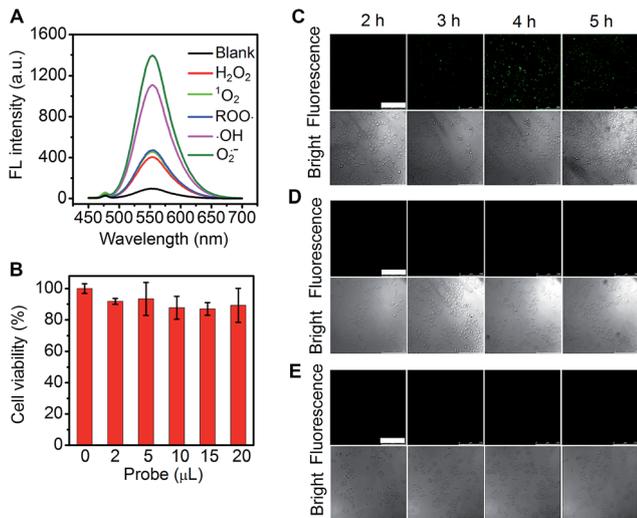


Fig. 8 (A) Fluorescence responses of *o*-phenylenediamine (0.14 mM) oxidized by different ROS for 15 min in the presence of Au-Hem-Net (10 nM equivalent hemin) at an excitation wavelength of 410 nm. (B) The viability of HeLa cells that were detected with MTT after treatment with 100 μL of culture medium containing *o*-phenylenediamine (0.14 mM) and different volumes of Au-Hem-Net (100 nM equivalent hemin) for 5 h at 37 °C. (C–E) Confocal fluorescence images of HeLa cells after incubation with *o*-phenylenediamine and Au-Hem-Net (C), *o*-phenylenediamine (D) and Au-Hem-Net (E) for different times at 37 °C. *o*-Phenylenediamine: 0.14 mM; Au-Hem-Net: 10 nM equivalent hemin. The scale bars represent 250 μm.

mixture, which maintained about 90% of the cell viability after 5 h of incubation (Fig. 8B). The confocal fluorescence imaging of intracellular ROS was performed by incubating the HeLa cells with Au-Hem-Net and *o*-phenylenediamine at 37 °C for different times. The fluorescence from the HeLa cells was noticeable after 2 h and increased with increasing incubation time, reaching a maximum fluorescence intensity at 4 h (Fig. 8C), suggesting the formation of a fluorescent product from the Au-Hem-Net catalyzed oxidation of *o*-phenylenediamine by intracellular ROS. In contrast, individual substrate or Au-Hem-Net treated HeLa cells did not show appreciable changes in the fluorescence images (Fig. 8D and E). Therefore, the cascade-assembled Au-Hem-Net possesses promising potential for application in the sensitive fluorescence imaging of intracellular ROS and/or other catalytic reaction-related biomolecules.

Conclusions

We present a highly catalytic network using axial coordination and coordination polymerization to achieve target-triggered cascade assembly based on the specific recognition of pyrimidinetrione moieties to the target. Two dual-functional pyrimidinetrione-containing derivatives of imidazole (MPT) and pyridine (PMPT) are designed to link the target Hg²⁺ and the Fe center of hemin that is covalently assembled on amino group modified AuNPs. The high catalytic activity of the network results from the triple amplification of the enzymatic activity during the cascade assembly: monomeric hemin loading on the

N-AuNPs, axial coordination of MPT or PMPT to the Fe center of hemin to form a five-coordinated hemin structure, and coordination polymerization *via* a 1 : 1 binding stoichiometry to form a “N–Hg^(II)–N” like binding motif for the ordered aggregation of the catalytic monomers. The catalytic activity of the network can be flexibly controlled by Hg²⁺, which leads to an extremely sensitive strategy for Hg²⁺ detection. The sub-attomolar detection limit is 7 orders of magnitude lower than those of conventional analytical methods. This catalytic system shows low cytotoxicity and has been successfully utilized in sensitive fluorescence imaging of intracellular ROS. The proposed catalytic network showed excellent affinity and high catalytic kinetics that are comparable to those of natural HRP. The cascade synthesis of the target-controllable catalytic system opens up new perspectives for not only the assembly of metalloporphyrin in the discovery and design of enzyme mimics, but also the development of biosensing protocols.

Acknowledgements

This study was supported by the National Natural Science Foundation of China (21375060, 21635005, 21605082, 21675084).

Notes and references

- J. P. Collman, N. K. Devaraj, R. A. Decréau, Y. Yang, Y. L. Yan, W. Ebina, T. A. Eberspacher and C. E. Chidsey, *Science*, 2007, **315**, 1565–1568.
- A. M. Beauchemin, *Nat. Chem.*, 2013, **5**, 731–732.
- M. Fischlechner, Y. Schaerli, M. F. Mohamed, S. Patil, C. Abell and F. Hollfelder, *Nat. Chem.*, 2014, **6**, 791–796.
- S. Ogi, K. Sugiyasu, S. Manna, S. Samitsu and M. Takeuchi, *Nat. Chem.*, 2014, **6**, 188–195.
- T. C. Bruice, *Acc. Chem. Res.*, 1991, **24**, 243–249.
- W. Auwärter, D. Écija, F. Klappenberger and J. V. Barth, *Nat. Chem.*, 2015, **7**, 105–120.
- S. J. Lee, C. D. Malliakas, M. G. Kanatzidis, J. T. Hupp and S. T. Nguyen, *Adv. Mater.*, 2008, **20**, 3543–3549.
- Z. C. Wang, Z. Y. Li, C. J. Medforth and J. A. Shelnutt, *J. Am. Chem. Soc.*, 2007, **129**, 2440–2441.
- M. Shema-Mizrachi, G. M. Pavan, E. Levin, A. Danani and N. G. Lemcoff, *J. Am. Chem. Soc.*, 2011, **133**, 14359–14367.
- P. Travascio, Y. F. Li and D. Sen, *Chem. Biol.*, 1998, **5**, 505–517.
- I. Willner, B. Shlyahovskiy, M. Zayats and B. Willner, *Chem. Soc. Rev.*, 2008, **37**, 1153–1165.
- F. Bedioui, *Coord. Chem. Rev.*, 1995, **144**, 39–68.
- L. G. Marinescu and M. Bols, *Angew. Chem., Int. Ed.*, 2006, **45**, 4590–4593.
- T. Hasobe, H. Imahori, P. V. Kamat and S. Fukuzumi, *J. Am. Chem. Soc.*, 2003, **125**, 14962–14963.
- W. Hieringer, *J. Am. Chem. Soc.*, 2011, **133**, 6206–6222.
- Y. M. Wang, X. F. Wang, S. K. Ghosh and H. P. Lu, *J. Am. Chem. Soc.*, 2009, **1**, 1479–1487.
- L. Zhang, J. P. Lei, F. J. Ma, P. H. Ling, J. T. Liu and H. X. Ju, *Chem. Commun.*, 2015, **51**, 10831–10834.



- 18 Y. Zang, J. P. Lei, P. H. Ling and H. X. Ju, *Anal. Chem.*, 2015, **87**, 5430–5436.
- 19 L. Chen, Y. Yang and D. L. Jiang, *J. Am. Chem. Soc.*, 2010, **132**, 9138–9143.
- 20 Q. G. Wang, Z. M. Yang, X. Q. Zhang, X. D. Xiao, C. K. Chang and B. Xu, *Angew. Chem., Int. Ed.*, 2007, **46**, 4285–4289.
- 21 T. Xue, S. Jiang, Y. Q. Qu, Q. Su, R. Cheng, S. Dubin, C. Y. Chiu, R. Kaner, Y. Huang and X. F. Duan, *Angew. Chem., Int. Ed.*, 2012, **51**, 3822–3825.
- 22 S. Jiang, R. Cheng, X. Wang, T. Xue, Y. Liu, A. Nel, Y. Huang and X. F. Duan, *Nat. Commun.*, 2013, **4**, 2225.
- 23 T. Xue, B. Peng, M. Xue, X. Zhong, C. Y. Chiu, S. Yang, Y. Q. Qu, L. Y. Ruan, S. Jiang, S. Dubin, R. B. Kaner, J. I. Zink, M. E. Meyerhoff, X. F. Duan and Y. Huang, *Nat. Commun.*, 2014, **5**, 3200.
- 24 O. K. Farha, A. M. Shultz, A. A. Sarjeant, S. T. Nguyen and J. T. Hupp, *J. Am. Chem. Soc.*, 2011, **133**, 5652–5655.
- 25 D. W. Feng, Z. Y. Gu, J. R. Li and H. C. Zhou, *Angew. Chem., Int. Ed.*, 2012, **51**, 10307–10310.
- 26 A. Fateeva, P. A. Chater, C. P. Ireland, A. A. Tahir, Y. Z. Khimyak, P. V. Wiper, J. R. Darwent and M. J. Rosseinsky, *Angew. Chem., Int. Ed.*, 2012, **51**, 7440–7444.
- 27 R. G. Cao, R. Thapa, H. Kim, X. D. Xu, M. G. Kim, Q. Li, N. Park, M. L. Liu and J. Cho, *Nat. Commun.*, 2013, **4**, 2076.
- 28 Y. H. Lin, L. Wu, Y. Y. Huang, J. S. Ren and X. G. Qu, *Chem. Sci.*, 2015, **6**, 1272–1276.
- 29 H. Li, Y. Y. Luk and M. Mrksich, *Langmuir*, 1999, **15**, 4957–4959.
- 30 T. Belser, M. Stçhr and A. Pfaltz, *J. Am. Chem. Soc.*, 2005, **127**, 8720–8731.
- 31 S. Debobrata, H. Partha and K. P. Tapan, *Angew. Chem., Int. Ed.*, 2013, **52**, 13314–13318.
- 32 E. M. Nolan and S. J. Lippard, *Chem. Rev.*, 2008, **108**, 3443–3480.
- 33 F. Zahir, S. J. Rizwi, S. K. Haq and R. H. Khan, *Environ. Toxicol. Pharmacol.*, 2005, **20**, 351–360.
- 34 C. H. Su, P. L. Wu and C. S. Yeh, *J. Phys. Chem. B*, 2003, **107**, 14240–14243.
- 35 K. K. Caswell, C. M. Bender and C. J. Murphy, *Nano Lett.*, 2003, **3**, 667–669.
- 36 E. S. Ryabova, *J. Biol. Inorg. Chem.*, 2004, **9**, 385–395.
- 37 Z. C. Chen, L. H. Wang and J. P. M. Schelvis, *Biochemistry*, 2003, **42**, 2542–2551.
- 38 F. A. Walker, M. W. Lo and M. T. Ree, *J. Am. Chem. Soc.*, 1976, **98**, 5552–5560.
- 39 J. P. Collman, T. Kodadek, S. A. Raybuck and B. Meunier, *Proc. Natl. Acad. Sci. U. S. A.*, 1983, **80**, 7039–7041.
- 40 Y. Miyake, H. Togashi, M. Tashiro, H. Yamaguchi, S. Oda, M. Kudo, Y. Tanaka, Y. Kondo, R. Sawa, T. Fujimoto, T. Machinami and A. Ono, *J. Am. Chem. Soc.*, 2006, **128**, 2172–2173.
- 41 X. J. Xue, F. Wang and X. G. Liu, *J. Am. Chem. Soc.*, 2008, **130**, 3244–3245.
- 42 J. N. Wang, L. Zhang, Q. Qi, S. H. Li and Y. B. Jiang, *Anal. Methods*, 2013, **5**, 608–611.
- 43 Q. B. Wang, N. Xu, Z. Gui, J. P. Lei, H. X. Ju and F. Yan, *Chem. Commun.*, 2014, **50**, 15362–15365.
- 44 N. L. Pacioni, M. González-Béjar, E. Alarcón, K. L. McGilvray and J. C. Scaiano, *J. Am. Chem. Soc.*, 2010, **132**, 6298–6299.
- 45 W. Yang, K. H. Liu, D. Song, Q. Du, R. N. Wang and H. M. Su, *J. Phys. Chem. C*, 2013, **117**, 27088–27095.
- 46 X. J. Yang, C. L. Fang, H. C. Mei, T. J. Chang, Z. H. Cao and D. H. Shangguan, *Chem.–Eur. J.*, 2011, **17**, 14475–14484.
- 47 J. S. Lee, M. S. Han and C. A. Mirkin, *Angew. Chem., Int. Ed.*, 2007, **46**, 4093–4096.
- 48 Y. K. Yang, K. J. Yook and J. S. Tae, *J. Am. Chem. Soc.*, 2005, **127**, 16760–16761.
- 49 S. Ando and K. Koide, *J. Am. Chem. Soc.*, 2011, **133**, 2556–2566.
- 50 R. Freeman, X. Q. Liu and I. Willner, *J. Am. Chem. Soc.*, 2011, **133**, 11597–11604.
- 51 P. W. Davidson, G. J. Myers, C. Cox, C. Axtell, C. Shamlaye, J. Sloane-Reeves, E. Cernichiari, L. Needham, A. Choi, Y. Wang, M. Berlin and T. W. Clarkson, *J. Am. Med. Assoc.*, 1998, **280**, 701–707.
- 52 Q. G. Wang, Z. M. Yang, M. L. Ma, C. K. Chang and B. Xu, *Chem.–Eur. J.*, 2008, **14**, 5073–5078.
- 53 H. Yamaguchi, K. Tsubouchi, K. Kawaguchi, E. Horita and A. Harada, *Chem.–Eur. J.*, 2004, **10**, 6179–6186.
- 54 Q. Q. Wang, S. Gonell, S. H. A. M. Leenders, M. Dürr, I. Ivanović-Burmazović and J. N. H. Reek, *Nat. Chem.*, 2016, **8**, 225–230.

