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A novel nanozyme based on selenopeptide-modified gold nanoparticles with a tunable glutathione peroxidase activity†

Dechen Zhang,^a Na Shen,^a Junrong Zhang,^a Jinming Zhu,^b Yi Guo^a and Li Xu^{id, a*}

In this study, we successfully prepared a selenium-containing pentapeptide (Sec-Arg-Gly-Asp-Cys)-modified gold nanozyme that exhibited glutathione peroxidase (GPx) activity. The nanozyme catalyzed the reduction of H_2O_2 by GSH, and its GPx activity was about 14 times that of free selenopeptide. Kinetic analysis indicated that the nanozyme changed the catalytic mechanism from the ping-pong mechanism of the peptide to an ordered mechanism. This indicated that the gold nanoparticle acts as a scaffold that may limit the mobility and constrain the conformation of the peptides, hence exposing the active center to the substrates, and allowing the multivalent selenopeptide to act cooperatively to increase the catalytic rate. Furthermore, upon addition of cysteamine to regulate the surface chemistry of gold nanoparticles and thereby modify the microenvironment of the active center, this nanozyme system could achieve a tunable GPx activity that was about 20 times that of free selenopeptide. Thus, the rational design of the nanozyme greatly improved and amplified the catalytic activity of the 'active unit' peptide. This study provides an alternative strategy to establish GPx mimics and provides new insights for the use of gold nanoparticles to develop nanozymes with biological applications.

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1. Introduction

Nanozymes are new artificial enzymes in the combined field of nanomaterials and enzymology research that aimed to design novel and efficient enzymatic mimics.^{1–3} Until now, a variety of nanomaterials have been used to mimic the activity of naturally occurring enzymes,^{4–8} such as nanocarbon materials,^{9–11} fullerene,¹² carbon nitride,¹³ Fe_3O_4 nanoparticles,¹⁴ mesoporous silica nanoparticles,¹⁵ dopamine nanoparticles,¹⁶ melanin nanoparticles,¹⁷ and metal–organic complexes.¹⁸ Noble metal-based nanomaterials have also shown significant advantages as enzyme mimetics;^{19,20} gold nanoparticles (AuNPs) possess excellent stability, good biocompatibility, ease of functionalization, and unique optical, and surface properties.^{21–24} The tailored surface functionalization of gold nanoparticles controls the interface between AuNPs and biomolecules,^{25–27} and it is crucial for development and design of tunable catalytic activities.^{28–30}

Glutathione peroxidase (GPx) is an important selenoenzyme in the cellular antioxidant defense system, which protects cells from oxidative damage by catalyzing the degradation of a variety

of hydroperoxides (ROOH) using glutathione (GSH) as the reductant.^{31–35} GPx contains an essential selenocysteine (Sec) residue at its active site,^{36–38} and this residue has clear catalytic characteristics.^{39,40} Considerable effort has been invested in constructing various GPx mimics,^{41,42} including small molecules,^{43,44} peptides,⁴⁵ cyclodextrin derivatives,^{46,47} abzymes,⁴⁸ bioimprinting proteins,^{49,50} and nanozymes.^{51,52} Small molecular mimics have the advantage of intrinsic structural stability, low immunogenicity, and facile membrane permeability. Ebselen is a well-known GPx mimic, which is a selenium-based organic complex with an anti-inflammatory, anti-oxidant and cytoprotective activity.^{53,54} However, its catalytic activity is much lower than that of macromolecule protein mimics,⁴¹ due to a weaker binding interaction of substrate. The improvement of the catalytic activity of small molecular mimic, while maintaining the above advantageous profile remains a challenge. We hypothesized that multivalent, functional systems anchored on a scaffold, and yet fully soluble, with limited mobility and conformational constraint could be a suitable system to act cooperatively in a catalytic process. Here, a self-assembled selenopeptide-anchored AuNP nanozyme as a GPx mimic was reported.

2. Experimental

Materials and reagents

Auric acid ($HAuCl_4 \cdot 4H_2O$) was purchased from Sinopharm Chemical reagent Co., Ltd. Trisodium citrate ($Na_3C_6H_5O_7$),

^aKey Laboratory for Molecular Enzymology and Engineering, The Ministry of Education, National Engineering Laboratory for AIDS Vaccine, School of Life Sciences, Jilin University, Changchun 130012, P. R. China. E-mail: xuli@jlu.edu.cn

^bChina-Japan Union Hospital of Jilin University, Changchun 130033, P. R. China

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cysteamine hydrochloride ($\text{HSCH}_2\text{CH}_2\text{NH}_2 \cdot \text{HCl}$), glutathione, glutathione reductase, and H_2O_2 were procured from Sigma-Aldrich. NADPH was purchased from Roche Applied Science. Pentapeptide Ser-Arg-Gly-Asp-Cys (SRGDC) and Sec-Arg-Gly-Asp-Cys (URGDC) were synthesized from GL Biochem (Shanghai) Ltd. Sample stock solution was prepared in Millipore (Milli-Q) water with a resistivity of 18.2 $\text{M}\Omega \text{ cm}$.

Apparatus

The kinetic measurements were recorded using a Shimadzu UV2450 spectrophotometer (Japan). TEM analysis was conducted on a JEOL JEM-1200EX model transmission electron microscope. The pH value of the buffer was determined with a Mettler Toledo 320 pH meter. Fourier transform infrared (FTIR) spectral studies were performed on a Nicolet 5700 spectrometer (Thermo Electron Corporation, USA) equipped with an infrared microscope. Spectra were acquired between 4000 and 400 cm^{-1} with a resolution of 2 cm^{-1} . All freeze-dried powder samples were tilted on the microstat for FTIR measurements.

Synthesis of AuNPs

AuNPs were prepared as described earlier.⁵⁵ Briefly, a 50 mL sample of aqueous HAuCl_4 (0.25 mM) was prepared in a 100 mL flask at room temperature. The solution was brought to a boil while being stirred, and the corresponding amount of aqueous sodium citrate with 3.5 : 1 molar ratio of citrate to Au^{3+} was added. The reaction was allowed to run until the solution reached a wine-red colour, which indicated completion of the reaction. These synthesized AuNPs were characterized using UV-vis absorption spectra and dynamic light scattering (DLS) with Zeta Potential Analyser (Brookhaven, USA) coupled to a BI-90Plus digital autocorrelator. According to Liu's method,⁵⁶ assuming a spherical shape and a uniform fcc structure, the extinction coefficient of the AuNP sample was determined according to the Lambert-Beer law, $A = \epsilon cl$. The AuNP concentration was determined using a Shimadzu UV2450 UV-visible spectrophotometer.

Preparation of nanozyme

Citrate-stabilized colloidal AuNPs were washed by centrifugation at 8000 rpm for 30 min, and re-dissolved in Millipore water. DLS analysis indicated that the hydrodynamic diameter of AuNPs was 20 nm, and the extinction coefficient was calculated to be $6.36 \times 10^8 \text{ L mol}^{-1} \text{ cm}^{-1}$. The concentration of AuNPs was adjusted to be 2 nM. For the surface functionalization of AuNPs, the different concentrations of URGDC were added dropwise to the AuNP solution, mixed by violent shaking, and then the homogenate was stored at 4 °C overnight. The resulting Au@SeH with the optimum AuNP : URGDC molar ratio and maximum enzyme activity was defined as MPC 1. Similar to the preparation of MPC 1, different concentrations of 2-aminoethanethiol was added to the MPC 1 system, mixed by violent shaking, and then the homogenate was stored at 4 °C overnight. The resulting Au@SeH/ NH_2 with the optimum

AuNP : cysteamine molar ratio and maximum enzyme activity was termed as MPC 2.

GPx activity assay

The GPx activity was assayed using H_2O_2 and GSH as substrates, as previously described.^{57,58} The reaction was carried out at 37 °C in 700 μL of solution containing 10 mM PBS (pH 7.0), 1 nM EDTA, 1 mM sodium azide, 1 mM GSH, 0.25 mM NADPH, 1 unit of glutathione reductase, and 1–50 μM samples. The mixture was pre-incubated for 5 min and then 0.25 mM NADPH solution was added and incubated for 3 min at 37 °C. The reaction was initiated by the addition of 0.5 mM H_2O_2 . The activity was determined based on the decrease in NADPH absorption at 340 nm. The nonenzymatic reaction affecting the measurement of the initial rate was subtracted to obtain the exact kinetic values.

One unit of activity was defined as the amount that utilizes 1 μmol of NADPH per minute. The activity was expressed in U μmol^{-1} .

Enzyme kinetics

The assay for determining enzyme kinetics was similar to that conducted for native GPx.^{59,60} To investigate the mechanism, assays were carried out under standard reaction conditions as described above by varying concentrations of GSH at a fixed concentration of H_2O_2 or *vice versa*. The initial rates were measured by observing the change of NADPH absorption at 340 nm. Data were analyzed using the Origin 8.0 software program. We determined the Kinetics constant by double-reciprocal plots (or Lineweaver-Burk plot) and fit to eqn (1) (ping-pong mechanism) or eqn (2) (ordered mechanism), where V is the reaction velocity, V_{\max} is the maximal velocity, $[\text{ROOH}]$ and $[\text{GSH}]$ are the concentrations of substrate H_2O_2 and GSH, K_S^{ROOH} is the dissociation constant of enzyme-ROOH, and K_m^{ROOH} and K_m^{GSH} are the Michaelis constants for H_2O_2 and GSH, respectively.⁶¹

$$\frac{1}{V} = \frac{K_m^{\text{ROOH}}}{V_{\max}} \frac{1}{[\text{ROOH}]} + \left(\frac{K_m^{\text{GSH}}}{V_{\max}} \frac{1}{[\text{GSH}]} + \frac{1}{V_{\max}} \right) \quad (1)$$

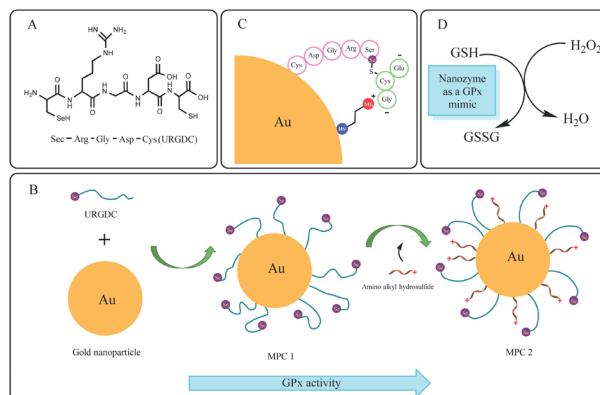
$$\frac{1}{V} = \left(\frac{K_m^{\text{ROOH}} K_m^{\text{GSH}}}{V_{\max}} \frac{1}{[\text{GSH}]} + \frac{K_m^{\text{ROOH}}}{V_{\max}} \right) \frac{1}{[\text{ROOH}]} + \left(\frac{K_m^{\text{GSH}}}{V_{\max}} \frac{1}{[\text{GSH}]} + \frac{1}{V_{\max}} \right) \quad (2)$$

3. Results and discussion

Design of nanozyme

Here, we propose a multivalent catalyst based on selenopeptide (URGDC) and cysteamine co-functionalized AuNPs, to construct a nanozyme with GPx-like activity (Scheme 1). This system provides unique biological functions, coupled with the original properties of AuNPs.





Scheme 1 Design of the nanozyme with GPx activity. (A) The structural formula of the URGDC with $-\text{SeH}$ at the N-terminus and $-\text{SH}$ at the C-terminus. (B) The construction of the nanozyme, with the GPx activity hypothesized to increase in order. This is a two-step process: first, URGDC was conjugated to gold nanoparticle (MPC 1); second, a mixed self-assembled monolayer was formed by further adding cationic amino alkyl hydrosulfide. (C) The schematic representation of the active site of MPC 2. Structural insights into the recognition and binding of the substrate GSH with MPC 2. The positively charged amino group directs the donor substrate GSH towards the catalytic center in such a way that its sulphydryl group must react with the selenium moiety. (D) The mechanism for nanozyme-catalyzed reduction of H_2O_2 by GSH.

Firstly, we designed a selenium-containing pentapeptide (Scheme 1A, URGDC, termed as 5SeP) with a delicate structure, in which the N-terminus was selenocysteine with $-\text{SeH}$ (catalytic group of GPx-like activity), and C-terminus was cysteine containing $-\text{SH}$ (binding group to AuNPs). Selenocysteine is the active site of natural GPx,⁶² which enables the pentapeptide GPx-like activity. The high affinity of thiol groups in the cysteine residue for gold atoms enables the pentapeptide binding activity through Au–S bond, resulting in the stable and oriented immobilization of peptides on the AuNPs.⁶³ The RGD motif is the recognition unit used by extracellular matrix proteins (vitronectin, fibrinogen, laminin, and collagen) to bind a variety of integrins that mediates cell attachment. It makes the selenopeptide more potent for identification to facilitate specific binding. Secondly, the selenopeptides were conjugated to AuNP through self-assembly to obtain one mimic (MPC 1). AuNP, as a scaffold, confines the mobility and conformation of randomly oriented URGDC to make most of the $-\text{SeH}$ groups exposed in a multivalent environment. Thus, GSH and H_2O_2 have ready access to the catalytic center, making the catalytic reaction more efficient. Thirdly, the nanozyme–substrate interface offers the opportunity to regulate enzyme activity. Cysteamine was employed to further decorate AuNPs by forming Au–S covalent bond (another mimic defined as MPC 2, Scheme 1B). It can adjust the distance and space structure of URGDC on the AuNP surface. Meanwhile, the electrostatic attraction between anionic GSH substrate and the positively charged monolayer facilitates its transport and binding to the active site (Scheme 1C). MPC 2 mimic provides an electrostatic architecture which, in each reductive step, directs the donor substrate GSH towards the

catalytic center in such a way that its sulphydryl group must react with the selenium moiety. Consequently, the MPC 2 mimic shows a stronger catalysis activity. The catalytic mechanism of the nanozyme is shown in Scheme 1D.

Characterization of the nanozyme

The synthesized AuNPs were investigated by TEM. It is apparent that the AuNPs are monodisperse, spherical in shape, and a narrow particle-size distribution with a mean size of about 20 nm (Fig. 1A). The characteristic surface plasmon resonance (SPR) band at 518 nm is shown in Fig. 1E. The calculated extinction coefficient of AuNP was $6.389 \times 10^8 \text{ L mol}^{-1} \text{ cm}^{-1}$.⁵⁵

The molecular simulation of URGDC was obtained by material studio (Fig. 1B), and the calculated cross-sectional diameter was 4 Å. Thus, it was calculated that about 8000 URGDC molecules are needed to completely cover the surface of one 20 nm AuNP.⁶⁴ Given that, the specific activities of Au@SeH with different Au : URGDC molar ratio were tested. It was found that Au@SeH with 1 : 1500 molar ratio of Au : URGDC (MPC 1) showed the maximum GPx activity of $5.05 \text{ U } \mu\text{mol}^{-1}$ (Fig. 1C, Table 1). Thus, the molar ratio of Au : URGDC was fixed to be 1 : 1500, and then different proportions of cysteamine were added to obtain Au@SeH/NH₂. The tunable specific activity of Au@SeH/NH₂ was measured, and we found that Au@SeH/NH₂

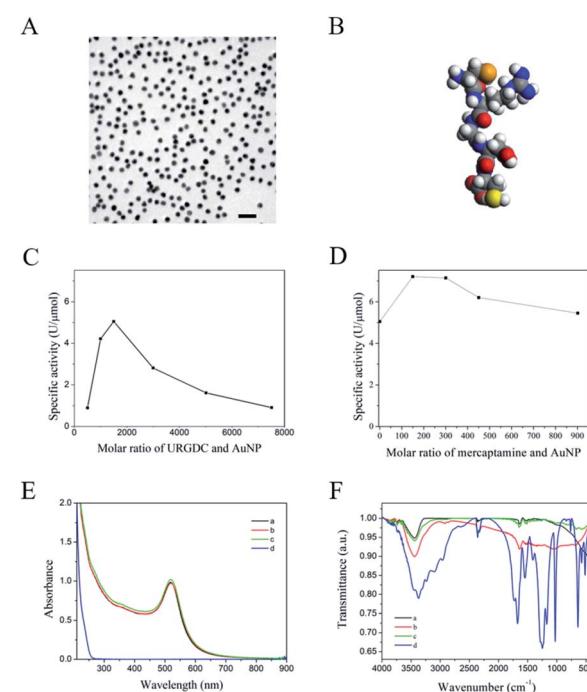


Fig. 1 Physical and enzyme activity characterization of nanozyme. (A) TEM image of gold nanoparticles. Scale bar is 50 nm. (B) The simulation structure of URGDC. Three-dimensional homology models of the peptides were generated, visualized and investigated with material studio. (C and D) The specific activity of different Au@SeH with various molar ratio of URGDC and AuNP (C) and Au@SeH/NH₂ with different proportion of cysteamine (D). (E) UV-vis spectra of AuNP (a), MPC 1 (b), MPC 2 (c) and URGDC (d) in 10 mM PBS (pH 7.0). (F) FTIR spectrum AuNP (a), MPC 1 (b), MPC 2 (c) and URGDC (d) at room temperature.



Table 1 The GPx activities of some mimics and the native GPx from rabbit liver

Catalyst	Substrate	Activity ^a (U μmol^{-1})
Ebselen ⁵⁴	H_2O_2 , GSH	0.99
2-SeCD ⁶⁷	H_2O_2 , GSH	7.40
SRGDC	H_2O_2 , GSH	0
URGDC	H_2O_2 , GSH	0.36
MPC 1	H_2O_2 , GSH	5.05
MPC 2	H_2O_2 , GSH	7.21
Native GPx ⁴¹	H_2O_2 , GSH	5780

^a The GPx activities were measured by a coupled enzymatic assay (as described) in sodium phosphate buffer (10 mM, pH 7.0) at 37 °C.

with 1 : 150 molar ratio of Au : cysteamine (MPC 2) showed the maximum GPx activity of 7.21 U μmol^{-1} (Fig. 1D, Table 1). Fig. 1E shows the UV-vis absorption spectra of MPC 1 and MPC 2. The 5SeP and cysteamine modification did not change the optical properties, since the shoulder band towards the longer wavelengths in the range of 600–700 nm did not appear, except that the peak absorbance of MPC 2 was slightly higher. This

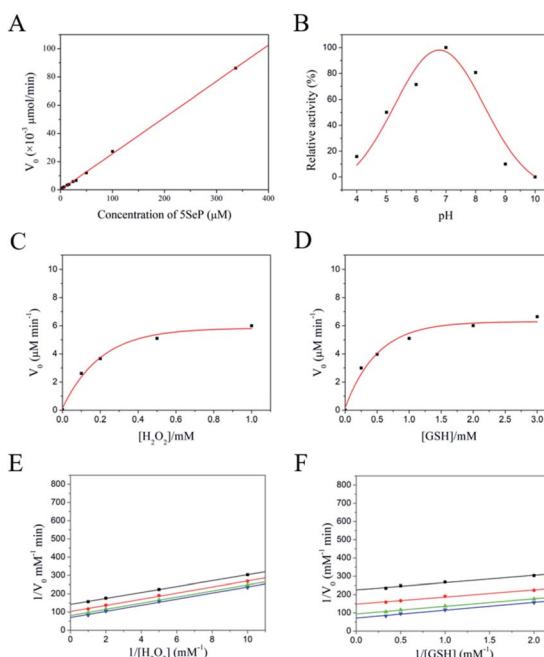


Fig. 2 Enzymatic reactive characteristics of 5SeP. (A and B) The influence of 5SeP concentration on initial velocity rate (A) and reactive system pH (B). The initial velocity rate was determined by changing the concentration of 5SeP between 0 and 400 μM , and pH from 4 to 10, respectively. (C and D) The effect of substrate H_2O_2 (C) or GSH (D) concentration on initial velocity rate for 5SeP. The absorbance of enzyme reactive system at 340 nm was scanned by varying H_2O_2 concentration at 1 mM GSH (C), and vice versa, varying GSH concentration at 0.5 mM H_2O_2 (D). (E and F) The double-reciprocal plots (or Lineweaver–Burk plot) of initial rate vs. H_2O_2 or GSH concentration for URGDC. $1/V_0$ versus $1/[\text{H}_2\text{O}_2]$ (L mmol^{-1}) of URGDC at $[\text{GSH}] = 0.5$ (■), 1 (●), 2 (▲) and 3 mmol L^{-1} (▼) (E). $1/V_0$ versus $1/[\text{GSH}]$ (L mmol^{-1}) of URGDC at $[\text{H}_2\text{O}_2] = 0.1$ (■), 0.2 (●), 0.5 (▲) and 1 mmol L^{-1} (▼) (F).

suggested that the AuNPs did not aggregate, and they are stable in the nanzyme system. It was also confirmed by TEM images (ESI Fig. S1†), there is no obvious change in the morphology of nanzyme. As shown in Fig. 1F, 5SeP exhibited S–H bond stretching vibration falls around 2357 cm^{-1} for free cysteine, and amide I and II regions showed two peaks located at 1674 and 1635 cm^{-1} , respectively. In the N–H stretching region, 5SeP showed a broader band located at 3373 cm^{-1} . Upon interaction with the surface of AuNP, the whole spectrum of 5SeP was strongly affected: amide I and II bands were weakened, and S–Au vibration appeared at 1644 cm^{-1} .⁶⁵ Furthermore, the N–H stretching band, which in in the free peptide located around 3373 cm^{-1} , became sharp and without any shoulder, probably forbidden by the surface selection rule.⁶⁶ This result illustrated

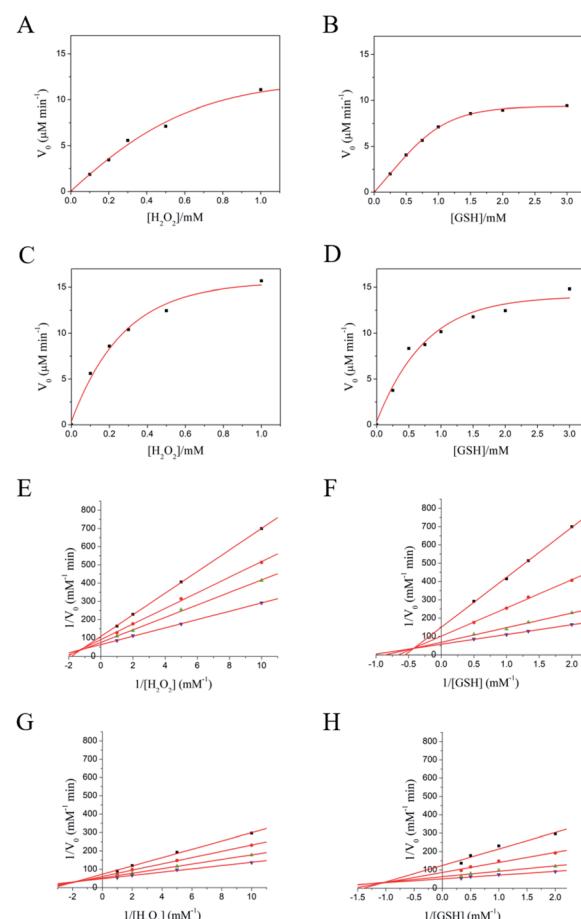


Fig. 3 Enzyme kinetics of nanzyme. (A and C) The influence on initial velocity rate of substrate H_2O_2 concentration for MPC 1 (A) and MPC 2 (C). The initial velocity rate was determined by varying H_2O_2 concentration at 1 mM GSH. (B and D) The effect on initial velocity rate of substrate GSH concentration for MPC 1 (B) and MPC 2 (D). The initial velocity rate was determined by varying GSH concentration at 0.5 mM H_2O_2 . (E–H) The double-reciprocal plots (or Lineweaver–Burk plot) of initial rate vs. H_2O_2 and GSH concentration for MPC 1 and MPC 2. (E and G) $1/V_0$ versus $1/[\text{H}_2\text{O}_2]$ (L mmol^{-1}) of MPC 1 and MPC 2 at $[\text{GSH}] = 0.5$ (■), 1 (●), 2 (▲) and 3 mmol L^{-1} (▼). (F and H) $1/V_0$ versus $1/[\text{GSH}]$ (L mmol^{-1}) of MPC 1 and MPC 2 at $[\text{H}_2\text{O}_2] = 0.1$ (■), 0.2 (●), 0.5 (▲) and 1 mmol L^{-1} (▼).



Table 2 Kinetic parameters of GPx mimic-catalyzed reduction of H_2O_2 by GSH

GPx mimic	K_{cat} (min $^{-1}$)	K_m^{GSH} (mM)	$K_m^{H_2O_2}$ (mM)	K_{cat}/K_m^{GSH} (M $^{-1}$ min $^{-1}$)	$K_{cat}/K_m^{H_2O_2}$ (M $^{-1}$ min $^{-1}$)
URGDC	0.85	0.72	0.28	1.18×10^3	3.04×10^3
MPC 1	14.83	0.64	0.22	2.32×10^4	6.90×10^4
MPC 2	15.65	0.32	0.18	4.97×10^4	8.89×10^4

the conjugation of 5SeP and/or cysteamine over the surface of AuNP.

GPx activity and kinetics of 5SeP

To study the GPx activity of 5SeP, we monitored the initial velocity under several concentrations of 5SeP (Fig. 2A). As a result, a significant liner correlation was observed between them. Analysis of the data revealed that the GPx activity of 5SeP was 0.36 U μmol^{-1} (Table 1). The SRGDC without -SeH catalytic group did not show GPx activity. Compared with nanzyme system, the GPx activity of MPC 2 was about 20 times that of free selenopeptide and 1.43 times that of MPC 1. Therefore, MPC 2 displayed a higher activity. In a multivalent environment, AuNP scaffold could efficiently facilitate the conjugation between substrate and multi active center. In addition, an electrostatic architecture (cysteamine) improved the reaction process in a more favourable action. For the other small molecular artificial GPx, the GPx activity is close to the same level of 2-SeCD mimic in the absent of hydrophobic environment for binding substrate.⁶⁷ It means that the multi active center system is more efficient for the artificial enzyme catalytic process.

Similar to the crude GPx, the catalytic activity of 5SeP was dependent on pH, GSH, and H_2O_2 concentration. We measured the GPx-like activity of 5SeP while varying the pH from 4 to 10, the H_2O_2 concentration from 0.1 to 1 mM, and the GSH concentration from 0.25 to 3 mM. The optimum pH was approximately 7.0 (Fig. 2B). The steady-state rate of catalysis increased towards a saturation plateau with increasing loads of substrate H_2O_2 or GSH, which was similar to the enzyme catalyzed reaction (Fig. 2C and D). Thus, it was confirmed that 5SeP had GPx-like activity and enzymatic reaction properties.

To investigate the mechanism of GPx activity of 5SeP, we determined the apparent steady-state kinetic parameters for the reaction. The double-reciprocal plots were obtained (Fig. 2E and F), and the parallel lines indicated that the GPx-like reaction of 5SeP comply with a ping-pong mechanism.⁶⁸

Kinetics study of the nanzyme

Next, we assessed the first-order kinetics to systematically study the property of nanzyme catalysis. As a result, a kinetic profile of the reaction towards saturation plateau was observed by varying the initial concentration of substrate H_2O_2 or GSH (Fig. 3A–D). Thus, it was revealed that both MPC 1 and MPC 2 possessed GPx-like activity and enzymatic reaction properties. To further investigate the mechanism of GPx activity of the nanzyme, we determined the apparent steady-state kinetic parameters for the reaction. The double-reciprocal plots were

obtained (Fig. 3E–H), and the crossed lines indicated that the GPx-like reaction of MPC 1 and MPC 2 abides by an ordered mechanism.⁶⁹

The kinetic parameters for URGDC, MPC 1, and MPC 2 are listed in Table 2. Their first order constants (K_m) for H_2O_2 diminished in order, and the formal second-order rate constant K_{cat}/K_m of MPC 1 and MPC 2 for H_2O_2 was 22.7 and 29.2 times higher than that determined under identical conditions for free 5SeP, respectively. The K_{cat}/K_m of MPC 1 and MPC 2 for GSH was 20.0 and 42.1 times higher than that determined under identical conditions for free 5SeP, respectively; this corresponds to the 'active unit' on AuNP surface. Thus, by adjusting with cysteamine, the MPC 2 could show more efficient catalysis activity.

Among the three mimics, the K_m value of both substrates showed a common profile, with $K_m^{GSH} > K_m^{H_2O_2}$ this demonstrated that H_2O_2 was a priority substrate, likely because H_2O_2 molecule is smaller than the GSH molecule, making it easier for it to approach the active site of the enzyme. However, while the access of H_2O_2 to the active site is not specific, the GSH shows specific/selective recognition; this is similar to the characteristics of natural GPx.

4. Conclusions

In conclusion, a selenopeptide functionalized AuNP was rationally designed in this study. The nanzyme-catalyzed reaction exhibited saturation kinetics: the reactive rate data complied with the Michaelis-Menten model of enzyme kinetics. The GPx activity of MPC 2 was 7.21 U μmol^{-1} , which is about 20 times of that of free selenopeptide. The kinetics analysis indicated that the catalytic mechanism of free peptide was the ping-pong mechanism, while the nanzymes (MPC 1, MPC 2) showed an ordered mechanism. Furthermore, the catalytic constant K_{cat}/K_m of both GSH and H_2O_2 were one order of magnitude higher than that of free selenopeptide. Thus, improved properties and synergistic activities could be achieved from an interface-regulating approach, as evident from the measured GPx activity and catalytic characteristics. We obtained not only a peptide-capped nanoparticle (MPC 1) with synergistic enzyme activity, compared with the 'active unit', but also a more efficient MPC 2 nanzyme by cysteamine adjustment. Thus, our study successfully demonstrates that AuNPs provide advantages of easily obtainable self-assembled monolayer and specific surface/interface chemistry, displaying a promising platform for improving and establishing delicate and efficient nanzyme catalytic system.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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