

Cite this: *RSC Adv.*, 2019, 9, 28814

One-pot synthesis of biomimetic glutathione peroxidase with temperature responsive catalytic behaviors

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Excessive reactive oxygen free radicals (ROS) are the main cause of various oxidative diseases. It is of great significance to develop antioxidant drugs that can intelligently regulate free radical concentrations. The biomimetic simulation of glutathione peroxidase (GPx) can provide an important theoretical basis for the development of antioxidant drugs. In order to explore a simple and efficient strategy for constructing biomimetic GPx, a microgel biomimetic GPx (PNTegel) with temperature responsive catalytic activity was prepared by a one-pot synthesis method. The PNTegel, with typical enzymatic catalytic characteristics, exhibited a maximum catalytic activity at 37 °C ($v_0 = 11.51 \text{ mM min}^{-1}$). The investigation of the catalytic mechanism of PNTegel suggested that the binding of different hydrophobic substrates to PNTegel was altered by the change of hydrophobicity of poly(*N*-isopropylacrylamide) (PNIPAM) in the microgel scaffold of PNTegel during the temperature response process. The change of hydrophobicity was the main factor for regulating the catalytic activity of PNTegel, which resulted in a temperature responsive catalytic behavior of PNTegel. This new strategy for the simple and efficient construction of biomimetic GPx by a one-pot method provides important theoretical support for exploring the preparation of highly effective antioxidant drugs.

Received 25th July 2019
Accepted 30th August 2019

DOI: 10.1039/c9ra05775g

rsc.li/rsc-advances

Introduction

Glutathione peroxidase (GPx) is an important selenoenzyme in the human body, and was first discovered by Mills *et al.*¹ Then, Epp *et al.* reported its crystal structure for the first time.² Commonly, GPx decomposes excess oxygen free radicals (ROS) in the human body using glutathione as a reducing agent and plays an important role in preventing various diseases such as cardiac cataracts, cardiovascular and cerebrovascular diseases, inflammation and Keshan disease.^{2–5} In order to provide a further theoretical basis for the development of antioxidant drugs, it is of great significance to study the catalytic mechanism of GPx and construct an effective biomimetic GPx.^{4,6–19} It is proved that selenocysteine in natural GPx is the catalytic center and plays an important role in maintaining high GPx catalytic activity.^{6,7,20,21} Through the biomimetic simulation of GPx catalytic center in natural GPx, researchers have prepared a series of biomimetic GPx. Generally, the scaffold materials for preparing biomimetic GPx mainly include small organic molecules,^{12,13,20–22} imprinted proteins,^{8,11} genetically engineered proteins,^{14,15,23}

supramolecular polymers,^{17,19,24,25} block polymers,^{26–29} microgels,^{30–32} and organic–inorganic hybrid materials.^{33–36} Compared with natural GPx, these biomimetic GPx were endowed with obvious advantages, such as high preparation yield, strong structural stability and good industrialization prospect. Among the scaffold materials of these biomimetic GPx, the microgel scaffold exhibits unique characteristics. Due to the three-dimensional structure of microgel is similar to that of natural GPx, the microgel is considered to be an efficient scaffold to construct biomimetic GPx and simulate the natural GPx catalytic behavior. Therefore, the construction of microgel-based biomimetic GPx is expecting, and our research team has carried out some exploration in this research field.^{30–32} Generally, the supramolecular microgel biomimetic GPxs were efficiently constructed by a host–guest supramolecular self-assembly, in which the host–guest molecules were modified to gain the catalytic function similar to the GPx. These biomimetic GPxs could successfully simulate the natural GPx by exhibiting a typical enzymatic catalytic behavior.

ROS are the by-products of normal human metabolism. Although a certain concentration of ROS plays an important role in normal physiological functions, the excessive free radicals can cause oxidative damage.^{5,37} Generally, natural GPx remove excess ROS and regulate the free radicals at an appropriate concentration to protect the normal physiological functions of the human body. Therefore, the development of

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biomimetic GPx that could intelligently regulate the concentration of ROS by changing its catalytic activity is desirable. It is of great significance for the development of intelligent antioxidant drugs. Using different stimuli-responsive materials, researchers have developed various smart biomimetic GPx with different responsive catalytic activity, such as temperature responsive,^{27–32,38–40} pH responsive,⁴¹ photo responsive,^{41–43} ATP responsive¹⁹ and so on. In particular, the construction of temperature responsive biomimetic GPx and the exploration of temperature responsive catalytic mechanisms have been systematically studied. For example, researchers prepared a series of temperature responsive biomimetic GPx based on block polymers,^{28,44} graft polymers,²⁹ microgels,³⁰ supramolecular microgels,^{31,32,38} *etc.* The construction of these biomimetic GPx not only efficiently expanded the new preparation strategy of biomimetic GPx but also intelligently regulated the catalytic activity of biomimetic GPx, which provided the basement for the development of antioxidant drugs. In our group, a series of temperature responsive biomimetic GPx based on microgel scaffold were constructed by the supramolecular self-assembly of temperature responsive materials. This strategy was environmentally friendly, while the biomimetic GPx were self-assembled only under a certain concentration range and special conditions.^{31,32,38,44} Moreover, the preparation process for supramolecular building blocks was complicated, which obstructed the large-scale industrial application. Therefore, exploring a new preparation strategy to simply and efficiently construct temperature responsive biomimetic GPx is still an important research work.

ATRP is one of the effective methods for preparing various structurally controllable polymers.^{45–47} In addition, the one-pot method presents a unique advantage in the simple and efficient preparation of functional materials.^{48–50} Inspired by these, a new strategy based on the combination of ATRP and one-pot method is worthwhile to be explored for the biomimetic GPx with high efficiency temperature response catalytic activity. In this work, a temperature responsive microgel biomimetic GPx was successfully constructed by a one-pot method using *N*-isopropylacrylamide, tellurium-containing polymerizable monomer (**Te monomer**) and acrylamide. These monomers provide the microgel with temperature response function, catalytic function and hydrophilic segment backbone, respectively. This work provides a new strategy and an important theoretical basis for the development of highly effective antioxidant drugs.

Experimental section

Materials

Sodium borohydride and 3-bromo-1-propanol were purchased from Sigma-Aldrich. Acrylic chloride and tellurium powder were purchased from Shanghai Reagent Company. Acrylic chloride was purchased from Anhui Wotu Co., Ltd. and used after being purified by vacuum distillation. *N,N'*-Methylene diacrylamide (MBA), acrylamide (AM), hexadecyl trimethyl ammonium bromide (CTAB), 5,5'-dithio-bis(2-nitrobenzoic acid), 4-nitrothiophenol (NBT), cumyl hydroperoxide, and hydrogen peroxide were used as received from J&K Scientific Ltd. *N*-

Isopropylacrylamide (NIPAM) was purchased from J&K Scientific Ltd., and used after cyclohexane/methanol recrystallization and drying. 2-Bromopropionyl bromide was purchased from Lancaster. Tris(2-dimethylaminoethyl)amine (ME₆TREN) was prepared according to the previously reported methods.⁵¹ Triethylamine and tetrahydrofuran were purchased from Nanning Lantian Reagent Co., Ltd., and used after rigorously dried with sodium. 3-Carboxyl-4-nitrobenzenethiol (TNB) was synthesized from 5,5'-dithiobis(2-nitrobenzoic acid) using the previously reported methods.²⁰

Instrumentations

The ¹H NMR spectra for the compounds and microgel were recorded on the Bruker 300 MHz spectrometer using a TMS proton signal as the internal standard. UV-vis spectra were obtained using a Shimadzu 2450 UV-vis-NIR spectrophotometer. Scanning electron microscopy (SEM) observations were carried out on a JEOL JSM-6700F scanning electron microscope. Transmission electron microscopy (TEM) observations were carried out on a JEOL JEM 3010 transmission electron microscope. The buffer pH values were determined with a METTLER TOLEDO 320 pH meter. Dynamic light scattering experiments were performed at Malvern-ZetaSizer Nano ZS instrument.

Synthesis of polymerizable small molecules (Te monomer)

The polymerizable small molecules (**Te monomer**) that contain catalytic centers were synthesized according to the reported literature.⁵² The NMR characterization data were as follows: ¹H NMR (300 MHz, D₂O) δ (ppm) 6.45–5.83 (6H, CH₂=CH–), 4.21 (t, 4H, COOCH₂), 3.31 (t, 4H, –CH₂Te), 2.09 (m, 4H, –CH₂–).

Preparation of PNTegel by one-pot method

The preparation of the microgel scaffold for **PNTegel** was carried out according to the previously reported literature.⁴⁷ The procedure was as follows: a magnetic stir bar, NIPAM (30 mmol), MBA (5 mmol), **Te monomer** (1 mmol), ME₆TREN (0.2 mmol), water (450 mL) and DMF (30 mL) were added to the reaction flask. The oxygen in the reaction system was removed by continuously purging nitrogen gas for 30 min. Then, CuBr (0.2 mmol) was added into the flask under the protection of nitrogen gas flow, followed by stirring for 40 minutes to form a complex of catalyst and ligand. The polymerization was then initiated by adding an initiator, benzyl 2-bromopropanoate (0.05 mmol, dissolved in 20 mL of DMF).²⁷ After reacting for 8 hours at 30 °C, the reaction flask was opened under a nitrogen stream and CTAB (0.06 mmol) was added into the flask. Subsequently, the temperature of the reaction system was raised to 70 °C, and the reaction was continued for 5 hours. Further, an aqueous solution (100 mL) of AM (30 mmol) was added to the reaction system under a nitrogen stream, and the reaction was continued for 24 hours at 70 °C. After that, the polymerization was terminated by the treatment of liquid nitrogen, and then the reaction solution was concentrated on a rotary evaporator. The resultant solution was dialyzed for five days to remove impurities. The target product (**PNTegel**) with the yield of 64.50% was obtained after lyophilization. The target



product was stored in refrigerator. The content of tellurium in the catalytic center of **PNTegel** was determined to be $1.49 \times 10^{-4} \text{ mmol mg}^{-1}$ according to the reported method.³⁸

Determination of the LCST for **PNTegel**

The optical transmissions (wavelength at 600 nm) of microgel polymer solution (**PNTegel**, 1 mg mL^{-1}) were measured at different temperatures using Shimadzu 2450 UV-vis-NIR spectrophotometer, which was employed to determine the LCST of **PNTegel**. All test samples were thermostated in a circulator bath with different temperatures prior to measurements. The LCST is defined as the temperature at which the change in transmittance of the polymer solution just occurs at the inflection point.

Determination of GPx activity

The catalytic activity of **PNTegel** was determined by the method that established by professor Hilvert with a slight modification.⁶ The enzyme catalytic reaction was performed in a 1 mL quartz cuvette. Typically, 700 μL of phosphate buffer solution ($\text{pH} = 7.0$), 100 μL of **PNTegel** solution ($10 \mu\text{M}$), and 100 μL of TNB solution (1.5 mM) were sequentially added to the quartz cuvette. After keeping at a fixed temperature for three minutes, 100 μL of cumene hydroperoxide (CUOOH) (2.5 mM) was added to the quartz cuvette to trigger the enzyme-catalyzed reaction. The measurement of the ultraviolet absorption at 410 nm for the aforementioned system was performed on Shimadzu 2450 UV-vis-NIR spectrophotometer to calculate the reaction rate. Simultaneously, the ultraviolet absorption of the system without **PNTegel** was also measured to determine the reaction rate ($\epsilon_{410} = 13\,600 \text{ M}^{-1} \text{ cm}^{-1}$, $\text{pH} = 7.0$). The catalytic rate of **PNTegel** was calculated by subtracting the reaction rate of the system that without **PNTegel** from the reaction rate of the system containing **PNTegel**.

Determination of repeatability of activity

The measurement of the repeatability was carried out when the temperature was converted repeatedly between 25 and 37 $^{\circ}\text{C}$. Firstly, the stock solution of **PNTegel** was maintained at 25 $^{\circ}\text{C}$, and the catalytic rate was determined at 25 $^{\circ}\text{C}$ according to the method mentioned above. Secondly, the temperature of the **PNTegel** stock solution was increased to 37 $^{\circ}\text{C}$, and the catalytic rate was also determined at 37 $^{\circ}\text{C}$. Thirdly, the **PNTegel** stock solution was then decreased to 25 $^{\circ}\text{C}$ for determining the catalytic rate at 25 $^{\circ}\text{C}$ using the identical method. The catalytic rates of the **PNTegel** stock solution were determined by converting the temperature repeatedly between 25 and 37 $^{\circ}\text{C}$ for several times.

Results and discussion

Construction of **PNTegel** based on one-pot method

Due to the important antioxidant physiological effects of glutathione peroxidase (GPx), Epp *et al.* first studied the crystal structure of GPx, indicating that the selenocysteine is the catalytic center of GPx.² In addition, by constructing biomimetic GPx with similar catalytic functions to the natural GPx,

researchers have further proved that the hydrophobic cavity consisted by hydrophobic amino acid residues also plays an important role in maintaining the high catalytic activity of natural GPx.^{28,44} Therefore, based on these researches, in this work, we designed a polymerizable monomer **Te monomer** as an alternative of the catalytic center of natural GPx (as shown in Fig. 1). Since PNIPAM is a polymer with temperature responsive property, the hydrophobic microenvironment formed by the hydrophobic polymer chain of PNIPAM in the temperature response process was employed to simulate the hydrophobic cavity of natural GPx. In addition, NIPAM and AM were the main functional monomers for constructing the microgel scaffold of biomimetic GPx. TNB and NBT were used as the thiophenol substrates, and CUOOH was employed as the peroxide substrate. These substrates were employed to determine the GPx catalytic activity of biomimetic GPx. MBA was a cross-linking molecule for preparing microgel. Except for the catalytic center, the **Te monomer** also acted as a cross-linking agent. A schematic diagram of the preparation of microgel scaffold for **PNTegel** by one-pot method is shown in Scheme 1.

As shown in Scheme 1, the preparation process of **PNTegel** by one-pot method was mainly divided into three stages. First, PNIPAM was synthesized by using ATRP, wherein the monomers of MBA and **Te monomer** were copolymerized in this process. Therefore, the catalytic center was anchored into the polymer scaffold of PNIPAM. Then, the temperature of the reaction system was raised to 70 $^{\circ}\text{C}$, and the reaction was carried out for about 5 hours. The solution of reaction system became turbid due to the poor solubility of PNIPAM polymer in the aqueous solution with 70 $^{\circ}\text{C}$. Under this condition, CTAB, a surfactant, was introduced into the reaction system to emulsify the reaction solution, which caused the reaction to be carried out mainly in the microemulsion. This therefore facilitated the formation of the main scaffold structure of the microgel. After a reaction of 5 hours, the monomer AM was added to the system. The polymerization reaction was continued for another 24 hours at 70 $^{\circ}\text{C}$, which fully anchored the polyacrylamide (PAM) into the scaffold of **PNTegel** and formed a shell layer of microgel. After that, the polymerization reaction was terminated by liquid nitrogen. After concentrating in a rotary evaporator, the resultant solution was dialyzed against a dialysis bag (molecular weight cutoff of 100 000), and freeze-dried to obtain the microgel material of **PNTegel**. In order to verify the successful preparation of microgels, the

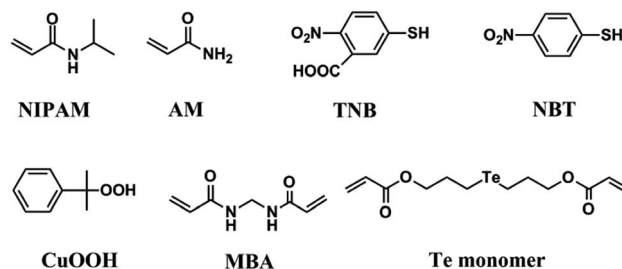
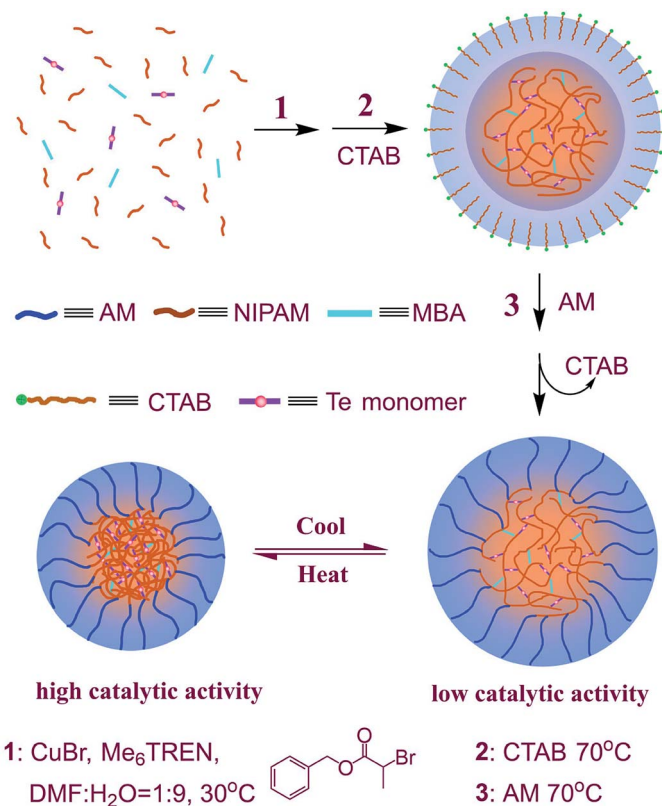


Fig. 1 The structures of the functional monomers NIPAM, AM, **Te monomer**, CUOOH, MBA and substrates (NBT, TNB).





Scheme 1 The graphical representation of the one-pot preparation process and reversible responsive behaviour of PNTegel.

microgel polymer scaffold of **PNTegel** was characterized by NMR, SEM, TEM, DLS and so on.

Firstly, as shown in Fig. 2, the structure of **PNTegel** was characterized by NMR. The proton signal of d was the hydrogen

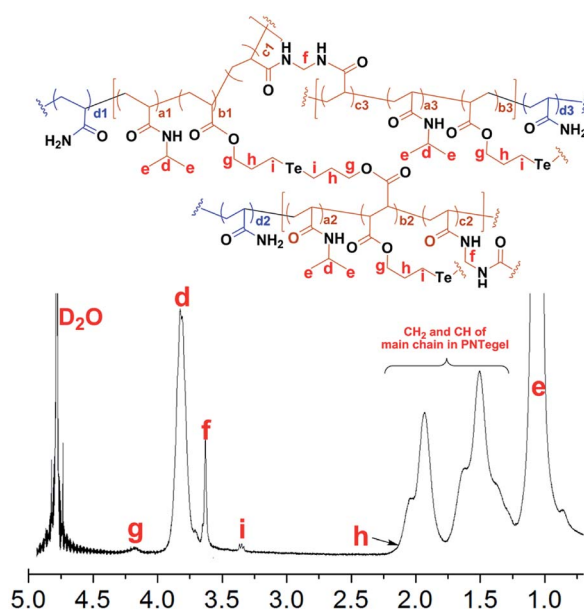


Fig. 2 ¹H NMR spectrum of PNTegel in D₂O.

signal peak of methine in PNIPAM. The signal peaks of chemical shift between 0.8 and 2.2 ppm were mainly attributed to the hydrogen connected with the main carbon chain of microgel and the methyl hydrogen e of PNIPAM. The proton signal of f was ascribed to the methylene hydrogen of MBA, and g was the hydrogen of methylene group attached to the ester group in **Te monomer**. The hydrogen signal peak of i was attributed to the hydrogen of the methylene group attached to the tellurium in **Te monomer**. And the hydrogen signal peak of the other methylene group of **Te monomer** was masked by the signal of the hydrogen in the polymer main chain. These results demonstrated that **Te monomer**, the catalytic center, was successfully introduced into the microgel scaffold.

Furthermore, the morphology of **PNTegel** was further characterized by SEM and TEM. As shown in Fig. 3, the microgel basically exhibited a spherical structure with the diameter of about 200 nm, indicating the successful preparation of a uniform microgel material by one-pot method. In addition, the average hydrodynamic diameter of the **PNTegel** microgel was also characterized by a dynamic light scattering (see Fig. 4a). It suggested that the average hydrodynamic diameter of **PNTegel** was about 250 nm at 25 °C. This result was slightly larger than the observation of SEM and TEM, which might be due to the swelling of the **PNTegel** during the measurement of the hydrodynamic diameter. Similar results have been reported by previous works.^{32,38} The characterization of the hydrodynamic diameter further suggested the successful preparation of microgel scaffold of biomimetic GPx materials.

Finally, the phase transition of PNIPAM in **PNTegel** was characterized by the change of optical transmittance of **PNTegel** solution (Fig. 5). The optical transmittance of **PNTegel** solution was higher when the temperature was lower than 33 °C, while the optical transmittance of **PNTegel** solution decreased as the increase of temperature and tended to be stable when the temperature was higher than 38 °C. The profile of the optical transmittance as a function of temperature indicated that the LCST of **PNTegel** was about 34.5 °C. This result suggested that the polymer chain of PNIPAM could be turned into a hydrophobic structure as the increase of temperature, which provided a hydrophobic microenvironment in **PNTegel**. This hydrophobic structure might play a significant role in simulating the hydrophobic cavity of natural GPx. The investigation of the hydrophobicity for the polymer scaffold in **PNTegel** provided

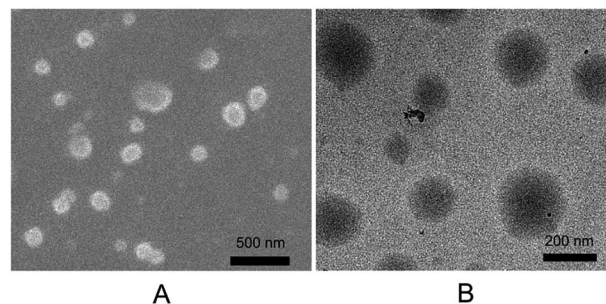


Fig. 3 SEM (A) and TEM (B) images of PNTegel.



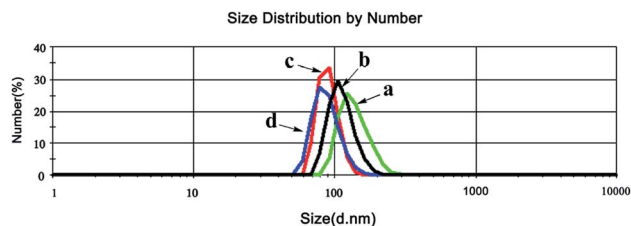


Fig. 4 Hydrodynamic diameters of PNTegel at varying temperature ((a) 25 °C; (b) 35 °C; (c) 37 °C; (d) 45 °C) determined using a Malvern-ZetaSizer Nano ZS instrument.

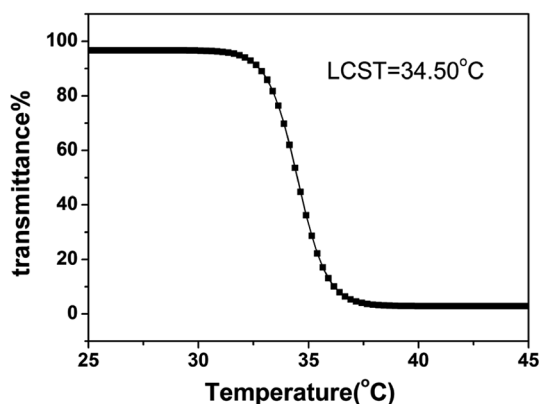


Fig. 5 Temperature dependences of optical transmittance at 600 nm obtained for PNTegel at the concentration of 1 mg mL⁻¹.

the basic information for the following discussion of the temperature responsive catalytic mechanism.

Taken together, the results of NMR, SEM, TEM and DLS proved that the microgel scaffold of PNTegel was successfully prepared based on one-pot method. These results also provided a support for studying the temperature responsive catalytic behavior and catalytic mechanism.

In order to construct an efficient biomimetic GPx, **Te monomer** was designed to be the catalytic center. The catalytic activity of PNTegel was determined according to the modified method reported by Hilvert *et al.*⁶ A typical enzyme catalytic reaction for the determination of catalytic activity was shown in

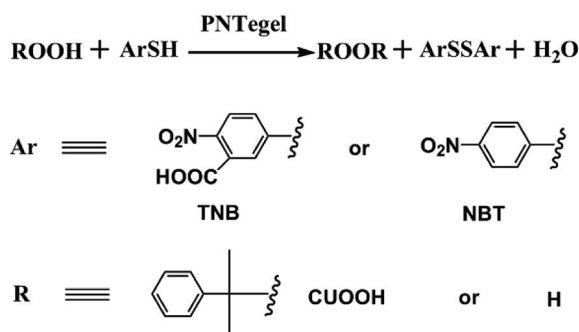


Fig. 6 Determination of GPx catalytic rate for the reduction of ROOH using ArSH as substrate.

Fig. 6. Among the catalytic substrates, TNB and NBT were employed to simulate the glutathione substrate, while CUOOH and H₂O₂ were used to mimic the peroxide substrate during the natural GPx catalytic process. Compared with H₂O₂ and TNB, CUOOH and NBT were the more hydrophobic substrates. Herein, using the hydrophobicity difference of these substrates, the catalytic mechanism and catalytic process for PNTegel could be investigated. The data of the catalytic activity for PNTegel under different conditions are listed in Table 1.

As shown in Table 1, the measurement data of the catalytic activity (v_0) for PNTegel in the temperature range of 25 to 45 °C was used to investigate the temperature responsive catalytic behavior of PNTegel. The results suggested that the catalytic rates of PNTegel exhibited a trend of typical temperature responsive catalytic behavior.

Aiming at evaluating the catalytic ability of PNTegel more objectively, the catalytic rates of PNTegel and the previously reported artificial GPx mimics were summarized in Table 2. Herein, the optimum catalytic activities of various tellurium-containing GPx mimics were determined under the identical conditions using CUOOH and TNB as catalytic substrates. It was observed that the slightly enhanced catalytic rate was obtained when the traditional small molecule artificial GPx (PhSeSePh) was employed. Compared with PhSeSePh, remarkable catalytic rates were achieved when other tellurium-containing artificial GPx in Table 2 were used. Typically, such artificial GPx can be divided into two categories. One of them is the artificial GPx anchored with merely two catalytic elements (catalytic center and hydrophobic environment), *i.e.* PNTegel, PNIPAM-CD-g-Te, Microgel_{max} and SM-Te_{max}. It was found that PNTegel exhibited the most excellent catalytic ability among such artificial GPx with two catalytic elements. Moreover, the other kind of artificial enzyme in Table 2 is the artificial GPx modified with three

Table 1 The catalytic rates (v_0) for the reduction of ROOH (250 μM) by ArSH (150 μM) in the presence of the PNTegel at pH 7.0 (50 mM PBS)

Temperature (°C)	ArSH	ROOH	v_0^a (mM min ⁻¹)
25	TNB	CUOOH	2.32 ± 0.04
28	TNB	CUOOH	2.45 ± 0.11
30	TNB	CUOOH	2.47 ± 0.09
32	TNB	CUOOH	3.51 ± 0.10
34	TNB	CUOOH	5.24 ± 0.11
36	TNB	CUOOH	9.66 ± 0.26
37	TNB	CUOOH	11.51 ± 0.29
38	TNB	CUOOH	11.24 ± 0.31
40	TNB	CUOOH	10.61 ± 0.27
42	TNB	CUOOH	10.07 ± 0.45
45	TNB	CUOOH	8.55 ± 0.48
37	TNB	H ₂ O ₂	2.34 ± 0.09
37	NBT	CUOOH	14.94 ± 0.44
37	NBT	H ₂ O ₂	5.86 ± 0.22

^a Mean ± SD, each sample was determined for three times, and the mean value was calculated to evaluate the catalytic ability of PNTegel. The rate of auto-oxidation of ArSH was deducted during the determination of the catalytic rate. It was assumed that one molecule of the tellurium was used as a catalytic center of enzyme to calculate the activity.



Table 2 The initial rates (v_0) for the reduction of ROOHs by ArSH in the presence of **PNTegel** and other tellurium-containing catalysts

Catalysts	Temperature (°C)	ArSH	ROOH	v_0^a (mM min ⁻¹)
PNTegel	37	TNB	CUOOH	11.51 ± 0.13
PhSeSePh	36	TNB	CUOOH	0.011 ± 0.001
PNIPAM-CD-g-Te^b	35	TNB	CUOOH	6.23 ± 0.25
Microgel_{max}^c	32	TNB	CUOOH	8.09 ± 0.8
SM-Te_{max}^d	38	TNB	CUOOH	5.60 ± 0.29
PN1^e	37	TNB	CUOOH	24.1 ± 1.8
SGPx^f	36	TNB	CUOOH	18.75 ± 0.29

^a Mean ± SD, each sample was determined three times, and the mean value was calculated to evaluate the catalytic ability of **PNTegel**. The rate of auto-oxidation of ArSH was deducted during the determination of the catalytic rate. It was assumed that one molecule of the tellurium was used as a catalytic center of enzyme to calculate the activity. ^b The supramolecular GPx mimic based on asupramolecular graft copolymer. ^c The microgel artificial GPx (Microgel) constructed in our previous report. ^d The supramolecular microgel artificial GPx constructed in our previous report. ^e The GPx mimic based on polystyrene nanoparticle *via* microemulsion polymerization. ^f The smart supramolecular artificial GPx with temperature responsive catalytic activity based on host-guest interaction and a blending process. ⁵⁴

catalytic elements (catalytic center, binding site and hydrophobic environment), *i.e.* **PN1** and **SGPx**. Noticeably, binding site was specially modified into the scaffold of such artificial GPx. Generally, such binding site endowed further enhanced catalytic rates with **PN1** and **SGPx**. And the catalytic ability of **PNTegel** was reasonable lower than that of **PN1** or **SGPx** since the binding site was absent in **PNTegel**. Herein, the main purpose of this work was to explore an efficient and simply method for constructing artificial GPx. Therefore, considering that the reasonable efficient catalytic ability was achieved based on one-pot synthesis, **PNTegel** was still a successful artificial GPx with temperature responsive catalytic behavior.

Moreover, in order to study the catalytic mechanism of **PNTegel**, the catalytic activity of **PNTegel** in reaction systems containing different substrates was also determined. Generally,

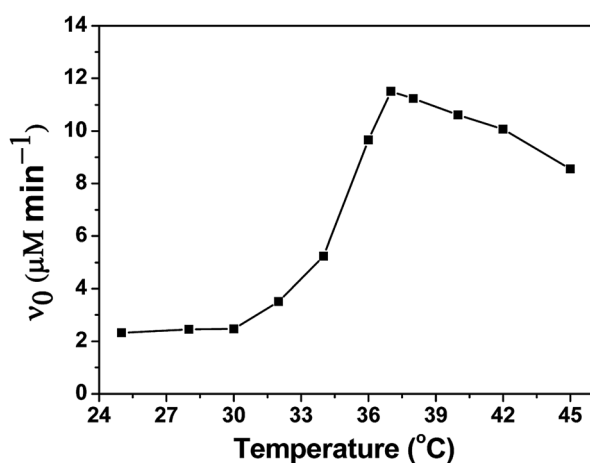


Fig. 7 The catalytic rate of **PNTegel** as a function of temperature for the catalytic reduction of CUOOH (0.25 mM) by TNB (0.15 mM) with a catalytic center of 1.00 μM.

the catalytic activity was determined by the combination of different substrates such as TNB and CUOOH, TNB and H₂O₂, NBT and CUOOH, NBT and H₂O₂, which were employed to study the influence of different hydrophobic substrates in the catalysis process of **PNTegel**. To more intuitively demonstrate the temperature responsive characteristics of the catalytic rate for **PNTegel**, the catalytic activity of **PNTegel** in the system of CUOOH and TNB as a function of temperature is provided in Fig. 7. Obviously, the catalytic rate of **PNTegel** was just slightly changed when the temperature was lower than 32 °C. Whereas, the catalytic rate of **PNTegel** increased significantly with the increasing of temperature when the temperature was above 32 °C. It was noticed that the maximum catalytic activity (11.51 μM min⁻¹) was achieved at 37 °C. However, the catalytic rates of **PNTegel** tended to be significantly decreased as the further promotion of temperature. The reason for these results was discussed in the following section.

Furthermore, to probe the catalytic mechanism of **PNTegel** promoting the reaction mentioned in Fig. 6, detailed kinetic

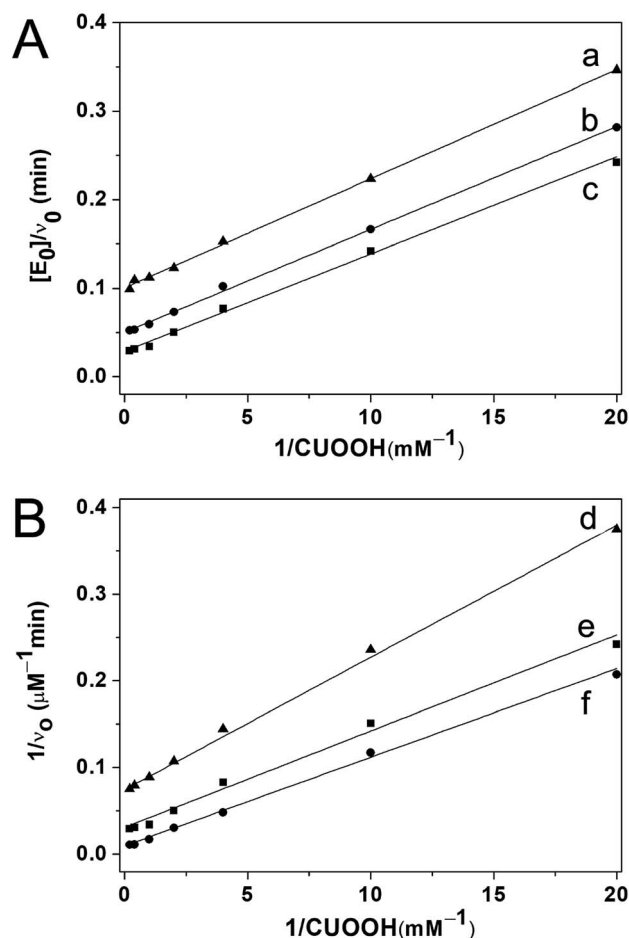


Fig. 8 (A) $[E_0]/v_0$ (min) vs. $1/[CUOOH]$ (mM⁻¹) for 1 μM **PNTegel** in PBS, pH 7.0 and 37 °C, at [TNB] = 0.04 mM (a), 0.10 mM (b), 0.15 mM (c). (B) Lineweaver-Burk plots obtained for the catalytic reaction in the presence of different concentration of **PNTegel** ([**PNTegel**] = 0.5 μM (d), 1 μM (e), 2 μM (f)) at pH 7.0 and 37 °C. The initial CUOOH concentration was fixed to 0.25 mM.



experiments were conducted. These experiments were designed and carried out according to the methods reported by Dong and coworkers.²⁰ They had confirmed that the catalyst–substrate complex existed during the catalytic cycle through detailed kinetic studies. Double-reciprocal plots of initial rate *versus* concentration of CUOOH at all the individual concentration was illustrated in Fig. 8. It suggested that the catalytic rates increased linearly with the concentration of CUOOH. And the rapidly increased catalytic rates were observed in the initial stages (see Fig. 8B and 9) while the concentration of CUOOH was increased (from 0 to 2.5 mM). However, the catalytic rates almost did not change when concentration of CUOOH was further increased. From this observation, it was clearly shown that this catalytic reaction featured the characteristic parallel lines of ping-pong mechanism. And it suggested that the covalent intermediate (–Te–S–Ar) was presented in the catalytic cycle.²⁰

In addition, the enzymatic properties of **PNTEgel** were characterized using TNB and CUOOH as substrates at 37 °C (Fig. 9). The **PNTEgel** exhibited a typical catalytic activity of enzyme. The apparent kinetic constants of **PNTEgel** in the test system containing TNB and CUOOH were as follows: $V_{\max} = 32.15 \mu\text{M min}^{-1}$, $k_{\text{cat}}^{\text{app}} = 32.15 \text{ min}^{-1}$, $K_{\text{m CUOOH}} = 356.91 \mu\text{M}$, $k_{\text{cat}}^{\text{app}}/K_{\text{m CUOOH}} = 9.01 \times 10^4 \text{ M}^{-1} \text{ min}^{-1}$, and the turnover number per catalytic center tellurium was calculated to be 84.75 min^{-1} .

Catalytic mechanism of **PNTEgel**

Based on the research on the crystal structure of GPx and the investigation of biomimetic GPx, it was indicated that hydrophobic cavity derived from the hydrophobic amino acid residues of ¹⁵⁰Phe, ¹⁴⁸Trp, and ³⁴Leu played an important role in maintaining the high catalytic activity.^{1,2} In this study, the hydrophobic structure of the natural GPx was simulated using the hydrophobic microenvironment derived from the PNIPAM polymer scaffold during the temperature responsive process. For natural enzymes, slight structural changes may lead to

significant alterations in catalytic activity. Therefore, we hypothesize that the hydrophobic change of PNIPAM polymer scaffold should be a key factor in regulating the temperature responsive catalytic activity of **PNTEgel**. In order to confirm this suppose, the catalytic mechanism of was discussed by the combination of the changes of optical transmittance, hydrodynamic diameter and catalytic activity with temperature in the catalytic process of **PNTEgel**.

The optical transmittance of **PNTEgel** solution was close to 100% when the temperature was lower than the typical temperature value of the LCST of PNIPAM (about 32 °C). Under this condition, the amide group in **PNTEgel** interacted with the surrounding water molecules by hydrogen bond, which swelled the microgel scaffold in aqueous solution. Therefore, the hydrodynamic diameter of **PNTEgel** was determined to be a relatively large value of about 250 nm (Fig. 4a). In addition, the **PNTEgel** presented a poor catalytic activity due to the absence of hydrophobic microenvironment for effectively gathering the substrates when the temperature was lower than 32 °C.

Subsequently, the temperature was elevated to above 32 °C, the hydrogen bonds between amide group in **PNTEgel** and surrounding water molecules was cleaved, which endowed the PNIPAM polymer in the **PNTEgel** a hydrophobic property. Hence, the PNIPAM polymer chains in the **PNTEgel** were shrunk, which resulted in a remarkable decrease of the optical transmittance of **PNTEgel** solution (Fig. 5) and a decline of the hydrodynamic diameter of **PNTEgel**. For instance, the hydrodynamic diameters of **PNTEgel** at 35 °C (Fig. 4b) and 37 °C (Fig. 4c) were about 230 and 170 nm, respectively, which are smaller than that of **PNTEgel** at 25 °C (about 250 nm). Obviously, the PNIPAM with a temperature responsive property offered a hydrophobic microenvironment for the **PNTEgel** under the temperature higher than the LCST. These hydrophobic microenvironments in the **PNTEgel** could readily gather the hydrophobic substrate and consequently enriched the enzyme-catalyzed substrate molecule around the catalytic center. Therefore, the catalytic activity of **PNTEgel** reached a maximum value of $11.51 \mu\text{M min}^{-1}$ at 37 °C, in which the appropriate hydrophobic microenvironment was formed.

Furthermore, as the further rise of temperature (above 37 °C), the optical transmittance of **PNTEgel** solution was basically stable at a low value. Meanwhile, the **PNTEgel** presented a smaller hydrodynamic diameter at 45 °C (Fig. 4d) in comparison to that of **PNTEgel** existed in the aqueous solution with a lower temperature. This may be attributed to the fact that the hydrophobicity of PNIPAM polymer was further enhanced, which resulted in a more compact structure of microgel scaffold at the high temperature. The tight aggregate structure of **PNTEgel** would encapsulate the catalytic center, which decreased the exposure of catalytic center to the catalytic substrate and thereby reduced the catalytic rate at the high temperature. Obviously, the catalytic activity of **PNTEgel** changed with the hydrophobicity of the polymer scaffold during the temperature responsive catalytic process. In other words, the structural change of **PNTEgel** played a key role in the regulation of catalytic activity.

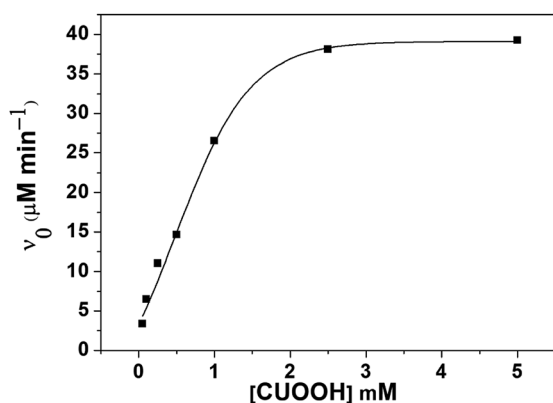


Fig. 9 The catalytic rate as a function of concentration of CUOOH in the presence of the **PNTEgel**. The initial concentration of TNB was fixed to be 0.15 mM, the concentration of CUOOH was 0.05, 0.10, 0.25, 0.5, 1, 2.5 and 5 mM, respectively.



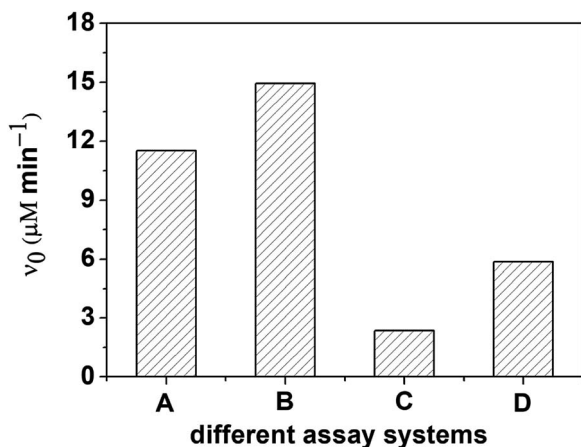


Fig. 10 The catalytic rates (v_0) of the reduction of hydroperoxides (250 μM) by TNB and NBT (150 μM) in the presence of the **PNTegel** at pH 7.0 (50 mM PBS) and 37 °C. (A) CUOOH and TNB; (B) CUOOH and NBT; (C) H_2O_2 and TNB; (D) H_2O_2 and NBT.

In order to further confirm this conclusion, the catalytic rates were measured in systems that contained different catalyzed substrates (Fig. 10). It was found from Fig. 9 that the catalytic activity of **PNTegel** in the system using NBT as glutathione substrate was higher than that in the system containing TNB when the same peroxide substrate was employed ($B > A$, $D > C$). This result indicated that the NBT, with a stronger hydrophobicity, could be more easily reacted as compared with the TNB. Similarly, **PNTegel** exhibited a higher catalytic activity ($A > C$, $B > D$) when a more hydrophobic CUOOH was employed as the peroxide substrate. These results suggest that **PNTegel** exhibits a higher catalytic activity in the reaction system that contains a more hydrophobic substrate. This might be due to the fact that PNIPAM polymer highlighted a hydrophobicity when the temperature was higher than the LCST of PNIPAM, which was likely to gather more hydrophobic substrates into the scaffold of **PNTegel** to a certain extent. Consequently, the contact of the catalytic center to the catalytic substrates was

increased, which promoted the catalytic rate of **PNTegel**. Therefore, the change trend of catalytic activity measured in different catalytic reaction systems further proved that the hydrophobicity of the polymer scaffold in **PNTegel** was an important factor for altering the catalytic activity.

For the temperature responsive biomimetic GPx, the reversibility of the catalytic behavior with temperature was also an important property. Therefore, the reversibility of catalytic activity for **PNTegel** was investigated during heating/cooling processes (see Fig. 11). It was found that the catalytic activity of **PNTegel** did not change substantially when the temperature was converted repeatedly between 25 and 37 °C.

Conclusions

In this work, a new strategy for construction of biomimetic GPx using a one-pot synthesis was successfully explored. The prepared biomimetic GPx, **PNTegel**, exhibited a typical temperature responsive catalytic activity. The highest catalytic activity ($11.51 \mu\text{M min}^{-1}$) in the system containing TNB and CUOOH was obtained at 37 °C during the temperature responsive process. The structure change of the polymer scaffold of **PNTegel** that induced by the temperature is mainly responsible for the change of the catalytic activity. Because of the absence of the hydrophobic microenvironment for gathering the reaction substrates, **PNTegel** presented a low catalytic activity at low temperature (<LCST). Whereas a high temperature (>37 °C) lead to a compact structure of **PNTegel**, which obstructs the contact of the catalytic center and substrates and also decreases the catalytic activity. This study may provide important theoretical supports for the preparation of highly effective antioxidant drugs.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (51663020), Natural Science Fund for Distinguished Young Scholars of Guangxi Province (2017GXNSFFA198007), the project of Guangxi Colleges and Universities for the Promotion of Foundation Ability of Young Teachers (2018KY0599 and 2019KY0469) and the Project of Guangxi Colleges and Universities Key Laboratory of Beibu Gulf Oil and Natural Gas Resource Effective Utilization (2017KLOG12). The authors also acknowledge the Guangxi Colleges and Universities Innovation Research Team.

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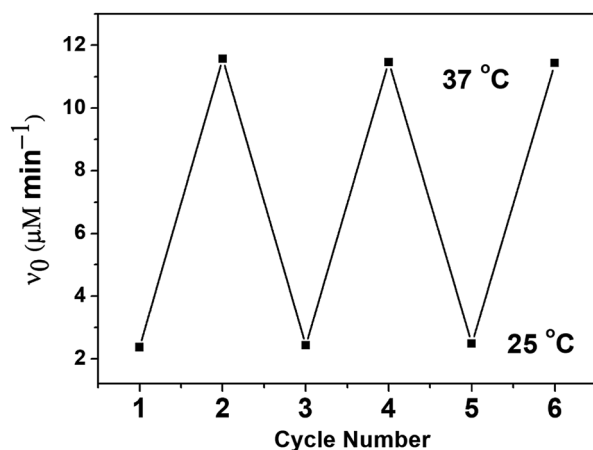


Fig. 11 Changes of the catalytic activity of **PNTegel** during heating/cooling cycles between 25 and 37 °C.



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