


Cite this: *RSC Adv.*, 2020, 10, 6647

An electrochemical aptasensor for lead ion detection based on catalytic hairpin assembly and porous carbon supported platinum as signal amplification†

Huali Jin,^a Di Zhang,^a Yong Liu^b and Min Wei^b*

An electrochemical aptasensor is fabricated for lead ion (Pb^{2+}) detection based on catalytic hairpin assembly as signal amplification. Biomass porous carbon derived from soybean straw (PCs) is used to load platinum nanoparticles (PtNPs), which are introduced as mimetic enzymes to catalyze the hydroquinone– H_2O_2 system. In the presence of Pb^{2+} , the complementary DNA (cDNA) releases from aptamer–cDNA and hybridizes with hairpin DNA1 (HP1) on the electrode surface. After bio-hairpin DNA2 (bio-HP2) is added, hybridization between HP1 and bio-HP2 further releases cDNA, which participates in the next cycle and triggers amplification, eventually forming a large number of bio-HP2/HP1 on the electrode surface. Then PtNPs@PCs immobilize on the electrode surface by specific binding of streptavidin with biotin, and catalyze the oxidation of hydroquinone in the presence of H_2O_2 . The produced electrochemical signal depends on the concentration of Pb^{2+} . The developed biosensor exhibits a wide linear range from 50 pM to 1000 nM with a detection limit of 18 pM, and high selectivity for Pb^{2+} over other environmentally relevant metal ions at concentration ratios of 100. The recoveries of Pb^{2+} in real samples were 93.5–108%. Thus, the proposed biosensor can provide an efficient method for sensitive and selective detection of Pb^{2+} in real samples.

Received 2nd January 2020

Accepted 6th February 2020

DOI: 10.1039/d0ra00022a

rsc.li/rsc-advances

1. Introduction

As one of the most hazardous heavy metal contaminants, lead ions (Pb^{2+}) can exist in the environment for a long time and accumulate in the body through skin absorption, the digestive tract or food chain, leading to nerve disorders, memory decay, or anemia even at very low level exposure, particularly in children.^{1,2} The World Health Organization (WHO) and the US Environmental Protection Agency (EPA) regulate that the maximum acceptable concentration of Pb^{2+} in drinking water is 48 nM and 72 nM, respectively.^{3,4} Recent studies reported that the adverse effects of Pb^{2+} are widely recognized as toxic at concentrations as low as 1 pM.^{5,6} Thus, it is of great value to construct simple, ultrasensitive and rapid methods for Pb^{2+} detection. Many methods such as atomic absorption spectrometry (AAS), inductively coupled plasma atomic emission spectrometry (ICP-AES), and inductively coupled plasma mass spectrometry (ICP-MS) have been reported for the detection of

Pb^{2+} .^{7–9} Although these conventional analytical methods are sensitive and accurate, they are labor-intensive, time-consuming, and expensive, which restrict their wider applications.¹⁰ At present, electrochemical sensors with different signal amplification strategies have been widely utilized in the detection of Pb^{2+} owing to its simple instrumentation, rapid response, low cost and high sensitivity.^{11–13}

As a kind of novel recognition element, aptamers exhibit many advantages, such as easy synthesis, excellent stability and strong specificity.^{14,15} Based on these unique properties, many electrochemical sensors based on aptamers have been fabricated for ultrasensitive detection of Pb^{2+} .^{16,17}

In order to further amplify response signal and enhance the sensitivity, different signal amplification techniques including palindromic molecular beacon (PMB),¹⁸ rolling circle amplification (RCA),¹⁹ DNA walker,²⁰ hybridization chain reaction (HCR)²¹ and catalytic hairpin assembly (CHA)²² have attracted increasing attention. Among them, as isothermal nucleic acid amplification strategy, with the simple reaction conditions, high sensitivity and the low costs, CHA is powerful for amplifying and transducing signals at the terminus of nucleic acid amplification reactions,^{23,24} and has been broadly applied in electrochemical sensing platforms.^{25,26} The presence of initiator DNA can trigger CHA process to achieve signal amplification.²⁷ DNzyme with the existence of Pb^{2+} can act as initiator for

^aCollege of Food Science and Technology, Henan Key Laboratory of Cereal and Oil Food Safety Inspection and Control, Henan University of Technology, Zhengzhou 450001, PR China

^bCollege of Chemistry and Chemical Engineering, Henan University, Kaifeng, 475004, PR China. E-mail: wei_min80@163.com

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ra00022a



triggering CHA process, and has been successfully used for Pb^{2+} detection.^{28,29} However, Pb^{2+} dependent DNzyme cleaves RNA irreversibly, and leads to the destroyed structure, high cost and hard reuse, which limit its wider applications.^{30,31} Instead, aptamer only changes the structure and can be reusable. To our best knowledge, aptamer-based CHA for Pb^{2+} detection have not been reported. Herein, CHA based on Pb^{2+} aptamer is designed to realize the amplification strategy.

In recent years, compared with natural enzymes, due to the advantages of broader reaction conditions, higher stability, lower production cost, and better catalytic activity, mimetic enzymes based on different nanomaterials have been developed to construct biosensors.^{32–34} Among them, with the advantages of good electrical conductivity, larger surface activity, favorable catalytic ability, and wider range of pH and temperature conditions, Pt-based nanomaterials such as Pt nanotubes and Pt nanoparticles (PtNPs) have been widely used in electrochemical biosensors to obtain the better performance.^{35–37} In order to avoid the aggregation of Pt-based nanomaterials and improve the catalytic activity, different nanomaterials such as metal–organic frameworks³⁸ and reduced graphene oxide³⁹ were used as carriers to enhance the dispersity. Due to the unique properties of pore structure, high surface area, good conductivity, low production cost and environment-friendly preparation, biomass porous carbons (PCs) derived from soybean straw can be a good choice to support PtNPs for electrochemical sensors.

Herein, a novel label-free electrochemical aptasensor is developed to detect Pb^{2+} . CHA based on Pb^{2+} aptamer is designed to realize the amplification strategy. PCs produced by soybean straw is introduced as the carrier of PtNPs to prepare PtNPs@PCs, which acts as peroxide mimetic enzyme to catalyze the oxidation of hydroquinone in the presence of hydrogen peroxide and realize the second signal amplification.

2. Experimental

The “Apparatus and electrochemical measurement” was provided in the ESI.†

2.1. Reagents and materials

Streptavidin (SA) was purchased from Solarbio Science and Technology Co., Ltd. (Beijing, China). 6-Mercapto-1-hexanol (MCH) was purchased from Aladdin Co., Ltd. (Shandong province, China). Tris(hydroxymethyl)aminomethane (Tris) was obtained from Shanghai Macklin Biochemical Co., Ltd. (Shanghai, China). Tris(2-carboxyethyl)phosphine (TCEP) was purchased from shanghai yuanye Bio-Technology Co., Ltd. (Shanghai, China). Hydroquinone (HQ) and lead nitrate ($\text{Pb}(\text{NO}_3)_2$) were ordered from Kermel Co., Ltd. (Tianjin, China). H_2O_2 (30%, w/w) was obtained from Haohua Co., Ltd. (Luoyang, Henan Province, China). Soybean straw comes from farmland (Qi County, Henan Province, P. R. China). All used reagents were of analytical grade. The ultrapure water was used throughout the experiments. All DNA sequences were synthesized by Sangon Biotechnology (Shanghai, China) and listed below:

Aptamer: 5'-GGG TGG GTG GGT GGG TAT-3'

Complementary DNA (cDNA): 5'-TCA TAC CCA CCC ACC-3';

Hairpin DNA 1 (HP1): 5'-TTT TGG GTG GGT ATG ACC ACC GCC CAC CCA-3'

Bio-Hairpin DNA 2 (bio-HP2): 5'-bio-TAT GAC CAC CTG GGT GGG CGG TGG TCA TAC CCA C-3'

Each HP was heated at 95 °C for 5 min and then slowly cooled to room temperature before using. All oligonucleotide solutions for DNA hybridization reaction were prepared by using 20 mM Tris-HCl (pH 7.4) containing 140 mM NaCl and 5 mM MgCl_2 . 0.1 M phosphate buffered solution (PBS, pH 7.4) was obtained by mixing the 0.1 M Na_2HPO_4 and 0.1 M KH_2PO_4 as working buffer.

2.2. Preparation of PtNPs@PCs-SA

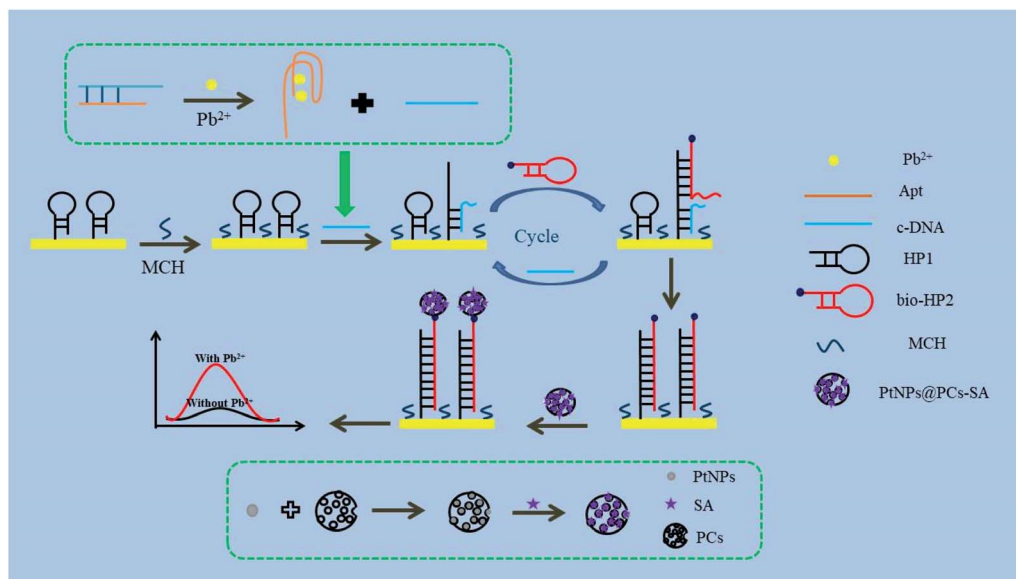
The honeycomb-like structure PCs was prepared by one-step pyrolysis using soybean straw as the precursor and magnesium oxide as the template. 1.0 g soybean straw was placed in the vibration mill together with 2.0 g magnesium oxide. After even grinding, the mixtures were carbonized at high temperature of 800 °C for 2 h with heating rate of 3 °C min^{-1} under the N_2 atmosphere in a tube furnace. The obtained black solid was treated with 1 M HCl in order to remove pore-forming agent magnesium oxide. Then, the examples were washed with distilled water until the solution pH reached 7, and were dried at 60 °C for 12 h in the oven. So, PCs was obtained. PtNPs@PCs were prepared using the ethylene glycol (EG) reduction method according to the reported literatures.⁴⁰ Finally, 2 mg of PtNPs@PCs was dispersed in PBS buffer, and then 2 mg of streptavidin was added with gently stirring for 12 h at 4 °C to sufficiently immobilize onto the surface of PtNPs@PCs *via* the strong Pt–N affinity.⁴¹

2.3. Fabrication of the electrochemical biosensor

Firstly, the mixture of 3 μL of 4 μM Apt and 3 μL of 4 μM cDNA were heated to 95 °C for 5 min then slowly cooled to room temperature to form double-stranded structure. Then Pb^{2+} with different concentrations was added into above solution for 40 min at 37 °C to obtain the mixture of Pb^{2+} /Apt and cDNA. 5 μL of 3.5 μM HP1 was incubated with 10 mM TCEP for 2 h to block the formed disulfide bonds, and then was added onto the surface of AuE at room temperature for 12 h to obtain the HP1/AuE. After washing with Tris-HCl buffer (pH 7.4), the HP1/AuE was blocked with MCH for 30 min at 37 °C. Then, the mixture of Pb^{2+} /Apt and cDNA were added to electrode for 1.5 h at 37 °C. Next, 5 μL of 4 μM bio-HP2 was dropped onto the electrode and incubated for 1.5 h at 37 °C. Finally, 5 μL of PtNPs@PCs-SA was added to the electrode and incubated at 37 °C for 2 h.

As shown in Scheme 1, the signal amplification strategy is conducted as follows: firstly, in the presence of Pb^{2+} , the combination of Pb^{2+} and aptamer forms stable G-quadruplex structure, leading to the release of cDNA. Then, the released cDNA hybridizes with HP1 on the AuE surface. After bio-HP2 is added onto the AuE surface, its hairpin structure opens and hybridizes with HP1. Thus, the cDNA is replaced by bio-HP2 and released from HP1 for the next cycle. When the cyclic processes





Scheme 1 Illustration of the electrochemical aptasensor for detection of Pb^{2+} .

complete, there are a large number of capture probe (bio-HP2/HP1) left on the electrode. Finally, PtNPs@PCs-SA are immobilized on the electrode by the specific recognition between biotin and streptavidin. Thus, in the presence of H_2O_2 , PtNPs@PCs acts as peroxide mimetic enzyme to catalyze the oxidation of HQ and realize the second signal amplification.

3. Results and discussion

3.1. Characterization of PtNPs@PCs

The prepared PCs and PtNPs@PCs were characterized by SEM and TEM, and the results were shown in Fig. 1. The SEM image of PCs (Fig. 1A) indicated that the obtained carbon materials were porous structure monolith after the removal of MgO template. As shown in Fig. 1B, it can be seen that PtNPs were uniformly spread on the surface of the porous of PCs.

Fig. 2 showed the XRD patterns of PtNPs@PCs . The diffraction peak at 2θ of 39.9° , 45.5° , 67.7° , and 81.4° correspond to

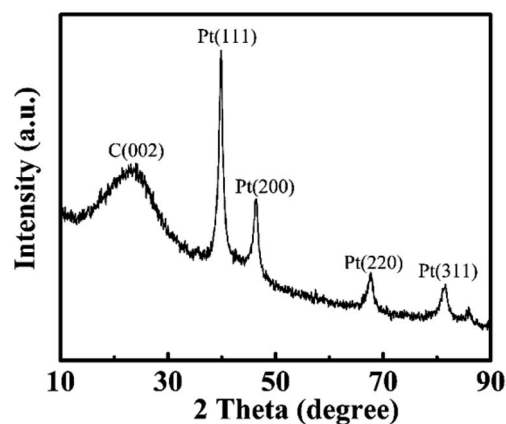


Fig. 2 XRD patterns of PtNPs@PCs .

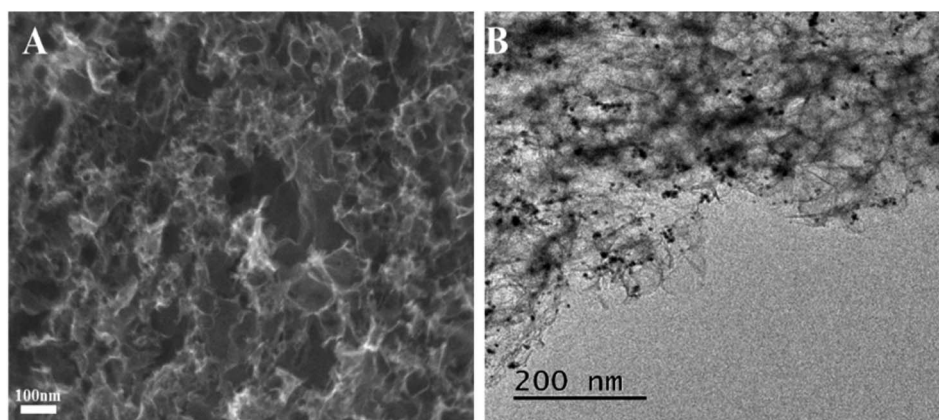


Fig. 1 (A) SEM image of the PCs and (B) TEM image of the PtNPs@PCs .

(110), (200), (220), and (311) crystalline planes of face-centered cubic structure of Pt, respectively. The diffraction peak at about 23.5° observed can be attributed to the C (002) facet of graphitic carbon.

In addition, DPV was used to characterize the mimic enzyme performance of PtNPs@PCs (Fig. S1†). When the bare electrode was placed in the PBS buffer containing HQ, the oxidation peak was measured at 0.136 V and the oxidation peak current was $22.09 \mu\text{A}$ (curve a). However, when the bare electrode was put into the PBS buffer with HQ and H_2O_2 , the oxidation peak potential shifted negatively, and the oxidation peak current increased to $32.63 \mu\text{A}$ (curve b), indicating that H_2O_2 can promote the oxidation of HQ. When the PtNPs@PCs-modified electrode was placed in PBS buffer with HQ, the peak current increased significantly to $91.54 \mu\text{A}$ and the peak potential shifted negatively to 0.064 V (curve c), indicating that PtNPs@PCs has the effect of accelerating electron transfer. When the PtNPs@PCs-modified electrode was immersed in the PBS buffer with HQ and H_2O_2 (curve d), the peak current further increased to $113.4 \mu\text{A}$ and the peak potential shifted negatively to 0.044 V, which was ascribed that larger surface activity and good electrical conductivity of PtNPs@PCs make it act as peroxidase-like catalyst to catalyze H_2O_2 and accelerate the oxidation of HQ. Moreover, due to its good stability, PtNPs@PCs can be reusable and retains its catalytic activity in a wider range of pH and temperature, making it a great potential application.

3.2. Characterization of the proposed biosensor

EIS is used to characterize the fabrication of biosensor. As shown in Fig. 3, the bare AuE showed a low R_{et} value (curve a). After HP1 was assembled on the AuE surface, the R_{et} value increased obviously (curve b), which is ascribed to the mutual repulsion between negatively charged phosphate group and redox probe. When the MCH was immobilized onto the HP1/AuE, the R_{et} value further increased (curve c), indicating that

MCH blocks nonspecific sites on the electrode surface. When cDNA was introduced, the R_{et} value increased due to the negative charge of DNA (curve d). Similarly, the incubation of bio-HP2 also led to further increase of R_{et} value (curve e). With the introduction of PtNPs@PCs-SA (curve f), the R_{et} value apparently decreased due to that the formation of PtNPs@PCs-SA on the electrode promote the electron transfer.

3.3. Electrochemical behavior of the biosensor

The prepared sensor is applied to the detection of Pb^{2+} , and the result is shown in Fig. 4. In the absence of Pb^{2+} , cDNA and Apt hybridizes to form the double-stranded DNA structure. Therefore, cDNA cannot reach the AuE surface to hybridize with HP1. The cycle amplification of CHA cannot be performed. So, the initial peak current was $0.65 \mu\text{A}$ (curve a). In the presence of $1 \mu\text{M}$ Pb^{2+} , Apt combines with Pb^{2+} to form a G-quadruplex structure, resulting in that cDNA detached from the double-stranded DNA structure and hybridized with HP1 on the AuE surface, thereby opening the hairpin structure of HP1. With the introduction of bio-HP2, cDNA released from HP1 due to the greater number of complementary bases between bio-HP2 and HP1. The released cDNA participated the next cycle and further hybridized with HP1 to realize the catalytic hairpin assembly and achieve the amplification strategy. When the cyclic processes completed, there were a large number of bio-HP2/HP1 formed on the electrode. By combining with bio-HP2 by the specific recognition between biotin and streptavidin, PtNPs@PCs-SA can catalyze the oxidation of HQ in the presence of H_2O_2 , thereby enhancing the peak current signal and realizing the detection of Pb^{2+} . So, the peak current of $5.84 \mu\text{A}$ was found (curve b). The produced electrochemical signal depends on the concentration of Pb^{2+} , which can be used for the detection of Pb^{2+} .

3.4. Optimization of the experimental conditions

The experimental conditions including bio-HP2 concentration, HQ concentration and PtNPs@PCs incubation time are optimized. As depicted in Fig. 5A, by increasing the bio-HP2

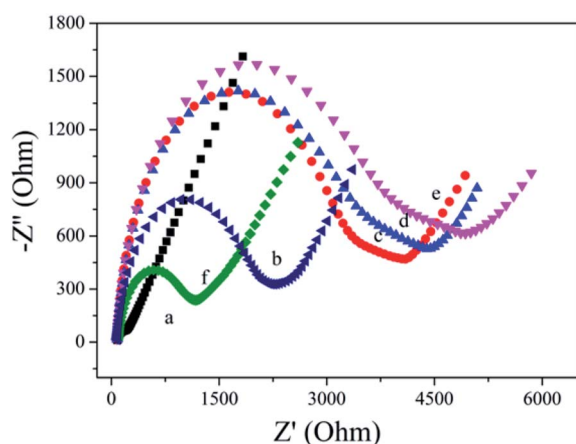


Fig. 3 EIS of different modified electrodes were conducted in 0.1 M PBS (pH 7.4) solution containing 5 mM $[\text{Fe}(\text{CN})_6]^{3-/4-}$ and 0.5 M KCl: (a) AuE; (b) HP1/AuE; (c) MCH/HP1/AuE; (d) cDNA/MCH/HP1/AuE; (e) bio-HP2/cDNA/MCH/HP1/AuE; (f) PtNPs@PCs-SA/bio-HP2/cDNA/MCH/HP1/AuE.

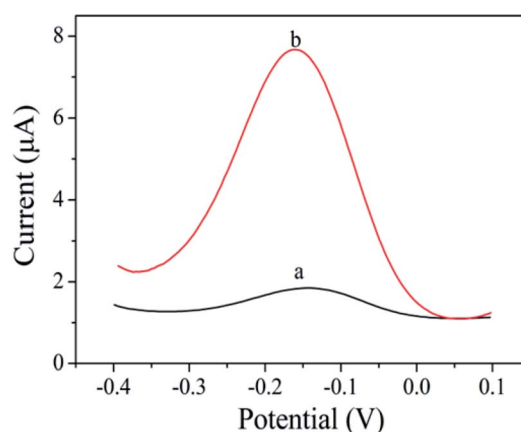


Fig. 4 DPV curves of without Pb^{2+} (a) and with Pb^{2+} ($1 \mu\text{M}$) (b) in PBS (pH 7.4) buffer containing 10 mM HQ and 10 mM H_2O_2 .



concentration, more PtNPs@PCs were modified on the electrode surface by the combination of biotin and streptavidin, and the current response gradually increased. When the concentration of bio-HP2 was 4 μM , the current response reached the maximum. As the concentration continued to increase, the current value became stable. Thus, 4 μM was chosen as the optimal concentration of bio-HP2 for the following experiments. As shown in Fig. 5B, with the concentration of HQ increased, the current response gradually

increased and tended to be steady after 10 mM. Thus, 10 mM was used as the optimal concentration of HQ. The incubation time of PtNPs@PCs was optimized, and the results were shown in Fig. 5C. As the incubation time increased, the current value gradually increased, and when the incubation time was 120 min, the current value reached the maximum. As the incubation time continued to increase, the current value slightly reduced. Therefore, 120 min was chosen as the optimal incubation time.

3.5. Analytical performance of biosensor

Different concentrations of Pb^{2+} are detected by the prepared biosensor. As shown in Fig. 6A, the peak current increased gradually with increasing Pb^{2+} concentration in the range of 0.05–5000 nM. From the results showed in Fig. 6B, a linear relationship between peak current and the logarithm of Pb^{2+} concentration from 0.05 nM to 1000 nM, and the linear regression equation was $I = 1.13 \lg C_{\text{Pb}^{2+}} + 12.47$ ($R^2 = 0.995$). The detection limit of 0.018 nM was calculated on the basis of 3

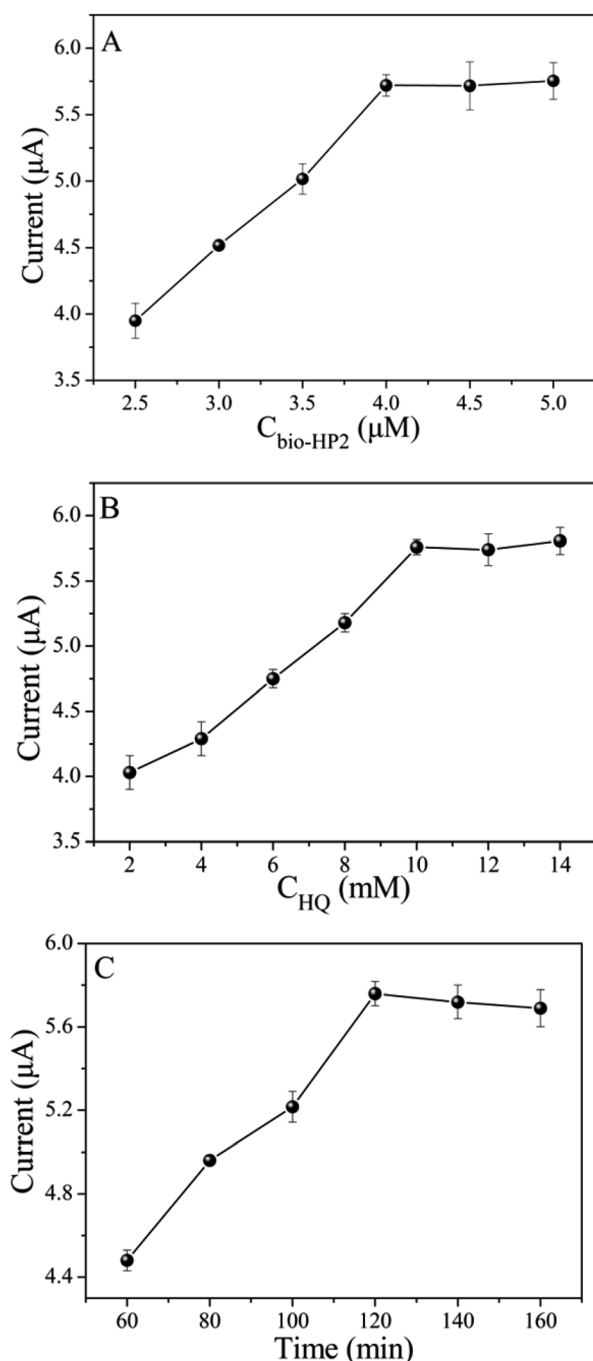


Fig. 5 The influence of (A) bio-HP2 concentration, (B) HQ concentration, and (C) the incubation time of PtNPs@PCs on the biosensor.

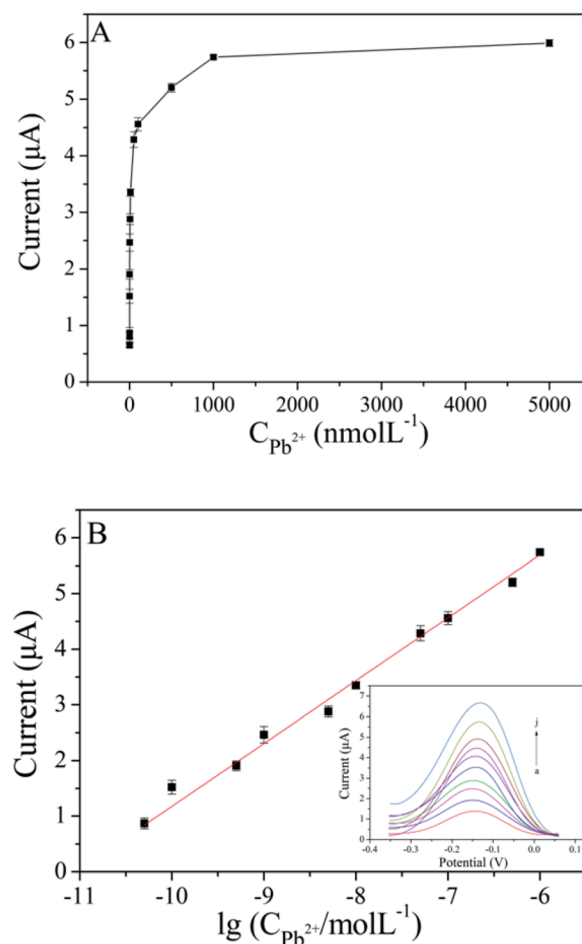


Fig. 6 (A) Relationship between the peak current and Pb^{2+} concentration (0 nM, 0.05 nM, 0.1 nM, 0.5 nM, 1 nM, 5 nM, 10 nM, 50 nM, 100 nM, 500 nM, 1000 nM, 5000 nM) ($n = 3$); (B) linear relationship between the peak current and the logarithm of Pb^{2+} concentration ranging from 0.05 nM to 1000 nM (inset: DPV curves of the aptasensor with Pb^{2+} concentration ranging from 0.05 nM to 1000 nM).

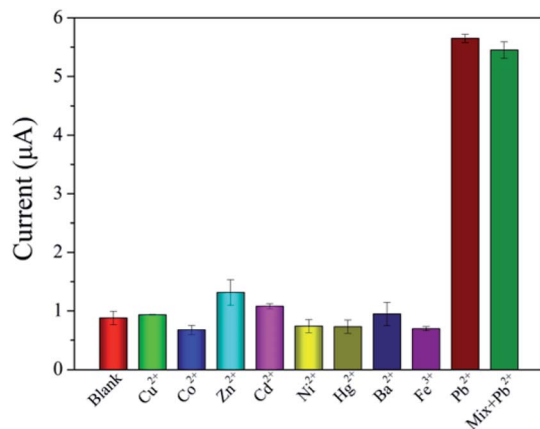


Fig. 7 Selectivity of the prepared biosensor for Pb²⁺ compared with other metal ions ($n = 3$). The concentration of Pb²⁺ is 1 μ M and that of other metal ions are 100 μ M.

times the standard deviation above the blank. As shown in Table S1,[†] the performance of the proposed aptasensor is better or comparable with that of previously reported Pb²⁺ sensors.

3.6. Selectivity, reproducibility and stability of the biosensor

Under the same experimental conditions, some metal ions such as Cu²⁺, Co²⁺, Zn²⁺, Cd²⁺, Ni²⁺, Hg²⁺, Ba²⁺ and Fe³⁺ are used to evaluate the selectivity of the biosensor, and the results are shown in Fig. 7. The concentration of other metal ions is 100 μ M, and that of Pb²⁺ is 1 μ M. In the absence of Pb²⁺, the current response of other metal ions had no significant change compared with that of the measured blank solution. The current response of the mixture with Pb²⁺ was similar to that with Pb²⁺ only. These results indicated that the biosensor had an excellent selectivity for Pb²⁺ detection.

The inter-assay precision was evaluated by five times parallel measurements of 1 μ M Pb²⁺, and relative standard deviation (RSD) value of 1.48% was obtained. The intra-assay precision was determined by five times repeated measurements on the same aptasensor, and RSD value of 4.69% was obtained. After a 15 day storage period at 4 °C, the aptasensor retained 88.1% of its initial current response, indicating the acceptable stability.

3.7. Analysis of Pb²⁺ in real samples

Different concentrations of Pb²⁺ at 500 nM, 10 nM and 0.1 nM were respectively added into tap water samples and lake

Table 1 Detection of Pb²⁺ in real samples with the proposed biosensor

Samples	Added (nM)	Found (nM)	Recovery (%)	RSD (%) ($n = 3$)
Tap water	500.0	496.9	99.4	5.02
	10.0	9.35	93.5	5.10
	0.1	0.105	104.9	4.71
Lake water	500.0	477.3	95.5	4.38
	10.0	10.8	108	3.11
	0.1	0.939	93.9	12.21

samples, which were detected using the proposed biosensor. As shown in Table 1, the recoveries were in the range of 93.5–108%, suggesting that proposed biosensor can be used for Pb²⁺ detection in real samples.

4. Conclusion

In summary, an electrochemical biosensor was fabricated for Pb²⁺ detection by catalytic hairpin assembly and PtNPs@PCs as signal amplification strategy. The developed biosensor exhibited higher sensitivity, good selectivity, and favorable recovery in water samples. Therefore, the proposed electrochemical biosensor has a potential application for Pb²⁺ detection in the environment, and can be expanded for the detection of other metal ions.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This study was funded by the National Key Research and Development Program of China (2018YFC1602800), Key Scientific and Technological Project of Henan Province (192102310255), and the Fundamental Research Funds for the Henan Provincial Colleges and Universities in Henan University of Technology (2016RCJH04).

References

- 1 J. N. Ding, Y. Liu, D. W. Zhang, M. L. Yu, X. J. Zhan, D. Zhang and P. Zhou, *Microchim. Acta*, 2018, **185**, 545.
- 2 L. Y. Wang, X. L. Peng, H. J. Fu, C. Huang, Y. P. Li and Z. M. Liu, *Biosens. Bioelectron.*, 2020, **147**, 111777.
- 3 F. L. Wang, Z. Wu, Y. X. Lu, J. Wang, J. H. Jiang and R. Q. Yu, *Anal. Biochem.*, 2010, **405**, 168–173.
- 4 L. Zhang, H. M. Deng, R. Yuan and Y. L. Yuan, *Microchim. Acta*, 2019, **186**, 709.
- 5 F. Long, A. N. Zhu and H. C. Wang, *Anal. Chim. Acta*, 2014, **849**, 43–49.
- 6 E. Rossi, *Clin. Biochem. Rev.*, 2008, **29**, 63–70.
- 7 Z. H. Hao, J. Z. Yao, R. L. Tang, X. M. Zhang, W. G. Li and Q. Zhang, *Spectrosc. Spectral Anal.*, 2015, **35**(2), 527–533.
- 8 Y. K. Wang, S. T. Gao, X. H. Zang, J. C. Li and J. J. Ma, *Anal. Chim. Acta*, 2012, **716**, 112–118.
- 9 S. Wang, X. Dong, B. Y. Dai, M. F. Pan, S. Y. He and J. P. Wang, *J. AOAC Int.*, 2015, **98**(1), 218–224.
- 10 Y. G. Wang, G. H. Zhao, Q. Zhang, H. Wang, Y. Zhang, W. Cao and Q. Wei, *Sens. Actuators, B*, 2019, **288**, 325–331.
- 11 J. J. Shi, J. C. Zhu, M. Zhao, Y. Wang, P. Yang and J. He, *Talanta*, 2018, **183**, 237–244.
- 12 B. Zhang, J. F. Chen, B. Q. Liu and D. P. Tang, *Biosens. Bioelectron.*, 2015, **69**, 230–234.
- 13 Q. Zhou, Y. X. Lin, Y. P. Lin, Q. H. Wei, G. N. Chen and D. P. Tang, *Biosens. Bioelectron.*, 2016, **78**, 236–243.



- 14 C. Y. Yang, Q. Wang, Y. Xiang, R. Yuan and Y. Chai, *Sens. Actuators, B*, 2014, **197**, 149–154.
- 15 F. L. Li, Y. M. Guo, X. Y. Wang and X. Sun, *Biosens. Bioelectron.*, 2018, **115**, 7–13.
- 16 S. M. Taghdisi, N. M. Danesh, P. Lavaee, M. Ramezani and K. Abnous, *Sens. Actuators, B*, 2016, **234**, 462–469.
- 17 F. Gao, C. Gao, S. Y. He, Q. X. Wang and A. Q. Wu, *Biosens. Bioelectron.*, 2016, **81**, 15–22.
- 18 R. J. Zeng, Z. B. Luo, L. S. Su, L. J. Zhang, D. P. Tang, R. Niessner and D. Knop, *Anal. Chem.*, 2019, **91**, 2447–2454.
- 19 H. X. Ji, F. Yan, J. P. Lei and H. X. Ju, *Anal. Chem.*, 2012, **84**, 7166–7171.
- 20 S. Z. Lv, K. Y. Zhang, Y. Y. Zeng and D. P. Tang, *Anal. Chem.*, 2018, **90**, 7086–7093.
- 21 X. Y. Xie, Y. Q. Chai, Y. L. Yuan and R. Yuan, *Anal. Chim. Acta*, 2018, **1034**, 56–62.
- 22 J. Y. Dai, Z. J. Duan, M. Z. Cao, M. R. Hao, H. F. He and D. Xiao, *Talanta*, 2018, **181**, 142–146.
- 23 S. S. Tang, Y. Gu, H. T. Lu, H. F. Dong, K. Zhang, W. H. Dai and X. J. Zhang, *Anal. Chim. Acta*, 2018, **1004**, 1–9.
- 24 Y. Jiang, B. L. Li, J. N. Milligan, S. Bhadra and A. D. Ellington, *J. Am. Chem. Soc.*, 2013, **135**(20), 7430–7433.
- 25 X. Y. Wu, Y. Q. Chai, P. Zhang and R. Yuan, *ACS Appl. Mater. Interfaces*, 2014, **7**(1), 713–720.
- 26 H. L. Shuai, K. J. Huang, L. L. Xing and Y. X. Chen, *Biosens. Bioelectron.*, 2016, **86**, 337–345.
- 27 Y. Wang, N. Gan, Y. Zhou, T. H. Li, F. T. Hu, Y. T. Cao and Y. J. Chen, *Biosens. Bioelectron.*, 2017, **97**, 100–106.
- 28 X. Y. Huang, J. L. Li, Q. Y. Zhang, S. Chen, W. Xu, J. Y. Wu, W. C. Niu, J. J. Xue and C. R. Li, *J. Electroanal. Chem.*, 2018, **816**, 75–82.
- 29 Y. D. Ma, C. Yu, Y. J. Yu, J. Chen, R. F. Gao and J. L. He, *Microchim. Acta*, 2019, **186**(10), 677.
- 30 H. Wang, Y. Liu and G. Liu, *Microchim. Acta*, 2018, **185**, 142.
- 31 D. Wang, C. C. Ge, K. P. Lv, Q. S. Zou, Q. Liu, L. P. Liu, Q. H. Yang and S. Y. Bao, *Chem. Commun.*, 2018, **54**, 13718–13721.
- 32 H. Wei and E. Wang, *Chem. Soc. Rev.*, 2013, **42**(14), 6060–6093.
- 33 D. P. Cormode, L. Gao and H. Koo, *Trends Biotechnol.*, 2018, **36**(1), 15–29.
- 34 S. Chen, P. Liu, K. W. Su, X. Li, Z. Qin, W. Xu, J. Chen, C. R. Li and J. F. Qiu, *Biosens. Bioelectron.*, 2018, **99**, 338–345.
- 35 R. J. Zeng, Z. B. Luo, L. J. Zhang and D. P. Tang, *Anal. Chem.*, 2018, **90**, 12299–12306.
- 36 Z. Z. Yu, Y. Tang, G. N. Cai, R. R. Ren and D. P. Tang, *Anal. Chem.*, 2018, **91**, 1222–1226.
- 37 Q. Yu, Y. M. Wu, Z. Liu, S. Lei, G. P. Li and B. X. Ye, *Biosens. Bioelectron.*, 2018, **107**, 178–183.
- 38 C. L. Zhang, J. L. He, Y. C. Zhang, J. Chen, Y. L. Zhao, Y. Z. Niu and C. Yu, *Biosens. Bioelectron.*, 2018, **102**, 94–100.
- 39 Y. M. Wu, L. N. Zou, S. Lei, Q. Yu and B. X. Ye, *Biosens. Bioelectron.*, 2017, **97**, 317–324.
- 40 Y. M. Zhang, Y. Liu, W. H. Liu, X. Y. Li and L. Q. Mao, *Appl. Surf. Sci.*, 2017, **407**, 64–71.
- 41 W. Cai, S. B. Xie, Y. Tang, Y. Q. Chai, R. Yuan and J. Zhang, *Talanta*, 2017, **163**, 65–71.

