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## Strategy for the detection of mercury ions by using exonuclease III-aided target recycling

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Herein, a simple, highly sensitive and selective exonuclease III (Exo III)-based sensor was developed for mercury ions  $Hg^{2+}$  detection without the need of complicated, high-cost and time-consuming labeling. In this strategy, two oligonucleotides were employed for the experiment design. DNA1 was prepared with one DNA strand which was designed as a quadruplex-forming oligomer, and which was constructed as 3'-protruding to ensure the probe cannot be digested by Exo III. When the target  $Hg^{2+}$  is introduced into the sensing system, it associates with DNA1 and DNA2 to form double strand DNAs (DNA1/DNA2 duplexes) which have a blunt 3'-terminus. Exo III then can degrade part of DNA1, releasing the DNA1 and  $Hg^{2+}$ , and ultimately liberating the G-rich oligomer. This released G-rich oligomer folds into a G-quadruplex structure and thus allows the formation of DNAzyme in the presence of hemin. The formed DNAzyme can effectively catalyze the  $H_2O_2$ -mediated oxidation of ABTS, giving rise to a change in solution color. The released DNA1 and  $Hg^{2+}$  are free to bind to another 3'-protruding terminus of DNA1 to trigger a new digestion reaction, leading to significant amplification of the signal. The present new strategy shows a limit of detection as low as 66 pM and excellent selectivity toward  $Hg^{2+}$  over a wide range of metal ions.

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## Introduction

Heavy metals are highly toxic and dangerous pollutants. While some heavy metal pollutants come from fertilizers and sewage, industrial waste is the main source of heavy metal pollution.<sup>1</sup> As one of the most toxic heavy metals, mercury with the feature of strong toxicity can cause serious human health problems even at very low concentrations.<sup>2</sup> Mercury can enter the food chain, posing a threat to human health. The exposure to mercury can cause a number of toxicological effects such as brain damage, kidney failure, and motion disorders.<sup>3</sup> Recognition of the toxic effects from minute concentrations of mercury has made it essential to monitor the level of mercury in aquatic environments. Traditional methods for  $Hg^{2+}$  quantification, include inductively coupled plasma mass spectrometry (ICP-MS), cold-vapor atomic fluorescence/absorption spectrometry (CV-AFS/CV-AAS) and ultraviolet-visible spectrometry, *etc.* are very sensitive and selective but require sophisticated and expensive instruments operated by professionals in a centralized laboratory. In order to overcome these drawbacks, it is still of great challenge to develop a new strategy for facile and sensitive detection of  $Hg^{2+}$ .

On the other hand,  $Hg^{2+}$  can bind directly to  $N_3$  of thymidine (T) in place of the imino proton and bridge two thymidine

residues to form a T-Hg-T base pair. The  $Hg^{2+}$ -mediated T-Hg<sup>2+</sup>-T pair is even more stable than the Watson-Crick A-T pair, without affecting the double-helical structure.<sup>4</sup> Because of this property, many DNA sensors for  $Hg^{2+}$  detection based on fluorescence, colorimetry, electrochemistry, and resonance scattering have been constructed by rational design of DNA sequences.<sup>4-8</sup>

To realize ultrasensitive detection of these trace amounts of  $Hg^{2+}$ , effective signal amplifications are commonly required. In recent years, the polymerase chain reaction (PCR)-like amplification strategies, including isothermal exponential amplification reaction (EXPAR), rolling circle amplification (RCA)<sup>9</sup> and ligase chain reaction,<sup>10</sup> have been incorporated into the biosensor strategy to achieve highly sensitive detection of targets. Although high sensitivity can be obtained by these PCR-like amplification strategies, these sensing strategies are limited due to the drawbacks of complex handing operation, easy contamination and longtime reaction.<sup>11-13</sup> On the other hand, a new class of signal amplification strategies based on nuclease (exonuclease or endonuclease)-assisted target recycling has been reported. The nuclease with sequence specific activity can degrade the specific probe to release the target DNA strands to achieve target recycling amplification.<sup>14-18</sup> This signal amplification strategy has been demonstrated to be useful in detection trace levels of proteins molecules or DNA targets. Unlike endonuclease which requires specific sequences for recognition, exonuclease has received increasing interest in target recycling-amplified DNA detection. For example,

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exonuclease III (Exo III), which catalyzed the stepwise removal of mononucleotides from 3'-termini, has been successfully used for amplifying detection of DNA targets<sup>13,15,19,20</sup> small molecules and proteins.<sup>21,22</sup> Despite the wide application of the nuclease-assisted recycling amplification in sensitive target detection, the extension of this technique for metal detection remains a major challenge because Exo III shows activity only on dsDNA strands.

Herein, a simple, highly sensitive and selective Exo III-based fluorescent sensor has been developed for  $Hg^{2+}$  detection without the need of complicated, high-cost and time-consuming labeling. Also, this method can be devised utilizing both the advantages of label-free architectures and signal-on architectures.

## Experimental

### Materials and chemicals

Oligonucleotides (guaranteed oligos, HPLC-purified) were purchased from Genscript Co., Ltd. (Nanjing, China). The sequence of the probe is as follows: DNA1: 5'-TTTGGGTAGGGCGGGTGGGTACCCAAAATTCTTCTT-3'; DNA2: 5'-TTGTTTGT-3'. Deionized (D. I.) water produced by a Milli-Q Millipore system (18.2 MΩ cm, Millipore Corp., Billerica, MA) was used for the preparation of all the solutions. ABTS (2,2-azinobis (3-ethylbenzothiazoline)-6-sulfonic acid), hemin,  $H_2O_2$ , and HEPES (4-(2-hydroxyethyl) piperazine-1-ethanesulfonic acid sodium salt) were purchased from Aladdin Reagents (Shanghai, China) and used as supplied. All other chemicals and solvents were obtained from the commercial sources and used directly without any further purification. Hemin stock solution (1 mM) was prepared in dimethylsulfoxide (DMSO) and stored at -20 °C. ABTS<sup>2+</sup> and  $H_2O_2$  solution were freshly prepared before used.

### Amplification reaction

Firstly, the mixture of DNA1 (2  $\mu$ M, 10  $\mu$ L), DNA2 (2  $\mu$ M, 10  $\mu$ L) and different concentration (0, 300, 500, 700, 800, 1000, 1500, 3000, and 5000 pM) of  $Hg^{2+}$  in 10 mM phosphate buffer solution (PBS, 0.1 M NaCl, pH 7.4) in a microtest tube incubated in room temperature for about 20 min to form the stem-loop structure DNA. Subsequently, the solution was incubated with 30  $\mu$ L of 10  $\times$  NEBuffer solution containing 20 U Exo III at 37 °C for 20 min. Finally, the resulting mixtures were heated at 90 °C for 10 min to deactivate the Exo III. For control experiment, the  $Pb^{2+}$ ,  $Cu^{2+}$ ,  $Cd^{2+}$ , and  $Zn^{2+}$  were used.

### Gel electrophoresis

Denaturing polyacrylamide gel electrophoresis (PAGE) (20%) was carried out in 1  $\times$  TBE buffer at a 100 V constant voltage for 1 h. The mixture of DNA1, DNA2 and  $Hg^{2+}$  was incubated in Exo III solution (20 U) at 37 °C for 1 h. Afterword, digestions were stopped by heating at 90 °C for 20 min. Then, the resulting mixtures were heated at 90 °C for 10 min to deactivate the Exo III. The gel was taken photograph under UV light after staining with EtBr for 15 min. The concentration of each DNA is 1  $\mu$ M.

### Colorimetric measurement

The above mentioned solutions were moved to 96-well microtiter plates. 4  $\mu$ L of hemin (10  $\mu$ M) and 126  $\mu$ L of HEPES buffer (25 mM HEPES, 20 mM KCl, 200 mM NaCl, 0.05% Triton X-100, and 1% DMSO, pH 7.4) were then added, and the mixture was incubated for 60 min at room temperature. ABTS<sup>2+</sup> and  $H_2O_2$  (20  $\mu$ L in total) were added to the mixture to reach final concentrations of 6 mM and 2 mM, respectively. Absorption intensity of the solutions were taken after 5 min of color development.

### Instrumentation

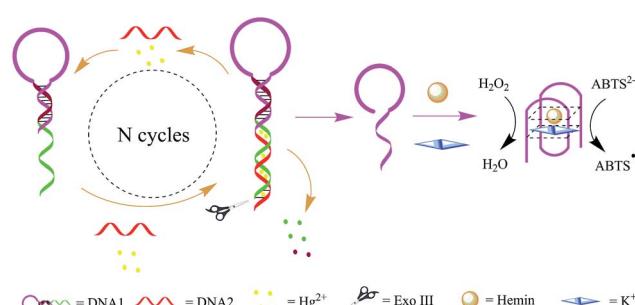
The absorption intensities were recorded by a SpectraMax M5 Multi-Mode Microplate Readers (Molecular Devices, USA), using a Costar 96 well microtiter plate (No. 3599, Corning, New York, USA).

## Results and discussion

### Mechanism of this strategy

The working principle of the designed Exo III amplification and DNAzyme canalization based colorimetric biosensor is illustrated in Scheme 1. The system mainly consists of DNA1 and DNA2. DNA1 includes three domains: the classical G-rich sequence which can form a G-quadruplex in the presence of hemin and  $K^+$ , the sequence that is complementary to part of G-rich sequence and the sequence that is complementary to DNA2 at the help of  $Hg^{2+}$ .

DNA1 was prepared with one DNA strand which designed as a quadruplex-forming oligomer, and which was constructed as 3'-protruding to ensure the probe cannot be digested by Exo III since Exo III only cleaves dsDNA strands with blunt of 3'-recessing termini. However, when the target  $Hg^{2+}$  is introduced into the sensing system, it associates with DNA1 and DNA2 to form double strand DNAs (DNA1/DNA2 duplexes) which have a blunt 3'-terminus. Exo III then can degrade part of DNA1, releasing the DNA1 and  $Hg^{2+}$ , and ultimately liberating the G-rich oligomer. This released G-rich oligomer folds into a G-quadruplex structure and thus allows the formation of DNAzyme in the presence of hemin. The formed DNAzyme can effectively catalyze the  $H_2O_2$ -mediated oxidation of ABTS ( $\lambda_{max} = 418$  nm), giving rise to a change in solution color. The

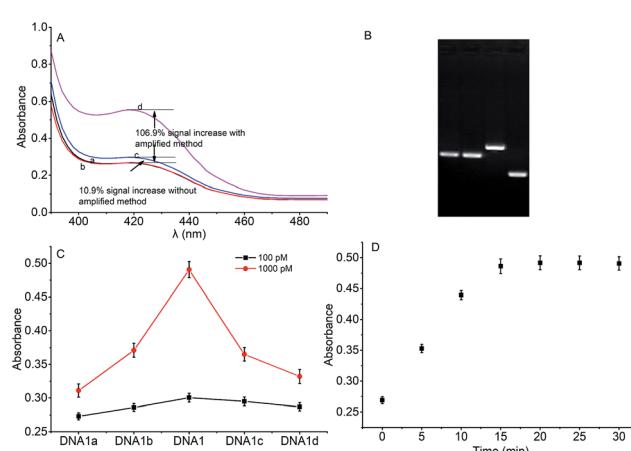


Scheme 1 Schematic illustration of the Exo III and G-quadruplex-based strategy for the detection of  $Hg^{2+}$ .

released DNA1 and  $\text{Hg}^{2+}$  are free to bind to another 3'-protruding terminus of DNA1 to trigger a new digestion reaction, leading to significant amplification of the signal.

### Feasibility study

To verify the feasibility of quantitative detection, the UV-vis absorption intensity changes under different conditions were investigated. As shown in Fig. 1, the absorption of the mixture of DNA1 and DNA2 (curve a), and the mixture of DNA1 and DNA2 with  $\text{Hg}^{2+}$  (3000 pM) (curve b) was relatively low. Upon addition of Exo III to the DNA1/DNA2 mixture without  $\text{Hg}^{2+}$ , the absorption intensity increase was only 10.9% (curve c). However, when both  $\text{Hg}^{2+}$  (3000 pM) and Exo III were present in the solution, we observed 106.9% increase in the absorption intensity (curve d). The amplification reactions were further verified by using denaturing polyacrylamide gel electrophoresis (PAGE), and the results showed that  $\text{Hg}^{2+}$  and Exo III can start amplification reaction and produce G-rich oligomer (line 1, product c).



**Fig. 1** (A) The UV-vis absorption intensity under different conditions: (a) DNA1 and DNA2; (b) DNA1, DNA2 and 3000 pM  $\text{Hg}^{2+}$ ; (c) DNA1 and DNA2 with Exo III; (d) DNA1, DNA2 and 3000 pM  $\text{Hg}^{2+}$  with Exo III. (B) Denaturing polyacrylamide gel electrophoresis characterization of the products by the Exo III assisted amplification method. Lanes (1) DNA1, (2) DNA1 and DNA2, (3) DNA1- $\text{Hg}^{2+}$ -DNA2 complex, and (4) DNA1, DNA2 and  $\text{Hg}^{2+}$  incubated with Exo III. Products (a) to (c): (a) DNA1; (b) DNA1- $\text{Hg}^{2+}$ -DNA2 complex; and (c) G-rich oligomer. DNA2 hard to be distinguished in the gel since the sort length (10 bp). (C) The absorption intensity difference of the method fabricated by five T-rich DNA sequences for the detection of  $\text{Hg}^{2+}$  at 100 pM (black points) and 1000 pM (red points), respectively. (D) Optimization of the incubation time of  $\text{Hg}^{2+}$ .

**Table 1** The designed DNA sequence

Define name	Sequence
DNA1	5'-TTTGGGTAGGGCGGGTTGGGTACCCAAA <u>ATTCTTTCTT</u> -3'
DNA2	5'- <u>TTGTTTGT</u> TT -3'
DNA1a	5'-TTTGGGTAGGGCGGGTTGGGTACCCAAA <u>ATTCTTTC</u> -3'
DNA2a	5'- <u>GTTTGTT</u> -3'
DNA1b	5'-TTTGGGTAGGGCGGGTTGGGTACCCAAA <u>ATTCTTTCT</u> -3'
DNA2b	5'- <u>TGTTTGT</u> TT -3'
DNA1c	5'-TTTGGGTAGGGCGGGTTGGGTACCCAAA <u>ATTCTTCTTT</u> -3'
DNA2c	5'- <u>TTTGTGT</u> TT -3'
DNA1d	5'-TTTGGGTAGGGCGGGTTGGGTACCCAAA <u>ATTCTTCTTTT</u> -3'
DNA2d	5'- <u>TTTGTTGT</u> TT -3'



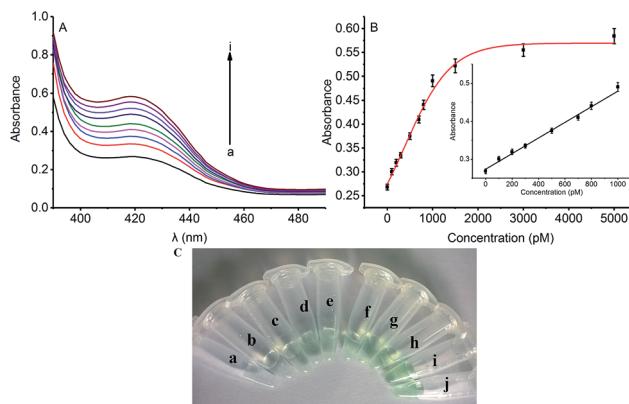


Fig. 2 (A) The UV-vis absorption intensity of the strategy for the assay of  $\text{Hg}^{2+}$  at different concentrations: from (a) to (i): 0, 300, 500, 700, 800, 1000, 1500, 3000, and 5000 pM  $\text{Hg}^{2+}$ , respectively. (B) The relationship between the absorption intensity and the concentrations of the  $\text{Hg}^{2+}$ . The insert is shown with the linear range from 0 to 1000 pM. All the data are taken from independent experiments with repetition for at least three times, and the presented data are the results of averaging. (c) Colorimetric analysis by naked eye with different concentrations  $\text{Hg}^{2+}$ : from (a) to (j): 0, 300, 500, 700, 800, 1000, 1500, 2000, 3000, and 5000 pM.

indicating that the release of the quadruplex-forming oligomer is highly dependent on the concentration of  $\text{Hg}^{2+}$ . Fig. 2B shows the corresponding calibration plot of the concentration of  $\text{Hg}^{2+}$  versus the UV-vis absorption intensity, in which the detection limit was calculated to be 66 pM according to the responses of the blank tests plus 3 times the standard deviation ( $3\sigma$ ). A good linear range from 0 to 1000 pM was shown in the insert of Fig. 2B. The fitting equation of the curve shown in the insert of Fig. 2B is  $Y = 0.00021X + 0.274$  ( $R^2 = 0.991$ ), where  $Y$  is the absorption intensity and  $X$  is the concentration of  $\text{Hg}^{2+}$ . Such low detection limit (66 pM) for  $\text{Hg}^{2+}$  detection is even lower to some existing signal amplification methods that using amplification strategy<sup>14</sup> and nearly 2 orders of magnitude lower to other previously reported  $\text{Hg}^{2+}$  detection systems based on catalytic DNAzyme.<sup>23</sup>

The color change in the presence of various concentrations of  $\text{Hg}^{2+}$  was also monitored by the naked eye (colorimetric analysis) to achieve visual detection. As illustrated in Fig. 2C, the green color of the reaction solution was gradually intensified with elevated concentration of  $\text{Hg}^{2+}$  from 0 to 5000 pM. Also, 500 pM (tube c)  $\text{Hg}^{2+}$  can be easily identified by the naked eye according to the distinct color difference between the sample and the blank test.

The selectivity of the proposed method is examined by monitoring the colorimetric signal of the solution with the presence of  $\text{Hg}^{2+}$  against other metal ions,  $\text{Pb}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Cd}^{2+}$ , and  $\text{Zn}^{2+}$ . As shown in Fig. 3, the presence of the control molecules (1000 pM) causes insignificant colorimetric signal change of the solutions while the addition of the target  $\text{Hg}^{2+}$  (1000 pM) results in a high colorimetric signal of the probe solution, revealing that the amplification method is highly selective. Such high selectivity can be related to the highly specific binding capability of the T- $\text{Hg}^{2+}$ -T. In other words, only the presence of  $\text{Hg}^{2+}$

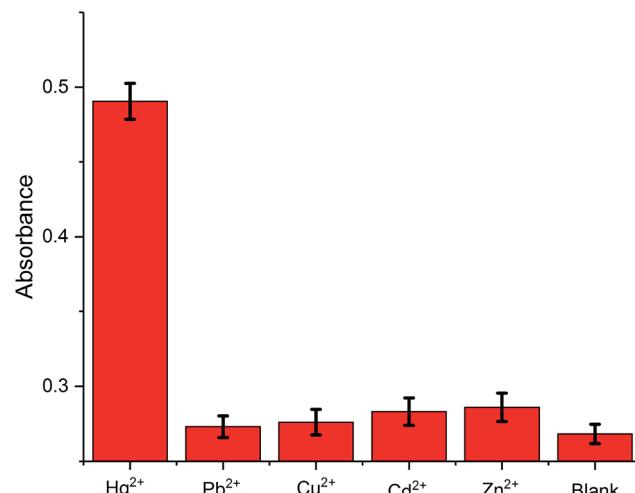


Fig. 3 Selectivity tests of this assay. The concentration of the ions is all 1000 pM.

Table 2 Results of the recovery test of  $\text{Hg}^{2+}$  in river water

Samples	$\text{Hg}^{2+}$ added (pM)	Found (pM)	Recovery (%)
1	200	197.6	$98.8 \pm 2.6$
2	500	492.3	$98.5 \pm 2.3$
3	1000	986.7	$98.7 \pm 3.2$
4	3000	2983.9	$99.5 \pm 1.8$

can trigger amplification cycles to achieve highly visual detection of target.

### The real applicability evaluation

It is critically important to evaluate the real applicability of the amplification method by challenging this strategy in various media. Given that the strategy is resistant to other ions influence, we challenged the assay by analyzing samples of complicated matrix to evaluate the applicability of the strategy to river water. Four concentration of  $\text{Hg}^{2+}$  (200 pM, 500 pM, 1000 pM, 3000 pM) were spiked into river water. Table 2 is shows the experimental results obtained in  $\text{Hg}^{2+}$ -spiked river water samples.  $\text{Hg}^{2+}$  concentration recoveries of 98.5–99.5% were achieved. These results showed that the interference of river water could be overcome. It means that, our proposed strategy could potentially be applied to realistic river water samples.

### Conclusions

In summary, we have developed a novel highly sensitive method to quantify  $\text{Hg}^{2+}$  as low as 66 pM by integrating a recycling cycles and demonstrated its feasibility in the application of realistic detection. We suggest that this assay offers two advantages. First, the assay does not involve any chemical modification of DNA, which makes it simple, low-cost and makes it a label-free strategy. Second, the method does not require troublesome separation procedures. We believe that, this method has the



potential to be used as a major tool for ultrasensitive quantitative analysis of  $Hg^{2+}$  or other heavy metal ions in river water.

## Conflicts of interest

There are no conflicts to declare.

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