



Nanopore sensor for copper ion detection using a polyamine decorated β -cyclodextrin as the recognition element

Yanli Guo,* Feifei Jian and Xiaofeng Kang*

A novel and simple nanopore sensing method has been developed for the detection of Cu^{II} ions using polyamine decorated cyclodextrin as the recognition element. The strong binding affinity between Cu^{II} and the amino groups of cyclodextrin inside an α -hemolysin pore causes the new current blockade events. The event frequency is linear for concentrations of Cu^{II} in the range 0.08–20 μM . The detection limit is as low as 12 nM. More significantly, the sensing system is highly specific for Cu^{II} and does not respond to other metal ions with concentrations up to 10 fold that of Cu^{II} . The applicability of this sensor has also been verified by the analysis of Cu^{II} ions in running water, suggesting the potential application of this sensing system.

Received 11th January 2017
 Accepted 3rd March 2017

DOI: 10.1039/c7ra00454k
rsc.li/rsc-advances

1. Introduction

Nanopore stochastic sensing based on single-molecule recognition is an emerging analytical technique on account of having high sensitivity and being rapid and low-cost.^{1,2} The principle of nanopore sensing is to monitor the ionic current modulation caused by analytes driving through nanopores under a fixed applied voltage.³ The characteristics of current modulation signatures, including frequency, amplitude and dwell time, depend upon the size, charged status and concentration of the analytes. In this way, nanopore sensing has been used successfully to detect a wide variety of substances, ranging from tiny metal ions^{4–6} to organic molecules^{7–9} and even biological macromolecules such as nucleic acids,^{10–12} protein^{13–15} and peptides.^{16,17} Furthermore, protein nanopores have shown attractive prospects as a next-generation DNA sequencing platform.¹⁸

So far, sensing metal ions with an α -hemolysin (α HL) nanopore can be realized in two ways. One is to mutate the protein to construct binding sites in the lumen of α HL. Choi and Mach used an α HL mutant pore with four cysteine mutations to detect Ag^{I} and Cd^{II} .¹⁹ Braha's group reported simultaneous detection of Zn^{II} , Co^{II} and Cd^{II} using an engineered pore containing histidine residues.²⁰ Similar research has been conducted by Kasianowicz *et al.*²¹ Another method is based on the interaction of metal ions and DNA or peptides. Pb^{II} was successfully detected by inducing the conformational change of the G4 DNA aptamer.²² Wen and colleagues designed specific T-

rich DNA oligomers as probes to detect Hg^{II} ions.⁴ Subsequently, they extended this strategy for the detection of Pb^{II} and Ba^{II} ions.⁵ Furthermore, a polyhistidine peptide chain was exploited to detect Cu^{II} based on the chelating reaction by Wang *et al.*²³ However, effective sensors with good selectivity, reproducibility and maneuverability are rare. There are still some problems that remain to be solved. First, it is difficult to operate and imitate for the method of modifying the nanopore interior due to the complexity of mutagenesis and separation of protein nanopore. Second, strong background disturbance exists in detecting low concentration metal ions, limiting the sensitivity of the method. Therefore, there is an upsurge of need in the development of a new approach to overcome these limitations.

In the present work, we used the polyamine decorated cyclodextrins as recognition element in α HL pore for the highly sensitive and selective detection of Cu^{II} ion, which is a well-known heavy metal and plays a vital role in many biological processes.^{24–26} Its concentration will directly affect people's health.^{27,28} As is well known, cyclodextrins could be lodged noncovalently within the lumen of the α HL pore and produced a substantial and incomplete channel block.²⁹ The host-guest inclusion of cyclodextrins with analytes could generate additional transient reductions in the current, which permitted the analytes to be identified and quantified.³⁰ Modified β -cyclodextrins (β CDs) have been also covalently attached to the pore to differentiate deoxynucleotides with over 99% confidence.³¹ Moreover, chiral discrimination was achieved by α HL pore equipped with the β CD adapter.³² However, previous studies involving cyclodextrins have been mainly focused on organic molecules,^{33,34} no attention has been paid to the applications for metal ions. Herein, we detect Cu^{II} for the first time with α HL nanopore containing a functionalized cyclodextrin. In this method, mutating the protein to construct binding sites for the

Key Laboratory of Synthetic and Natural Functional Molecular Chemistry, College of Chemistry & Materials Science, Northwest University, Xi'an 710127, P. R. China.
 E-mail: guoyl@nwu.edu.cn; kangxf@nwu.edu.cn; Fax: +86-029-81535026; Tel: +86-029-81535026



Cu^{II} is not necessary, which simplifies the fabrication of nanopore sensors. More importantly, we validated the practicality of this method for the detection of Cu^{II} in environmental samples through analyses of running water.

2. Experimental section

2.1. Reagents and materials

The (WT)₇, (M113F)₇ and (M113R)₇ α HL protein were synthesized and purified according to the methods reported from documents.^{7,35} All the metal salts, CuCl_2 (>99%), HgCl_2 (>99%), CuCl_2 (>99%), ZnCl_2 (>99%), MgCl_2 (>99%), CdCl_2 (>99%), $\text{Co}(\text{NO}_3)_2$ (>99%), $\text{Ni}(\text{NO}_3)_2$ (>99%), $\text{Mn}(\text{NO}_3)_2$ (>99%), TbCl_3 (>99%), GdCl_3 (>99%), $\text{Dy}(\text{CH}_3\text{COO})_2$ (>99%), purchased from Aladdin and were prepared at concentrations of 10.0 mM each. Heptakis-(6-deoxy-6-amino)- β -cyclodextrin (am₇ β CD) (>99%) was obtained from Cycloab (Budapest, Hungary). Both NaCl (>99%) and Tris (>99%), used to prepare electrolyte solution, were obtained from Kermel Chemical Reagents Co., Ltd. (Tianjin, China). It should be noted that the pH of the electrolyte solutions adjusted by using hydrochloric acid. All reagents were dissolved in ultrapure water. Teflon film (25 μm) was ordered from Goodfellow Corp. (Malvern, PA, USA). The 1,2-diphytanoyl-sn-glycero-3-phosphocholine (DPhPC) used for planar bilayer lipid formation was gained from Avanti Polar Lipids (Alabaster, AL, USA). Running water sample was collected from Northwest University (Xian, China). The sample was filtered through a 0.22 μm membrane (Shanghai Xin Ya Purification Equipment Co., Ltd., China) prior to the detection.

2.2. Single channel recording

Single channel electrical measurements were carried out by using traditional methods. For simplify, planar lipid bilayer membranes of DPhPC were typically created using the method of Montal and Mueller³⁶ on an aperture 120–150 μm in diameter in a Teflon film which separates two identical compartments. Each compartment was filled with 1500 μL of buffer solution (*cis* buffer and *trans* buffer). Both sides of aperture were pre-treated with a 10% v/v hexadecane/pentane mixture before the addition of buffer solution. Then the compartments were both injected with 900 μL buffer solution to the level just below the aperture and lipid solution was added to buffer. When the pentane was evaporated, solvent-free lipid monolayer formed at the solution-air interface. The remaining 600 μL buffer solution was introduced to compartments until the level of buffer rose above the aperture. Unless otherwise stated, (M113F)₇ pores were added to the grounded *cis* compartment (0.05–0.2 ng mL⁻¹). After the successful insertion of a single α HL pore, copper ions and am₇ β CD were added to the *cis* and *trans* side respectively. The single channel current was detected with two freshly prepared Ag/AgCl electrodes, collected with a patch clamp amplifier (Axopatch 200B, Axon Instruments, Foster city, CA, USA) and filtered with a low-pass Bessel filter with a corner frequency of 5 kHz and then digitized with a Digidata 1440A A/D converter (Axon Instruments) at a sampling frequency of 20 kHz.

2.3. Data analysis

Single channel current recordings were performed and analyzed with pClamp 10.3 (Axon Instruments). Origin 8.5 (Microcal, Northampton, MA) was employed for histogram construction, curve fitting and graph presentation. Both the values of τ_{on} (the mean interevent interval) and τ_{off} (the mean dwell time) for am₇ β CD– Cu^{II} complexes, were obtained from the dwell time histograms by fitting the distributions to single exponential functions by the Levenberg–Marquardt procedure.¹⁶ The current blockades were produced by the fitted Gaussian distributions.

3. Results and discussion

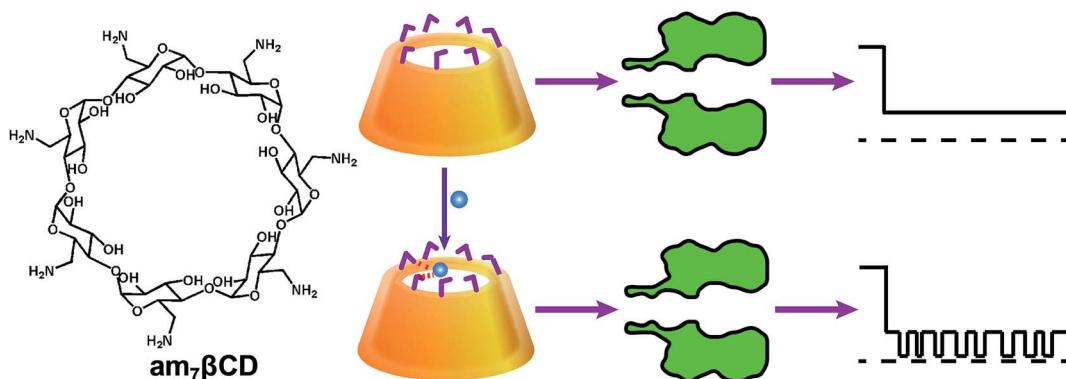
3.1. The principle of the Cu^{II} detection

Inspired by the successful applications of copper complexes of polyamines decorated cyclodextrin³⁷ and the wealth of information about cyclodextrins interacted with α HL nanopore,³⁸ we designed that a simple but efficient Cu^{II} ions sensor which could be operate easily under mild conditions. As shown in Scheme 1, we used β -cyclodextrin with the seven primary hydroxyls replaced with amino groups (heptakis-(6-deoxy-6-amino)- β -cyclodextrin; am₇ β CD) as recognition element, which possess a range of nitrogen donor atoms and are very effective ligands for Cu^{II} ions with high specificities. Without Cu^{II} ions, am₇ β CD adapter which entered into the lumen of the α HL pore only produced one kind of events. In sharp contrast, upon addition of Cu^{II} ions to electrolyte solution, they would coordinate with am₇ β CD molecules and form am₇ β CD– Cu^{II} complexes, which resulted in new additional current blockade events having significantly different signatures from those in the absence of Cu^{II} ions. It permitted the Cu^{II} ions to be readily recognized.

To demonstrate this hypothesis, our initial experiments were carried out at pH 8 under identical conditions with different α HL nanopore containing an am₇ β CD adapter as the detector. As displayed in Fig. 1, significant differences in event signature were observed before and after addition of Cu^{II} ions for three protein nanopores. It is clear that dwell time of am₇ β CD was quite different (3.97 ± 0.18 ms) in (WT)₇ and (1.03 ± 0.09 ms) in (M113R)₇. In the case of (M113F)₇ pore, the am₇ β CD adapter almost permanently locked into a state of $77.2 \pm 0.5\%$ block. It has been reported that sensing with molecular adapter would be enhanced if the adapter did not every so often dissociate from the pore.³⁹ As a result, the (M113F)₇ protein could provide an enhanced resolution for Cu^{II} ions recognition compared with that observed in other α HL pore.

To further prove that the new type of events are related to the formation of am₇ β CD– Cu^{II} complexes, control experiments were examined. Firstly, when EDTA was introduced to the solution, we found that these events originated from am₇ β CD were eliminated, which demonstrated a strong association between Cu^{II} ions and these events (Fig. 2a). Further, am₇ β CD in nanopore stochastic sensing was replaced with natural β CD. The results showed that no additional events was observed in





Scheme 1 Cartoon showing Cu^{II} ions recognition with a protein nanopore using $\text{am}_7\beta\text{CD}$ as recognition element. The $\text{am}_7\beta\text{CD}$ and $\text{am}_7\beta\text{CD}-\text{Cu}^{II}$ complexes with the pore produced events having significantly different signatures, thus permitting them to be readily distinguished. Chemical structure of $\text{am}_7\beta\text{CD}$ molecule was shown in the left.

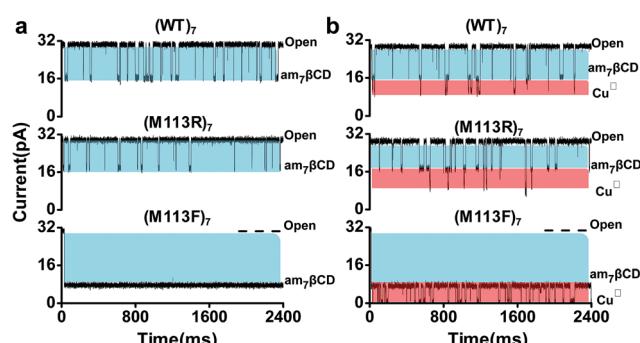


Fig. 1 Typical single-channel current recording traces showing the effect of αHL protein pore on Cu^{II} ions detection. The experiments were performed at 40 mV in three different αHL nanopores containing an $\text{am}_7\beta\text{CD}$ molecule before (a) and after (b) addition of Cu^{II} ions. Conditions: 20 μM Cu^{II} ions added from the *cis* side and 40 μM $\text{am}_7\beta\text{CD}$ from the *trans* side, 1 M NaCl, 10 mM Tris, pH 8.

the current trace, confirming the role that $\text{am}_7\beta\text{CD}-\text{Cu}^{II}$ complexes play in the Cu^{II} ions detection (Fig. 2b).

3.2. Optimization of detection conditions

In order to achieve high sensitivity and selectivity of detection of Cu^{II} ions, the optimal detection conditions are essential. To begin with, we compared the signals where Cu^{II} ions were added from the *trans* side of the bilayer and the case where Cu^{II} ions were added from the *cis* side of the bilayer (Fig. 3). In general, the transport of the charged analytes through the nanopore was dominated by electrophoresis. Since Cu^{II} ions have positive charges, we anticipated the Cu^{II} ions to traverse more easily the nanopore from *trans* to *cis* side instead of from *cis* to *trans* when positive voltage was applied to the *trans* chamber. However, the experimental results showed that the event frequency of Cu^{II} ions from *trans* side was 10 times than from *cis* side, in apparent contradiction with the assumption. In fact, there is a combined action of diffusion, electrophoresis, and electroosmosis in the transport of charged molecules through nanopore. A few experimental studies have demonstrated that the electroosmotic flow (EOF) provided an

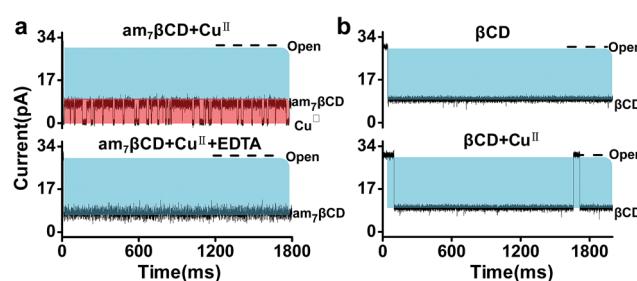


Fig. 2 (a) Typical single-channel current recording traces from a single (M113F)₇ pore at 40 mV before (top) and after (bottom) addition of EDTA. (b) Representative single-channel current traces of βCD in the absence (top) and presence (bottom) of Cu^{II} ions. Conditions: 40 μM $\text{am}_7\beta\text{CD}$ or βCD added from the *trans* side, 20 μM Cu^{II} ions and 20 μM EDTA added from the *cis* side, 1 M NaCl, 10 mM Tris, pH 8.

important or even dominant, contribution to the analytes transport, especially for small neutral molecules^{40,41} or molecules with a low net charge,^{42–44} under appropriate experimental conditions. Our experiment phenomenon also was interpreted as EOF being the main force driving electrically. The αHL protein nanopore exhibits a selectivity to anions⁴⁵ and $\text{am}_7\beta\text{CD}$ molecule is positively charged⁴⁶ at pH 8. Previous studies have demonstrated that anion selective of αHL becomes more anion selective when natural or the positively charged βCD adapter is lodged within the channel lumen.⁴⁷ Therefore, the (M113F)₇ pore equipped with a cationic adapter, $\text{am}_7\beta\text{CD}$, was still anion selective. EOF is consist with the direction of anion flow under applied positive voltages. As a consequence, when Cu^{II} ions were introduced into the *cis* chamber, current blockade events occurred preferentially. While following the direction of cation flow and opposing to EOF, only a few blockades were observed for Cu^{II} ions added from the *trans* side of the bilayer.

As previously documented, the transmembrane voltage is an important factor for determining the translocation of analytes.^{48,49} In addition to providing a novel approach for sensing Cu^{II} ions, our sensor permits the determination of association and dissociation rate constants, separately, and formation constants for $\text{am}_7\beta\text{CD}-\text{Cu}^{II}$ complexes. The rate constants of



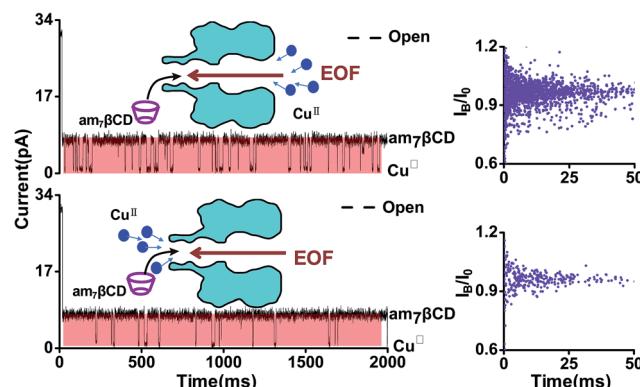


Fig. 3 Representative single-channel current recording traces at 40 mV, in presence of 20 μM Cu^{II} ions added from the *cis*-side (top) or from the *trans*-side (bottom) of the bilayer. The scatter plots of the events (dwell time vs. current blockage (I_8/I_0)) for the corresponding current traces showing in the right. Conditions: 40 μM am₇βCD added into the *trans* chamber, (M113F)₇, 1 M NaCl, 10 mM Tris, pH 8.

am₇βCD–Cu^{II} for association and dissociation were calculated by corresponding formula, that is, $k_{\text{on}} = 1/(\tau_{\text{on}}[\text{Cu}^{\text{II}}])$ and $k_{\text{off}} = 1/\tau_{\text{off}}$. To better illustrate voltage with Cu^{II} ions binding events, we did the experiment in which Cu^{II} ion was kept constant at 20 μM , while transmembrane voltage was changed. As expected, the frequency of event decreased with increasing applied positive voltage, which corresponded to decreases in the association rate constants (Fig. 4a, left). On the contrary, the dissociation rate constant k_{off} increased with voltage, suggesting that the rather high voltage was harmful for the association of Cu^{II} ions (Fig. 4a, middle). Equilibrium formation constants were calculated by using $K_f = k_{\text{on}}/k_{\text{off}}$ for am₇βCD–Cu^{II} and differed by over 10-fold under the current experimental conditions (Fig. 4a, right). The most extreme values were produced at the voltage of 40 mV and 160 mV. For instance, Cu^{II} ions were bound 10.7 times more strongly at 40 mV ($K_f = 1.5 \pm 0.1 \times 10^4 \text{ M}^{-1}$) than at

160 mV ($K_f = 1.4 \pm 0.1 \times 10^3 \text{ M}^{-1}$). These data implied that Cu^{II} binding event was favored at lower values of voltage. So as to improve the Cu^{II} ions detection sensitivity, we used +40 mV as the applied voltage in the remaining experiments.

Based on the fact that certain properties of the α HL protein pore are dependent, to a great extent, on pH, including the ion selectivity, conductance, charge state and magnitude of single-channel noise.^{49–51} Thus, we carried out the Cu^{II} ion detection experiments with am₇βCD at three different pH values (Fig. 4b). The results showed that, in the absence of Cu^{II} ions, only one current blockade level was observed at pH 8 and 10. And there was no additional background noise arised from the am₇βCD state, indicating that am₇βCD was firmly held in the β barrel of protein nanopore under these conditions. In contrast, there were a lot of substates during occupied by am₇βCD in pH 6 buffer solution, which might be attributed to the conformational changes of am₇βCD in the lumen of the α HL pore.³⁸ On the other hand, the addition of 20 μM Cu^{II} ions to the buffer solution of pH 8 produced a large number of markedly new blockade events having a mean residual current of $0.49 \pm 0.12 \text{ pA}$ and a mean dwell time of $14.82 \pm 1.13 \text{ ms}$. By contrast, no additional current block was detected when Cu^{II} ions was applied to the buffer solution, whatever the buffer solution pH 6 or 10. Apart from the interference from the background noise, the possible reason why the event signatures of Cu^{II} ions were not detected at pH = 6 was that amino groups of the cyclodextrin were completely protonated.⁴⁶ When the buffer solution pH increased to 10, Cu^{II} ions were almost hydrolysis entirely. Therefore, the electrolyte solution of pH 8 was selected for the following experiments.

3.3. Detection sensitivity and selectivity for Cu^{II}

To test the sensitivity of this sensor, Cu^{II} ions at various concentrations were examined under the optimum experimental conditions. As illustrated in Fig. 5 and 6a, the event frequency increased with increasing concentration of added

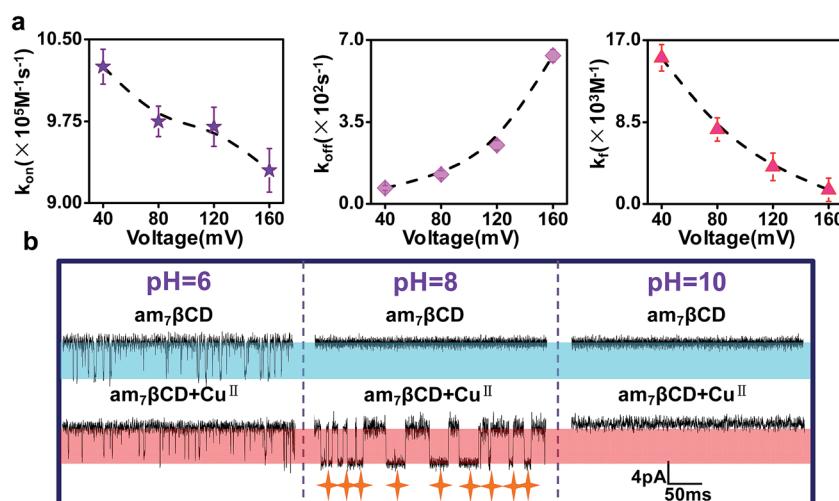


Fig. 4 (a) Dependence of the kinetic constants for the interaction of Cu^{II} with am₇βCD on voltage at pH 8, (left) k_{on} ; (middle) k_{off} ; (right) K_f . (b) Typical single-channel current recording traces at 40 mV, showing the effect of solution pH on Cu^{II} ions detection. The experiments were performed with (M113F)₇ nanopore in a buffer solution comprising 1 M NaCl and 10 mM Tris. Conditions: 20 μM Cu^{II} ions added from the *cis* side and 40 μM am₇βCD from the *trans* side.



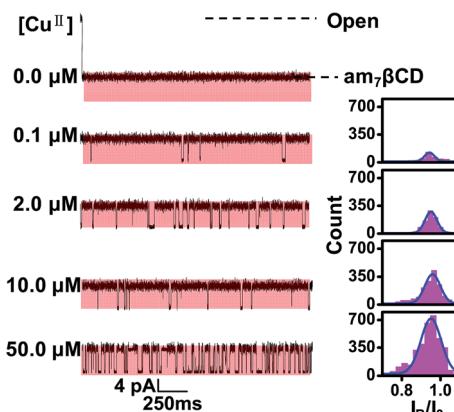


Fig. 5 Nanopore detection of Cu^{II} ions using $\text{am}_7\beta\text{CD}$ molecule as recognition element. Left panel: representative single channel recordings with the $(\text{M113F})_7$ nanopore of Cu^{II} ions at various concentrations in the presence of 40 μM $\text{am}_7\beta\text{CD}$. Right panel: the corresponding concentration-dependent event amplitude histograms. Conditions: Cu^{II} ions added from the *cis* side and $\text{am}_7\beta\text{CD}$ from the *trans* side, 1 M NaCl, 10 mM Tris, pH 8, +40 mV.

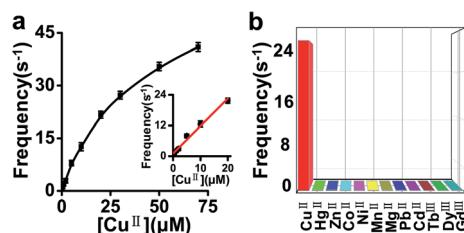


Fig. 6 (a) Dose response curve and (b) selectivity for the Cu^{II} ions nanopore sensor system. The inset of figure (a) shows an enlarged portion of the dose response curve at a range of low Cu^{II} ions concentrations. The experiments were performed at +40 mV in the presence of 40 μM $\text{am}_7\beta\text{CD}$. Conditions: 200 μM interfering ions and various concentration Cu^{II} ions added from the *cis* side and 40 μM $\text{am}_7\beta\text{CD}$ from the *trans* side, $(\text{M113F})_7$, 1 M NaCl, 10 mM Tris, pH 8.

Cu^{II} ions. Linear regression analysis showed good linearity between the frequency and Cu^{II} concentration ranging from 80 nM to 20 μM with a correlation coefficient of 0.98 (Fig. 6a, inset). The detection limit of Cu^{II} ions could be as low as 12 nM ($S/N = 3$), which was lower than previously reported values ($LOD = 40$ nM).²³ Subsequently, under the same conditions, we evaluated the selectivity of this sensing platform toward Cu^{II} (20 μM) relative to other kinds of metal ions including Pb^{II} , Cd^{II} , Mg^{II} , Co^{II} , Mn^{II} , Ni^{II} , Zn^{II} , Hg^{II} , Dy^{III} , Gd^{III} and Tb^{III} (each 200 μM). These metal ions, especially Co^{II} , Ni^{II} , Zn^{II} ions, were well known to be able to interact with amino groups. Apparently, the results revealed that no obvious response could be observed upon the addition of other ions (Fig. 6b). Compared with other nanopore sensors for metal ions, our sensor is more selective toward Cu^{II} ions over the tested interference ions.

3.4. Detection of Cu^{II} in running water

To the end, after a number of screenings of experimental conditions, the applicability of this methodology for detecting

Table 1 Results of the detection of Cu^{II} in real sample

Sample	Added Cu^{II} (μM)	Measured (μM)	Recovery (%)
Running water 1	0	0.35	—
Running water 2	0.50	0.82	91.4
Running water 3	2.00	2.37	105.7
Running water 4	4.00	4.38	108.6

Cu^{II} ions in a real sample was further evaluated. The content of Cu^{II} ions in the sample of running water were detected and the concentration is $0.36 \pm 0.05 \mu\text{M}$. What's more, as shown in Table 1, the measured values for samples with known amounts of Cu^{II} ions showed good recoveries of 91.4–108.6%. The results demonstrated that this nanopore sensor had strong anti-interference ability and could be applied for the detection in real samples.

4. Conclusions

In summary, we have developed a simple, selective and sensitive nanopore sensor for detection of Cu^{II} ions by employing $\text{am}_7\beta\text{CD}$ molecule as recognition element. Based on the chelating reaction between $\text{am}_7\beta\text{CD}$ and Cu^{II} ions, this sensor has been successfully applied to determination of a wide range of Cu^{II} ions from 80 nM to 20 μM . Compared with other nanopore sensors, our sensing system exhibits relatively low detection limit and better selectivity for Cu^{II} ions. More importantly, analyses of running water samples revealed that this approach has potential for detection of Cu^{II} ions in real environmental samples.

Acknowledgements

This work was supported by the National Nature Science Foundation of China (No. 21175105, 21375104 and 21327806), the Specialized Research Fund for the Doctoral Program of Higher Education of China (No. 20126101110015) and the Natural Science Foundation of Shaanxi Province of China (No. 2014JM2050).

References

- 1 J. E. Reiner, A. Balijepalli, J. W. Robertson, J. Campbell, J. Suehle and J. J. Kasianowicz, *Chem. Rev.*, 2012, **112**, 6431–6451.
- 2 B. M. Venkatesan and R. Bashir, *Nat. Nanotechnol.*, 2011, **6**, 615–624.
- 3 J. J. Kasianowicz, E. Brandin, D. Branton and D. W. Deamer, *Proc. Natl. Acad. Sci.*, 1996, **93**, 13770–13773.
- 4 S. Wen, T. Zeng, L. Liu, K. Zhao, Y. Zhao, X. Liu and H. C. Wu, *J. Am. Chem. Soc.*, 2011, **133**, 18312–18317.
- 5 C. Yang, L. Liu, T. Zeng, D. Yang, Z. Yao, Y. Zhao and H. C. Wu, *Anal. Chem.*, 2013, **85**, 7302–7307.



6 G. Liu, L. Zhang, D. Dong, Y. Liu and J. Li, *Anal. Methods*, 2016, **8**, 7040–7046.

7 S. Cheley, L. Q. Gu and H. Bayley, *Chem. Biol.*, 2002, **9**, 829–838.

8 A. J. Boersma, K. L. Brain and H. Bayley, *ACS Nano*, 2012, **6**, 5304–5308.

9 H. C. Wu and H. Bayley, *J. Am. Chem. Soc.*, 2008, **130**, 6813–6819.

10 M. Ayub, S. W. Hardwick, B. F. Luisi and H. Bayley, *Nano Lett.*, 2013, **13**, 6144–6150.

11 Y. L. Ying, J. Zhang, R. Gao and Y. T. Long, *Angew. Chem., Int. Ed.*, 2013, **52**, 13154–13161.

12 A. Meller, L. Nivon, E. Brandin, J. Golovchenko and D. Branton, *Proc. Natl. Acad. Sci.*, 2000, **97**, 1079–1084.

13 J. Nivala, L. Mulroney, G. Li, J. Schreiber and M. Akeson, *ACS Nano*, 2014, **8**, 12365–12375.

14 M. Pastorizagallego, M. F. Breton, F. Discala, L. Auvray, J. M. Betton and J. Pelta, *ACS Nano*, 2014, **8**, 11350–11360.

15 D. Rotem, L. Jayasinghe, M. Salichou and H. Bayley, *J. Am. Chem. Soc.*, 2012, **134**, 2781–2787.

16 L. Movileanu, J. P. Schmittschmitt, J. M. Scholtz and H. Bayley, *Biophys. J.*, 2005, **89**, 1030–1045.

17 H. Y. Wang, Y. L. Ying, Y. Li, H. B. Kraatz and Y. T. Long, *Anal. Chem.*, 2011, **83**, 1746–1752.

18 S. Cornelis, Y. Gansemans, L. Deleye, D. Deforce and N. F. Van, *Sci. Rep.*, 2017, **7**, 41759.

19 L. S. Choi, T. Mach and H. Bayley, *Biophys. J.*, 2013, **105**, 356–364.

20 O. Braha, L. Q. Gu, L. Zhou, X. Lu, S. Cheley and H. Bayley, *Nat. Biotechnol.*, 2000, **18**, 1005–1007.

21 J. Kasianowicz, B. Walker, M. Krishnasastri and H. Bayley, *MRS Online Proc. Libr.*, 1993, **330**, 217–223.

22 H. Y. Wang, Z. Y. Song, H. S. Zhang and S. P. Chen, *Microchim. Acta*, 2016, **183**, 1003–1010.

23 G. Wang, L. Wang, Y. Han, S. Zhou and X. Guan, *Biosens. Bioelectron.*, 2014, **53**, 453–458.

24 A. Badarau and C. Dennison, *J. Am. Chem. Soc.*, 2011, **133**, 2983–2988.

25 R. V. Rathod, S. Bera, S. Man and D. Mondal, *RSC Adv.*, 2016, **6**, 34608–34615.

26 E. Gaggelli, H. Kozlowski, D. Valensin and G. Valensin, *Chem. Rev.*, 2006, **106**, 1995–2044.

27 J. Liu and Y. Lu, *J. Am. Chem. Soc.*, 2007, **129**, 9838–9839.

28 S. Lutsenko, A. Gupta, J. L. Burkhead and V. Zuzel, *Arch. Biochem. Biophys.*, 2008, **476**, 22–32.

29 L. Q. Gu, O. Braha, S. Conlan, S. Cheley and H. Bayley, *Nature*, 1999, **398**, 686–690.

30 Y. Astier, O. Braha and H. Bayley, *J. Am. Chem. Soc.*, 2006, **128**, 1705–1710.

31 J. Clarke, H. C. Wu, L. Jayasinghe, A. Patel, S. Reid and H. Bayley, *Nat. Nanotechnol.*, 2009, **4**, 265–270.

32 X. F. Kang, S. Cheley, X. Y. Guan and H. Bayley, *J. Am. Chem. Soc.*, 2006, **128**, 10684–10685.

33 L. Q. Gu, S. Cheley and H. Bayley, *Science*, 2001, **291**, 636–640.

34 J. Gupta, Q. Zhao, G. Wang, X. Kang and X. Guan, *Sens. Actuators, B*, 2013, **176**, 625–631.

35 L. Wang, Y. Han, S. Zhou, G. Wang and X. Guan, *ACS Appl. Mater. Interfaces*, 2014, **6**, 7334–7339.

36 M. Montal and P. Mueller, *Proc. Natl. Acad. Sci.*, 1972, **69**, 3561–3566.

37 G. Impellizzeri, G. Maccarrone, E. Rizzarelli, G. Vecchio, R. Corradini and R. Marchelli, *Angew. Chem., Int. Ed.*, 1991, **30**, 1348–1349.

38 L. Q. Gu, S. Cheley and H. Bayley, *J. Gen. Physiol.*, 2001, **118**, 481–494.

39 H. C. Wu, Y. Astier, G. Maglia, E. Mikhailova and H. Bayley, *J. Am. Chem. Soc.*, 2007, **129**, 16142–16148.

40 F. Piguet, F. Discala, M. F. Breton, J. Pelta, L. Bacri and A. Oukhaled, *J. Phys. Chem. Lett.*, 2014, **5**, 4362–4367.

41 S. P. Bhamidimarri, J. D. Prajapati, D. B. B. Van, M. Winterhalter and U. Kleinekathöfer, *Biophys. J.*, 2016, **110**, 600–611.

42 L. Mereuta, M. Roy, A. Asandei, J. K. Lee, Y. Park, I. Andricioaei and T. Luchian, *Sci. Rep.*, 2014, **4**, 3885.

43 A. Asandei, M. Chinappi, J. K. Lee, S. C. Ho, L. Mereuta, Y. Park and T. Luchian, *Sci. Rep.*, 2015, **5**, 10419.

44 A. Asandei, I. Schiopu, M. Chinappi, H. S. Chang, Y. Park and T. Luchian, *ACS Appl. Mater. Interfaces*, 2016, **8**, 13166–13179.

45 B. Egwolff, Y. Luo, D. E. Walters and B. Roux, *J. Phys. Chem. B*, 2010, **114**, 2901–2909.

46 B. Hamelin, L. Jullien, F. Guillo, J. M. Lehn, A. Jardy, L. D. Robertis and H. Driguez, *J. Phys. Chem.*, 1995, **99**, 17877–17885.

47 L. Q. Gu, S. M. Dalla, J. B. Vincent, G. Vigh, S. Cheley, O. Braha and H. Bayley, *Proc. Natl. Acad. Sci.*, 2000, **97**, 3959–3964.

48 O. Braha, J. Webb, L. Q. Gu, K. Kim and H. Bayley, *ChemPhysChem*, 2005, **6**, 889–892.

49 L. Q. Gu and H. Bayley, *Biophys. J.*, 2000, **79**, 1967–1975.

50 P. G. Merzlyak, M. F. Capistrano, A. Valeva, J. J. Kasianowicz and O. V. Krasilnikov, *Biophys. J.*, 2005, **89**, 3059–3070.

51 O. V. Krasilnikov, M. P. Capistrano, L. N. Yuldasheva and R. A. Nogueira, *J. Membr. Biol.*, 1997, **156**, 157–172.

