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ACS Sens., **Just Accepted Manuscript** • Publication Date (Web): 04 May 2017

Downloaded from <http://pubs.acs.org> on May 10, 2017

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ACS Publications

ACS Sensors is published by the American Chemical Society, 1155 Sixteenth Street N.W., Washington, DC 20036

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Peptide-mediated nanopore detection of uranyl ions in aqueous media

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ABSTRACT

Uranium is one of the most common radioactive contaminants in the environment. As a major nuclear material in production, the environmental samples (like soil and groundwater) can provide signatures on uranium production activity inside the facility. Thus, developing a new and portable analytical technology for uranium in aqueous media is significant not only for environmental monitoring, but also for non-proliferation. In this work, a label-free method for the detection of uranyl (UO_2^{2+}) ions is developed by monitoring the translocation of a peptide probe in a nanopore. Based on the difference in the number of peptide events in the absence and presence of uranyl ions, nanomolar concentration of UO_2^{2+} ions could be detected in minutes. The method is highly selective; micromolar concentrations of Cd^{2+} , Cu^{2+} , Zn^{2+} , Ni^{2+} , Pb^{2+} , Hg^{2+} , Th^{4+} , Mg^{2+} and Ca^{2+} would not interfere with the detection of UO_2^{2+} ions. In addition, simulated water samples were successfully analyzed.

KEYWORDS

Nanopore sensing; Uranyl ion; Peptide; Biosensor; Chelation

Because of uranium mining, nuclear power production, nuclear weapon development, as well as other industrial and medical application, uranium has been one of the most common radioactive contaminants in the environment, which raises concerns about its environmental impact and risk for human health.^{1,2} For example, UO_2^{2+} can disturb organ function by accumulating in the skeleton, kidneys, lungs, and liver.³⁻⁷ In aqueous solutions, uranium can exist in four different oxidation states such as +3, +4, +5, and +6, but uranyl (UO_2^{2+}) and its complexing ions are the most stable and common species. Thus far, various analytical techniques have been utilized to detect uranyl ions, including radiospectrometry,⁸ inductively coupled plasma mass spectrometry,⁹ fluorescence,¹⁰ colorimetric^{11,12} and complexometric titration¹³. However, most of these techniques are laborious, time-consuming, and require the use of sophisticated instruments or fluorescent labels / dyes. In addition, environmental samples like soil and groundwater near a uranium processing facility can provide significant signatures about its production activities inside the facility, and to detect such early signatures at declared or undeclared areas is significant for preventing nuclear materials from wrong people.¹⁴ Therefore, development of other fundamentally different techniques, which are especially label-free, easy to operate, and potentially field-deployable, for uranyl detection is highly desirable not only for environmental monitoring but also for non-proliferation of nuclear materials or weapons.

Nanopore stochastic sensing has attracted substantial interest as an emerging label-free and amplification-free technique for measuring single molecules.¹⁵⁻¹⁸ Under an applied voltage bias, the movement of an analyte in a nanopore produces a measurable ionic current blockage. The identity and the concentration of the analyte could be revealed from its characteristic current signatures such as the event residence time, amplitude, frequency and even the shape of the blockage.¹⁹ In addition to its biosensing application,²⁰⁻²⁹ nanopore sensing technology has been successfully applied to study a variety of other research areas, for example, covalent and non-covalent bonding interactions,^{30,31} biomolecular folding and unfolding,³²⁻³⁵ and enzyme kinetics.³⁶⁻³⁸ It should be noted that nanopore biosensing is generally achieved by modifying the nanopore interior to introduce binding sites for molecular recognition of target analytes.³⁹⁻⁴³ Recently, Long et al. utilized the effect from electrochemical confined-space to efficiently convert the single DNA/peptide characteristics into measurable electrochemical signatures with high temporal and current resolution, which has been successfully achieved in the study of the heterogeneous structure-function relationship of biomolecules.⁴⁴⁻⁴⁶ More recently, Wang and coworkers reported a coordination chemistry-based stochastic nanopore sensing method for the detection of Cu^{2+} ions by using a peptide as a ligand probe.⁴⁷ The detection was based on the effect of Cu^{2+} on peptide translocation in a nanopore. Briefly, in the absence of Cu^{2+} , the translocation of the copper-chelating agent in the nanopore produced only one major type of events. In contrast, in the presence of Cu^{2+} ions, they interacted with the

copper-chelating agent to form copper chelates, which produced a new type of events in the nanopore. By taking advantage of these new events, quantitative detection of Cu^{2+} ions could be achieved.

In this work, based on the similar metal ion-chelating agent interaction principle but with a different detection mechanism, we developed a new nanopore method for the detection of UO_2^{2+} in aqueous media. We also investigated the effect of metal ions on the detection of UO_2^{2+} . The results demonstrated that this nanopore detection method is highly sensitive and selective to UO_2^{2+} in aqueous media, and the presence of Ni^{2+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Pb^{2+} , Hg^{2+} , Th^{4+} , Ca^{2+} , and Mg^{2+} showed little impact on its detection and quantification.

EXPERIMENTAL SECTION

Chemicals and Reagents. Peptide HH_{14} , a 14-amino acid peptide with a sequence of HHHHHHKHHHYHHH, was obtained from WatsonBio sciences (Houston, TX). Other chemicals such as $\text{UO}_2(\text{NO}_3)_2$ (99.999%), $\text{Ca}(\text{NO}_3)_2$ (99.999%), $\text{Mg}(\text{NO}_3)_2$ (99.999%), $\text{Ni}(\text{NO}_3)_2$ (99.999%), $\text{Zn}(\text{NO}_3)_2$ (99.999%), $\text{Cu}(\text{NO}_3)_2$ (99.999%), $\text{Cd}(\text{NO}_3)_2$ (99.999%), $\text{Pb}(\text{NO}_3)_2$ (99.999%), $\text{Hg}(\text{NO}_3)_2$ (99.999%), $\text{Th}(\text{NO}_3)_4$ (99.999%), NaCl (99.999%), HCl (ACS reagent, ≤ 1 ppm heavy metals), NaH_2PO_4 (BioXtra grade, $\geq 99.5\%$), H_3PO_4 (ACS reagent, $\leq 0.002\%$ heavy metals), $\text{C}_2\text{H}_3\text{NaO}_2$ (BioXtra grade, $\geq 99.0\%$), $\text{CH}_3\text{CO}_2\text{H}$ (ACS reagent, $\geq 99.7\%$), $\text{NaC}_6\text{H}_7\text{O}_7$ ($\geq 99\%$), $\text{C}_6\text{H}_8\text{O}_7$ (FG, $\geq 99.5\%$), and Trizma base (BioXtra grade, $\geq 99.9\%$) were bought from Sigma (St. Louis, MO). All the chemicals, including the HH_{14} peptide, were dissolved in HPLC-grade water (ChromAR, Mallinckrodt Baker). The stock solutions of the peptide and metal salts were prepared at concentrations of 10 mM each, and were kept at -20°C before and after use. The buffer solutions used in this study included: 1) 1.0 M NaCl and 10 mM tris with pH values adjusted to 6.5 and 7.5 using HCl ; 2) 1.0 M NaCl and 10 mM NaH_2PO_4 with pH values adjusted to 4.5 and 5.5 using H_3PO_4 ; 3) 1.0 M NaCl and 10 mM CH_3COONa with pH values adjusted to 4.5 and 5.5 using CH_3COOH ; and 4) 1.0 M NaCl and 10 mM $\text{NaC}_6\text{H}_7\text{O}_7$ “sodium citrate” with pH values adjusted to 4.5 and 5.5 using $\text{C}_6\text{H}_8\text{O}_7$ “citric acid”. Lipid 1,2-diphytanoylphosphatidylcholine was purchased from Avanti Polar Lipids (Alabaster, AL). Teflon film was obtained from Goodfellow (Malvern, PA). The α -hemolysin (αHL) (M113F)₇ protein pores was made according to our previous work.⁴⁸

Electrical Recording and Data Analysis. Single channel recordings were carried out at $24 \pm 1^\circ\text{C}$ in a two-compartment chamber, which is separated by a Teflon septum having a 150 μm diameter aperture (refer to Supporting Information, Fig. S1, for a schematic illustration of the nanopore sensor system). Briefly, the planar bilayer was formed on the aperture of the Teflon film using 1,2-

diphytanoylphosphatidylcholine. Unless otherwise noted, the experiments were performed under symmetrical buffer conditions, with the α HL proteins added to the grounded *cis* compartment, while metal ion salts and the peptide probe were introduced to the *trans* side of the chamber device. Currents were recorded with an Axopatch 200B amplifier (Molecular Devices, Sunnyvale, CA, USA), filtered with a built-in four-pole low-pass Bessel filter at 10 kHz, and then sampled at 50 kHz with a Digidata 1440 A/D converter (Molecular Devices).

The signatures of current blockage events were obtained using Clampfit 10.5 software (Molecular Device). Specifically, the conductance values and the mean residence time (τ_{off}) for the HH₁₄ peptide were derived from the amplitude and the residence time histograms by fitting the distributions to Gaussian and single exponential functions, respectively.³¹ The change (Δn) in the number of peptide HH₁₄ events after addition of metal ions, including UO_2^{2+} , to the solution was calculated by using the equation: $\Delta n = n_0 - n_1$, where n_0 represented the number of HH₁₄ events in the absence of metal ions, while n_1 depicted the number of peptide HH₁₄ events in the presence of metal ions. Therefore, a positive value of Δn indicated a reduction in the number of peptide events after addition of metal ions to the solution. Each single-channel current trace was recorded for 10 minutes. At least three separate experiments, in each of which a new protein nanopore was used, were performed for each sample.

RESULTS AND DISCUSSION

Detection of UO_2^{2+} ions using peptide HH₁₄. Peptides possess a range of potential donor atoms, and are very effective ligands for a variety of metal ions with high specificities. As a noted example, the amyloid beta (A β) peptides, which is crucially involved in Alzheimer's disease, binds Cu^{2+} ions in vitro, and binding results in aggregation of the A β peptide. In particular, it is well documented that the histidine and cystine residues in the peptide display strong affinity for divalent or trivalent metal ions due to the chelation/coordination interaction.⁴⁹ Since uranyl ions themselves could not produce any current modulation events in the nanopore (Supporting Information, Fig. S2), in order to detect UO_2^{2+} , we utilized a 14-amino acid peptide (i.e., HH₁₄) as the ligand probe. The three histidines in the 6-, 13- and 14-positions of the peptide HH₁₄ sequences were designed based on the finding that, in the Cu(II)-A β complex, the Cu^{2+} ions were coordinated by three histidine amino acids (i.e., His-6, His-13 and His-14) in the A β peptide.⁴⁹ The other nine histidines were introduced to increase the coordination possibility between the peptide ligand and the target metal ion. Similar to the Cu^{2+} sensor reported previously, the peptide probe HH₁₄ produced only one major type of events (Fig. 1a). However, unlike the Cu^{2+} sensor, no new events were observed after addition of UO_2^{2+} to the peptide solution. Instead, the number of peptide events decreased. Furthermore, we noticed that, with an increase in the concentration of added

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3 UO_2^{2+} , the peptide events become fewer and fewer. Specifically, when $0.5 \mu\text{M}$ UO_2^{2+} ions were added to
4 the peptide HH_{14} ($40 \mu\text{M}$) solution, the number of peptide events decreased by $56.5 \pm 2.4 \%$ (Fig. 1b). As
5 the concentration of uranyl ions increased to $10 \mu\text{M}$, $92.8 \pm 2.2 \%$ of the HH_{14} peptide events disappeared
6 (Fig. 1c). Since the I-V Curves of HH_{14} , UO_2^{2+} , and their mixtures showed that the existence of uranyl in
7 the nanopore didn't rectify ionic current (Supporting Information, Fig. S3), one possible reason for our
8 observation that addition of UO_2^{2+} to the peptide HH_{14} solution did not produce new types of events, but
9 only decreased the peptide event count is because the interactions between peptide HH_{14} and uranyl ions
10 led to formation of UO_2^{2+} - HH_{14} complexes, which passed through the nanopore too rapidly to be captured
11 by the nanopore sensor ($\sim 200 \mu\text{s}$ resolution). Note that the isoelectric point of histidine is around 7.5,
12 while that of lysine is ~ 9.7 . Therefore, under our experimental conditions, peptide HH_{14} was positively
13 charged. After chelation with UO_2^{2+} , the net positive charge of the peptide-uranyl ion complex increased,
14 and hence, the complex would be electrophoretically driven through the nanopore more rapidly than the
15 uncomplexed peptide. Alternatively, the metal ion-peptide complexes might have larger molecular sizes
16 than the nanopore opening so that they could not enter and pass through the pore. However,
17 stoichiometric consideration of the UO_2^{2+} - HH_{14} interaction could not explain such a large (56.5%)
18 reduction in the peptide HH_{14} events after addition of $0.5 \mu\text{M}$ to $40 \mu\text{M}$ HH_{14} . Furthermore, dynamic
19 light scattering experiment (data not shown) demonstrated that uranyl would not induce HH_{14} aggregation.
20 Therefore, the most likely mechanism behind our finding was that the binding of uranyl to the peptide
21 HH_{14} enabled other uncomplexed peptide molecules to undergo conformational change. It is worth
22 mentioning that disappearance of the biomolecule events or change in its event signatures due to
23 conformational change has been reported previously.^{35,44,50}

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26 **pH effect on the sensitivity of the nanopore sensor.** Nanopore sensor usually has an extremely low
27 background, and hence increasing the peptide HH_{14} event frequency in the nanopore offers the potential
28 to greatly improve the detection limit for UO_2^{2+} quantitation. It has been well documented that the
29 electrolyte pH affects the properties such as conductance and ion selectivity of the protein pores.⁵¹⁻⁵³
30 Therefore, a change in the pH of the electrolyte solution may influence peptide translocation in the
31 nanopore, thus affecting the sensor sensitivity and resolution. Note that, nanopore experiments are usually
32 carried out at/near the physiology pH. In our previous studies, we demonstrated that, as the electrolyte pH
33 decreased, not only the event frequency and residence time of target analytes (e.g., biomolecules and
34 terrorist agents) increased, but the contrast in the event signatures between the target analyte and other
35 matrix components also improved.⁵⁴⁻⁵⁵ On the other hand, it is well known that the coordination properties
36 of a histidine residue within a peptide sequence depend enormously on its position in a peptide chain;
37 further, the metal-peptide complexes formed can exist in a variety of conformations that are dependent
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not only on the concentrations of both the peptide and metal ion but also on the pH of the reaction medium.⁵⁶⁻⁵⁸ To achieve highly sensitive detection of UO_2^{2+} , translocation of peptide HH_{14} in the αHL nanopore was carried out in a series of electrolyte solutions with different pH values (from pH 4.5 to 7.5) and different buffer components. Our experimental results (Fig. 2 and Supporting Information, Table S1) showed that, as the pH value of the electrolyte solution decreased from pH 7.5 to pH 4.5, the frequency and the mean residence time of the peptide events decreased by ~ 10 -folds, and ~ 60 -folds, respectively. The results were not unreasonable considering the net charges of peptide HH_{14} at these various pH values. As discussed in the previous section, the isoelectric point of histidine is ~ 7.5 , while that of lysine is ~ 9.7 . Therefore, in our various investigated buffer solutions of different pH values (from pH 4.5 to pH 7.5), peptide HH_{14} had net positive charges. By systematic calculation of the charge state of HH_{14} (Supporting Information, Table S2), we found that peptide HH_{14} had a +1.05, +4.05, +10.31, +12.69 charge at pH 7.5, pH 6.5, pH 5.5, and pH 4.5, respectively. Therefore, under a positively applied potential bias, in theory, a decrease in the pH of the electrolyte solution would lead to a decrease in the peptide event residence time and an increase in the peptide event frequency. However, due to the resolution of the single channel recording setup, most of the rapid peptide events (e.g., at pH 4.5 and pH 5.5) were missed under our experimental conditions. The electrolyte buffer solution of pH 6.5 rather than pH 7.5 was used in the remaining experiments because a larger percent reduction in the number of peptide events after addition of uranyl ion ($1\ \mu\text{M}$) to the peptide solution ($10\ \mu\text{M}$) was obtained at this pH. Specifically, after addition of uranyl, the number of peptide events was reduced by 83.0% in the pH 6.5 solution compared to 6.2% for the pH 7.5 solution, again suggesting that the net charge of the uranyl-peptide complex played an important role in the disappearance of the biomolecule events.

The effect of voltage bias and peptide concentration on UO_2^{2+} detection. To identify the optimum conditions needed to achieve the maximum nanopore resolution for the detection of UO_2^{2+} , we further investigated the translocation of peptide HH_{14} (without/with UO_2^{2+}) in the nanopore at different voltages, ranging from +60 mV to +140 mV. Our experimental results (Fig. 3) showed that, in the absence of UO_2^{2+} , both the frequency and the blockage amplitude of the peptide events increased, while the peptide event residence time decreased as the applied potential bias increased. After addition of UO_2^{2+} to the peptide HH_{14} solution, the percent reduction in the number of peptide events first increased and then did not change significantly with an increase in the voltage. Although the percent event reduction at +100 mV was slightly smaller than that of +140 mV ($78.9 \pm 5.8\%$ vs. $79.4 \pm 5.7\%$), +100 mV was chosen as the optimum applied potential, and this voltage was used in the remaining experiments because the bilayer at +140 mV was not as stable as that at +100 mV. Note that, the principle for our nanopore sensor to detect UO_2^{2+} was based on the effect of uranyl on the frequency of the peptide HH_{14} events. In order to achieve

highly sensitive detection of UO_2^{2+} , two conditions need to be satisfied: one is a large number of peptide events in the absence of UO_2^{2+} ; and the other is a large percent peptide event reduction in the presence of UO_2^{2+} .

In addition, the effect of the peptide HH_{14} concentration on the nanopore sensor resolution was examined. We found that the number of peptide events was linearly proportional to the peptide concentration, suggesting that the concentration of the peptide would not affect the sensitivity of the nanopore significantly (Supporting Information, Fig. S4). A concentration of 40 μM HH_{14} was used in the remaining experiments since it produced enough events for statistical data analysis within a relatively short recording time.

Sensitivity and selectivity of the UO_2^{2+} nanopore sensor. Utilizing the current physical condition (i.e., pH 6.5, +100 mV applied potential bias, and 40 μM HH_{14} peptide), dose response curve for UO_2^{2+} detection was constructed by monitoring the interaction between peptide HH_{14} and the nanopore in the presence of UO_2^{2+} ions at various concentrations, ranging from 25 nM to 500 nM. Our experimental results showed that the percent peptide event reduction was linearly correlated with the UO_2^{2+} concentration from 25 nM to ~ 200 nM (Fig. 4a). It was found that the detection limit (which is defined as the UO_2^{2+} concentration corresponding to three times the standard deviation of blank signal) in a 10-minute electrical recording was 10 nM, which is more than good enough for analyzing uranyl ion in natural water (note that the maximum contamination level for UO_2^{2+} in drinking water defined by U.S. Environmental Protection Agency is 130 nM).

Nine metal ions, including Ni^{2+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Pb^{2+} , Hg^{2+} , Th^{4+} , Ca^{2+} , and Mg^{2+} , were selected as potential interfering species to examine the selectivity of the nanopore UO_2^{2+} sensor because of their similar chemical properties and/or abundances in water. With the exception of Th^{4+} (5 μM) and Ca^{2+} (500 μM), the concentrations of all the other metal ions used in this investigation were 20 μM each. Our single-channel recording experimental results suggested that these nine metal ions did interact with peptide HH_{14} to form metal-peptide complexes. However, the existence of these cationic species would not affect uranyl ion detection significantly. As shown in Fig. 4b, in the presence of Mg^{2+} , Cd^{2+} , Pb^{2+} , Hg^{2+} , and Th^{4+} , the number of peptide events increased by $0.5 \pm 3.1\%$, $9.7 \pm 1.9\%$, $13.9 \pm 3.8\%$, $7.0 \pm 1.2\%$, and $11.3 \pm 2.6\%$, respectively. Similar to UO_2^{2+} , the existence of Ca^{2+} , Ni^{2+} , Cu^{2+} , or Zn^{2+} ions in the solution led to a decrease in the peptide event count. However, considering that only small event count decreases ($5.1 \pm 1.9\%$, $5.6 \pm 0.9\%$, $2.0 \pm 1.1\%$, and $9.6 \pm 3.7\%$ for Ca^{2+} , Ni^{2+} , Cu^{2+} , and Zn^{2+} , respectively) were obtained in the presence of relatively large (20 to 500 μM) concentrations of

interfering metal ions, the effect is negligible (note that, in comparison, 56.5 ± 2.4 % decreases in the number of peptide events were observed after addition of $0.5 \mu\text{M}$ uranyl ions to the solution). Taking together, the combined results suggest that our nanopore sensor is highly selective to UO_2^{2+} . It should be noted that, histidines can bind to a variety of divalent and trivalent metal ions, and especially display high affinity to Ni^{2+} , Co^{2+} , Cu^{2+} , and Zn^{2+} .^{47,49} The high selectivity of the nanopore to UO_2^{2+} based on unselective histidine-ion coordination suggests that stoichiometry and the strength of the binding affinity between metal ions and peptide HH_{14} were less likely to be the sensing mechanism of the nanopore uranyl sensor, thus favoring our interpretation that addition of UO_2^{2+} to the peptide HH_{14} solution caused significant reduction in the peptide event count was because the binding of uranyl to the peptide HH_{14} enabled other uncomplexed peptide molecules to undergo conformational change.

Effect of the salt gradient on the sensitivity of the UO_2^{2+} nanopore sensor. Use of an asymmetric electrolyte gradient instead of the conventional symmetric electrolyte solution is a well-established approach to significantly increase the event frequency for the translocation of DNA/RNA molecules through a nanopore, thus improving the sensor sensitivity for nucleic acid analysis.⁵⁹ To examine whether this strategy can be employed to improve the sensitivity for UO_2^{2+} detection, two experiments were performed to investigate the translocation of peptide HH_{14} in the nanopore under asymmetric electrolyte conditions. In one experiment, the *cis* chamber compartment contained a solution comprising 1.5 M NaCl and 10 mM Tris (pH 6.5), while the *trans* compartment contained a solution comprising 0.5 M NaCl and 10 mM Tris (pH 6.5). In the other experiment, we increased the salt concentration in the *cis* compartment to 3 M, while maintained the 0.5 M NaCl in the *trans* compartment. Our experimental results (Fig. 5) showed that, with a salt gradient solution instead of the conventional symmetric electrolyte solution, there was indeed a significant increase in the number of peptide translocation events. Specifically, compared with the symmetric electrolyte solution of 1.0 M NaCl (*cis*) / 1.0 M NaCl (*trans*), a salt gradient of 1.5 M NaCl (*cis*) / 0.5 M NaCl (*trans*), and a salt gradient of 3 M NaCl (*cis*) / 0.5 M NaCl (*trans*) resulted in ~ 1.75-fold, and 10-fold increase in the number of peptide events, respectively. It is not unreasonable to expect tens- and even hundreds- fold increase in the peptide event count if a larger salt gradient than that of 3 M NaCl (*cis*) / 0.5 M NaCl (*trans*) is used. As discussed in the voltage effect on HH_{14} translocation, with an increase in the applied potential bias, the number of peptide events increased. Under the symmetric electrolyte solution of 1.0 M NaCl (*cis*) / 1.0 M NaCl (*trans*) and at +100 mV, the detection limit of our nanopore sensor for UO_2^{2+} was 10 nM. By taking advantage of the salt gradient effect and using a larger voltage bias (> 100 mV), the detection limit for UO_2^{2+} would be greatly improved. As a noted example, with a salt gradient of 3 M NaCl (*cis*) / 0.5 M NaCl (*trans*), the detection limit of the nanopore uranyl sensor was 2 nM at +100 mV (Supporting Information, Fig. S5). Using the same salt

gradient but with the applied potential increased to +140 mV, another 4-fold enhancement in the sensor sensitivity could be achieved (Supporting Information, Fig. S6). This sensitivity is comparable with those of other reported highly sensitive electrochemical, optical, and radiospectrometry methods,^{8, 10-12} with detection limits ranging from ~100 pM to 50 nM.

Simulated water sample analysis. To demonstrate the potential application of our nanopore sensor in real-world sample analysis, three simulated uranyl ion-contaminated water samples were created by spiking 100 nM uranyl ions to the tap water (obtained from our life science building), lake water (from Lake of Michigan), and Ice Mountain brand bottled spring water. The simulated water samples were analyzed by our nanopore sensor under the symmetrical buffer conditions. Our experimental results (Fig. 6) showed that the percent event reduction values (22.7 ± 2.3 %, 19.9 ± 0.2 %, and 19.7 ± 1.4 %) of the three simulated water samples were similar to that (19.3 ± 1.8 %) of the control sample (i.e., uranyl ion standard solution), suggesting the matrix component in the water would not affect uranyl ion detection significantly.

CONCLUSIONS

In summary, a highly selective and sensitive nanopore sensor was successfully developed to detect UO_2^{2+} ions by using a peptide molecule as a chelating agent and taking advantage of peptide translocation in the nanopore. Although the formation of UO_2^{2+} -peptide complex did not produce new types of events in the nanopore, the percent reduction in the uncomplexed peptide translocation events could be utilized for UO_2^{2+} quantitation. The high selectivity for UO_2^{2+} of our nanopore sensor was supported by two experiments, i.e., the interference study and simulated water analysis. Our study showed that, in spite of their similar chemical properties and/or large concentration in the real-world samples, metal ions such as Cd^{2+} , Th^{4+} , Cu^{2+} , Zn^{2+} , Ni^{2+} , Pb^{2+} , Hg^{2+} , Mg^{2+} and Ca^{2+} , and other matrix components would not interfere with UO_2^{2+} detection significantly. Our developed nanopore sensor may find useful applications in detection of uranyl ions in natural water for environmental monitoring or for signatures on nuclear material production activity inside a processing facility.

ASSOCIATE CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website. Additional tables and figures, including effect of buffer solution on peptide translocation in the nanopore, net charges of peptide HH_{14} at different pH values, schematic illustration of the uranyl nanopore sensor, typical single-channel recording trace segment of uranyl ions, I-V curves of peptide HH_{14} , UO_2^{2+} and their

mixtures in the nanopore, dose response curve for peptide HH₁₄, dose response curve for uranyl in asymmetric electrolyte conditions, and voltage effect on the sensitivity of the nanopore uranyl sensor.

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Notes

The authors declare no competing financial interests.

ACKNOWLEDGMENT

This work was financially supported by the National Institutes of Health (1R15GM110632).

FIGURES LEGEND

Figure 1. Nanopore detection of UO₂²⁺ ions using peptide HH₁₄. (a) 40 μM HH₁₄; (b) 40 μM HH₁₄ + 0.5 μM UO₂²⁺; and (c) 40 μM HH₁₄ + 10 μM UO₂²⁺. (*Left*) Typical single-channel current recording trace segments; and (*Right*) the corresponding event amplitude histograms. The experiments were performed at +100 mV with the (M113F)₇ αHL pore in an electrolyte solution containing 1.0 M NaCl and 10 mM Tris•HCl (pH 6.5). Both the peptide and uranyl ions were added to the *trans* compartment of the nanopore sensing chamber. Dashed lines represent the levels of zero current.

Figure 2. The effect of electrolyte pH on the mean residence time of the peptide HH₁₄ events. The experiments were performed at +80 mV with the (M113F)₇ αHL nanopore in a series of electrolyte solutions with different pH values and in the presence of 10 μM peptide HH₁₄.

Figure 3. The effect of the applied potential bias on the (a) number of occurrences; (b) blockage amplitude; and (c) residence time of the peptide HH₁₄ events; as well as (d) percent reduction in the number of HH₁₄ events after addition of uranyl ions. The experiments were performed with the (M113F)₇ αHL protein nanopore in an electrolyte solution containing 1 M NaCl and 10 mM Tris (pH 6.5) and in the presence of 10 μM peptide HH₁₄ at various voltages ranging from +60 mV to +140 mV. The concentration of UO₂²⁺ used in Fig. 3d was 0.5 μM.

Figure 4. Sensitivity and selectivity of the UO₂²⁺ nanopore sensor. a) Dose-response curve; and (b) interference study. The experiments were performed at +100 mV with the (M113F)₇ αHL protein

nanopore in an electrolyte solution containing 1 M NaCl and 10 mM Tris (pH 6.5) and in the presence of 40 μ M peptide HH₁₄. With the exception of Th⁴⁺ (5 μ M), Ca²⁺ (500 μ M), and UO₂²⁺ (0.5 μ M), the concentrations of all the other metal ions shown in Fig. 4b were 20 μ M each. In Figs. 4a and 4b, the change (Δn) in the number of peptide HH₁₄ events after addition of UO₂²⁺ to the solution was calculated by using the equation: $\Delta n = n_0 - n_1$, where n_0 represented the number of HH₁₄ events in the absence of uranyl, while n_1 depicted the number of peptide HH₁₄ events in the presence of UO₂²⁺.

Figure 5. Effect of salt gradient on UO₂²⁺ detection. (a) Symmetric electrolyte solution of 1.0 M NaCl and 10 mM Tris•HCl (pH 6.5) (*cis*) / 1.0 M NaCl and 10 mM Tris•HCl (pH 6.5) (*trans*), (b) salt gradient of 1.5 M NaCl and 10 mM Tris•HCl (pH 6.5) (*cis*) / 0.5 M NaCl and 10 mM Tris•HCl (pH 6.5) (*trans*), and (c) salt gradient of 3 M NaCl and 10 mM Tris•HCl (pH 6.5) (*cis*) / 0.5 M NaCl and 10 mM Tris•HCl (pH 6.5) (*trans*). The experiments were performed at +100 mV with the (M113F)₇ α HL nanopore in the presence of 10 μ M peptide HH₁₄, which was added to the *trans* compartment of the nanopore sensing chamber. The event counts in Figs. 5a-c were calculated based on 5 min single channel recording trace segments.

Figure 6. Simulated water sample analysis. The experiments were performed at +100 mV with the (M113F)₇ α HL protein nanopore in an electrolyte solution containing 1 M NaCl and 10 mM Tris (pH 6.5) and in the presence of 40 μ M peptide HH₁₄.

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Figure 1

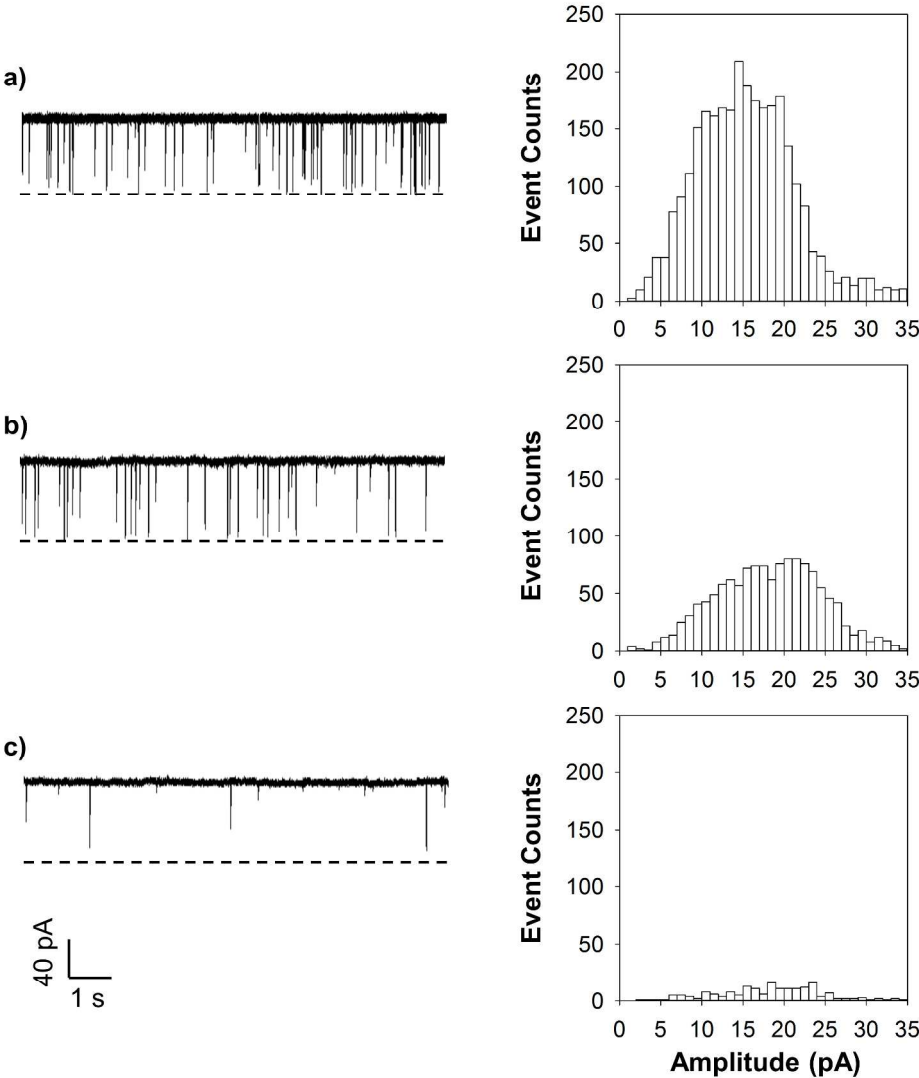


Figure 2

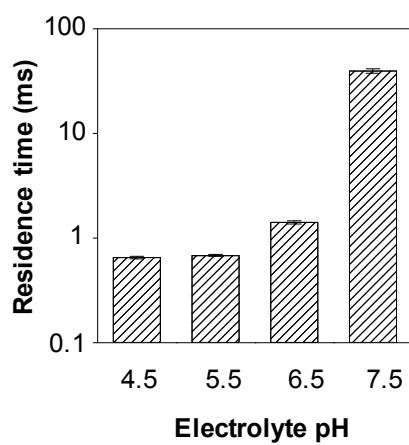


Figure 3

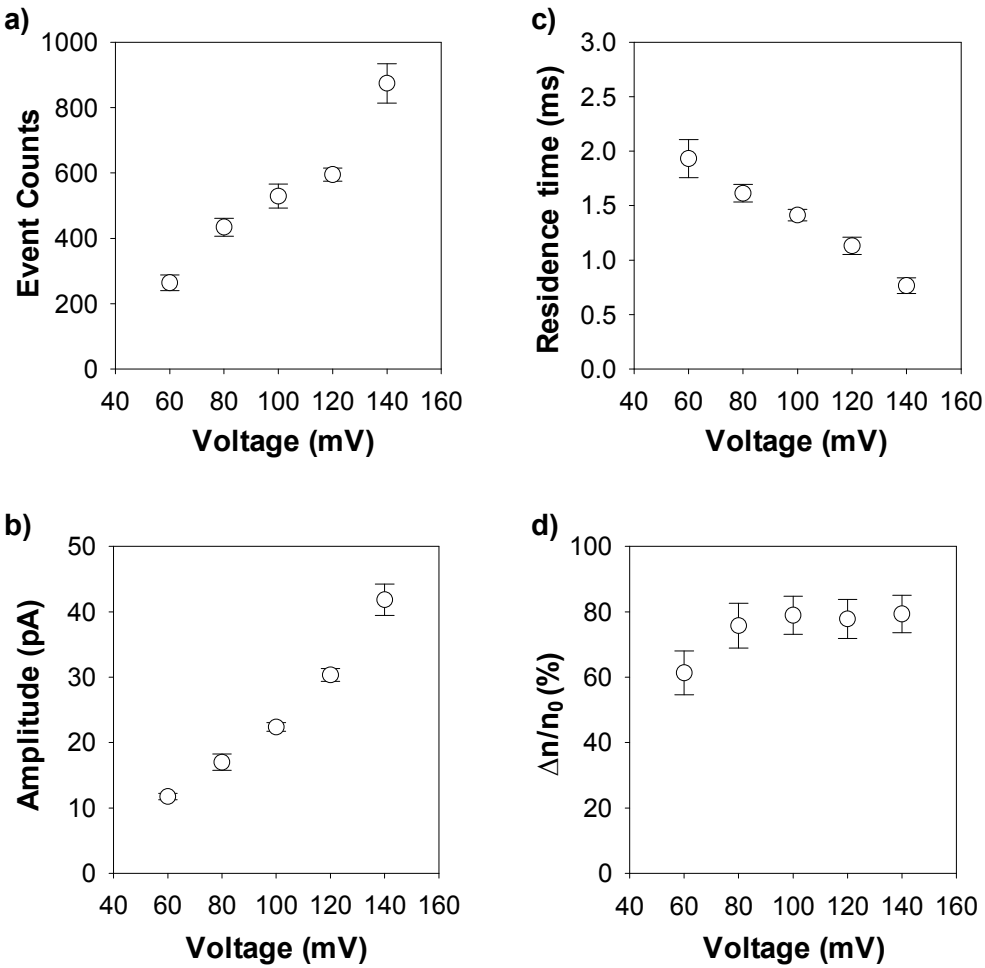


Figure 4

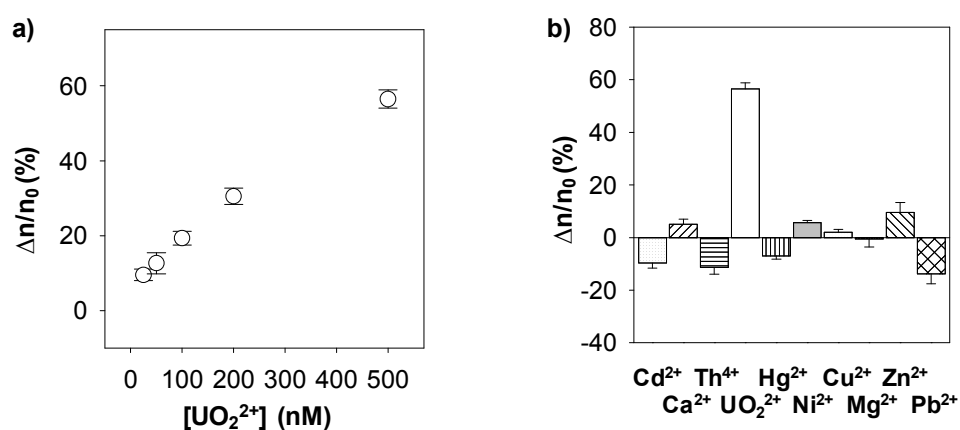


Figure 5

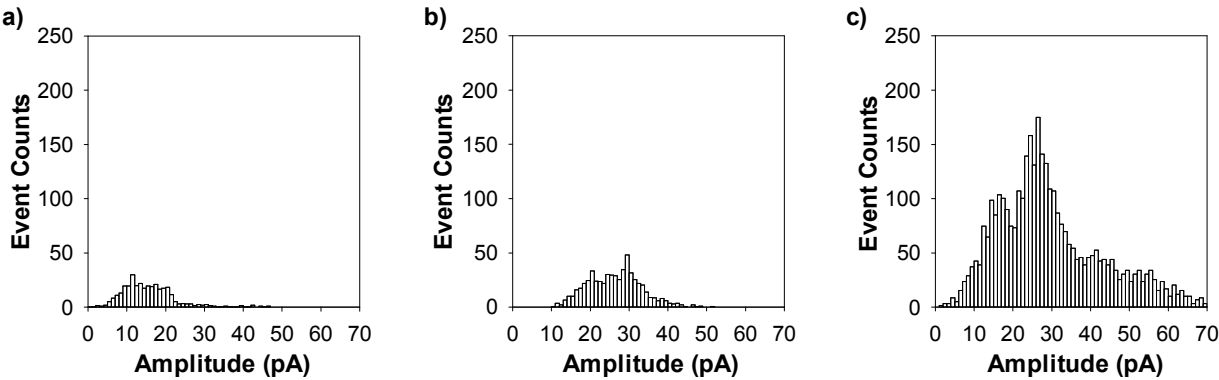
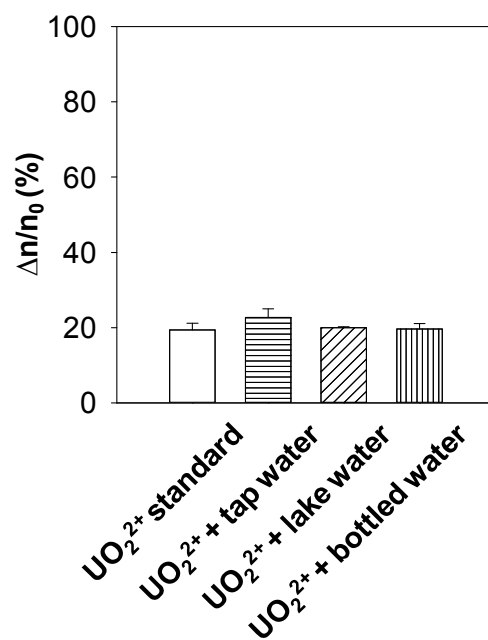


Figure 6



“For TOC Only”

