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Recyclable nanoparticles based on a boronic acid–diol complex for the real-time monitoring of imprinting, molecular recognition and copper ion detection†

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Molecularly imprinted polymers (MIPs) have now become one of the most remarkable materials in the field of molecular recognition. Although many efforts have been made to study the process and mechanism of molecular imprinting, it has not been possible to monitor the interactions between the template and the growing polymer chains under real-time experimental conditions. The behavior of the template–monomer complex during the whole polymerization process has remained largely unknown. In this work, we introduce a fluorescence technique that allows monitoring of the template–functional monomer complex during an actual imprinting process, as well as the real-time signaling of template binding and dissociation from the imprinted polymer. For the first proof-of-principle, we select Alizarin Red S (ARS) and 4-vinylphenylboronic acid as the template and functional monomer, respectively, to synthesize MIP particles *via* precipitation polymerization. As the formation of the template–functional monomer complex leads to strong fluorescence emission, it allows the status of the template binding to be monitored throughout the whole reaction process in real time. Using the same fluorescence technique, the kinetics of template binding and dissociation can be studied directly without particle separation. The hydrophilic MIP particles can be used as a scavenger to remove ARS from water. In addition, the MIP particles can be used as a recyclable sensor to detect Cu ions. As the Cu ion forms a stable complex with ARS, it causes ARS to dissociate from the MIP nanoparticles, leading to effective fluorescence quenching. The non-separation analytical method based on fluorescence measurement provides a convenient means to study molecular imprinting reactions and the kinetics of molecular recognition using imprinted polymers. The recyclable nanoparticle sensor allows toxic Cu ions to be detected directly in water in the range of 0.1–100 μM with a recovery of 84–95%.

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1 Introduction

The molecular imprinting technique provides an effective approach to preparing tailor-made polymer materials with a functional performance similar to antibodies and enzymes.^{1,2} Over the past years, the configurations of molecularly imprinted polymers (MIPs) have shifted from bulk polymers to nanoparticles,^{3,4} and the compatibility of MIPs with aqueous solvents has been improved greatly.^{5,6} These developments, together with advances in post-imprinting modifications,^{7,8}

have enabled MIPs to be applied not only in solid-phase extraction⁹ and chemical sensing¹⁰ but also as substitutes for biological receptors,¹¹ enzyme-like catalysts¹² and drug-delivery vehicles.^{13,14} Fundamental studies on molecular imprinting have contributed to a better understanding of the formation mechanisms of molecular-recognition sites in MIPs, and have provided useful guidance for the design of better-performing MIPs.¹⁴ Most of the studies have been based on *in silico* strategies, electronic structure methods and molecular dynamics simulations to study molecular interactions in pre-polymerization mixtures.^{15,16} In some cases, the results of computational methods have been validated using experimental analysis of the molecular imprinting system.^{17–19} Due to the complicated nature of the radical polymerization process, almost all *in silico* studies have focused on the pre-polymerization mixture, and it is difficult to use computer simulation to study molecular complexes during a real molecular imprinting polymerization process. Except for a few attempts to study the status of a

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that has a temperature controlling system to maintain the cuvette temperature at 40 °C. The fluorescence spectrum was recorded continuously every 1 min for 180 min.

2.4 Binding of ARS by MIP particles

MIP particles were suspended in 20 mM phosphate buffer (pH 8.5) *via* sonication to give a final concentration of 1 mg mL⁻¹. In a series of 2 mL microcentrifuge tubes, the particle suspension (0.5 mL) was mixed with ARS (0.5 mL) of different concentrations prepared in 20 mM phosphate buffer (pH 8.5). The samples were shaken gently using a rocking table at room temperature for 1 h. After this step the samples were centrifuged. The free ARS in the supernatant was measured using UV-vis spectrometry. For comparison, the NIP particles were used as the control and were tested using the same procedure.

2.5 Monitoring the binding and release of ARS

MIP and NIP particle suspensions (0.5 mL, 1 mg mL⁻¹) were transferred to a cuvette equipped with a magnetic stirrer. The cuvette was put into a spectrofluorometer. After adding 0.5 mL of ARS solution (0.2 mM, prepared in 20 mM phosphate buffer, pH 8.5), the fluorescence spectrum of the particle suspension was recorded every 0.5 min for 60 min. After this step, 180 mg of fructose was added to the cuvette, and the fluorescence spectrum was recorded every 1 min for 180 min.

2.6 Detection of Cu²⁺ ions using MIP particles

MIP particles were suspended in 20 mM phosphate buffer (pH 8.5) *via* sonication to give a final particle concentration of 1 mg mL⁻¹. In a series of 2 mL microcentrifuge tubes, 0.5 mL of the particle suspension was mixed with 0.5 mL of CuSO₄ solution. After the samples were shaken gently using a rocking table at room temperature for 2 h, the fluorescence spectra of the samples were recorded.

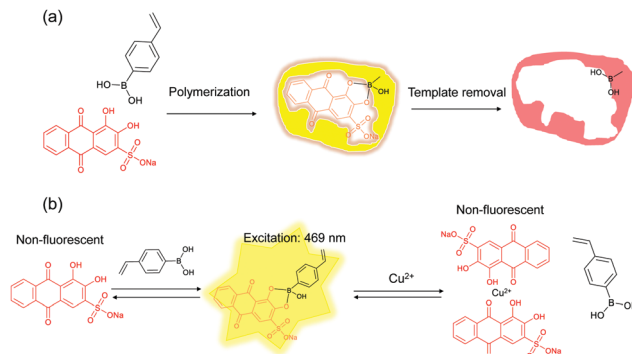
The concentration of Cu²⁺ in spiked rainwater (collected on March 15, 2021 in Lund, Sweden) and in tap water was determined using the above fluorescence measurement method. For comparison, the concentrations of Cu²⁺ in the spiked samples were also measured using the commercial copper assay kit.

2.7 Detection of Cu²⁺ ions using recycled MIP particles

After their first use for the detection of Cu²⁺, the MIP particles were mixed with 20 mL of 0.5% EDTA to remove the Cu²⁺. The particles were then washed with water thoroughly and dried in a vacuum desiccator. The MIP particles were re-suspended in 20 mM phosphate buffer (pH 8.5) to measure the capacity of ARS binding. To evaluate the regenerated MIP particles for the detection of Cu²⁺, the MIP particles (20 mg) were added to 20 mL of 1 mM ARS solution and shaken overnight to load ARS. The particles were washed thoroughly with phosphate buffer (pH 8.5) and dried in a vacuum desiccator before being used to detect Cu²⁺.

3 Results and discussion

The general principle of molecular imprinting is based on the formation of a template–functional monomer complex using a



Scheme 1 (a) Synthesis of ARS-imprinted polymer particles. (b) Formation of fluorescent adduct from the reaction between ARS and VPBA, and quenching of the fluorescence caused by the displacement of VPBA by Cu²⁺.

cross-linking polymerization system.^{38,39} The imprinting effect mainly depends on the strength of the molecular interaction between the functional monomer and the template.²¹ Recently, Piletsky and co-workers proposed imprinting as a result of the dynamic interaction of oligomers and protoparticles with molecular template.⁴⁰ To gain further detailed insight into the mechanism of molecular imprinting, it is essential to have a simple and practical method for monitoring, in real time, the complexation between the template and the functional monomer/oligomer/cross-linked particles. Scheme 1a shows the route for preparing ARS-imprinted polymer particles. VPBA was used as the functional monomer and MBA as the cross-linking monomer. A redox initiator APS/TEMED was used to initiate the polymerization in phosphate buffer containing ethanol. Scheme 1b shows the formation of a fluorescent adduct caused by the template–functional monomer binding, and quenching of the fluorescence caused by the Cu²⁺ ion due to dissociation of the ARS–boronic acid complex.

3.1 Characterization of MIP particles

DLS was used to measure the hydrodynamic diameter of the MIP and NIP particles. DLS measurements were carried out

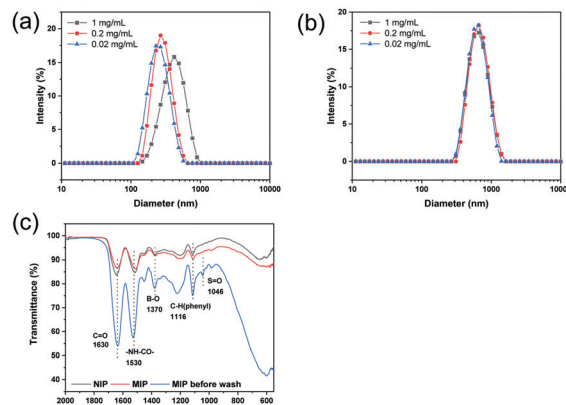


Fig. 1 Hydrodynamic particle size of MIP (a) and NIP (b) measured using DLS in deionized water. (c) FT-IR spectra of NIP particles, and MIP particles before and after washing.



using different particle concentrations. As shown in Fig. 1a and Table S1 (ESI[†]), the diameter of the MIP particles decreased from 428 nm to 263 nm, and the polydispersity index (PDI) decreased from 0.360 to 0.126 when the particle concentration was changed to 0.2 and 0.02 mg mL⁻¹. For the NIP particles in Fig. 1b, the particle diameter was found to be around 650 nm with a PDI of 0.11. From the SEM images (Fig. S1, ESI[†]), it was difficult to determine the size of MIP and NIP particles due to serious aggregation.

In Fig. 1c, the FTIR spectra of the MIP and NIP particles are presented. It is easy to find the characteristic bands corresponding to VPBA and MBA in the polymer samples. The IR band at 1370 cm⁻¹ was assigned to B–O stretching, and the band at 1116 cm⁻¹ (C–H stretching) was caused by the stretching of the benzene ring in VPBA. The characteristic absorbance of the amide group from MBA was found at 1630 cm⁻¹ (C–O stretching) and 1530 cm⁻¹ (–NH–CO– stretching). The above results confirmed that VPBA and MBA have been copolymerized to form the particles. For the MIP particles before washing, an IR band at 1046 cm⁻¹ arising from the S=O group in ARS was observed. This IR band disappeared after the template ARS was removed from the MIP particles.

3.2 Confirmation of molecular imprinting effect

Fig. 2a shows the binding of ARS by MIP and NIP particles measured using different initial ARS concentrations (from 0.02 to 0.1 mM). At all the ARS concentrations, the amount of ARS binding on the MIP particles was higher than on the NIP particles. The maximum capacity of the MIP toward ARS (~50 μmol g⁻¹) was higher than for the NIP (~20 μmol g⁻¹). These results indicate that more molecular recognition sites were created in the MIP by involving ARS in the polymerization process. Using AFA as a structural analogue of ARS and NADH as a *cis*-diol compound, the selectivity of the target binding by the MIP was tested. AFA has a dihydroxyanthraquinone structure and its molecular size is almost the same as ARS. This fact makes AFA a useful reference molecule for evaluating the importance of the *cis*-diol structure for molecular binding to the imprinted cavities. NADH contains a *cis*-diol moiety that is similar to ARS but NADH lacks the dihydroxyanthraquinone structure. Using NADH as a reference molecule allows us to study any non-specific binding caused by random boronic acid



Fig. 2 (a) ARS binding on MIP and NIP particles measured in 20 mM phosphate buffer (pH 8.5). Concentration of MIP particles: 0.5 mg mL⁻¹; incubation time: 1 h. (b) Binding of ARS, AFA and NADH on MIP particles measured in 20 mM phosphate buffer (pH 8.5). Initial concentration of the test compounds: 0.02 mM; concentration of MIP particles: 0.5 mg mL⁻¹.

outside the imprinted cavities. As shown in Fig. 2b, the MIP exhibited no binding for AFA and NADH, indicating that the MIP demonstrates high selectivity for ARS.

3.3 Kinetics of ARS binding studied *via* fluorescence measurements

Based on the fact that the fluorescence intensity increases when ARS forms an adduct with boronic acid, we expected that the kinetic recognition of ARS by MIP particles can be studied using a separation-free fluorescence assay. Therefore, the intensity of the fluorescence of the MIP particles after mixing with ARS was measured at different times, and the result was used to calculate the rate of ARS binding. As shown in Fig. 3a, the fluorescence intensity of the MIP particles increased with time, and eventually reached equilibrium after 60 min of incubation with ARS. Fig. 3b shows clearly that the intensity at 560 nm increased with time and reached equilibrium after 60 min. In the initial stage, the amount of ARS bound increased quickly. After 5 min, the binding of ARS by the MIP increased more slowly. As a comparison, the NIP was measured using the same procedure, and the fluorescence emission also increased after the addition of ARS, and reached equilibrium within 60 min (Fig. 3c and d).

To study the kinetics of ARS binding with the MIP and NIP, the experimental data in Fig. 3 were fitted using a two-phase exponential association model (eqn (1)):^{41,42}

$$I = I_0 + aB_1(1 - e^{-k_{\text{obs}1}t}) + aB_2(1 - e^{-k_{\text{obs}2}t}) \quad (1)$$

where I and I_0 are the fluorescence intensity measured at time t and at 0 min, respectively, a is a constant that relates to the fluorescence intensity to bound ARS, B_1 and B_2 are the amount of ARS bound to the two types of sites, and $k_{\text{obs}1}$ and $k_{\text{obs}2}$ are the observed rate constants for ARS binding. The results of the fitting are presented in Table 1. In the two-phase exponential

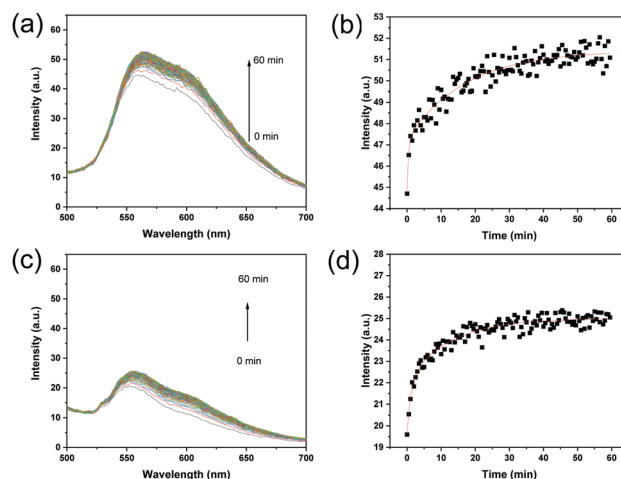


Fig. 3 (a) Fluorescence emission spectra of MIP particles measured at different times after mixing with ARS. (b) Kinetic binding of ARS with MIP particles measured in 20 mM phosphate buffer. (c) Fluorescence emission spectra of NIP particles measured at different times after mixing with ARS. (d) Kinetic binding of ARS with NIP particles measured in 20 mM phosphate buffer. Concentration of MIP particles: 0.5 mg mL⁻¹; concentration of ARS: 0.1 mM.



Table 1 Results of the fitting of kinetic data of ARS binding to polymer particles. Constant a was calculated from the fluorescence emission caused by ARS binding (Fig. 2a)

Polymer	$k_{\text{obs}1}$ (min^{-1})	B_1 ($\mu\text{mol g}^{-1}$)	$k_{\text{obs}2}$ (min^{-1})	B_2 ($\mu\text{mol g}^{-1}$)	a ($\text{g } \mu\text{mol}^{-1}$)
MIP	2.27	19.5	0.058	31.5	0.132
NIP	0.91	9.45	0.0793	11.2	0.262

association model, both the observed rate constant $k_{\text{obs}1}$ (for fast binding site) and the amount of ARS bound to the MIP are much larger than for the NIP, and the ARS for the slow binding site (B_2) of the MIP is also significantly larger than for the NIP.

In Fig. 3c, the fluorescence signal observed from the NIP at the very beginning was caused by the very fast binding of ARS to the polymer particles. As shown in Fig. S2a (ESI[†]), before the addition of ARS, the NIP particles themselves were not fluorescent. The nearly instantaneous emission of fluorescence from the NIP particles after the addition of ARS indicates that the surface-exposed boronic acid on the NIP particles reacted with ARS very rapidly (within a few seconds while the sample cuvette was transferred to the fluorometer). For the MIP particles exposed to ARS, the initial fluorescence intensity was actually slightly lower than MIP particles alone (Fig. S2b, ESI[†]), due to the UV-vis absorption of ARS in the range of 400–600 nm. The background fluorescence of the MIP particles was due to the residual template that has not been completely removed during the particle purification.

3.4 Displacement of ARS from MIP by fructose

Considering the dynamic nature of the boronate ester bond, it should be possible to release the ARS that has been loaded on the MIP particles by introducing a *cis*-diol compound, *e.g.*, fructose. When fructose is added, it is able to form a boronate ester bond with boronic acid because fructose contains a similar *cis*-diol structure to ARS. The formation of the boronic acid–fructose complex causes ARS to dissociate from boronic acid, thereby leading to fluorescence quenching. The displacement of ARS can be monitored by measuring the fluorescence intensity change. In this regard, after the ARS–MIP binding has reached equilibrium, fructose at a high concentration was added to the MIP particle suspension. Fig. 4a shows the fluorescence spectra of

the particles after the addition of fructose. It is obvious that the fluorescence intensity decreased with time and eventually reached equilibrium. While the process of ARS binding to the MIP particles took 60 min to reach equilibrium, the displacement of ARS from the MIP by fructose lasted 180 min. When the binding and displacement curves are overlaid, as shown in Fig. S3a (ESI[†]), it becomes clear that addition of fructose caused the fluorescence intensity to decrease even after the original signal from the MIP particles was reached. The additional ARS released from the MIP particles is due to the residual template that has not been removed during the purification of the initial MIP particles. For the NIP particles, addition of fructose also caused the fluorescence intensity to decrease (Fig. 4b). However, after the original fluorescence intensity was reached, no further signal reduction was observed (Fig. S3b, ESI[†]).

3.5 Monitoring the molecular imprinting process in real time

In a previous study, the nucleation and growth of imprinted polymer particles in precipitation polymerization was monitored using DLS.²¹ The DLS analysis revealed that the nucleation of propranolol-imprinted polymer particles and the formation of imprinted sites occurred in the early stages of the imprinting reaction. In this work, because the fluorescence signals from the ARS–VPBA adduct and the ARS-oligomers containing VPBA can be easily detected, we conceived that it should be possible to monitor in real time the status of the complexation of the template–functional monomer and template–polymer chains during the molecular imprinting process. As shown in Fig. S4 (ESI[†]), when measured separately in the reaction solvent, MBA, VPBA and ARS themselves did not have a fluorescence emission (excitation at 469 nm). A mixture of VPBA and MBA did not show any fluorescence either. As expected, only the mixture composed of VPBA, MBA and ARS showed a clear fluorescence emission at around 620 nm. This result indicates that the fluorescence emission was caused by the ARS–VPBA adduct, and was not affected by MBA. Interestingly, the emission spectrum of the VPBA–ARS complex (with a maximum emission at 620 nm) was different from the results of MIP particles treated with a low concentration of ARS (Fig. 3a), where the maximum emission occurred at 560 nm. In Fig. 3a, the concentration of ARS was 0.1 mM while the concentration of ARS in Fig. S4 (ESI[†]) was 2 mM, the same as used in the actual molecular imprinting reaction. To understand the reason for the longer wavelength emission of the VPBA–ARS complex at the higher concentration, we measured the emission spectra of the VPBA–ARS solution after series dilution with the imprinting solvent. As shown in Fig. S5 (ESI[†]), in the low concentration range, the emission of the VPBA–ARS solution centered around 560 nm and the emission intensity increased with concentration. In the high concentration range, however, the emission band was red-shifted to 620 nm, accompanied by a reduced intensity. The concentration-dependent red shift of the fluorescence most likely resulted from a transition of VPBA–ARS complex from monomer to dimer or aggregate structures.^{43,44} The fluorescence bands at 560 nm and 620 nm therefore represented the monomer and dimer/aggregate

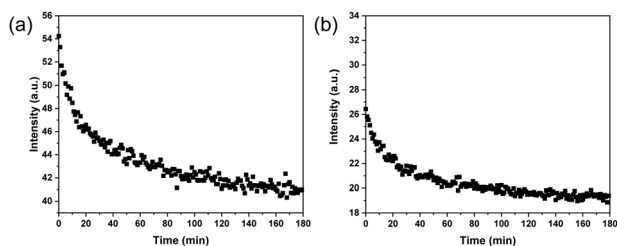


Fig. 4 (a) Change in fluorescence intensity of ARS-loaded MIP particles (0.5 mg mL^{-1}) measured after the addition of 1 M fructose. (b) Change in fluorescence intensity of ARS-loaded NIP particles (0.5 mg mL^{-1}) measured after the addition of 1 M fructose. The fluorescence intensity was measured at 560 nm in 20 mM phosphate buffer.



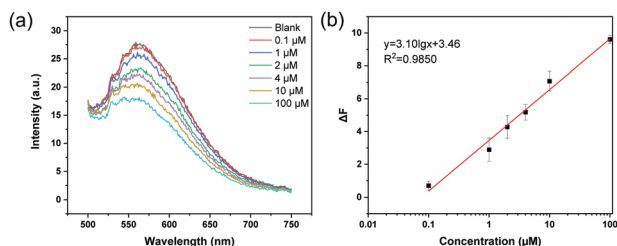


Fig. 7 (a) Fluorescence emission spectra of ARS-loaded MIP particles (0.5 mg mL^{-1}) mixed with Cu^{2+} solution at different concentrations in phosphate buffer (20 mM, pH 8.5). (b) Linear relationship between the fluorescence intensity change and the logarithm of the Cu^{2+} concentration.

probe, the metal ions can be measured in the same concentration range as was achieved using the original MIP particles.

3.8 Detection of Cu^{2+} in spiked water samples

In order to evaluate the selectivity of the MIP-based assay, the fluorescence response of MIP nanoparticles to Cu^{2+} , Na^+ , Mg^{2+} , Fe^{3+} , Ca^{2+} , K^+ , Ag^+ and Zn^{2+} was investigated. In Fig. S9 (ESI[†]), the change in the fluorescence intensity of the MIP particle suspension (0.5 mg mL^{-1}) caused by $10 \text{ } \mu\text{M}$ of different metal ions in water is presented. It is clear that only Cu^{2+} exhibited an obvious quenching of the fluorescence. Therefore, the MIP-based fluorescent probe has a satisfactory specificity for the detection of Cu^{2+} in water.

3.9 Detection of Cu^{2+} in water samples

The reliability of the proposed fluorescence method for detecting Cu^{2+} in real samples was studied by measuring Cu^{2+} in tap water and rain water. The analytical results were compared with those obtained using a commercial copper assay kit. As shown in Table 2, the recoveries of Cu^{2+} in the different samples were in the range of 84.0% to 126%, with a relative standard deviation (RSD) of 0.58% to 3.2%. Using the commercial copper assay kit, a chromogen was utilized to form a colored complex with the copper ions. The commercial assay kit was not able to detect Cu^{2+} in tap water and rainwater at the $1 \text{ } \mu\text{M}$ level. By comparing with the commercial copper assay kit, we conclude that the MIP-based fluorescence assay can be used for the fast detection of Cu^{2+} in water.

3.10 Repetitive adsorption of ARS and detection of Cu^{2+} with MIP nanoparticles

To investigate the use of MIP nanoparticles for the repetitive adsorption of ARS and the detection of Cu^{2+} , the uptake of ARS

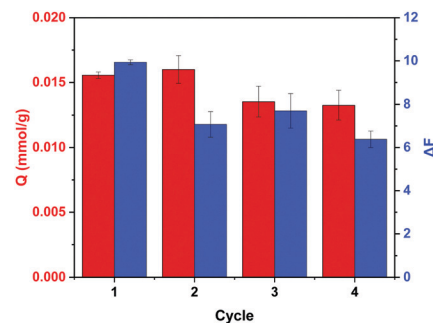


Fig. 8 Recycling and reuse of MIP nanoparticles for the adsorption of ARS and the detection of Cu^{2+} in buffer. Particle concentration: 0.5 mg mL^{-1} ; ARS concentration: 0.02 mM . Cycle 1 used the original MIP particles, and cycles 2–4 used MIP particles regenerated from the previous cycle followed by reloading with 1 mM ARS and washing.

by regenerated MIP nanoparticles and the detection of Cu^{2+} in aqueous solution were further studied. In the first cycle, MIP particles were used to bind ARS in water. At the same time, the same MIP particles were exposed to Cu^{2+} to measure the fluorescence intensity change. The uptake of ARS and the Cu^{2+} -induced fluorescence intensity change were recorded. After the first cycle, the MIP particles that had been exposed to Cu^{2+} were washed with EDTA solution, and then loaded with ARS before the particles were used to measure the change in the fluorescence intensity caused by Cu^{2+} . The uptake of ARS and the Cu^{2+} -induced fluorescence intensity change were recorded as the results of cycle 2. Fig. 8 shows the results of ARS binding and the Cu^{2+} -induced fluorescence intensity change obtained in the different cycles. It is obvious that ARS binding in cycle 2 was almost the same as in cycle 1, which indicates that the residual ARS in the polymer matrix did not affect the ARS binding at the 0.02 mM concentration level. The change on fluorescence for cycle 1 (ΔF_1) caused by Cu^{2+} was due to the complexation of Cu^{2+} with the residual ARS in the original MIP, whereas the change in fluorescence for cycle 2 (ΔF_2) was caused by the complexation of Cu^{2+} with the ARS re-loaded in the MIP. As the amount of residual ARS in the original MIP may not be the same as the regenerated and reloaded MIP, the measured ΔF_1 and ΔF_2 values are different. Nevertheless, the ΔF values measured in cycles 2–4 for the regenerated MIP appeared to be similar. The slight reduction in ARS binding and fluorescence response after the second cycle were most likely caused by the loss of polymer particles during the multiple centrifugation and decantation steps. Generally, the MIP particles prepared in this

Table 2 Recovery of Cu^{2+} at different concentration levels in tap water and rainwater

Sample	Cu^{2+} conc. spiked (μM)	This method			Copper assay kit		
		Detected (μM)	Recovery (%)	RSD (%)	Detected (μM)	Recovery (%)	RSD (%)
Tap water	0	4.81	—	0.9	3.43	—	0.11
	1	5.96	101	0.58	2.67	—	1.1
	10	17.6	126	3.16	12.7	93.1	1.26
Rainwater	0	0.44	—	2.7	0.7	—	0.9
	1	1.29	84	1.99	−0.74	—	0.65
	10	11.1	94.7	1.95	10.3	96.6	0.63



work displayed satisfactory recyclability and reusability for the adsorption of ARS and the detection of Cu^{2+} .

4 Conclusions

In this work, we demonstrated a simple, sensitive, and straightforward method to study the process of molecular imprinting using real-time fluorescence measurements. Using ARS as a model template and VPBA as the functional monomer, we have shown that the template–functional monomer complex plays an important role in realizing the specific molecular recognition of the final imprinted polymer. The molecularly imprinted sites in the model system start to form as soon as 20 min after the start of free radical polymerization. The method of mechanistic study in the present work can be extended to other molecular imprinting reactions. Besides monitoring molecular imprinting reactions, the analytical method also allowed us to study template binding and dissociation in real time, providing a convenient means of understanding the molecular recognition kinetics. The ARS-imprinted polymer can be used as a scavenger to remove ARS directly from water, and as a fluorescent probe to detect Cu^{2+} without any tedious sample preparation.

Author contributions

Xiaoting Xue designed, performed, and validated all the experiments, and wrote the manuscript. Man Zhang and Haiyue Gong provided discussion and feedback. Lei Ye designed and supervised the project, and carried out the project administration and funding acquisition. All authors contributed to discussion of the results and approved the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

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