Enhancing the Oil-Fouling Resistance of Polymeric Membrane Ion-Selective Electrodes by Surface Modification of a Zwitterionic Polymer-Based Oleophobic Self-Cleaning Coating

Longbin Qi,* Tianjia Jiang, Rongning Liang, and Wei Qin*

ABSTRACT: Due to the frequent oil spill accidents and pollution of industrial oily wastewater, oil fouling has become a great challenge to polymeric membrane ion-selective electrodes (ISEs) for applications in oil-contaminated areas. Herein, a simple approach is proposed to enhance the oil-fouling resistance of polymeric membrane ISEs by surface modification of a zwitterionic polymer-based underwater oleophobic coating. As a proof-of-concept, a classical poly(vinyl chloride) membrane-based calcium ion-selective electrode (Ca2+-ISE) is chosen as a model sensor. The zwitterionic polymer-based coating can be readily modified on the sensor’s surface by immersion of the electrode into a mixture solution of dopamine and a zwitterionic acrylate monomer (i.e., sulfobetaine methacrylate, SBMA). The formed poly(SBMA) (PSBMA) coating alters the oleophilic membrane surface to an oleophobic one, which endows the surface with excellent self-cleaning properties without loss of the sensor’s analytical performance. Compared to the pristine Ca2+-ISE, the PSBMA-modified Ca2+-ISE exhibits an improved analytical stability when exposed to oil-containing wastewater. The proposed approach can be explored to enhance the oil-fouling resistance of other polymeric membrane-based electrochemical sensors for use in the oil-polluted environment.

1. INTRODUCTION

As a well-established analytical tool, potentiometric polymeric membrane ion-selective electrodes (ISEs) play an important role in environmental monitoring. Due to the frequent oil spill accidents and increasing discharges of industrial oily wastewater, especially in coastal areas and oilfields, oil fouling are becoming commonplace, which would induce a great challenge for plasticized polymeric membrane sensors to analyze environmental samples with high accuracy and stability. Indeed, oil fouling are found to be much more intractable compared to other foulants (e.g., proteins and microorganisms) owing to their easy deposition and deformation on polymer membranes via strong hydrophobic interactions. When the oil droplets adhere onto a polymeric membrane, they can spread all over the membrane surface in a short time. Moreover, the lipophilic components of the oil droplets could be extracted into the organic membrane phase resulting in a drift in the potential response and loss of the sensor’s selectivity. To date, considerable efforts have been devoted to improve the sensors’ biocompatibilities. However, research on the sensors’ environmental compatibilities are still rather rare. Considering the negative effects of oil fouling, it is crucial to develop a polymeric membrane ISE configuration with oil-fouling resistance for reliable detection in environmental samples.

Many approaches, such as sample pretreatments (e.g., solvent extraction) and advanced sampling techniques, have been developed to mitigate the interferences of lipophilic compounds. However, these methods are usually time-consuming and unsuitable for the real-time analysis. Surface modification has been proven as an effective method to prevent fouling. Hydrophilic materials, such as poly(ethylene glycol) (PEG)-based polymers, hydrogels, and zwitterionic materials, have been employed to protect sensors from direct contact with foulants. Among these, zwitterionic materials are recognized as an excellent candidate for construction of fouling-resistance surfaces due to their electric neutrality, excellent hydrophilicity, underwater oleophobicity, and good stability. The unique structures of zwitterionic materials containing equivalent positively and negatively charged groups allow binding with water molecules via strong electrostatic interactions. Moreover, compared with the PEG-based materials, the zwitterionic units (e.g., sulfobetaine methacrylate, SBMA) could bind to more...
A constant current of 0.014 mA was applied for the sensor cleaning ability against oil foulants but does not in surface of the Ca\textsuperscript{2+}-ISE is modified before. 20, 24 A poly(3,4-ethylenedioxythiophene) coating endows the modified Ca\textsuperscript{2+}-ISE with an excellent self-cleaning ability against oil foulants but does not influence the sensor’s analytical performance.

2. EXPERIMENTAL SECTION

2.1. Membranes and Electrodes. The calcium-ion-selective membranes (Ca\textsuperscript{2+}-ISMs) contained 2.3 wt % N,N,N’,N’-tetracyclohexyl-3-oxapentanediamide (ETH 129), 2.2 wt % sodium tetrakis [3,5-bis(trifluoromethyl)phenyl] borate (NaTFPB), 31.8 wt % poly(vinyl chloride) (PVC), and 63.7 wt % 2-nitropbenzoyl octyl ether (o-NPOE). The membrane cocktail solution was prepared by dissolving 360 mg of the membrane components into 3.6 mL of tetrahydrofuran (THF). After being stirred for 2 h, the membrane cocktail solution was poured into a glass ring (i.d. 36 mm) fixed on a glass plate to obtain the Ca\textsuperscript{2+}-ISM.

The solid-contact Ca\textsuperscript{2+}-ISEs were prepared as described before.\textsuperscript{20,24} A poly(3,4-ethylenedioxythiophene) (PEDOT):poly(sodium 4-styrenesulfonate) (PSS) layer was used as the ion-to-electron transducer of the solid-contact Ca\textsuperscript{2+}-ISEs. This layer was electrodeposited onto the surface of a glassy carbon electrode (GCE, i.d. 3.0 mm) using a mixture solution with 10\textsuperscript{-2} M 3,4-ethylenedioxythiophene and 10\textsuperscript{-1} M PSS. A constant current of 0.014 mA was applied for the polymerization of 714 s.\textsuperscript{41,42} The obtained GCEs were rinsed with deionized water and dried at room temperature. Finally, 80 μL of the membrane cocktail solution was drop-cast onto the GCE to generate a sensing membrane with a thickness of approximately 100 μm.

2.2. Surface Modification of the Ca\textsuperscript{2+}-ISEs. A one-step deposition method was used to modify the zwitterionic polymer coating onto the surface of the Ca\textsuperscript{2+}-ISE.\textsuperscript{43} Briefly, 2 mg mL\textsuperscript{-1} of dopamine and 30 mg mL\textsuperscript{-1} of zwitterionic acrylate monomer SBMA were dissolved into a Tris-HCl buffer solution (pH = 8.5, 50 mM). Subsequently, the membrane electrodes were immersed into the obtained solution and incubated for 12 h at room temperature. During this process, the self-polymerization of dopamine can trigger the polymerization of the acrylate monomer SBMA to form PSBMA. Simultaneously, the generated polydopamine assists PSBMA to deposit onto the electrode surface. After modification, the obtained electrodes were thoroughly washed with deionized water.

2.3. Electrochemical Measurements. The electromotive force (EMF) values were measured by using an electrochemical workstation (CHI 760D, Shanghai Chenhua Apparatus Corporation, China) at room temperature in the galvanic cell: Ag/AgCl/3 M KCl/sample solution/ISM/ PEDOT-PSS/GCE. For Ca\textsuperscript{2+} detection, the electrodes were conditioned in 10\textsuperscript{-3} M CaCl\textsubscript{2} solution overnight before use. Selectivity measurements were carried out by using the separate solution method,\textsuperscript{46} and the electrodes were conditioned in 10\textsuperscript{-3} M NaCl solution overnight before measurements. The measured EMF values were corrected for the liquid junction potentials, and the ion activities were calculated according to the Debye–Hückel equation.

2.4. Evaluation of the Anti-Oil-Fouling Properties of the PSBMA-Modified Ca\textsuperscript{2+}-ISEs. To evaluate the oil-fouling resistance of the PSBMA-modified Ca\textsuperscript{2+}-ISEs, the potentiometric responses of the electrodes after exposure to oil foulants were tested. The artificial wastewater samples containing 5% (v/v) of the oils were prepared. The diesel oil and crude oil samples (collected from the Bohai Sea, China) were selected as the model foulants, since they are typical oil foulants in the environment. The conditioned membrane electrodes were immersed in the oil-containing wastewater samples for 1 h under continuous stirring. Then, the fouled electrodes were rinsed with deionized water and transferred to the cell for potentiometric measurements. To observe the oil fouling of
the membrane electrodes, the surfaces of the Ca\(^{2+}\)-ISMs were photographed. For easy visualization, the diesel oil sample was stained with Sudan III.

3. RESULTS AND DISCUSSION

Due to oil spill accidents and industrial oil pollution, oil fouling is becoming a great challenge for polymeric membrane ISEs to be applied in the environmental fields. Since the plasticized polymeric membrane matrices are hydrophobic, oil foulants can easily deposit on the sensing membrane via strong hydrophobic interactions, which may cause measurement errors and shorten the sensors’ lifetimes. In this study, an underwater oleophobic self-cleaning coating based on a zwitterionic polymer was developed to enhance the oil-fouling resistance of the Ca\(^{2+}\)-ISE. As shown in Scheme 1, the proposed surface coating can be formed simply by immersion of the electrode into a solution containing dopamine (2 mg mL\(^{-1}\)) and zwitterionic acrylate monomer SBMA (30 mg mL\(^{-1}\)) at pH 8.5. After the one-step modification, the oleophobic membrane surface can be changed to an oleophobic one. In this case, the oil droplets adjacent to the sensor surface can be cleaned by the PSBMA coating, thus effectively eliminating the oil-fouling problems (Scheme 1, right). In contrast, for the pristine sensor, the oil droplets could rapidly be adsorbed on the sensor surface and cause sensor failures (Scheme 1, left).

The images of scanning electron microscopy (SEM) show that numerous nanoparticles can be found on the surface of the PSBMA-modified Ca\(^{2+}\)-selective membrane (Figure S1). The results of the water contact angle (WCA) measurements illustrate that the WCA of the Ca\(^{2+}\)-selective membrane decreases from 86.3 ± 1.7° to 38.1 ± 4.2° after PSBMA modification, which indicates that the hydrophilicity of membrane surface is significantly improved by the PSBMA coating (Figure S2). The spectra of attenuated total reflectance-Fourier transform infrared spectroscopy (ATR-FTIR) show the new peaks at 1726 and 1033 cm\(^{-1}\) after PSBMA coating (Figure S2), which are attributed to the O=C=O and S=O symmetric stretching vibrations, respectively.\(^{43}\) These results indicate that the SBMA units have been incorporated into the polymer. Under alkaline conditions, dopamine can be oxidized into dopamine quinone and then form 5,6-dihydroxyindole and 5,6-indolequinone.\(^{47}\) These intermediates can be transferred into the semiquinone radical species via the single-electron exchange reaction and electron resonance, and then further integrated into polydopamine via the radical coupling reactions.\(^{47,48}\) Simultaneously, these radicals can also act as reactive species to trigger the free-radical polymerization of SBMA to form PSBMA.\(^{37,43}\) Due to the intrinsic adhesive nature of polydopamine, the PSBMA can be codeposited onto the surface of the substrate via noncovalent interactions, such as hydrogen bonding and π-π interactions.\(^{37,45,49}\)

Since the deposition time plays a vital role in the oleophobicity of the prepared PSBMA coating, the influence of deposition time was investigated. As shown in Figure S4, the pristine Ca\(^{2+}\)-ISE has a small underwater contact angle (UWOCa) of 25.4 ± 2.1° for dichloromethane, indicating its high lipophilicity. As the deposition time increases from 4 to 12 h, the UWOCa of the pristine Ca\(^{2+}\)-ISE increases rapidly from 42.0 ± 1.3 to 141.5 ± 3.4°, which is mainly due to the increase in the thickness of the oleophobic PSBMA coating. A further increase in deposition time would not obviously improve the UWOCa. Thus, 12 h was selected as the optimal modification time. Moreover, several common oils including diesel oil, petroleum ether, and peanut oil were also used to measure the UWOCAs of the PSBMA modified Ca\(^{2+}\)-ISE. As shown in Figure 1, the oleophilic membrane surface becomes oleophobic with significantly increased UWOCa values for all of these oils. As a comparison, the UWOCAs for the polydopamine-modified Ca\(^{2+}\)-ISE were also investigated (see Figure S5), which are much smaller than those of the PSBMA-modified Ca\(^{2+}\)-ISE. These results indicate that the oleophobicity of PVC-based Ca\(^{2+}\)-ISE can be significantly enhanced by the proposed PSBMA coating. In addition, the PSBMA coating has an excellent long-term stability. Experiments show that the PSBMA-modified Ca\(^{2+}\)-ISE can still maintain a UWOCa value of 142° for diesel oil after storage at 4 °C for one month.

The zwitterionic polymers have been recognized as effective materials with excellent underwater oleophobicity to resist oil fouling.\(^{35,50}\) The self-cleaning properties of the PSBMA-modified Ca\(^{2+}\)-ISE were characterized by investigating the underwater-oil adhesion on the membrane surface. As illustrated in Figure 2A, the diesel oil droplet can quickly adhere to the surface of the pristine Ca\(^{2+}\)-ISE once it contacts the membrane surface, indicating that the pristine Ca\(^{2+}\)-ISE has a high oil-fouling tendency. In contrast, no obvious oil adhesion is observed for the PSBMA-modified Ca\(^{2+}\)-ISE (Figure 2B). In addition, the free oil droplet can quickly slide away from the surface of PSBMA-modified Ca\(^{2+}\)-ISE (Figure 2C). Indeed, these data prove that the PSBMA-modified membrane has an excellent self-cleaning ability. After PSBMA modification, a zwitterionic membrane surface can be obtained which has excellent fouling resistance due to the formation of a compact and continuous hydration layer on the membrane surface via electrostatically induced hydration.\(^{13,39}\) The hydration layer could strongly weaken the hydrophobic interactions between the oil droplets and membrane surface. Under the conditions of slight hydraulic stirring, the oil droplets can be easily expelled from the membrane surface without any adhesion. In this case, the coalescence of oil droplets and formation of an oil layer on the membrane surface

![Figure 1. Underwater-oil contact angles (UWOCAs) for the pristine and PSBMA-modified Ca\(^{2+}\)-selective membranes. Dichloromethane, diesel oil, petroleum ether, and peanut oil were used to measure the UWOCAs. Experimental conditions: dopamine, 2 mg mL\(^{-1}\); SBMA, 30 mg mL\(^{-1}\); pH, 8.5; immersion time, 12 h.](https://doi.org/10.1021/acs.analchem.1c01116)
can be effectively inhibited. Therefore, the oil-fouling resistance of the PSBMA-modified membrane electrode can be significantly enhanced.

It should be noted that the improvement in sensor’s fouling-resistance properties should not sacrifice the analytical performance. Therefore, the potentiometric responses of the pristine and PSBMA-modified Ca²⁺-ISEs were tested in the range from 10⁻⁸ to 10⁻² M. As shown in Figure 3A, the pristine Ca²⁺-ISE exhibits a Nernstian response slope of 29.2 ± 0.2 mV decade⁻¹ over the range from 10⁻⁶ to 10⁻² M with a detection limit of 0.3 × 10⁻⁵ M. Similar to the pristine Ca²⁺-ISE, the PSBMA-modified Ca²⁺-ISE shows a response slope of 28.7 ± 0.9 mV decade⁻¹ with a detection limit of 5.5 × 10⁻⁷ M. These results indicate that the PSBMA coating cannot impair the thermodynamic extraction equilibrium of Ca²⁺. As shown in Figure 3B, the response time of the PSBMA-modified Ca²⁺-ISE is longer than that of the pristine Ca²⁺-ISE. After modification, the response time of the Ca²⁺-ISE increases from 2.9 to 13.9 s when the concentration of Ca²⁺ is changed from 10⁻⁶ to 10⁻⁵ M. Such a change in response time is more pronounced for lower concentrations of Ca²⁺ but negligible for concentrations higher than 10⁻³ M. This may be attributed to the fact that the PSBMA layer could prevent the diffusion of Ca²⁺ between the sensing membrane and the aqueous layer. The selectivity coefficients of the PSBMA-modified Ca²⁺-ISE were measured by using the separate solution method, and Na⁺, K⁺, and Mg²⁺ were selected as the interfering ions. The results shown in Table S1 demonstrate that the selectivity coefficients of the PSBMA-modified Ca²⁺-ISE are comparable to those of the pristine Ca²⁺-ISE.

In order to evaluate the feasibility of the proposed self-cleaning Ca²⁺-ISE for practical applications, the potentiometric responses of the pristine and PSBMA-modified Ca²⁺-ISEs before and after immersion of the electrodes in the oil-containing wastewaters were measured. As illustrated in Figure 4A, after immersion in the wastewater containing 5% (v/v) diesel oil for 1 h under stirring, the response slope of the pristine Ca²⁺-ISE decreases by 56.2% (from 29.2 to 12.8 mV decade⁻¹). In contrast, no obvious change is observed for the PSBMA-modified Ca²⁺-ISE (from 28.7 to 28.4 mV decade⁻¹).

Figure 2. Self-cleaning properties of the PSBMA-modified Ca²⁺-selective membranes: (A) dynamic images of a diesel oil droplet approaching and adhering to the pristine Ca²⁺-selective membrane, (B) dynamic images of a diesel oil droplet approaching and leaving the PSBMA-modified Ca²⁺-selective membrane, and (C) dynamic images of a free diesel oil droplet sliding away from the PSBMA-modified Ca²⁺-selective membrane. The red arrows represent the moving directions of the diesel oil droplets.

Figure 3. Influence of the surface modification on the sensor’s analytical performance: (A) time-dependent potentiometric response traces of the pristine and PSBMA-modified Ca²⁺-ISEs and (B) response times of the pristine and PSBMA-modified Ca²⁺-ISEs. The response times were calculated from the potentiometric response traces when the concentration of Ca²⁺ was changed from 1 × 10⁻⁶ to 1 × 10⁻⁵ M. Error bars represent one standard deviation for three measurements.

Figure 4. Typical calibration curves of the pristine and PSBMA-modified Ca²⁺-ISEs after underwater contacts with 5% (v/v) diesel oil (A) and crude oil (B), respectively, for 1 h under stirring.
The similar results are also observed for the crude oil fouling. As shown in Figure 4B, the pristine Ca\textsuperscript{2+}-ISE even loses its original potential response, while the PSBMA-modified Ca\textsuperscript{2+}-ISE still maintains an acceptable response slope of 260 mV decade\textsuperscript{-1}. These results suggest that the proposed self-cleaning Ca\textsuperscript{2+}-ISE exhibits a significantly improved stability when exposed to oil foulants. Such remarkable difference in sensor’s potentiometric response is probably due to the fact that the pristine PVC-based oleophilic membrane has a tendency to adsorb oil droplets resulting in the accumulation of oils on the membrane surface.\textsuperscript{13} The adsorbed oil foulants could be dissolved in the membrane phase to change the membrane components and/or may form an oil layer to insulate the ISE membrane from the sample solution. For the PSBMA-modified Ca\textsuperscript{2+}-ISE, the self-cleaning oleophobic coating can effectively prevent the adhesion of oil foulants on the electrode surface and thus improve the sensor’s analytical stability when exposed to oil foulants.

Furthermore, the Ca\textsuperscript{2+}-ISMs were examined visually after contact with the oil-containing wastewaters. The diesel oil sample was stained using Sudan III to improve visualization. Indeed, as shown in Figure S6, after contacts with the oil-containing wastewaters, a large amount of the tested oils with the dense and uniform distributions can be observed on the unmodified Ca\textsuperscript{2+}-ISMs, indicating the high fouling tendency of the unmodified membrane. Figure S6 also indicates that, for the PSBMA-modified Ca\textsuperscript{2+}-ISMs, negligible oil adhesion is observed after contacts with the oil-containing wastewaters, thus further confirming the enhanced fouling-resistance of the proposed sensor.

■ CONCLUSIONS

We demonstrate here for the first time that the oil-fouling resistance of polymeric membrane Ca\textsuperscript{2+}-ISEs can be enhanced by surface modification of a zwitterionic PSBMA coating. Experiments show that the analytical performance of the Ca\textsuperscript{2+}-ISE in terms of response slope, detection limit, and selectivity is not influenced by the PSBMA coating. The PSBMA coating increases the UWOCA of the sensor’s surface from 25.4 ± 2.1 to 141.5 ± 3.4° and endows the sensor with excellent self-cleaning properties against oil foulants. After fouling with 5% (v/v) diesel oil or crude oil, the response of the pristine Ca\textsuperscript{2+}-ISE decreases by 56.2% or even completely lost in terms of the response slope. In contrast, the PSBMA-modified Ca\textsuperscript{2+}-ISEs can still maintain acceptable response slopes after being fouled by oils. The proposed strategy has a promising potential to develop polymeric membrane ISEs with excellent oil-fouling resistance for reliable potentiometric detection in the oil-polluted environment.

■ ASSOCIATED CONTENT

1 Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.analchem.1c01116.

Experimental details including reagents and characterization; SEM images (Figure S1) for the PSBMA-modified membrane; water contact angles (Figure S2) for the PSBMA-modified membrane; ATR-FTIR spectra for the polydopamine and the product from polymerization of dopamine and SBMA (Figure S3); influences of the modification time on the UWOCA for PSBMA-modified Ca\textsuperscript{2+}-selective membrane (Figure S4); UWOCA for the polydopamine-modified Ca\textsuperscript{2+}-selective membrane (Figure S5); photographs of the pristine and PSBMA-modified Ca\textsuperscript{2+}-selective membranes before and after contact with oils (Figure S6); and potentiometric selectivity coefficients of Ca\textsuperscript{2+}-ISEs before and after modification (Table S1) (PDF)

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Notes

The authors declare no competing financial interest.

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