Application of a New Macrocyclic -PVC Electrode to Potentiometric Studies of Fe(III) Ion

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ABSTRACT

A PVC membrane Fe(III) ion selective electrode has been constructed using a new macrocyclic as membrane carrier. The electrode exhibits a good potentiometric response for Fe(III) over a wide concentration range 1.0×10^{-4} to 1.0×10^{-1}M with a slope 19.4±0.5 mV/decade and a working pH range of 6.5-9.0. It has a fast response time ≤30s. The best performance was observed with the membrane having the 30.5% PVC, 3.0% ligand, 61.0% acetophenone and 5.5% oleic acid composition.

Keywords: Ion selective electrode; PVC membrane; Fe(III) determination; Macrocyclic ionophore

INTRODUCTION

Ion-selective electrodes (ISEs), especially those with neutral carrier-based solvent polymeric membranes, have been studied for more than three decades, and are now routinely employed for direct potentiometric measurements of various ionic species in environmental, industrial and clinical samples [1,2,3]. The increasing use of ion sensors in the fields of environmental, agricultural and medicinal analysis is stimulating analytical chemists to develop new sensors for the fast, accurate, reproducible and selective determination of various species. In the past few decades, considerable efforts have led to the development of selective sensors for alkali and alkaline earth metals and for heavy metals. Among heavy metals, trivalent metals have received less attention in spite of their widespread occurrence in rocks, alloys, food products, sea- and fresh-water, plants and animals [4].

A significant number of macrocyclic compounds including crown ethers, cryptands, aza-crowns and thiacrowns, which have been synthesized in various cavity sizes and shapes have already been
exploited for the fabrication of poly(vinyl chloride) membrane electrodes for transition and heavy metal ions [5-12].

Fabrication of a new ion specific ISE with high selectivity and sensitivity, wide linear concentration range, long lifetime, good reproducibility and low cost is always in need. A literature survey revealed that a large number of ISE based on PVC membrane were reported for many inorganic ions [13], but to the best of our knowledge, very little reports were appeared on PVC-based trivalent ions.

In this paper, we report the use of new ligand as a neutral carrier in the construction of a PVC membrane electrode selective to Fe(III) ion.

**EXPERIMENTAL**

**Reagents**

All of reagents and solvents involved in synthesis were of analytically grade a received without further purification. Tetrahydrofuran (THF), acetophenone (AP), oleic acid (OA), high relative molecular weight PVC were obtained from Aldrich and Merck. a new macrocyclic ionophore (Fig.1).

Chloride and nitrate salts of all other cations and detergent used (all from Merck or Fluka) were of the highest purity available and used without any further purification.

**Fig. 1.** Structure of H3L.

**Apparatus**

Potentiometric and pH measurements were carried out using a metrohm digital pH/mV meter ion analyzer in stirred solution. In all instances, an Ag-AgCl / KCl (sat.) electrode (Azar electrode company, Urmia, Iran) was used in conjucation with the respective indicator electrode. A Haoke model FK2 circulation water bath was used control the temperature of the test solution.

**Electrode preparation and potential measurement**

A mixture of PVC, oleic acid, acetophenone and ionophore to give a total mass of 105 mg, was dissolved in about 2 mL of THF and the solution was mixed well. The resulting clear mixture was transferred into a glass dish of 2 cm diameter. The solvent was evaporated slowly until an oily concentrated mixture was obtained.

A pyrex tube (5 mm o.d.) was dipped into the mixture for about 10 s so that a nontransparent membrane of about 0.2 mm thickness was formed. The tube was then pulled out from the mixture and kept at room temperature for about 2 h. The tube was then filled with internal solution 1.0 \times 10^{-3} M Fe(III) with pH = 3. The electrode was finally conditioned for 24 h by soaking in a 1.0 \times 10^{-2} M Fe(III) solution with pH = 3.

The potential measurements were carried out with the following assembly:

\[ \text{Ag/AgCl} / 3 \text{ M KCl} / \text{internal solution (1.0 } \times 10^{-3} \text{ M FeCl}_3, 1.0 \times 10^{-3} \text{ M HCl}) / \text{PVC membrane} / \text{test solution} / 3 \text{ M KCl} / \text{Ag-AgCl} \]

**RESULTS AND DISCUSSION**

Effect of membrane composition on the electrode response

The potential responses of various ion-selective electrodes based on H3L are in Fig. 2. Expect for the Fe(III) ion selective electrode, in all other cases the slope of the
corresponding potential-pM plots is much lower than the expected Nernstian slopes.

Besides the critical role of the nature of the ion carrier in preparing membrane-selective sensors, some other important features of the PVC membrane are known to significantly influence the sensitivity, linearity range and selectivity of ion-selective electrodes. These include the amount of ionophore, the nature of solvent mediator, the plasticizer/PVC ratio and especially the nature of additives used [14-20].

Thus, based on the results obtained on the optimization of the membrane composition, the membrane with the optimized composition of PVC : ionophore : acetophenone : oleic acid ratio of 10:1:22:2 was selected for the preparation of the polymeric membrane electrode. The characteristic parameters of the optimized membrane are summarized in Table 1.

Reversibility of the electrode
To evaluate the reversibility of the electrode, a similar procedure in the opposite direction was adopted. The measurements were performed in the sequence of high-to-low (from 1.0×10^{-2} to 1.0×10^{-3} M) sample concentration and the results are shown in Fig. 3. Fig. 3. shows that the potentiometric response of the electrode is reversible, although the times needed to reach equilibrium values were longer than that of low-to-high sample concentration [21].

![Fig. 2. Response at pH= 3 against some of cations for ion-selective electrodes containing H3L as ionophore.](image)

<table>
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<th>Table 1. Specification of the Fe(III)-ISE based on H3L</th>
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The response time
The response time of the electrode was measured after successive immersion of the electrode in a series of Fe solution, in each of which the Fe(III) concentration was increased tenfold, from $1.0 \times 10^{-6}$ to 1.0 M. At lower concentrations, however, the response time was longer and reached 30 s for a Fe(III) concentration of $1.0 \times 10^{-4}$ M. The actual potential versus time traces is shown in Fig. 4 for Fe(III) concentration of $1.0 \times 10^{-4}$ M.

CONCLUSION
The membrane sensor incorporating H$_3$L as the electroactive phase can be used to determine Fe(III) in the wide concentration range. The sensor exhibited good reproducibility over a useful lifetime about 2 months. This electrode is superior to the existing electrodes with regard to the slope, pH range, response time and selectivity over a number of cation.

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