Ion-Exchange Applications of Newly Synthesized of Aminoacetophenone, Biuret, Formaldehyde Derivative as New Chelating Terpolymer Resin

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A new chelating terpolymer resin (o-AABF) was synthesized through copolymerization of o-amino acetophenone, biuret, formaldehyde in (1:1:3) mol ratio in acidic medium (HCl) by condensation reaction. The synthesized copolymer resin was characterized by UV-visible, FTIR and 1H NMR techniques. The empirical structure and molecular weight of the resin were analyzed by elemental analysis. Morphological structure of the synthesized resin was studied by scanning electron microscopy. Thermal properties and stability of the resin was determined by DSC and TGA. The intrinsic viscosity and the viscosity average molecular weight were calculated. Selectivity and binding capacity of the terpolymer resin studied by batch equilibrium method toward Ni2+, Co2+, Cd2+, Cu2+, Zn2+ and Cr3+ in different pH and treatment time.

Keywords: Chelating, Terpolymer, Intrinsic viscosity, Biuret, o-amino acetophenone.

INTRODUCTION

The heavy metal ion toxicity has increased strongly because of many industries catalyst contains metal ions. Many techniques have been improved for the pre-concentration and removal of metal ions such as electrodeposition coprecipitation techniques have been improved for the pre-concentration and because of many industries catalyst contains metal ions. Many industry processes require the removal of heavy metal ions, which are toxic to human health. If the toxic metal ion can be recovered, the energy and material require of the waste water treatment process can be simple. The use of chelating resins for separation and removal metal ions is the method choice due to its highly separation efficiency, high degree of selectivity, high loading capacity [1-3]. Chelating resins are widely used for the treatment of radioactive wastes from nuclear power stations [4,5]. The aromatic compound with substituents like –OH, –COOH and –NH, with urea and formaldehyde shows selective ion-exchange properties, thermal resistance properties and coordinating properties [6]. The metal ion uptake increase with increasing mole proportions of the copolymer synthesized from substituted benzoic acid [7]. Salicylic acid-formaldehyde-resorcinol resin has been synthesized and explored its use for removal and separation of heavy metal ions from their binary mixtures [8]. The metal ion-binding properties of terpolymer resins derived from 2,4-dihydroxyacetophenone-biuret-formaldehyde and 2,4-dihydroxy benzaldehyde-oxamide and formaldehyde, respectively have been investigated. In this research, it was synthesized chelating terpolymer resin using an ecofriendly technique and reported for its good binding capacity for Ba2+ and Zn2+ ions [9]. In the present paper, we describe the synthesis of terpolymer resin derived from o-aminoacetophenone and biuret with formaldehyde (o-AABF). The resin has been studied for sorption of Ni2+, Co2+, Cd2+, Cu2+, Zn2+ and Cr3+ from aqueous solutions. For this purpose, various factors effecting the chelating such as pH and treatment time. The regenerating efficiency has been studied by using nitric acid.

EXPERIMENTAL

o-Amino acetophenone (98 %, B.D.H), Biuret (99 %, Fluka) and formaldehyde (37 %, B.D.H), sodium hydroxide (98 %, B.D.H), hydrochloric acid (99 %, B.D.H), nitric acid (99 %, B.D.H). The stock standard solutions of Ni(II), Cd(II), Co(II), Cu(II), Zn(II) and Cr(III) were prepared by dissolving appropriate amount of metal chlorides in deionized water, N,N-dimethyl formamide (DMF) (Acros, 99.6 %), dimethyl sulfoxide-d6 (DMSO-d6). Elemental analysis is recorded by Microestimation technique Euro EA 3000 Single. FTIR spectrum of synthesized resin carried out in Shimadzu FT-IR 8300 series spectrophotometer. 1H NMR (300 MH2) spectra was recorded using (Ultra Shield, Bruker, Switzerland (DMSO-d6)). UV-visible spectrum of result terpolymer resin was recorded by (Double beam Shimadzu 1800, Japan). Amount of metal ions was estimated by atomic absorption spectrophotometer technique (Emission spectrophotometer Shimadzu (A-A680).
Thermal analysis of the resin was recorded using DSC (STA PT-1000 linseis model) linseis Company-Germany. Viscometer [k = 0.005232] Schott-Gerate made in Germany.

**Synthesis of o-AABF terpolymer resin:** The new terpolymer resin *ortho*-amino acetonaphone biuret formaldehyde (o-AABF) was synthesized by the condensation polymerization. In 3 necks, volumetric flask, mixed of *o*-amino acetonaphone (13.5 g, 0.1 mol) and biuret (10.30 g, 0.1 mol) with 37 % formaldehyde (9.0 g, 0.3 mol) in HCl medium at 130 °C in an oil bath for 5 h. After cooling, red brown solution was formed and next day poured in the ice water to obtain yellowish precipitate then washed with distilled water (Scheme-I). The dried resin was further purified by dissolving in 8 % NaOH and regenerated in 1:1(v/v) HCl/ water [10-12]. This process was repeated twice to separate the pure terpolymer resin and dried vacuum at room temperature.

The purified terpolymer resin was finally grinded and sieved to obtain uniform particles and kept in desiccator over silica gel. The sieved resin was used for further characterization. The FT-IR spectrum of (CO-CH$_3$)$_2$, 1599 cm$^{-1}$ of the (Ar-NH$_2$, 1429 cm$^{-1}$ of methylene bridges (CH$_2$), 1697 cm$^{-1}$ of (C=O) in biuret. UV-visible absorption bands in 313 nm. $^1$H NMR (400 MHz, DMSO-$d_6$, δ ppm): 10.97-10.92 (s, 1H, C$_6$), 7.12-5.91 (s, 2H, Cp,d), 5.91-5.32 (s, 2H, C$_{1697}$, t, 1H, C$_6$), 3.21-2.64 (s, 6H, C$_2$). Anal. calcd. for C$_{56.31}$ %, H 6.22 %, N 20.26 %, O 17.25 %.

**Preparation of metal solutions:** Set of standard metal solutions (Ni$^{2+}$, Co$^{2+}$, Cd$^{2+}$, Cu$^{2+}$, Zn$^{2+}$ and Cr$^{3+}$) were prepared via weighting of certain amount of metal salts (NiCl$_2$·6H$_2$O, CoCl$_2$·6H$_2$O, CdCl$_2$·2H$_2$O, CuCl$_2$·2H$_2$O, ZnCl$_2$ and CrCl$_3$·6H$_2$O), respectively in the deionized water [13] in concentrations 1000 and 100 ppm with different pH (2-6) by using (1 M HCl) and (1 M NaOH) to control the pH.

**Chelation efficiency of o-AABF:** The chelation efficiency of o-AABF terpolymer resin was evaluate by batch equilibration method.

The finely group terpolymer was used to determine its metal ion uptake capacity for specific metal ion such as Ni$^{2+}$, Co$^{2+}$, Cd$^{2+}$, Cu$^{2+}$, Zn$^{2+}$ and Cr$^{3+}$ in the form of their chloride salt aqueous solution, metal ion binding capacity for the terpolymer was studied under two different experimental conditions.

The procedure adopted on removes of metal ions by shaking of the terpolymer (0.1 g) with (10 mL) solution having (100 ppm) metal ion concentration for 24 h at higher pH (pH = 6) and 25 °C where the loading capacity will be in maximum range [14]. The mixture was then filtered off and washing were collected and then the amount of metal ion was estimated by using atomic absorption spectrophotometer

**Recovery of metals ions from o-AABF:** Reactivation of the dry loading resin was carried out by shaking with 10 mL (3 M HNO$_3$) at different times. After filtration, the metal ions concentrations in the acid solution were calculated via atomic absorption using standard solution curve for the metal in same acid concentration. From recovery metal ions concentration can be calculate the recovery percentage [15].

**RESULTS AND DISCUSSION**

**Terpolymer resin synthesis:** The synthesized resin (o-AABF) was dark brown colour. The resin are soluble in polar solvent like dimethyl sulfoxide, N,N-dimethyl formamide and tetrahydrofuran, also soluble in mixture of aqueous sodium, potassium hydroxide solution. The synthesized resin did not show sharp melting point but undergo decomposition above 500 °C.

The dilute solution of the synthesized terpolymer in concentration of 10$^{-5}$ M in dimethyl sulfoxide exhibited strong UV-visible absorption bands in 313 nm assignable to the (n-$\pi^*$) transitions resulting from the conjugation between the aromatic rings and nitrogen atom (Fig. 1).

![Fig. 1. UV-visible spectrum of o-AABF terpolymer](image)

The FT-IR spectrum of o-AABF shows the sharp band at 1682 cm$^{-1}$ due to the stretching vibration of carbonyl group of (CO-CH$_3$) and $-NH$ stretching vibration of Ar-NH$_2$ group

![Scheme-I: The synthesis routes of (o-AABF) terpolymer resin](image)
which is further confirmed by –NH bending vibration appearing at 1599 cm\(^{-1}\) [16]. A band appearing in the region of 1429 cm\(^{-1}\) shows the presence of methylene bridges (CH\(_2\)) in the polymer chain [17]. A sharp and weak peak obtained at 2939 cm\(^{-1}\) indicates the presence of stretching vibration of methylene group (-CH\(_2\)) in the terpolymer chain and the sharp band obtained at 1697 cm\(^{-1}\) due to presence of carbonyl group (C=O) in biuret in the terpolymer chain (Fig. 2).

The DSC thermogram of the synthesized o-AABF terpolymer showed a distinct first glass transition (\(T_g\)) appearing at 1599 cm\(^{-1}\) (-NH-CO) (f). The sharp peak is attributed to the protons of –NH- (h) and –NH\(_2\) groups. The signal in the range of (\(\delta_H = 10.97-10.92\)) ppm due to the amide proton (\(\delta_H = 7.12-5.91\)) ppm attributed to the protons (b,d) of the aromatic ring. The peak in the range (\(\delta_H = 3.21-2.64\)) ppm is attributed to protons of amide proton linkage (-NH-CO) (f). The sharp peak is attributed to the protons of –CH\(_2\) (a) and CH\(_3\) (c) groups.

Effect of acidic function on the analytical efficiency of o-AABF ions ((Ni\(^{2+}\), Cd\(^{2+}\), Cu\(^{2+}\), Zn\(^{2+}\), Cr\(^{3+}\)) have been studied. After shaking 10 mL of corresponding metal ions, a solution concentration (100 ppm) with 0.1 g of polymer for a period of 24 h has shown a great response ion towards polymer.

Analytical chelating resin efficiency: The study of the analytical efficiency of the chelating resins is one of the most important applications in analytical chemistry, where the selective resins for specific elements of the presence of other elements will be based on different active groups in the resin, as per the nature of the atom-giving and this leads to the use of resin as selective separation of different ions to be capture.

There are several techniques used in the study of analytical efficiency of chelating resins. The best of these techniques used for this purpose are atomic absorption spectroscopy which is used in this study. The analytical efficiency of the polymer [o-AABF] towards several ions (Ni\(^{2+}\), Cd\(^{2+}\), Co\(^{2+}\), Cu\(^{2+}\), Zn\(^{2+}\), Cr\(^{3+}\)) have been studied. After shaking 10 mL of corresponding metal ions, a solution concentration (100 ppm) with 0.1 g of polymer for a period of 24 h has shown a great response ion towards polymer.

Effect of treatment time at maximum capacity to load o-AABF terpolymer: The effect of treatment time on the efficiency of loaded polymer [o-AABF] to extract metal ions as it was noted that increasing treatment time for the solutions of the studied ions increased loading of the polymer capacity, though the ions reach equilibrium after about (10 h) and the amount of load capacity becomes a little bit until they reach 24 h of treatment. Table-1 and Fig. 5 show the effect of treatment time on the load capacity of the polymer [o-AABF] ions (Ni\(^{2+}\), Cd\(^{2+}\), Co\(^{2+}\), Cu\(^{2+}\), Zn\(^{2+}\), Cr\(^{3+}\)) in acidic studied functions.

Effect of acidic function on the analytical efficiency of o-AABF: The study showed that the acidic function was with significant effect in the loading capacity of the resin for all studied ions. It has been observed that in general, a higher load capacity of the resin for these ions are at the highest acidic function used in the study of each ion (Fig. 6).

Intrinsic viscosity of o-AABF: The viscosity of the resin solution was measured using capillary tube called (Ubbelohde viscometer) by measure the flow time of the solvent used in the composition of the solution polymer (\(t^*\)) and then measuring
TABLE-1

<table>
<thead>
<tr>
<th>Ions</th>
<th>pH</th>
<th>Capacity resin mg/g loading</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0.5 h</td>
<td>1 h</td>
</tr>
<tr>
<td>Co²⁺</td>
<td>1.0</td>
<td>2.0</td>
</tr>
<tr>
<td>Ni²⁺</td>
<td>1.3</td>
<td>4.0</td>
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<td>6.0</td>
</tr>
<tr>
<td>Ni²⁺</td>
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<td>2.0</td>
</tr>
<tr>
<td>Ni²⁺</td>
<td>2.5</td>
<td>4.0</td>
</tr>
<tr>
<td>Ni²⁺</td>
<td>3.2</td>
<td>6.0</td>
</tr>
<tr>
<td>Ni²⁺</td>
<td>5.1</td>
<td>7.0</td>
</tr>
<tr>
<td>Cu²⁺</td>
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<td>2.0</td>
</tr>
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<td>Cu²⁺</td>
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<td>Cd²⁺</td>
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</tr>
<tr>
<td>Cr³⁺</td>
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<td>4.0</td>
</tr>
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<td>6.0</td>
</tr>
<tr>
<td>Cr³⁺</td>
<td>8.5</td>
<td>7.0</td>
</tr>
</tbody>
</table>

the same solution time (t), across these measurements can calculate viscosity as follows:

Relative viscosity:  \[ \eta_{rel} = \frac{t}{t_0} \]

Specific viscosity:  \[ \eta_{sp} = \eta_{rel} - 1 \]

Reduced viscosity:  \[ \eta_{red} = \frac{\eta_{sp}}{c} \]

where c is grams of polymer in (100 mL) of solvent.

Intrinsic viscosity:  \[ [\eta] = K M^a \]

The fact that the viscosity of an experimental relationship with the molecular weight of the several polymeric systems so Stodnker predicted in 1930 that the viscosity is about equal the molecular weight with a slight modification where it compensates intrinsic viscosity instead of reducing viscosity as shown in equation mark reported by Houwnik [20,21].

\[ [\eta] = K M^a \]

where a, K are constants depend on the type of polymer and solvent and solution temperature.

The viscosity of the prepared polymer in this study was measured using viscometer type (Ubbelohde viscometer) in DMSO-as solvent.

Fig. 5. Number of milligrams ions (Co²⁺, Ni²⁺, Cu²⁺, Zn²⁺, Cd²⁺, Cr³⁺) drawn by (0.1 g) of the polymer [o-ABBF] as a function of time in different acidic functions
The relative viscosity is obtained for the resin solution and converted into a relative and reduce viscosity (Table-2) then intrinsic viscosity (0.563 dL/g) was practically determined via graph between reducing viscosity and concentration of the prepared resin, the intersection of the curve with the y-axis at zero value of concentration as shown in Fig. 7.

<table>
<thead>
<tr>
<th>TABLE-2</th>
<th>THE RELATIVE VISCOSITY, SPECIFIC VISCOSITY, REDUCED VISCOSITY</th>
</tr>
</thead>
<tbody>
<tr>
<td>C¹</td>
<td>T²sec</td>
</tr>
<tr>
<td>0.03</td>
<td>113</td>
</tr>
<tr>
<td>0.05</td>
<td>116</td>
</tr>
<tr>
<td>0.07</td>
<td>120</td>
</tr>
<tr>
<td>0.10</td>
<td>128</td>
</tr>
</tbody>
</table>

1: Resin concentration; 2: Solution time (sec.); 3: The relative viscosity (dL/g); 4: Specific viscosity (dL/g); 5: Reduced viscosity (dL/g)

Fig. 6. Effect of acidic function on the loading capacity (LC) of polymer [o-AABF] of metal ions (Co²⁺, Ni²⁺, Cu²⁺, Zn²⁺, Cd²⁺, Cr³⁺) in different treatment times

Conclusions
(1) Polymeric resin prepared from o-amino acetophenone, biuret and formaldehyde proved the ability to work as chelating
resin where its extractive efficiency depends on the number of chelating groups or groups capable to form the coordinating links of the group -NH$_2$.

(2) The ability of resin in the extraction of metal ions depends on several factors, including the nature of metals and ions and nature of the link between chelating resin and ions.

(3) The ability of chelating resins composition increased with increasing ionic valance.

(4) In this study, it is found the highest extractive efficiency of the resin is when the values pH = 8 where the basic midfield lead to a partial melting of the resin, leading to the difficulty of separating the resin loaded metal ions.

REFERENCES


