An Eco-friendly Synthesis, Characterization, Morphology and Ion Exchange Properties of Tercopolymer Resin derived from 2,4 dihydroxyacetophenone

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Abstract – An effectual chelating resin was synthesized for the detoxification of harmful metal ions using a synthesized resin involving 2,4-dihydroxyacetophenone and catechol with formaldehyde (2,4-HACF-II) in 2:1:3 mol ratio using hydrochloric acid by condensation technique. The synthesized tercopolymmer resin was characterized by elemental analysis, FTIR and 1H NMR spectroscopy. The average molecular weight of resin was determined by nonaqueousconductometric titration method. The physico-chemical parameters have been evaluated for the terpolymer resin. The semicrystalline nature of the synthesized tercopolymer was characterized by scanning electron microscopy (SEM). Batch equilibrium method was employed to study the selectivity and binding capacity of the tercopolymner resin toward certain metal ions such as Cu2+, Co2+, Ni2+, Zn2+ and Pb2+ in different electrolyte concentrations, wide pH ranges and time interervals.

Keywords-Morphology, Resin, Ion exchange, pH range and Batch equilibrium.

1- INTRODUCTION

Ion-exchange may be defined as the reversible exchange of ions between the substrate and surrounding medium. Ion exchange technique can remove traces of ion impurities from water process liquors and given out a product of ultra pure quality in a single efficient and technoeconomically viable manner. Ion exchangers are widely used in analytical chemistry, hydrometallurgy, antibiotics, purification and separation of radioisotopes and find large application in water treatment and pollution control [1, 2]. Lutforet al. [3] prepared a chelating ion exchange resin containing amidoxime functional group. The chelating poly (amidoxime) resin was characterized by FT-IR spectra, TG and DSC analyses. Samir et. al. [4] synthesized ion exchange resin from 8-quinolinyl methacrylate and characterized by conventional methods. The metal ion uptake capacities of synthesized copolymers were estimated by batch equilibration method using different metal ion solutions under different experimental conditions. Three phenol-formaldehyde chelating resins, poly (8-hydroxyquinoline-5, 7 diylmethene) and poly (2-aminophenol-5,7diylmethene) were synthesized and characterized by Ebraheem [5]. The chelating characteristics of these polymers were studied by a batch equilibrium technique. The ion exchange capacity, effect of electrolyte on metal ion up take, rate of metal uptake and distribution of metal ion at different pH with resin copolymer derived from thiosemicarbazone derivatives of phenol compound shows higher order than the resin copolymer derived from semicarbazone derivatives [6]. Recently much work has been carried out to study the ion exchange properties of anchoring functional chelating groups on the polymeric network. The synthesized polymer resins showing versatile applications and properties attracted the attention of scientists and introduce the recent innovations in the polymer chemistry. These polymers can be used as high energy material [7], ion-exchanger [8], semiconductors [9], antioxidants [10], fire proofing agent [11], optical
storage data [12], binders [13], molding materials, [14] etc. Pal et al. [15] has reported electrical conductivity of salicylic acid-biuret/dithio-oxamide / dithiobiurettioxane polymer resins. Ion-exchange technique can remove traces of ionic impurities from water/process liquors and gives out a product of ultra pure quality in a simple efficient and technoeconomically viable manner. Ion-exchangers are widely used in water treatment and pollution control [16,17]. Various Hydroxybenzoic acid-formaldehyde and 4-hydroxyacetophenone-biuret-formaldehyde copolymers have been reported and found to be used as ion-exchangers [18,19]. The polymer resin under investigations is found to be cation exchanger having both ion-exchange group and chelating group in the same polymer matrix therefore the resin can be used selectively for the purpose of purification of waste water. One of the important applications of this chelating and functional polymer is its capability to recover metal ions from waste solution. Hence the chelating ion-exchange property of the 2,4-HACF-II polymer resin was also reported for specific metal ions.

II-EXPERIMENTAL METHOD

Chemicals and reagent

The chemicals such as 2,4-dihydroxyacetophenone and catechol were purified by rectified spirit, formaldehyde (37%), metal chlorides and nitrates (AR grade, Merck) were used. All other solvents and the indicators were of the analytical grade procured from India. Standardized disodium salt of EDTA was used as a titrant for all the complexometric titrations.

Synthesis of 2,4-HACF-II tercopolymer resin

The 2,4-HACF tercopolymer resin was prepared by the condensation polymerization of 2,4-dihydroxyacetophenone (2.72gm, 0.2mol) and catechol (1.10gm, 0.1mol) with formaldehyde (11.25ml, 0.3mol) in hydroxychloric acid medium at 126±2°C in an oil bath for 5h under refluxed condition with occasional shaking. The solid product obtained was immediately removed from the flask as soon as the reaction period was over. It was washed with cold water, dried and powdered. The powder was repeatedly washed with hot water to remove excess of 2,4-dihydroxyacetophenone-catechol-formaldehyde resin, which might be present along with the 2,4-HACF-II tercopolymer resin.

The dried resin was further purified by dissolving in 8% NaOH and regenerated in 1:1 (v/v) HCl/H₂O. This process was repeated twice to separate the pure polymer. The resulting polymer resin washed with boiling water and dried in vacuum at room temperature. The purified tercopolymer resin was finally ground well and kept in a vacuum over silica gel. The yield of the polymer was found to be 84 % and the reaction route for the synthesis is shown in Fig. 1.

![Fig. 1 Formation of 2,4-HACF Tercopolymer Resin](image)

III-CHARACTERIZATION OF TERCOPOLYMER RESIN

Physicochemical and elemental analysis

The tercopolymer resin was subject to micro analysis for C, H and N on an ElementerVario EL III Carlo Ebra 1108 elemental analyzer. The number average molecular weight \( M_n \) was determined by conductometric titration in DMSO medium using ethanolic KOH as the titrant by using 25 mg of sample. A plot of the specific conductance against the milliequivalents of KOH requires for neutralization of 100 g of polymer was made. Inspection of such a plot revealed that there were many breaks in the plot. From this plot, The first break and the last break were noted. The calculation of \( M_n \) by this method is based on the following consideration. On the basis of average degree of polymeration, \( DP \) the average molecular weight has to be determined by following equation.
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\[ \text{Mn} = \text{DP x Repeat unit weight.} \]

The intrinsic viscosity was determined using a Tuan-Fuoss viscometer [20] at six different concentrations ranging from 0.3 to 0.05 wt% of resin in DMF at 30°C. Intrinsic viscosity (\( \eta \)) was calculated by the Huggin’s equation [21] and Kraemer’s equation. [22].

\[
[\eta_{sp}/c] = [\eta] + K_1[\eta]^2C
\]

\[
\ln [\eta_{sp}/c] = [\eta] - K_2[\eta]^2C
\]

**Spectral Analysis**

Electron absorption spectra of tercopolymer (2,4-HACF-II) were recorded in 200 to 800 nm range by using Shimadzu automatic recording double beam spectrophotometer (UV-VIS-NIR Spectrometer) UV-240 at Sophisticated Analytical Instrument Facility, Punjab University, Chandigarh having 10 nm optical path supplied with the instruments were used. An infra-red spectrum of 2,4-HACF-II polymer resin was recorded in the region 4000 to 400 cm\(^{-1}\) were scanned in KBr pellet on Perkin Elmer Model 983 spectrophotometer at Sophisticated Analytical Instrument Facility, Punjab University, Chandigarh. A Nuclear Magnetic Resonance (\(^1\)H NMR) spectrum of newly synthesized polymer resin has been scanned on 90 MHz for proton using BrukerAvance II 400 NMR Sepctrometer in deuterated dimethyl sulphoxide (DMSO-\(d_6\)) at Sophisticated Analytical Instrumentation Facility, Panjab University, Chandigarh. Scanning electron micrograph of polymer has been scanned and magnification by scanning electron microscope at Sophisticated Test and Instrumentation Centre, STIC, Cochin University, Cochin.

**Ion-exchange properties**

The ion-exchange property of the 2,4-HACF-II terpolymer resin was determined at three different variations given below.

**Determination of metal uptake in the presence of electrolytes of different concentrations**

The 2,4-HACF terpolymer (25 mg) was taken in a pre-cleaned glass bottles and each of the electrolytes (25 mL) such as NaClO\(_4\), NaCl, Na\(_2\)SO\(_4\) and NaNO\(_3\) in different concentrations \( v_i \) 0.01, 0.1 and 0.5 M. The pH of the suspension was adjusted to the required value either by adding 0.1 M HCl or 0.1 M NaOH and mechanically stirred for 24 h at 25 ºC for swelling of the terpolymer. Then exactly 0.1 M of the metal ion solution (2 mL) was added and vigorously stirred for 24 h at room temperature. The mixture was then filtered off and washed with distilled water. The filtrate and the washings were collected and then the amount of metal ion adsorbed was estimated by titrating against standard Na\(_2\)EDTA solution. A blank experiment was also performed following the same procedure without the polymer sample. The amount of metal ions taken up by the polymer in the presence of a given electrolyte can be calculated from the difference between the actual value and that from the blank.

**Evaluation of the Distribution of Metal Ions at Different pH**

The distribution of each of the metal ions at various pH ranging from 1.5 to 6 between the polymer phase and the aqueous phase were determined in the presence of 1 M KNO\(_3\) at 25 ºC. The distribution ratio D, may be defined as

\[ D = \frac{\text{Weight (in mg) of metal ions taken up by 1 g of the resin sample}}{\text{Weight (in mg) of metal ions present in 1 mL of the solution}}. \]

**Evaluation of Rate of Metal Ion Uptake**

The time required for attaining the state of equilibrium under the experimental conditions was evaluated by a series of experiments carried out to determine the amount of metal ion adsorbed by the terpolymer at specific time intervals. 25 mg of the polymer sample was mechanically stirred with 25 mL of 1 M KNO\(_3\) to allow the polymer to swell. The pH of the suspension was adjusted to the required value by adding either 0.1 M HCl or 0.1 M NaOH. It was observed that under the given experimental conditions the state of equilibrium established within 24 h at 25 ºC. The rate of metal ion uptake is expressed as the percentage of the metal ion uptake after the specific time related to the state of equilibrium. It is given as,

\[ \text{Metal ion uptake} = \frac{\text{Amount of metal ion adsorbed}}{\text{Amount of metal ion adsorbed at equilibrium}} \times 100 \]

**IV- RESULTS AND DISCUSSION**

The resin sample was dark brown in color and soluble in solvents like N,N-dimethylformamide (DMF),
tetrahydrofuran (THF), dimethylsulphoxide (DMSO), aqueous sodium and potassium hydroxide solutions and insoluble in alcohol, chloroform and carbon tetrachloride. The melting point of resin is 499K and empirical formula of the terpolymer resin is found to be C_{25}H_{22}O_{8} which is in good agreement with the calculated values of C, H and O. The molecular weight of terpolymer resin was also determined by conductometric titration. The calculated molecular weight for 2,4-HACF resin is 3600.

Electronic Spectral Analysis

The UV-Visible spectra of the 2,4-HACF tercopolymer resin in pure DMF were recorded in the region 200-800 nm. The spectra have shown in Fig 2. The UV-Visible spectra of 2,4-HACF tercopolymer resin gave rise to two characteristic bands at about 230 to 270 nm and 310 to 370 nm. This observed positions for the absorption bands clearly indicate the presence of carbonyl group in the sample belonging to 2,4-dihydroxyacetophenone moiety. The former band (more intense) appeared as a result of \( \pi \rightarrow \pi^* \) [23] transition while the latter band (less intense) may be due to \( n \rightarrow \pi^* \) [24] electronic transition.

Infra-Red Spectral Analysis

The IR spectra of 2,4-HACF is presented in Fig. 4. The singlet obtained in the region 2.26 (\( \delta \)) ppm may be due to the methylene proton of Ar-CH\(_2\)-Ar bridge [27]. The weak multiple signals (unsymmetrical pattern) in the region of 7.71 (\( \delta \)) ppm may be attributed to aromatic proton (Ar-H) [28]. The signals in the 8.19 (\( \delta \)) ppm may be due to phenolic hydroxyl protons. The much downfield chemical shift for phenolic –OH indicates clearly the intramolecular hydrogen bonding of -OH group [29]. The signal appeared in the region at 6.32 (\( \delta \)) ppm is due to the methylene proton of Ar-CH\(_2\) bridge [30]. The methyl protons of the Ar - CO - CH\(_3\) moiety may be identified by the intense peak at 3.28 (\( \delta \)) ppm.

Scanning electron microscopy (SEM)

The scanning electron micrograph of 2,4-HACF-II tercopolymer resin have been scanned and magnification...
by analytical scanning electron microscope at Sophisticated Test and Instrumentation Centre, STIC, Cochin University, Cochin as shown in the Fig.5. Surface analysis has found great use in understanding the surface features of the polymers. SEM gives the information of surface topology and defect in the structure. The semi crystalline nature of 2,4-HACF II tercopolymer resin was established by scanning electron microscopy. The morphology of resin sample thus identified by SEM as crystalline as well as amorphous or transition between crystalline and amorphous, showing more or less good ion capacity. The polymerization reaction proceeds by introducing amorphous character in the tercopolymer sample.

Fig. 5 Scanning electron microscopy of 2,4-HACF TercopolymerResin

Ion-exchange properties

Batch equilibrium technique developed by De Geiso et al. [31] and Gregor et al. [32] was used to study ion-exchange properties of 2,4-HACF-II polymer resin. The results of the batch equilibrium study carried out with the polymer 2,4-HACF-II are presented in Tables 1–3. Five metal ions Cu\(^{2+}\), Ni\(^{2+}\), Co\(^{2+}\), Zn\(^{2+}\) and Pb\(^{2+}\) in the form of aqueous metal nitrate solution were used. The ion-exchange study was carried out using three experimental variables such as a) electrolyte and its ionic strength b) uptake time and c) pH of the aqueous medium.

Effect of Electrolytes on Metal Ion Uptake

Generally as concentration increases of the electrolyte, the ionization decreases, the number of ligands decreases in the solution which form the complex with less metal ions and more ions are available for adsorption. The variable metal ions uptake capacity of 2,4-HACF tercopolymer resin may be due to the strong and weak complex formation between electrolyte ligands and metal ions. Examination of the data given in Table 1 reveals that the amount of metal ions taken up from a given amount of tercopolymers depends on the nature and concentration of the electrolyte present in the solution. In presence of perchlorate, chloride and nitrate ions, the uptake of Cu\(^{2+}\) and Ni\(^{2+}\) ions increases with increasing concentration of the electrolytes, whereas in presence of sulphate ions the amount of the above mentioned ions taken up by the tercopolymers decreases with increasing concentration of the electrolytes [33]. Moreover, the amount of Co\(^{2+}\), Zn\(^{2+}\) and Pb\(^{2+}\) ions taken up by the tercopolymer samples decreases with increasing concentration of the chloride, nitrate, perchlorate and sulphate ions [33]. This may be explained in terms of the stability constants of the complexes which Cu\(^{2+}\), Ni\(^{2+}\), Co\(^{2+}\), Zn\(^{2+}\) and Pb\(^{2+}\) ions from with these anions. SO\(_4^{2-}\) might form rather strong complexes with Ni\(^{2+}\) and Cu\(^{2+}\) ions, while ClO\(_4^-\), NO\(_3^-\) and Cl\(^-\) might form weak complexes and, therefore, might not be expected to influence the position of the Ni\(^{2+}\) and Cu\(^{2+}\) chelates equilibrium as much as SO\(_4^{2-}\). Sulphate, perchlorate, nitrate and chloride might form rather strong chelates with Co\(^{2+}\), Zn\(^{2+}\) and Pb\(^{2+}\) and therefore might be expected to influence the position of the Co\(^{2+}\), Zn\(^{2+}\) and Pb\(^{2+}\) chelates equilibrium.

Table 1: Evaluation of the uptake of metal ions by 2,4-HACF-II resin

| Metal ions | Concentration of electrolytes (mol/L) | Weight of metal ion uptake in presence of electrolytes (mg) |
|------------|--------------------------------------|----------------------------------------------------------|
| Cu\(^{2+}\) | 0.01 | 2.54 | 2.42 | 3.59 |
|           | 0.05 | 2.67 | 2.54 | 2.78 |
|           | 0.10 | 2.94 | 2.79 | 2.11 |
|           | 0.50 | 3.23 | 3.46 | 1.45 |
|           | 1.00 | 3.32 | 3.54 | 0.83 |
| Ni\(^{2+}\) | 0.01 | 1.56 | 1.32 | 2.78 |
|           | 0.05 | 2.13 | 1.67 | 2.04 |
|           | 0.10 | 2.96 | 2.03 | 1.45 |
|           | 0.50 | 3.43 | 2.23 | 0.89 |
|           | 1.00 | 4.11 | 2.56 | 0.52 |
| Co\(^{2+}\) | 0.01 | 1.59 | 1.95 | 1.65 |
|           | 0.05 | 1.42 | 1.72 | 1.38 |
|           | 0.10 | 1.18 | 1.49 | 1.27 |
|           | 0.50 | 0.84 | 1.11 | 1.11 |
|           | 1.00 | 0.47 | 0.72 | 0.82 |
| Zn\(^{2+}\) | 0.01 | 1.75 | 2.39 | 1.89 |
|           | 0.05 | 1.52 | 2.07 | 1.52 |
|           | 0.10 | 1.24 | 1.67 | 1.21 |
|           | 0.50 | 0.92 | 1.24 | 1.17 |
|           | 1.00 | 0.67 | 0.75 | 0.81 |
| Pb\(^{2+}\) | 0.01 | 1.46 | 1.83 | 1.92 |
|           | 0.05 | 1.29 | 1.69 | 1.64 |
|           | 0.10 | 1.17 | 1.34 | 1.32 |
|           | 0.50 | 1.03 | 1.12 | 1.03 |
|           | 1.00 | 0.82 | 0.68 | 0.75 |
Evaluation of the Rate of Metal Ion Uptake

The data of metal ion uptake at various shaking time intervals is given in Table 2. These results indicate that the time taken for the uptake of the different metal ions at a given stage depends on the nature of metal ion under given conditions. It is found that Cu^{2+}, Ni^{2+}, Co^{2+} and Zn^{2+} ions require about 5h for the establishment of the equilibrium where as Pb^{2+} ions required almost 6h for equilibrium. Thus, the rate of metal ion uptake follows the order Cu^{2+}, Ni^{2+}, Co^{2+}, Zn^{2+}> Pb^{2+} for all the terpolymers. The trend is good agreement with earlier workers [34, 35].

Table 2 Evaluation of rate metal ion uptake by 2,4-HACF-II resin

| Metal ions | Equilibrium attainment (%) |
|------------|-----------------------------|
|            | Time (h) 1 2 3 4 5 6        |
| Cu^{2+}    | 62.6 72.7 81.4 86.3 94.1 -  |
| Ni^{2+}    | 69.4 75.3 83.3 89.4 95.3 -  |
| Co^{2+}    | 47.4 63.4 77.3 85.5 95.9 -  |
| Zn^{2+}    | 53.2 63.2 72.2 85.3 94.2 -  |
| Pb^{2+}    | 32.4 49.1 65.2 75.1 85.4 95.2 |

Distribution Ratio of Metal Ions at Different pH

The effect of pH on the amount of metal ion distributed between two phases can be explained by the result shown in the Table 3. The result indicate that the relative amount of metal ion adsorbed by the tercopolymer resin increase with increasing pH of the medium. The study was carried out up to definite pH value for the particular metal ion to prevent hydrolysis of the metal ions at higher pH. Cu^{2+} and Ni^{2+} have highest distribution ratio at pH 2.5 to 6, therefore these ions can be taken up more selectively between the ranges of pH 2.5-6. The other three ions Co^{2+}, Zn^{2+} and Pb^{2+} have lower distribution ratio over pH range of 2.5 to 6. The high distribution ratio may be due to higher stability constant of metal ligand complexes and vice versa. Cu^{2+} and Ni^{2+} may form more stable complex, indicate that these ions adsorbed more selectively to the higher extent, rather than Co^{2+}, Zn^{2+} and Pb^{2+} which may form rather weak complexes with ligand during the course of time, therefore these ions adsorb to a least extent over an entire pH range study [36]. Thus, the order of selectivity of metal ions by all the terpolymer is found to be Cu (II) > Ni (II) > Co (II) > Zn (II) > Pb (II). The results are in good agreement with earlier co-workers [36]. Hence, the results of this type of study are helpful in selecting the optima pH for a selective uptake of a metal ion from a mixture of different ion.

Table 3 Evaluation of distribution ratio D, at different pH

| Meta 1 ions | Distribution ratio (KD) of the metal ions |
|-------------|------------------------------------------|
|             | pH of the medium                          |
|             | 1.5 2 2.5 3 3.5 4 5 6                     |
| Cu^{2+}     | - 74.8 92.5 212.3 489.7 865.4 1134.4     |
| Ni^{2+}     | - 63.4 79.3 145.3 356.1 567.1 874.3      |
| Co^{2+}     | - 48.2 57.4 93.4 143.1 268.5 453.3       |
| Zn^{2+}     | - 57.4 62.4 89.3 165.5 244.6 343.2       |
| Pb^{2+}     | - 42.4 62.4 79.8 112.3 156.2 278.4       |

V- CONCLUSION

A tercopolymer 2,4-HACF-II based on the condensation reaction of 2,4dihydroxyacetophenone and catechol with formaldehyde in the presence of acid catalyst was prepared. The 2,4-HACF-II resin is a selective chelating cation exchange terpolymer resin for certain metals. The uptake capacities of metal ions by the polymer resin were pH dependent. The uptake capacities of metal ions by the tercopolymer resin were carried out by batch equilibrium technique. Due to the considerable difference in the adsorption capacity at different pH media, the rate of metal uptake and distribution ratio at equilibrium, it may be possible to use the tercopolymer for separation of particular metal ions. Also from the FTIR and 3H NMR spectral studies the proposed structure of the 2,4-HACF-II polymer resin has been determined.

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