Preparation and Characterization of (PVA+ZnO) Polymer Films Doped With Cu$^{2+}$ Ions

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Abstract: 5 mol% (PVA+ZnO): (x mol%) Cu$^{2+}$ ions doped polymer films were prepared by solution casting technique. The obtained films are analyzed using different characterization techniques like SEM, FTIR, optical and EPR. The SEM micrographs shows the improvement in the crystallinity of the Cu$^{2+}$ doped PVAZn polymer films compared with the pure PVAZn film. The Fourier transform infrared spectroscopy (FTIR) spectrum shows a characteristic peaks of C-H, C=C, CH$_2$ and C-O bonds. The optical absorption spectrum exhibits broad bands at 412 nm and 692 nm, which were corresponds to the $^3B_{18} \rightarrow ^2E_g$ and $^3B_{18} \rightarrow ^2B_{2g}$ transition. The EPR spectra of all the polymer samples exhibits resonance signals characteristic of Cu$^{2+}$ ions. The spin-Hamiltonian parameter indicate that the ground state of Cu$^{2+}$ ions is $d^9$ and the site symmetry around Cu$^{2+}$ ions is tetragonally distorted octahedral site. The number of spins participating in resonance (N) and paramagnetic susceptibility ($\chi$) were calculated from the EPR data for various concentrations of Cu$^{2+}$ ions.

I. Introduction

Polymers have potential applications as permanent and transient data storage materials or as basic materials to the fabrication of active and passive light guides. In the holographic recording, “ion” doped organic polymers are highly efficient compared with other polymer films [1]. The main advantage of the polymer films are simplicity of their preparation. The present system consists of simple combinations of poly (vinyl alcohol) and zinc acetate as host materials. PVA is important due to its enhanced properties in industrial, commercial, medical and food applications [2]. PVA is the most widely produced water-soluble synthetic polymer having a wide range of dopant dependent properties. It has a good resistance and excellent physical properties [3]. Surface capping by PVA is useful in achieving a higher and relatively steady UV sensitivity in ZnO materials. Zinc is one of the best promising materials for electrical devices, including transparent conductive films, light emitting diodes and photo-catalyst [4]. In particular, introducing ZnO into polymeric matrices can modify the optical, electrical and mechanical properties of polymer films [5]. The doping of copper ions into the host material will influence the packing efficiency, which change the thermal properties of the polymer material [1].

II. Experimental

Analytical grade of poly (vinyl alcohol), Zinc Acetate and copper sulphate (CuSO$_4$·5H$_2$O) purchased from Himedia, Mumbai, India and double distilled water is used as solvent throughout the experiment. The 5 mol% of each PVA and zinc acetate were dissolved in double distilled water separately and stirred with magnetic stirrer until the chemicals completely dissolved and forms a clear viscous solution. Then the desired volume of both solutions were mixed and stirred for 30 min. The desired concentration of CuSO$_4$·5H$_2$O (0.2, 0.4, 0.6 & 0.8 mol %) solutions were added drop wise to the prepared 5 mol% of (PVA+ZnO) solution and stirred thoroughly with magnetic stirrer for another 30 min. Then the solution was casted on petri dishes and the films were collected after 7 days. Here after the combination of (PVA+ZnO) polymer films were referred as PVAZn polymer films. The surface and cross sectional morphology of the pure PVAZn and copper doped PVAZn polymer films were recorded using Hitachi make model, S-2830N, Japan made Scanning Electron Microscope (SEM).

The infrared absorption spectra of the polymer films were measured at room temperature in the wave number range 2000-600 cm$^{-1}$ by a Jasco, FT-IR130 Japan make Fourier Transform computerized infrared spectrophotometer. The optical absorption spectra of the folded polymer samples (thickness 1mm ±0.1mm) recorded at room temperature using a dual-beam spectrophotometer JASCO V-670, Japan in the wavelength range 400-800 nm. X-Band EPR spectra were recorded on a Bruker EMX X-band EPR spectrometer working at 9.15 GHz using a rectangular cavity operating at a 100 kHz modulation frequency at room temperature.

III. Results and discussions

3.1 Scanning Electron Microscope (SEM)

The surface morphology of the PVA+ZnO films are studied with SEM images. The SEM images of pure and Cu$^{2+}$ doped PVAZn polymer films are shown in Fig.1 (a) and (b). The prepared films have homogeneous distribution of Zn particles. When copper ions introduced in to the PVAZn matrix and compared
with the pure polymer matrix. The PVAZn:Cu²⁺ film shows spherical grain distribution with a uniform density. The clear change in surface morphology was observed with the Cu²⁺ ion doped in PVAZn polymer film.

3.2 Fourier transform infrared studies

The FTIR spectra of pure and Cu²⁺ doped PVAZn polymer films were shown in Fig.2. The FTIR spectra consist of several groups of bands for the undoped and copper doped polymer films. The main bands observed in the range of 610-691 cm⁻¹ has been assigned to Zn-O stretching of the polymer complex [6]. The main band at 837 cm⁻¹ has been assigned to C-C stretching mode of the host polymer. The bands observed at 1000-1100 cm⁻¹ has been assigned to C-O-C stretching mode formed by the cross linking reaction of PVA with ZnO and the bands around 1450-1558 cm⁻¹ has been assigned to bending vibrations of CH₂ of PVA group [7]. The band at 1642 and 1702 cm⁻¹ has been assigned to stretching mode of C-O. The main change for the undoped and copper doped polymer is shifting of the absorption bands. The Zn-O bands at 616 and 667 cm⁻¹ for pure PVAZn polymer shifts towards the 641 and 691 cm⁻¹ in the Cu²⁺ PVAZn polymer film. This indicates the cross linking of Cu-O with Zn-O in the host polymer [8]. The band at 837 cm⁻¹ shifts towards the lower region 796 cm⁻¹, which has been assigned to C-C mode indicating the amorphous nature of the sample. The band at 1702 cm⁻¹ became broaden in the doping of Cu²⁺ ions in PVAZn polymer due to the alteration of PVA chains with Zn-O and Cu-O. Therefore introducing the Zn-O and Cu-O in the PVA structure modifies the semi crystalline structure of PVA confirming the cross linking of metal ions with the main chain of the PVA. The observed band positions and their corresponding band assignments presented in Table 1.

3.3 Optical absorption studies

The room temperature optical absorption spectra were recorded for Cu²⁺ ions doped PVAZn polymer film in the wavelength range 400-800 nm. The absorption spectra exhibit two absorption bands at 412 and 692 nm respectively. There is no specific change in the band positions except variation in intensity of the bands with the concentration. Hence as a remarkable spectrum, the absorption spectrum of (0.2 mol%) Cu²⁺ doped PVAZn polymer film is given as Fig.3. The broad band at 692 nm has been assigned to 3¹B₁g → 1¹B₁g transition of Cu²⁺ ion in a distorted octahedral site and the band at 412 nm has been assigned to 3¹B₁g → 2¹E₁g transition [1].

The investigations of optical transitions and electronic band structure can be analyzed using the fundamental absorption edge in the UV region of the optical absorption spectrum. The main feature of the absorption edge is an exponential increase of the absorption coefficient α (ν) with photon energy hν [9]. The absorption coefficient α (ν) can be calculated using the formula,

\[
α (ν) = (A/d) 2.303
\]

where A is the absorbance at frequency 'ν' and 'd' is the thickness of the sample. The optical band gap for direct and indirect transitions can be determined by using the relation,

\[
E_{\text{opt}} = hν - (α hν/B)^{1/2}
\]

where B is a constant. The direct and indirect band gaps were calculated from the plots of (αhν)² and (αhν)¹/₂ as a function of photon energy (hν) and are shown in Fig.4(a) and (b). The corresponding values are taken by extrapolating the linear portion of absorption coefficient meet at x=0.

Urbach plots are the natural logarithm of absorption coefficient ln (α) is plotted against photon energy (hν). In the present study such an Urbach plots were plotted and are shown in Fig.5. The direct, indirect optical band gaps and Urbach energies were calculated with respect to the copper concentration in the PVAZn polymer films and are presented in Table 2. From the Table, it is observed that the direct and indirect band gaps energy decrease with the increase of copper concentration. This may be due to the loose packing of the host complex with the copper.

3.4 Electron paramagnetic resonance (EPR)

The EPR spectrum of the undoped polymer film confirming that the starting materials used in the present work were free from transition metal impurities or any other paramagnetic contamination. When various mol% of CuSO₄ were added to PVAZn films, the EPR spectra of all the investigated samples exhibits a well resolved resonance signals as shown in Fig.6. The EPR spectrum exhibits four parallel and one perpendicular line for the samples under study. For Cu²⁺ ions, a regular octahedral site may not exist because the cubic symmetry is disturbed by electronic-hole in the degenerate dₓ²−y² orbital and this produces the tetragonal distortion. The EPR spectra of Cu²⁺ ions in the present polymer film can be well interpreted in terms of the spin-Hamiltonian for d⁷ ions in an axial symmetry [10],

\[
\mathcal{H} = β[g_{∥}H_S + g_{⊥} (H_I, S_x + H_y, S_y)] + A_{∥} I_z S_z + A_{⊥} (I_x S_x + I_y S_y)
\]

where the terms have their usual meanings given in our previous papers [11].
In EPR spectrum, one can get four parallel and four perpendicular signals for copper ions. However, in the present study, we observed only four parallel and one perpendicular signal due to $^{65}$Cu and $^{63}$Cu isotopes. The spectral parameters for all the samples are well resolved with four parallel components ($g_1 \approx 2.38$) and a perpendicular component around ($g_\perp \approx 2.05$). The observed $g$ and A values are presented in Table 3. From the table, it can be observed that, the $g$ values follow the trend $g_1 > g_\perp > 2.0023$. We consider that the ground state for the paramagnetic electrons is $d_{x^2-y^2}$ orbital and the Cu$^{2+}$ ions being located in the distorted octahedral sites $(D_{4h})$ [12-15].

3.4 (a) Calculation of spin concentration (N)

The number of spins participating in resonance can be calculated by comparing the present system with standard known concentration of CuSO$_4$ 5H$_2$O. The well – known expression given by Weil et al., [16] was used to calculate the spin concentration in the present work and is given by,

$$ N = \frac{A_x(\text{Scan})^2 \ G_{\text{std}} (B_{\text{m, std}}) \ (P_{\text{std}})^{1/2} \ [S(S+1)]_{\text{std}}}{A_{\text{std}} (\text{Scan})^2 G_x (B_{\text{m}}) (g_x)^2 (P_x)^{1/2} [S(S+1)]_x} $$

where the terms have their usual meanings [11].

The spin concentration (N) participating in resonance for the present system was calculated for different concentrations of copper ions (0.2 to 0.8 mol%) and a plot was drawn between copper concentrations against spin concentration for this polymer system and is shown in Fig.7. From the graph, it can be observed that the spin concentration participating in resonance (N) has a non-monotonic (i.e., increases then decrease) behavior with the increase in concentration. This may be due to the presence of other copper states existing in the polymer complex [17].

3.4 (b) Paramagnetic susceptibility

The paramagnetic susceptibility of the polymer films at room temperature was calculated from the EPR data using the formula [18],

$$ \chi = \frac{N g^2 \beta^2 J(J+1)}{3K_B(T)} $$

where the terms have their usual meanings [11]. The paramagnetic susceptibilities were calculated for PVAZn: Cu$^{2+}$ (0.2 - 0.8 mol%) polymer films and a plot was drawn for the reciprocal of susceptibility $(1/\chi)$ as a function of concentrations is shown in Fig.8. From the graph, the susceptibility of the samples decreases with the concentrations of copper in the host matrix.

IV. Conclusions

- PVAZn:Cu$^{2+}$ polymer films were prepared by solution casting technique at room temperature.
- The clear and uniform surface structure was observed from the Scanning electron microscopy (SEM).
- From the EPR and optical absorption spectral studies, the Cu$^{2+}$ ions occupied in tetragonally distorted octahedral sites in the host matrix.
- The decrease in paramagnetic susceptibility indicates the increase in amorphous nature of the sample.
- The FT-IR spectrum exhibits the Zn-O, CH$_2$, C-H, C=O, C=C groups of stretching and bending vibrations.

References


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Table 1. FTIR band positions and their corresponding band assignments of PVAZn and Cu$^{2+}$ ions doped PVAZn polymer films.

<table>
<thead>
<tr>
<th>S.No</th>
<th>Cu$^{2+}$ (mol%)</th>
<th>PVAZn:Cu$^{2+}$ (mol%)</th>
<th>Band assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.2</td>
<td>616 619 623 647 641 641</td>
<td>Zn-O</td>
</tr>
<tr>
<td>2</td>
<td>0.4</td>
<td>667 676 679 680 693 750</td>
<td>Zn-O</td>
</tr>
<tr>
<td>3</td>
<td>0.6</td>
<td>837 833 799 789 750 837</td>
<td>CH$_2$</td>
</tr>
<tr>
<td>4</td>
<td>0.8</td>
<td>1080 1019 1022 1101 1147</td>
<td>CO</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1137 - 1093 1146 -</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td></td>
<td>1459 1443 1444 1431 1463</td>
<td>CH$_2$</td>
</tr>
<tr>
<td>6</td>
<td></td>
<td>1521 1532 1517 1517 1516</td>
<td>C=C</td>
</tr>
<tr>
<td>7</td>
<td></td>
<td>1541 1558 1561 1558 1558</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td></td>
<td>1642 1619 1632 1640 1621</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td></td>
<td>1702 1700 1686 1692 1682</td>
<td></td>
</tr>
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</table>

Table 2. The direct, indirect band gaps and Urbach energies of Cu$^{2+}$ ions doped PVAZn polymer films.

<table>
<thead>
<tr>
<th>S.No</th>
<th>Cu$^{2+}$ (mol %)</th>
<th>Direct (eV)</th>
<th>Indirect (eV)</th>
<th>Urbach energy ($\Delta E$, eV)</th>
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<tr>
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<td>3.67</td>
<td>3.40</td>
</tr>
<tr>
<td>2</td>
<td>0.4</td>
<td>3.82</td>
<td>3.40</td>
<td>2.78</td>
</tr>
<tr>
<td>3</td>
<td>0.6</td>
<td>3.51</td>
<td>3.31</td>
<td>1.83</td>
</tr>
<tr>
<td>4</td>
<td>0.8</td>
<td>3.29</td>
<td>3.13</td>
<td>1.34</td>
</tr>
</tbody>
</table>

Table 3. The g values and the hyperfine splitting constant ($A$) values of Cu$^{2+}$ ions doped in PVAZn polymer films calculated using EPR spectra.

| S.No | Cu$^{2+}$ (mol %) | $g_{||}$ | $g_{\perp}$ | $A \times 10^{9}$ (cm$^{-1}$) | No. of spins (10$^{25}$/kg) |
|------|------------------|----------|-------------|-------------------------------|------------------------------|
| 1    | 0.2              | 2.45     | 2.08        | 138.4                         | 1.214                        |
| 2    | 0.4              | 2.39     | 2.05        | 138.4                         | 2.169                        |
| 3    | 0.6              | 2.39     | 2.07        | 138.4                         | 1.9039                       |
| 4    | 0.8              | 2.38     | 2.07        | 130.9                         | 1.4726                       |

Fig. 1(a) & (b)

Fig. 2
Fig. 3

Absorbance (arb.units)

Wavelength (nm)

Fig. 4 (a)

Fig. 4 (b)
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Fig.5

![Graph showing ln(α) vs. hv (eV)]

Fig.6

![Graph showing first derivative of absorption vs. Magnetic field (mT)]

Fig.7

![Graph showing No of spin concentration vs. Concentration (mol%)]
Caption for figures:
Fig.1: SEM images of (a) pure PVAZn and (b) (0.4 mol%) Cu$^{2+}$:PVAZn polymer film.
Fig.2: FTIR spectra of a pure PVAZn and Cu$^{2+}$:PVAZn polymer films recorded at room temperature.
Fig.3: Optical absorption spectrum of (0.2 mol %) Cu$^{2+}$:PVAZn polymer film.
Fig.4: A plot of (a) $(\alpha h \nu)^2$ as a function of $h \nu$ and (b) $(\alpha h \nu)^{1/2}$ as a function of $h \nu$ for Cu$^{2+}$:PVAZn polymer films.
Fig.5: A plot of ln(α) as a function of $h \nu$ for PVAZn:Cu$^{2+}$ polymer films.
Fig.6: EPR spectra of Cu$^{2+}$:PVAZn films recorded at room temperature.
Fig.7: A plot of no.of spins (N) against concentration of copper in Cu$^{2+}$:PVAZn polymer films.
Fig.8: A plot of reciprocal of susceptibility (1/$\chi$) as a function of copper concentration in Cu$^{2+}$:PVAZn polymer films.