SYNTHESIS, CHARACTERIZATION AND ANTIBACTERIAL STUDIES ON DIVALENT TRANSITION METAL COMPLEXES OF HYDRAZINE WITH ARYL SUBSTITUTED ACETIC ACIDS

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ABSTRACT

Metal phenylacetate Sesquihydrazinate hydrates of the formula M(PhAc)\textsubscript{2}.1.5N\textsubscript{2}H\textsubscript{4}.H\textsubscript{2}O where M=Co,Ni or Cd and Metal phenoxyacetate Sesquihydrazinate hydrates M(PhOAc)\textsubscript{2}.1.5N\textsubscript{2}H\textsubscript{4}.H\textsubscript{2}O Where M=Co, Ni, Cd or Mn have been prepared and characterized by analytical, IR spectral and thermal studies. The composition of complexes have been determined by analytical studies. Infrared spectral data indicate that the bidentate bridging by hydrazine molecules and monodentate coordination by carboxylate ions to the central metal ion. Thermogravimetry (TG) and differential thermal analyses (DTA) in air have been used to study the thermal behaviour of the complexes. The simultaneous TG-DTA curves of all the complexes in air resulted in the formation of respective metal or metal oxide as final residue. These complexes decompose exothermically either in single step or decompose through respective metal carboxylate intermediates. The antibacterial activity of the prepared complexes screened against *Staphylococcus aureus, Escherichia coli, Pseudomonas aeruginosa and Proteus mirabilis.

Keywords: Phenylacetate Sesquihydrazinate hydrates, phenoxyacetate Sesquihydrazinate hydrates.

1. INTRODUCTION

Hydrazine is a versatile ligand and forms a wide variety of complexes with various metal ions. The monodentate and bridging bidentate coordination of the hydrazine molecule on complexation has been well documented in the literature (Braibanti et al., 1968). The emerging interest in these hydrazine complexes is mainly due to their structure and thermal behaviour. Hydrazine carboxylates of the transition metal ions with variety of acids have been reported. These include simple aliphatic monocarboxylic acid (Ravindranathan and Patil, 1983; Sivasankar and Govindrajan, 1994a; Sivasankar and Govindrajan, 1995a; Sivasankar and Govindrajan, 1997; Vikram and Sivasanka, 2007) aliphatic dicarboxylic acids (Gajapathy et al., 1983; Sivasankar and Govindrajan, 1994b; Govindrajan et al., 1995; Yasodhai and Govindrajan, 2000; Sivasankar, 2006) aromatic mono and dicarboxylic acids (Kuppasamy and Govindrajan, 1996; Sivasankar, 2006). Most of these formed as mono, bis-hydrazine or hydrazinium carboxylates and few tris-hydrazine carboxylates are known with aliphatic substituted acetic acid (Kuppasamy and Govindrajan, 1995) and other Inorganic anions (Sivasankar and Govindrajan, 1995; Nicholls et al., 1996). Generally all phenolic derivatives show antibacterial property. Particularly 2,4-dichlorophenoxycetic acid altered envelope properties of the bacteria Escherichia coli, such as hydrophobic index (Blague et al., 2001). This prompted us to make antibacterial study of hydrazinium salt of dichlorophenoxycetic acid against Escherichia coli.

In this context, we present some new metal hydrazine complexes with aromatic carboxylic acids namely phenylacetic acid and phenoxyacetic acid.

2. EXPERIMENTAL

2.1. Preparation of M(PhAc)\textsubscript{2}.1.5N\textsubscript{2}H\textsubscript{4}H\textsubscript{2}O where M=Co,Ni or Cd

The Cobalt, Nickel and Cadmium complexes were prepared by the addition of an aqueous solution (50 ml) of hydrazine hydrate (0.2 ml, 0.004 m) and phenylacetic acid (0.5 g, 0.0036 m) to the corresponding aqueous solution (50 ml) of metal nitrate hydrates (Co(NO\textsubscript{3})\textsubscript{2}.6H\textsubscript{2}O,0.5 g, 0.0017 m, Ni(NO\textsubscript{3})\textsubscript{2}.6H\textsubscript{2}O,0.5 g, 0.0017 m, Cd(NO\textsubscript{3})\textsubscript{2}.4H\textsubscript{2}O, 0.5 g, 0.0016 m). The mixture was stirred well to get a clear solution. This solution was concentrated on a water bath to 20ml and it was kept for complexation. After 15 minutes complex was formed. It was filtered and washed by using water, alcohol and diethyl ether and air dried.

2.2. Preparation of Mn(PhAc)\textsubscript{2}.1.5N\textsubscript{2}H\textsubscript{4}H\textsubscript{2}O

The manganese complex was also prepared by the same procedure with aqueous solution (50 ml) of phenylacetic acid (0.5 g, 0.0036 m), hydrazine
hydrate (0.4 ml, 0.008 m) and aqueous solution of manganese acetate hydrate (0.5 g, 0.002 m).

2.3. Preparation of M(PhOAc) \( \cdot \frac{1}{2} \text{SN} \ H \cdot \text{H O} \) Where \( M=\text{Co, Ni, Cd or Mn} \)

The same procedure was followed with aqueous solution of Phenoxoacetic acid (0.5 g, 0.0032 m), hydrazine hydrate (0.2 ml, 0.004 m) and metal nitrate hydrates (0.5 g, 0.0017 m) (or) Manganese acetate hydrate (0.5 g, 0.002 m).

2.4. Quantitative methods

The hydrazine content in the complexes was determined by titration using KI\( \text{O}_3 \) as the titrant (Von Burg and Stout, 1991). The percentage of metals in the complexes was estimated by the standard methods given in the Vogel’s textbook (Von Burg and Stout, 1991).

2.5. Physico-chemical techniques

2.5.1. Infrared spectrum

The infrared spectrum of the solid precursor sample was recorded by the KBr disc technique using a Perkin Elmer 597/1650 spectrophotometer.

2.5.2. Thermal analysis

The simultaneous TG-DTA experiment was carried out in Shimadzu DT40, Stanton 781 and STA 1500 thermal analyzer. Thermal analysis was carried out in air at the heating rate of 10\(^\circ\)C per minute using 5-10 mg of the sample. Platinum cups were used as sample holders and alumina as reference. The temperature range was ambient to 700\(^\circ\)C.

2.5.3. Biological assay

The antibacterial activities of the prepared complexes were determined by the disc diffusion method. The bacteria were cultured in nutrient agar medium and used as inoculum for the study. The antibacterial activity of the synthesized compounds of 25\(\mu\)g, 50\(\mu\)g, 100\(\mu\)g and 200\(\mu\)g concentrations were tested against Staphylococcus aureus, Escherichia coli, Pseudomonas aeruginosa and Proteus mirabilis. The inhibition zones were calculated and recorded.

3. RESULTS AND DISCUSSION

3.1. Chemical formula determination of the complexes

The chemical formula \([M(\text{PhOAc})]_2 \cdot \frac{1}{2} \text{SN} \text{H}_2 \text{O} \) (M=Ni, Co, Mn, Cd) and \([\text{M(PhOAc)}]_2 \cdot \frac{1}{2} \text{SN} \text{H}_2 \text{O} \) (M = Co, Ni, Cd, Mn) has been assigned to the prepared complexes, based on the observed and calculated percentage of hydrazine and metals, which are found to match closely with the calculated values (Table 1).

### Table 1. Compositional analysis data of the prepared complexes.

| Compound          | Molecular weight (gm) | Hydrazine (%) | Metal (%) |
|-------------------|-----------------------|---------------|-----------|
|                   | Obsd value | Calcd value | Obsd value | Calcd value |
| Co(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 397.35 | 12.25 | 12.16 | 14.55 | 14.92 |
| Ni(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 395.08 | 12.66 | 12.17 | 14.50 | 14.86 |
| Cd(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 449.31 | 9.93 | 10.70 | 26.2 | 25.24 |
| Mn(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 391.3 | 12.25 | 12.28 | 14.00 | 14.03 |
| Co(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 427.39 | 10.09 | 11.25 | 13.00 | 13.80 |
| Ni(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 427.1 | 10.65 | 11.25 | 13.00 | 13.74 |
| Cd(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 481.33 | 9.40 | 9.99 | 24.46 | 23.46 |
| Mn(PhOAc)\(\cdot\)2.1.5N\text{H}_2\text{O} | 423.32 | 10.65 | 11.36 | 12.00 | 12.79 |

PhAc - Phenylacetate

3.1.1. PhOAc – Phenoxoacetic

These are in good agreement with proposed formulae of the complexes.

3.2. FT-IR spectral analysis

The hydrated derivatives displayed a broad band in the region 3625-3282 cm\(^{-1}\) due to O-H stretching shows the presence of water molecule. The absorption band in the region 3246-3224 cm\(^{-1}\).This is due to the N-H stretching frequency of N\(\text{H}_2\text{O}\). The complexes show asymmetric and symmetric stretching frequencies of COO in the region 1610-156cm\(^{-1}\) and 1392-1338 cm\(^{-1}\) respectively. The \(\Delta\gamma(\gamma_{\text{sym}}-\gamma_{\text{sym}})\) of COO in range >222 cm\(^{-1}\) confirms the monodentate coordination of carboxylate anion. In the complexes, the N-N stretching is seen in the range 983-943 cm\(^{-1}\) confirming the bridging bidentate coordination of hydrazine (Sivasankar and Govindrajan, 1996). The IR spectra of the prepared complexes are displayed in Fig. 1-8.

### Table 2. FT-IR spectral data of the prepared complexes.

| Compound          | \(\gamma(\text{O-H})\) | \(\gamma(\text{N-S})\) | \(\gamma_{\text{asy}}(\text{COO})\) | \(\gamma_{\text{sym}}(\text{COO})\) | \(\Delta\gamma\) |
|-------------------|-----------------------|-----------------------|-------------------------------|-------------------------------|-----------------|
| Co(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 3294 | 3226 | 1606 | 1382 | 224 | 970 |
| Ni(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 3292 | 3224 | 1604 | 1382 | 222 | 975 |
| Cd(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 3303 | 3246 | 1610 | 1384 | 226 | 958 |
| Mn(PhAc)\(\cdot\)1.5N\text{H}_2\text{O} | 3625 | - | 1566 | 1392 | -943 |
| Co(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 3282 | 3228 | 1598 | 1342 | 256 | 948 |
| Ni(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 3282 | 3244 | 1585 | 1344 | 247 | 983 |
| Cd(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 3326 | 3242 | 1608 | 1346 | 262 | 968 |
| Mn(PhOAc)\(\cdot\)2.15N\text{H}_2\text{O} | 3330 | 3240 | 1583 | 1338 | 245 | 964 |

\(\Delta\gamma = (\gamma_{\text{asy}}(\text{COO}) - \gamma_{\text{sym}}(\text{COO}))\)
Fig. 1. Infrared spectrum of Co(PhAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 2. Infrared spectrum of Ni(PhAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 3. Infrared spectrum of Cd(PhAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 4. Infrared spectrum of Mn(PhAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 5. Infrared spectrum of Co(PhOAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 6. Infrared spectrum of Ni(PhOAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 7. Infrared spectrum of Cd(PhOAc)$_2$.15N$_2$H$_4$.H$_2$O.

Fig. 8. Infrared spectrum of Mn(PhOAc)$_2$.15N$_2$H$_4$.H$_2$O.

Table 3. The thermal decomposition pattern of some complexes are given.

| Compound                          | DTA - Peak Temp/ °C | TG - Temp Rang/ °C | Mass Loss % | Decomposition product |
|-----------------------------------|---------------------|--------------------|-------------|-----------------------|
|                                   | Found               | Calcd              |
| Co(PhAc)$_2$.15N$_2$H$_4$.H$_2$O  | 220(+), 440(+)      | 175-245, 245-500   | 14.55, 80.92 | Co(PhAc)$_2$, CoO     |
|                                   |                     |                    | 16.71, 81.03|
| Cd(PhOAc)$_2$.15N$_2$H$_4$.H$_2$O | 140(-), 205(-)      | 105-160, 170-230   | 3.6, 12          | Cd(PhOAc)$_2$.15N$_2$H$_4$, Cd(PhOAc)$_2$, CdO |
|                                   |                     |                    | 3.7, 13.72      |
|                                   |                     |                    | 73.46, 73.21    |
| Ni(PhAc)$_2$.15N$_2$H$_4$.H$_2$O  | 230(+), 465(+)      | 195-250, 370-480   | 16.37, 83.32    | Ni(PhAc)$_2$, NiO     |
|                                   |                     |                    | 16.72, 81.08    |

Exothermic - (+)  Endothermic - (-)
3.3. Thermal analysis

3.4. Co(PhAc)$_2$.1.5N$_2$H$_4$.H$_2$O

This complex undergoes two step decomposition. The TG curve shows 14% mass loss in the temperature range 175-245°C which coincides with the calculated mass loss for the formation of Co(PhAc)$_2$ with loss of hydrazine and water molecule. The higher temperature decomposition indicates that coordinated water molecule. This intermediate further decomposes in the temperature range 245-500°C to give CoO as the final product. DTA shows exotherm corresponding to the above two stages at 220 and 440°C, respectively. The TG-DTA pattern of this complex is given in Fig. 9.

Fig. 9. Co(PhAc)$_2$.1.5N$_2$H$_4$.H$_2$O

3.5. Cd(PhOAc)$_2$.1.5N$_2$H$_4$.H$_2$O

This DTA curves reveal three peaks corresponding to three step decomposition of the complex as shown by TG. The first endothermic peak at 140°C is assigned to the loss of molecule of water. The second step corresponds to the decomposition of the intermediate, Cd(PhOAc)$_2$.1.5N$_2$H$_4$ to yield Cd(PhOAc)$_2$ which is observed as an endotherm at 205°C in DTA. The Cd(PhOAc)$_2$ further decomposes exothermally at 520°C in the final step to form CdO as the end product. The thermogram of the complex is given in Fig. 10.

Fig. 10. Cd(PhOAc)$_2$.1.5N$_2$H$_4$.H$_2$O

3.6. Ni(PhAc) .1.5N .H .H .O

This complex undergoes two step decomposition. The TG curve shows 16% mass loss in the temperature range 195-250°C which coincides with the calculated mass loss for the formation of Ni(PhAc)$_2$ as an intermediate. This intermediate further decomposes in the temperature range 370-480°C to give NiO, as the final product. DTA shows exotherm corresponding to the above two stages at 230 and 465°C, respectively. The TG-DTA pattern of this complex is given in Fig. 11.

Fig. 11. Ni(PhAc)$_2$.1.5N$_2$H$_4$.H$_2$O

3.7. Coordination geometry

The analytical and physico chemical studies suggest that, in these complexes, the hydrazine molecules are present as a bidentate bridging ligand. The phenylacetate and phenoxyacetate ions are seen to present as a monodentate ligand (Figs. 12 and 13) as evident from IR spectra. From TG-DTA analysis we confirmed that water molecule is present as a coordinated one. The complexes are isolated only as polycrystalline powders. Hence, without crystal structure, it is very difficult to predict the environment of the metal in the complexes. Six-coordination has been tentatively proposed for all the complexes with octahedral stereochemistry. The insoluble nature of these complexes conforms to the polymeric structure.

Fig. 12. Polymeric structure M(PhAc)$_2$.1.5N$_2$H$_4$.H$_2$O where M = Co, Ni, Cd and Mn

Fig. 13. Polymeric structure M(PhOAc)$_2$.1.5N$_2$H$_4$.H$_2$O where M = Co, Ni, Cd and Mn

3.7. Antibacterial studies
The complexes of phenylacetic and phenoxyacetic acid have been screened to evaluate their antibacterial activities against (A) \textit{Staphylococcus aureus}, (B) \textit{Escherichia coli}, (C) \textit{Pseudomonas aeruginosa}, (D) \textit{Proteus mirabilis}, respectively at two different concentrations (Fig. 14). The radius of the zone of inhibition was measured in millimeter. \textit{Cefepimezobactum} were used as a standard control and is tabulated as follows.

Table 4. Antibacterial activities of the complexes are given. (the test solutions were prepared in dil HCl).

| S. No | Compound        | \textit{S. aureus} | \textit{E.coli} | \textit{P. aeruginosa} | \textit{P. mirabilis} |
|-------|----------------|-------------------|----------------|-----------------------|----------------------|
|       |                | Diameter of inhibition zone (mm) |
|       |                | 50 | 100 | 50 | 100 | 50 | 100 | 50 | 100 | 50 | 100 | 50 | 100 |
| 1     | Co(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | 17 | 18 | - | - | 18 | - | 17 | - | - | - | - | - | - |
| 2     | Ni(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | - | 12 | 12 | - | - | - | - | - | - | - | - | - |
| 3     | Cd(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 4     | Mn(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | 12 | 15 | - | - | - | - | - | - | - | - | - | - | - |
| 5     | Co(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | 17 | 8 | 18 | - | 20 | - | 14 | - | - | - | - | - | - |
| 6     | Ni(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | - | - | - | - | - | 8 | 10 | - | - | - | - | - | - |
| 7     | Cd(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | 7 | - | 8 | - | 6 | 12 | 12 | - | - | - | - | - | - |
| 8     | Mn(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 9     | \textit{Cefepimezobactum} | 20 | 24 | 20 | - | 20 | - | - | - | - | - | - | - | - | - |

(9) – standard
(-) – no activity

From the result, it has been observed that there is a concentration dependent percentage inhibition in the tested compounds. The activities of the complexes have been compared with the activity of standard antibiotics (\textit{Cefepimezobactum}) and it has been found that all the complexes showed good activities but compound Mn(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O has no activity against all the four microorganisms.

Fig. 14. Screening for the antibacterial activity at 100% and 50% concentration

From the Table 3.4, it can be seen that the compounds Co(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O showed remarkable activity against \textit{Pseudomonas aeruginosa}, and \textit{Escherichia coli}. Compound Co(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O showed remarkable activity against \textit{Pseudomonas aeruginosa}, and \textit{Staphylococcus aureus}.

The results suggest that the antibacterial activity of complexes prepared from both phenylacetic acid phenoxyacetic acid are almost same. This may be due to structural relativity between them.

4. CONCLUSION

Transition metal hydrazine complexes of the type M(PhAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O where M = Co, Ni, Cd (or) Mn, M(PhOAc)$_2$ \cdot 1.5N$_2$H$_4$H$_2$O where M = Co, Ni, Cd (or) Mn were prepared in aqueous medium by using respective metal nitrate hydrate or metal acetate hydrate, hydrazine hydrate and phenylacetic acid or phenoxyacetic acid. Among the hydrazine derivatives, sesquihydrinates are unique complexes, and also it is appropriate to mention here that these types of complexes are separated only with these acids except cadmium oxalate sesquihydrinate$^{126}$.

The prepared complexes were characterized by IR and thermogravimetric analysis. The antibacterial
activity of the complexes against *Staphylococcus aureus*, *Escherichia coli*, *Pseudomonas aeruginosa* and *Proteus mirabilis* were also carried out.

The IR spectral data indicates that the binding of hydrazine to a metal ion is a bidentate fashion. Carboxylate ligands are monodentatively coordinated to the central metal ion. The broad peak around 3625-3282 cm⁻¹ indicates the presence of water molecule. The prepared complexes undergo two or three step decomposition to form metal oxide as the final product. The higher temperature dehydration indicates that the presence of water molecule as coordinated one.

The antibacterial screening shows that Co(PhAc)₂L₅N-H₄H₂O, Cd(PhOAc)₂L₅N-H₄H₂O are active against *Staphylococcus aureus*, *Escherichia coli*, *Pseudomonas aeruginosa* and *Proteus mirabilis*. Co(PhAc)₂L₅N-H₄H₂O active against *Staphylococcus aureus*, *Pseudomonas aeruginosa* and *Proteus mirabilis*. Ni(PhOAc)₂L₅N-H₄H₂O active against *Staphylococcus aureus*, *Pseudomonas aeruginosa* and *Proteus mirabilis*. Mn(PhAc)₂L₅N-H₄H₂O active against *Staphylococcus aureus*, and *Escherichia coli*. Ni(PhAc)₂L₅N-H₄H₂O and Cd(PhAc)₂L₅N-H₄H₂O shows active against only *Escherichia coli* and *Proteus mirabilis* respectively. Mn(PhOAc)₂L₅N-H₄H₂O does not show any antibacterial activity.

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