STRUCTURAL AND THERMAL CHARACTERISTICS OF HYBRID Co(II) PEROVSKITE CRYSTALS

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ABSTRACT
The single crystals of benzyltributylammonium tetrachloro cobaltate (II) monohydrate (BTBA-Co) crystals were grown by slow evaporation solution growth technique at room temperature. The structural characteristics of the grown crystals were studied using Fourier transform infrared (FT-IR) spectroscopy, nuclear magnetic resonance (NMR - 1H and 13C) spectroscopy, powder X-ray diffraction (PXRD) and unit cell parameter analyses. Phase transitions were studied through thermal analyses (TG-DTA and DSC).

Keywords: perovskite, hybrid crystals, XRD, thermal analyses

INTRODUCTION
Semiconducting metal halide perovskites have attracted considerable attention in optoelectronics, including photovoltaics and light emission, because of their excellent photophysical properties1-5. The organic-inorganic perovskite crystals are the materials used for solar cells, piezoelectric, dielectric, semiconducting and related potential applications6,7. Perovskite is the class of crystalline compounds described with the crystal structure of A2BX4, ABX3 etc., where A and B are the cations of organic and inorganic compounds, respectively and X is a halide. Recently, there are several methods were adopted to synthesize perovskite Single crystals like slow evaporation solution growth, slow cooling method, hydrothermal growth, gel growth and vapor growth. In this, slow evaporation solution growth technique is one of the simple and versatile routes to grow single crystals. Ono et al8 reported organometal halide perovskites as solar cells with the reasonable percentage of efficiency. Several researchers are reported this kind of crystals for alternate energies3,9,10.

The perovskite single crystals, for instance, provide an excellent platform to characterize a range of materials properties that could not be achieved by studying polycrystalline thin films. Hybrid perovskite single crystals exhibit superior luminescent performance than their thin film counterparts, enabling potential applications like large area display. Understanding the mechanism of the crystal growth provides insights into the growth process and enhancement of quality. The main objective of the present investigation is to the synthesis of benzyltributylammonium tetrachloro cobaltate (II) monohydrate single crystals. Also, the structural and thermal behavior of the grown crystals were studied using physio-chemical characterizations.

EXPERIMENTAL
The Growth of Single Crystals (BTBA-Co)
The single crystals of benzyltributylammonium tetrachloro cobaltate (II) monohydrate (BTBA-Co) crystals are grown by slow evaporation of saturated aqueous solutions at room temperature. Aqueous solutions containing benzyl tributyl ammonium chloride and cobalt (II) chloride in 2:1 molar ratio,
respectively, are prepared by using triply distilled water. The two solutions are mixed together well and the resulting solution is filtered through a Whatman No.4 filter paper into a clean beaker. The beaker is covered with aluminum foil. Care is taken to minimize the temperature gradient and mechanical shock. Bright and blue colored crystals of BTBA-Co are obtained within 35-45 days. The grown crystals are collected from the mother liquor by using well-cleaned forceps. The harvested crystals are recrystallized repeatedly to get crystal of good quality.

\[
\begin{align*}
2 \left[ (C_6H_5)N^+CH_2-C_6H_5Cl^- \right] + & \text{CoCl}_2 \xrightarrow{\text{Medium}} \left[ (C_6H_5)N^+CH_2-C_6H_5 \right]_2\text{CoCl}_4\cdot\text{H}_2\text{O} \\
\text{BTBA-Co}
\end{align*}
\]

Scheme-1: Formation of BTBA-Co single crystals

**Characterization Techniques**

The elemental analysis and FT-IR spectrum are carried out in ELEMENTAR VARIO EL III instrument and THERMO NICOLET AVTAR370 DTGS instrument, respectively. The NMR (\(^1\)H and \(^{13}\)C) spectrum of BTBA-Co crystals are carried out in AMX 400 spectrometer. The simultaneous TG-DTA study is done on a NETZSCH STA 409C/CD thermal analyzer and the curves are obtained under a nitrogen atmosphere at a heating rate of 10ºC per minute from room temperature to 800ºC. The cell parameters of grown crystals are recorded using BRUKER SMART APEX CCD single crystal diffractometer using MoK\(\alpha\) radiation. A METTLER TOLEDO DSC822e instrument is used to record the low-temperature DSC curves for the samples at STIC, Cochin. The cooling run is carried out from 30 ºC to -100ºC and the heating run from -100 ºC to 30ºC both at a rate of 10ºC per minute under a nitrogen atmosphere.

### RESULTS AND DISCUSSION

**Elemental Analysis and FT IR Spectrum**

The elemental analysis of dark blue coloured grown crystals shows that the compound contains carbon: 58.3% (59.15%), hydrogen: 10.4% (9.08%) and nitrogen: 3.7% (3.63%). The results indicate that both experimental and theoretical values (given in the brackets) are found to be in good agreement with each other and conform to the molecular formula of the compound \([C_6H_5CH_2-N-(C_4H_9)_3]_2[\text{CoCl}_4].\text{H}_2\text{O}, \text{BTBA-Co}\). The proposed crystal structure of BTBA-Co crystals are shown in Scheme-2.

![FTIR Spectrum and XRD pattern of BTBA-Co Crystal](image)

Further, the crystal formation can be confirmed through FTIR spectrum, shown in the Fig.-1. The frequencies observed at 3491 cm\(^{-1}\) is due to the asymmetric O-H stretching vibration mode. The absorption frequency at 3063 cm\(^{-1}\) is due to symmetric O-H stretching vibration mode as well as C-H stretching of an aromatic nucleus. The frequencies at 2965 and 2875 cm\(^{-1}\) are due to the asymmetric and symmetric C-H stretching vibrational mode. The absorption frequencies at 1644, 1584, 1479, 1455, 1381 and 1213 cm\(^{-1}\) are due to C=C stretching (skeletal) of phenyl nucleus, CH\(_2\) asymmetric bending, CH\(_3\) asymmetric bending, CH\(_2\) symmetric bending, CH\(_3\) symmetric bending and C-H in-plane bending of phenyl ring respectively. The CH\(_3\) rocking, asymmetric C-C stretching, asymmetric C-N stretching, C-H
out-plane bending in phenyl ring, CH$_2$ rocking mode, C-C out-plane bending in aromatic nucleus and C-H out-plane bending in mono substituted benzene ring observed at 1145, 1066, 1032, 868, 803, 726 and 702 cm$^{-1}$ respectively. The C-N-C skeleton vibrational modes are observed at 627, 546 and 459 cm$^{-1}$.

**Powder X-Ray Diffraction and Cell Parameters**

The powder X-ray diffraction pattern of the BTBA-Co crystal is shown in the Fig.-1b. The sharp and well defined Bragg peaks in the powder XRD pattern confirm the crystallinity of the compound. The unit cell parameters of the crystal are determined by a least-squares technique using reflections. The compound crystallizes in triclinic system. The unit cell parameters for the crystal are $a = 11.24 \text{ Å}; b = 11.35 \text{ Å}$ and $c = 17.26 \text{ Å}$. $\alpha = 76.41^\circ$, $\beta = 82.42^\circ$ and $\gamma = 84.29^\circ$ ($\alpha \neq \beta \neq \gamma$). The volume of the unit cell is $2116 \text{ Å}^3$.

**NMR Spectroscopy**

The $^1$H and $^{13}$C NMR spectrum of BTBA-Co crystal are shown in the Fig.-2(a) and 2(b).

In $^1$H NMR (Fig.-2a), a singlet peak is observed at $\delta 7.5$ ppm which is due to the presence of aromatic protons. The two singlet peaks are observed at $\delta 4.9$ and $\delta 3.2$ ppm and are due to the water and methylene group protons which are attached with the aromatic ring respectively. A multiplet peak is observed at $\delta 1.4$ ppm which may due to the n-butyl group protons in the compound.

In $^{13}$C NMR (Fig.-2b), a peak observed at $\delta 132.528$ ppm is due to the C9 carbon in the benzyl group. The peaks observed at $\delta 130.882$, $\delta 129.487$ and $\delta 127.382$ ppm are due to the C8, C7 and C6 carbons respectively. C5 methylene carbon peak is observed at $\delta 61.765$ ppm. The butyl group carbons C4, C3, C2 and C1 are observed at $\delta 58.005$, $\delta 23.553$, $\delta 19.286$ and $\delta 13.160$ ppm respectively.

**Thermal Characteristics**

Thermogravimetry (TG) and differential thermal analysis (DTA) curves of BTBA-Co are shown in the Fig.-3a. When the compound is heated from 30 to 800ºC, it remains stable up to 170ºC. Decomposition is observed between 170 ºC and 550 ºC with weight loss of 61%. The remaining 39% residue is unaffected even after heating up to 800 ºC. The above weight losses can be explained by formulating the following decomposition pattern.

\[
[\text{C}_6\text{H}_5\text{CH}_2\text{-N-(C}_4\text{H}_9)_3\text{]}_2[\text{CoCl}_4]\text{H}_2\text{O} \xrightarrow{\text{170 ºC} - \text{550ºC}} (\text{C}_7\text{H}_7\text{)}_2\text{CoCl}_2 + 2\text{N(C}_4\text{H}_9)_3\text{↑} + \text{Cl}_2\text{↑} + \text{H}_2\text{O}↑
\]

<table>
<thead>
<tr>
<th>Theoretical Loss</th>
<th>59.4%</th>
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<tbody>
<tr>
<td>Experimental Loss</td>
<td>61%</td>
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The above pattern suggests the formation of the tropylium cobalt chloride as residue. Two moles as tributylammonia, 1 mole of chlorine gas and 1 mole of water are eliminated as a gaseous product at this temperature range. The proposed decomposition pattern amounts to a calculated weight loss of 59.4 %, whereas the experimental weight loss is 61%. The weight loss difference of 1.6% is well within the experimental error limits.
In DTA thermogram, the broad endothermic peak is observed at 130°C shows the removal of surface moisture in the compound at this temperature. The broad band at 320°C is due to the first stage decomposition. The DTA results correlate well with the corresponding TG results. The low-temperature DSC curve is shown in the fig. 3b. No thermal anomalies are observed in both cooling and heating runs. It clearly shows that there is no structural phase transition in the compound at temperatures below room temperature.

CONCLUSION

BTBA-Co single crystal with good optical quality is grown using slow evaporation technique at room temperature. XRD studies reveal that the grown crystal belongs to a triclinic system. From the elemental composition results, the presence of carbon, hydrogen, and nitrogen are exactly coincidence with the theoretical molecular formula of the compound. The various functional groups present in the BTBA-Co crystal is evaluated using FT-IR studies. Further, the molecular structure of the complex crystal was analyzed using $^1$H and $^{13}$C NMR spectroscopy studies and TG-DTA analyses show BTBA-Co is thermally stable up to 130°C.

REFERENCES

7. P. Umari, E. Mosconi, and F. De Angelis, Scientific Reports, 4, 4467 (2014), DOI:10.1038/srep04467

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