ISSN : 0974 - 7486

Volume 11 Issue 11



Materials Science An Indian Journal FUIL Paper

MSAIJ, 11(11), 2014 [349-352]

Cobalt levo-tartrate crystals;

Gel growth;

Thermo gravimetric

analysis;

Kinetic parameters;

Thermodynamic parameters.

Thermal study of gel-grown cobalt levo-tartrate crystals

H.O.Jethva

Maharaja Shree Mahendrasinhji Science College, Morbi – 363642, Gujarat, (INDIA) E-mail:hojethva@rediffmail.com

ABSTRACT

Metal tartrate compounds find various applications. Cobalt tartrate finds applications in semiconductors, medicine optics, gold industries and veterinary drugs. In the present study, cobalt levo-tartrate crystals were grown using silica hydro-gel as growth medium. Spherulitic reddish crystals were obtained. The crystals were characterized by Thermo gravimetric analysis and kinetic as well as thermodynamic parameters were evaluated for the different stage of decomposition.

© 2014 Trade Science Inc. - INDIA

INTRODUCTION

position stage.

The metal tartrate compounds find various applications in the different fields of science and technology, for example, ferroelectric applications of calcium tartrate^[1], piezoelectric application of cadmium tartrate^[2], addition of lead tartrate in gasoline to prevent knocking in motors [3] and application of carbonate solutions containing Co(II) tartrate complexes in an electrochemical procedure of anodic deposition of cobalt oxhydroxide film on a glassy carbon substrate in an alkali medium^[4]. The gel growth technique is found to be suitable to grow tartrate compound crystals, which is elaborately described by Henisch^[5]. The growth of several tartrate compound crystals by the gel technique and their characterization were reported, for instance, cadmium tartrate ^[6-11], iron tartrate ^[12] and lead tartrate ^[13,14]. In the present study, the present author has attempted to grow the crystals of cobalt levo-tartrate and characterized them by TGA and evaluated the kinetic as well as thermodynamic parameters for the dehydration and decom-

EXPERIMENTAL

KEYWORDS

In the present study, the single diffusion method^{[14-} ^{17]} was employed for the growth of cobalt levo-tartrate crystals. The silica hydro gel was used as a growth medium. To prepare the gel, a solution of sodium metasilicate of 1.05 specific gravity and 1 M solution of levo tartaric acid were mixed in such a manner that the pH of the mixture was set at 4.5. The gel solution was poured in to glass test tubes of 15 cm length and 2.5 cm diameter and allowed to set in the gel form. The supernatant solution containing cobalt nitrate solutions was poured on the set gel carefully without damaging the gel.

All the chemicals were of AR grade and obtained from Sigma Aldrich. The following reaction is expected to take place in the formation of cobalt levo-tartrate crystals.





Figure 2: 1 G curve for cobait levo-tartrate crystals

$Co(NO_3)_2 \cdot 6H_2O + H_2C_4H_4O_6 + nH_2O \rightarrow$ $CoC_4H_4O_6 \cdot nH_2O + 2HNO_3 + 6H_2O$ (1)

The amount of HNO₃ produced is very less in comparison to the nutrients being supplied to the growing crystals and hence no major limitation is imposed [5,14-17]

Photograph of the grown crystals is shown in the figure 1.

The grown crystals were characterized by TGA. The TGA was recorder on Perkin Elmer make instrument, model Pyris-1 DSC, Pyris-1 TGA, DTA-7. The data were recorded from room temperature to 700°C at a heating rate of 10°C min⁻¹ in atmosphere of air.

RESULT AND DISCUSSION

There are reports available in literature on thermal studies on pure and mixed metal tartrate systems, for example, cadmium tartrate ^[18], mercuric iodate crys-tals ^[19], iron-manganese levo-tartrate tartrate compound ^[20], ternary iron-manganese-cobalt tartrate compound ^[15], ternary iron-manganese-nickel tartrate compound ^[16] and lead-cadmium mixed levo-tartrate crystals ^[17].

The TGA curve for cobalt tartrate crystals is shown in figure 2.

There are three stages of decomposition starting from room temperature of 45 °C. The first stage of decomposition starts from 45 °C and continues up to 250 °C where the weight loss is about 18%. The second stage of decomposition starts from 250 °C and continues up to 390 °C where the weight loss is about 65%. The third stage of decomposition starts from 390 °C and continues up to 500 °C where the mass loss is about 70%. Comparing the observed and calculated percentage weight losses suggests chemical formula for the given crystal to be $CoC_4H_4O_62.6H_2O$. The TGA result for cobalt levo-tartrate is shown TABLE 1.

Many researchers have used thermo gravimetric data to calculate the kinetic parameters of solid state reaction including mass ^[21-25]. By using kinetic param-

Stage	Temperature	Mass loss (%)		Reaction	
	range (°C)	Observed	Calculated		
Ι	45 - 250	17.6	18.5	$CoC_4H_4O_62.6H_2O \rightarrow CoC_4H_4O_6$	
II	250 - 390	64.21	64.57	$CoC_4H_4O_6 \rightarrow CoCH_3O$	
III	390 - 500	70	70.48	$CoCH_3O \rightarrow CoO$	

TABLE 1 : TGA result for cobalt levo-tartrate crystals

Materials Science An Indian Journal eters such as order of reaction, frequency factor and energy of activation, the shape of curve is determined. With the help of Coats and Redfern^[23] relation the kinetic parameters were calculated. In the present study, the kinetic and thermodynamic parameters have been calculated for dehydration as well as for decomposition of crystals.

Coats and Tedfern relation is given by

$$\log_{10}\left[\frac{1-(1-\alpha)^{1-n}}{T^2(1-n)}\right] = \left\{\log_{10}\left[\frac{AR}{\alpha E}\right]\left[1-\frac{2RT}{E}\right] - \frac{E}{2.3RT}\right\}$$
(2)

where $\alpha = \frac{W_o - W}{W_o - W_f}$, W_o the initial weight, W the

weight at time t, W_t the final weight, n the order of reaction, A the frequency factor, E the activation energy of the reaction, R a gas constant and α the heating rate in °C/min.

To determine the the value of activation energy and

order of reaction, a plot of $\log_{10}\left[\frac{1-(1-\alpha)^{1-n}}{T^2(1-n)}\right]$ versus $\frac{1}{T}$

is drawn for different values of n and the best linear plot gives the correct value of n. This equation is valid for all values of *n* except n = 1.

Figure 3(a) and 3(b) is a plot of Coats and Redfern equation for the cobalt levo-tartrate crystals for the dehydration as well as decomposition stage, respectively.

The values of order of reaction, activation energy and frequency factor for the dehydration as well as decomposition stage are listed in the TABLE 2.



Figure 3: A plot of Coats and Redfern relation for cobalt levotartrate crystals

The thermodynamic parameters have been evaluated for the dehydration as well as decomposition stages by using the standard relations [26]. The standard enthalpy of activation "#H° was calculated by using the relation 3.

$$\Delta^{\#} H^{o} = E - 2RT$$
(3)
The standard entropy of estivation $A^{\#} S^{o}$ use call

The standard entropy of activation $\Delta^{\#}S^{\circ}$ was calculated by using relation 4.

$$\Delta^{\#}S^{o} = 2.303 \times R \times \log_{10}\left[\frac{Ah}{kT}\right]$$
(4)

here, k is the Boltzman constant, h the Planck constant, T the temperature and A is the frequency factor.

The standard Gibbs free energy of activation $\Delta^{\#}G^{\circ}$ is estimated by using the relation 5.

An Indian Journal

TABLE 2 : Kinetic parameters for cobalt levo-tartrate crystals								
tage	n	Activation energy (kJ mol ⁻¹)		Frequency factor				
Dehydration	2	122.85		1.26×10^{29}				
Decomposition	0	33.35		1.52×10^{16}				
	TABLE 3 : Therm	odynamic par	ameters for cobalt levo-1	artrate	crystals			
Stage	Standard enthalpy ∆ [#] H ^o (kJ mol ⁻¹)		Standard entropy ∆ [#] S ^o (kJ mol ⁻¹)		Standard entropy $\Delta^{\#}G^{0}$ (kJ mol ⁻¹)			
Dehydration	120.69		0.310		120.38			
Decomposition	25.98	;	0.062		25.93			
					D Materials Scien			

ABLE 2 : Kineti	c parameters fo	or cobalt levo	-tartrate crystals
-----------------	-----------------	----------------	--------------------

Full Paper -

$\Delta^{\#}\mathbf{G}^{\mathsf{o}} = \Delta^{\#}\mathbf{H}^{\mathsf{o}} - \mathbf{T} \,\Delta^{\#}\mathbf{S}^{\mathsf{o}}$

(5)

The values of standard enthalpy, standard entropy and standard Gibbs free energy for the dehydration as well as decomposition stage are listed in the TABLE 3.

Positive values of the standard enthalpy and the standard entropy of activation suggest that the process is spontaneous at high temperatures and the positive values of standard Gibbs free energy suggest that the process is thermodynamically unstable ^[15].

CONCLUSION

Pure cobalt levo-tartrate crystals were grown in silica gel by using 1 M cobalt nitrate solution as supernatant solution. From TG curves, it was found that the crystals were thermally unstable. Upon heating, they become anhydrous and decomposed into metal oxide through a single stage. The kinetic parameters were evaluated for both the stages by using Coats and Redfern relation. The thermodynamic parameters were evaluated for both the stages by using the standard relations. The presence of water molecules was detected and calculated. The exact chemical formulation for the pure cobalt levo-tartrate crystals were suggested.

ACKNOWLEDGEMENT

The author is thankful to the Principal and the Management of Maharaja Shree Mahendrasinhji Science College, Morbi for their encouragement.

REFERENCES

- [1] H. B.Gon; J. Cryst. Growth, 102, 501 (1990).
- [2] J.S.Hopwood, A.W.Nicol; Crystallogr, J. Online, 1600, 5767 (1972).
- [3] N.J.Rahway; The Merck index of chemicals and drugs, 6th ed. Merck and Co., (1952).
- [4] I.G.Casella; J. Electroanal. Chem., 520(1-2), 119 (2002).
- [5] H. K.Henisch; Crystal Growth in Gels, Dover Publication: New York (1993).
- [6] M.E.Torres, T.Lopez, J.Peraza, J.Stockel, A.C.Yanes; J.Appl. Phy., 84(10), 5728 (1998).

- [7] R.M.Dabh, M.J.Joshi; Indian J. Phys., 77, 481 (2003).
- [8] K.Sangwal, A.Kothari, S.K.Arora; Surf. Sci., 600, 1475 (2006).
- [9] A.Kothari, S.K.Arora; ICMAT, Singapore (2009).
- [10] N.S.Patil, P.A.Savale, S.K.Bachhav, S.T.Pawar; Archives of Phys Res., 2, 39 (2011).
- [11] P.Vera-Cruz, R.A.Toscano, J.Balmaseda, M.Basterrechea, N.Nino, L.F.del Castillo; Cryst. Eng. Comm., 14, 8608 (2012).
- [12] S.Joseph, H.S.Joshi, M.J.Joshi; Cryst. Res. Technol., 32(2), 339 (1997).
- [13] M.Abdulkadhar, M.A.Ittyachen; J. Cryst. Growth, 39, 365 (1977).
- [14] H.O.Jethva, M.V.Parsania; Asian J. Chem., 22(8), 6317 (2010).
- [15] S.J.Joshi, K.P.Tank, B.B.Parekh, M.J.Joshi; Cryst. Res. Technol., 45, 303 (2010).
- [16] S.J.Joshi, K.P.Tank, B.B.Parekh, M.J.Joshi; J. Therm. Anal. Calorim., (2012).
- [17] H.O.Jethva, P.M.Vyas, K.P.Tank, M.J.Joshi; J. Therm. Anal. Calorim., (2014).
- [18] S.K.Arora, A.Kothari, B.Amin, B.Chudasama; Cryst. Res. Technol., 42(6), 589 (2007).
- [19] A.B.Patil, K.B.Saraf; ISRJ, 1(3), 213 (2011).
- [20] S.J.Joshi, B.B.Parekh, K.D.Parikh, K.D.Vora, M.J.Joshi MJ; Bull. Mater. Sci., 29(3), 307 (2006).
- [21] S.Joseph, M.J.Joshi; Indian J. Phys., 71A, 183 (1997).
- [22] R.M.Dabhi, M.J.Joshi; Indian J. Phys., 76A, 481 (2003).
- [23] A.W.Coats, J.P.Redfern; Nature, 201, 183 (1964).
- [24] H.H.Horowitz, G.Metzger; Anal. Chem., 35, 1464 (1963).
- [25] P.N.Kotru, K.K.Raina, M.L.Koul; J. Mater. Sci., 21, 3933 (1986).
- [26] K.J.Laidler; Chemical kinetics, Harper and Row, New York (1987).

Materials Science Au ^qudian gournal