

## Synthesis and Characterization of Cadmium Substituted Copper Nano – Ferrites

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### ABSTRACT

A series of copper substituted cadmium nano ferrites with the compositional formula  $Cu_{1-x}Cd_xO_4$  (where  $x=0.1, 0.3, 0.5$  and  $0.7$ ) were prepared by the Citrate gel auto combustion technique and sintered at 873 K for 4 hr. The X-ray diffractograms (XRD) clearly exhibited the existence of single phase cubic spinel structure. The crystallite size was found in the range of 10-42nm. The values of lattice parameter were calculated by using d spacing. The calculated values of lattice parameter and X-ray density were explained on the basis of composition.

**Keywords:** Cu-Cd Nano Ferrites; X-Ray Density; lattice parameter

### I. INTRODUCTION

Spinel ferrites have been studied extensively due to easy to synthesis and abundant uses in technological and industrial applications [1-3]. The useful properties of the spinel ferrites mostly depend upon the chemical composition, preparation methods, sintering temperature, nature of the additives and their distribution i.e. tendency to occupy tetrahedral (A) or octahedral (B) site [4]. Cd has a strong preference octahedral site (B-site) [5].  $Cu^{2+}$  is the divalent ions which occupy essentially tetrahedral A-site depending on the sample preparation [6] when substituted in ferrites.

Among the spinel structures, Cu-Cd nano-ferrites ferrite has been widely used in different kinds of magnetic devices, such as inductors, magnetic heads, and magnetic devices, such as inductors, magnetic heads, magnetic refrigeration and magnetic resonance imaging. Thus, the magnetic and electric properties of nickel ferrite have been researched and improved [7]. Cu-containing ferrites form an interesting group of ferrites because of their typical electrical and magnetic properties and change in crystal structure on thermal treatment [8]. Cu-Cd nano-ferrites are low cost materials and have important magnetic and electrical properties for technological applications.

### II. EXPERIMENTAL METHOD

Cu-Cd Nano Ferrite samples with the chemical formula  $Cu_{1-x}Cd_xO_4$  (where  $x=0.1, 0.3,$

$0.5$  and  $0.7$ ) were synthesized using citrate gel auto combustion technique. The molar quantity of AR grade of Cadmium Nitrate ( $Cd(NO_3)_2 \cdot 6H_2O$ ), Copper Nitrate ( $Cu(NO_3)_2 \cdot 3H_2O$ ), Ferric Nitrate ( $Fe(NO_3)_3 \cdot 9H_2O$ ) and Citric Acid-Citrate ( $C_6H_8O_7 \cdot H_2O$ ) Ammonia ( $NH_3$ ) raw materials were taken as starting material. Metal nitrates and citric acid were dissolved in deionized water. Metal nitrate solutions were mixed with citric acid solution in 1:3 molar ratio of nitrate to citric acid. The  $p^H$  of the solution was adjusted to 7 using ammonia. The solution was first heated at 80°C to transform into gel and then ignited in a self-propagating combustion manner to form a fluffy loose powder. The as-burnt ferrite powders were grained by agate motor then calcined at 600°C for 4hr. the calcined ferrite powders were again grained by agate motor. As this method is a chemical route it requires no ball milling hence little scope of contamination, and better homogeneity.

The structural characterization was carried out using X-Ray Diffractometer Bruker (Karlsruhe, Germany) D8 advanced system with a diffracted beam monochromatic  $Cu K_\alpha$  radiation ( $\lambda = 1.5405 \text{ \AA}$ ) radiation source between the Bragg Angles 20° to 80° in steps of 0.04°/Sec.

The X-ray density  $\rho_x$  of the prepared samples was calculated by the relation

$$\rho_x = \frac{8M}{Na^3}$$

### III. RESULTS AND DISCUSSION

The X-ray diffraction patterns for  $\text{Cu}_{1-x}\text{Cd}_x\text{O}_4$  (where  $x=0.1, 0.3, 0.5$  and  $0.7$ ) was sintered at  $600^\circ\text{C}$  are shown in Fig.1. The X-ray patterns show all sample are existence of the single phase of cubic spinel structure [13]. It can be seen from the Table 1 the values of the particle size varies from 10 nm to 42 nm. Though all the samples were prepared under identical condition, the crystallite size was not the same for all Cd concentrations. This was probably due to the preparation condition followed here which gave rise to different rate of ferrite formation for different concentrations of Cu, favoring the variation of crystallite size. Fig.2. shows the lattice parameter versus composition. It can be seen from the figure that the values lattice parameter increases with the increase of composition. Plot of X-ray density with composition is shown in Fig 3. It can be seen from the figure the value of X-ray density is maximum for 0.7 composition.

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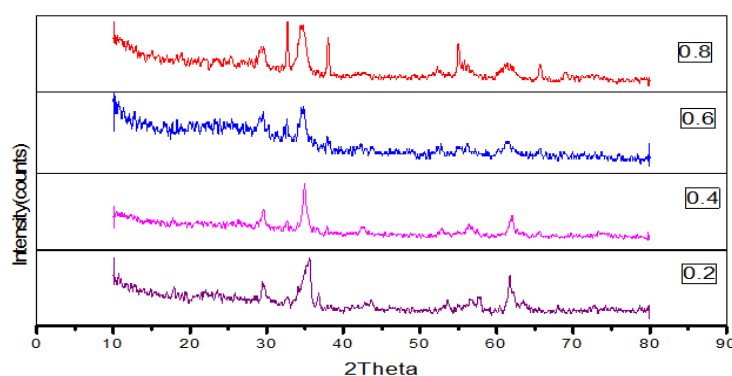


Fig. 1. XRD patterns for Cu-Cd nano-ferrites

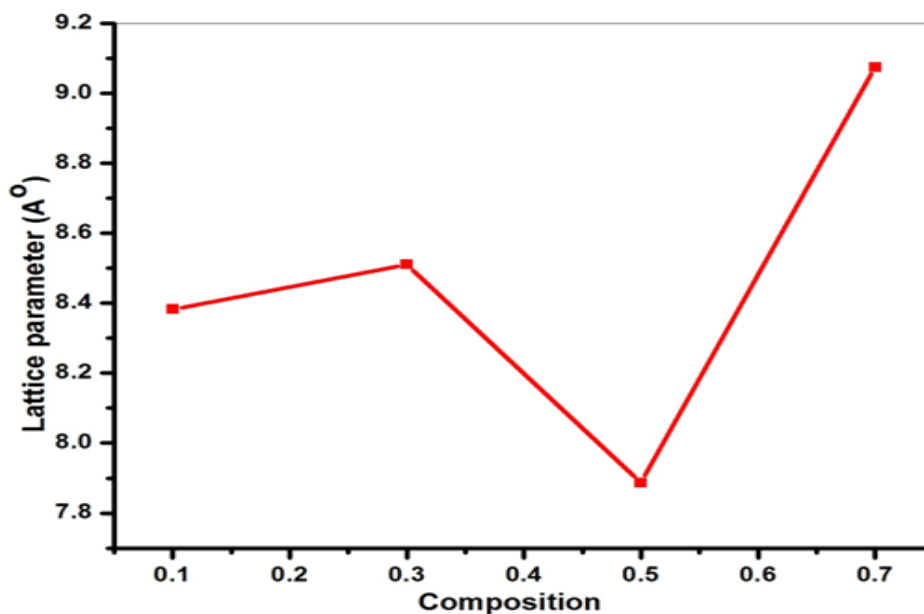


Fig.2. Plot of lattice parameter with composition.

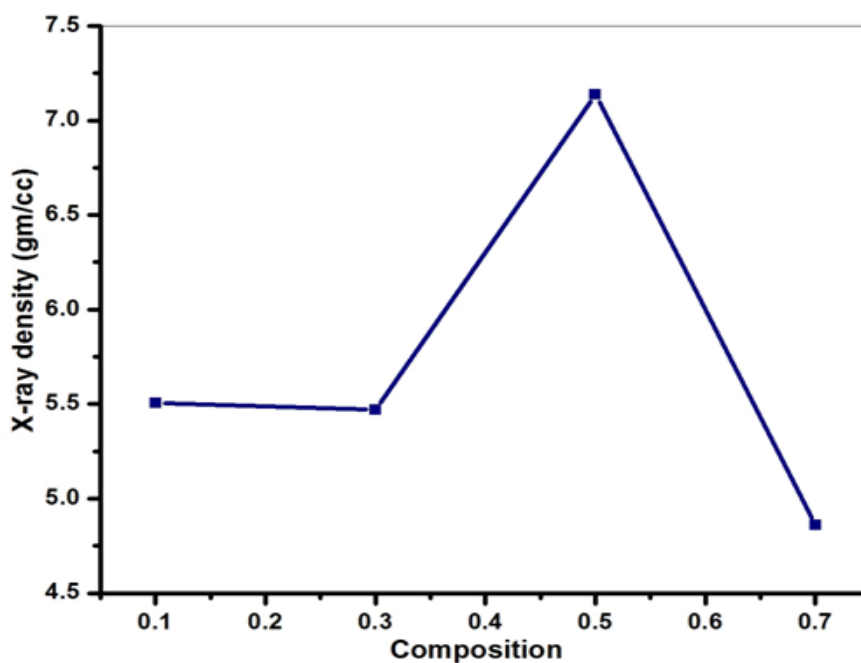


Fig.3. Plot of X-ray density with composition.

| Cu-Cd | 2theta  | fwhm   | d-space (Å) | D-crystalline size (nm) | A-lattice parameter (Å) | x-density gm/cm <sup>3</sup> | M-weight gm |
|-------|---------|--------|-------------|-------------------------|-------------------------|------------------------------|-------------|
| 0.1   | 35.5183 | 0.9446 | 2.52752     | 22.32                   | 8.3828                  | 5.5049                       | 244.103     |
| 0.3   | 34.9646 | 0.3936 | 2.56627     | 10.21                   | 8.5113                  | 5.4699                       | 253.877     |
| 0.5   | 37.8300 | 0.0667 | 2.37822     | 12.06                   | 7.8876                  | 7.1374                       | 263.65      |
| 0.7   | 32.7288 | 0.2362 | 2.73629     | 42.79                   | 9.0752                  | 4.8598                       | 273.425     |

Table.1. Lattice parameter, X-ray densities and crystal Size for Cu-Cd nano-ferrites.