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6.039(IIJIF) Peer-Reviewed International Journal Issue-60, Vol-03 or fraud callers and hackers for increase the trust on cashless transaction in India.

- 3. Government should make to increase speed of internet connectivity and reduce the network connecting problems while making cashless transaction.
- 4. After failing transaction immediately provides settlement by banks and service providers

Conclusion

After demonetization in India increased number of cashless transaction. Transparent economic development is need today therefore cashless transaction is secure and transparent. Government of India has to promote cashless transaction through various discount or incentive cash back offers are one of the important factor that Induce the common propels for the use of cashless transaction and discourages physical cash payments by the way of proper supervision of restrictions for using cash based transaction.

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Structural and Infra-red Analysis Of Ni_{0.7+x}Zn_{0.3} Zr_xFe_{2.2x}O₄ System.

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Abstract:- The structural properties and IR absorption spectra of Zr4* substituted Ni-Zn ferrites having general formula Nigges Znas $Zr_x Fe_{2-2x}O_4(x = 0.0, 0.1, 0.2, 0.3, 0.4, 0.5)$ were prepared by standard double sintering ceramic technique. The X-ray analysis confirmed the single phase cubic spinal structure formation of the samples. The lattice constant was found increases with increase of zr4+ ion substitution. which is attributed to ionic size difference of cations involved. IR spectra of all samples have been used to locate band positions, which are observed to lie near 400cm⁻¹ and 600cm⁻¹. The bond length $R_{_{\rm A}}$ and $R_{_{\rm B}}$ were found to increase with composition.

Ferrites, Structural Keywords:-Properties, Cation distribution, Infra-red, force constant.

I. INTRODUCTION

Ferrites are widely used in many electronic devices. These are preferred because of their high electrical resistivity, chemical stability, mechanical hardness and reasonable constant,[1-4]. Ni-Zn ferrites are well known technological magnetic materials finding their applications in various electronic devices. The physical properties of Ni-Zn ferrites are very Printing Area: Interdisciplinary Multilingual Refereed Journal

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sensitive to the method of preparation, the amount and the type of substitution [5]. The ferrites are commonly prepared by ceramic technique which involves high temperature sintering and prolong heating. This method

produces particles of coarse nature. Spinel ferrites are compounds of iron oxides and some transition metal oxides and they exhibits

important electrical and magnetic properties, which made them extensively useful in technological and industrial applications such

as magnetic storage in microwave devices [6,7].

Nickel ferrite is a well known hard magnetic material with inverse spinel structure. The saturation magnetization and coercivity of nickel ferrite is higher than the other nickel, manganese spinel ferrites [8]. However, to our knowledge zirconium has not been substituted in nickel ferrite. Zirconium is a lustrous, grayishwhite, soft, ductile and malleable metal which is solid at room temperature, though it becomes hard and brittle at lower purities. Zirconium is highly resistant to corrosion by alkalis, acids, salt water and other agents. Ni-Zn ferrites have high resistivity and low eddy currents. Therefore they could be considered as the most versatile ferrites. Ni-Zn ferrites have been commercially used in radio frequency circuits, high quality filters, rod antennas, transformer cores, read/ write heads for high speed digital tape and operating devices [9]. Introducing small amount of foreign ion can change the electrical and magnetic properties of the ferrites [10]. Addition of tetravalent ions like Ti4+, Ge4+, Si4+, Mn4+ influences the structural, magnetic and transition properties of the system.

In the present study, the effect of zirconium substitution in nickel ferrite formula $Ni_{0.7+x}Zn_{0.3}Zr_{x}Fe_{2.2x}O_{4}(x = 0.0, 0.1, 0.2, 0.3, 0.4,$ 0.5,0.6), on structural properties of nickel ferrite synthesized by standard double sintering ceramic method are reported. Zirconiumcontaining compounds are used in many biomedical applications [11].

Experimental Details:- Polycrostolines Ni-Zn ferrites having the chemical formula $Ni_{0.7+x}Zn_{0.3}Zr_xFe_{2.2x}O_4(x = 0.0, 0.1, 0.2, 0.3, 0.4)$ 0.5,0.6) were prepared by standard double sintering ceramic method using the appropriate quantities of A.R. grade ferric oxide (Fe,0.). Nickel oxide (Nio), Zinc Oxide (ZnO) and Zirconium dioxide (ZrO₂) . The samples were pre-sintered for 24 hours in furnace at 500°C Final sintering of the ferrite specimen was carried out for 24 hours at 1000°C. The X-rav diffraction patterns of all the samples were obtained on Philips diffractometer operated at 25KV and 20mA. IR spectra of all the samples were recorded on Perkin-Elmer IR spectrometer (Model No. 783) in the range 200-800cm⁻¹ in kBr medium.

II. RESULT AND DISCUSSION 1.X-ray diffraction studies:

X-ray diffraction patterns of all the samples are recorded at room temperature represented in Fig 1.1. These figures show well defined peaks and (311) reflection appears to be more intense in all the samples. All the planes are allowed planes, which confirm the formation of single phase cubic spinel structure without appearance of any extra peaks. The Inter-planer spacing (d) values were calculated for the recorded peaks using Bragg's law and the lattice constant 'a' was calculated for each plane using the following relation 1.1.

$$a = d_{hkl}\sqrt{(h^2 + k^2 + l^2)}$$
 1.1

The average lattice constant 'a' may be obtained by calculating the average of the 'a' values for each diffraction plane. The variation of lattice constant with Zr4+ ion content is shown in figure 1.2. It is found that the lattice constant increases with increasing Zr4+ ion substitution. In the present system, the increase in a lattice constant of the composition with increase in Zr4+ content x is attributed to replacement of smaller ionic radius of Fe3+ (0.67Å) by lager ionic radius of Zr4+ (0.80Å) obeying Vegard's Law [12,13]. The

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ISSN: 2394 5303 Impact Factor Peer-Reviewed International Journal Issue-60, Vol-03 results of variation of lattice constant with Zr4 substitution are in good agreement with other Zr4+ substituted spinel ferrite [14, 15].

2. Infra-Red Spectroscopy (IR):-

The IR spectra of the present ferrite system recorded at room temperature for the samples are shown in Fig. 1.3. The spectra show two distinct bands as observed in other ferrites [16, 17]. The IR spectra of all the samples have been used to locate the band positions. The band positions are observed to lie near 400 cm⁻¹ and 600 cm⁻¹. The values of band positions n, and n, are given in Table 1.2. The bands are found to be in the reported range. The higher frequency band n, is assigned to the intrinsic vibrations of the tetrahedral groups and the low frequency band n, is assigned to the octahedral groups [18, 19].

It is observed from Table 1.2 that there is a difference in the band positions. The difference in the band positions is expected because of the difference in the Fe3+-O2- Fe3+ distances for the tetrahedral and octahedral complexes. The IR spectra have been used to determine force constant K, and Ko, which depends upon vibrational frequency, cation mass etc. In order to find out force constant K and K_n, the analysis of IR given by Waldron [18] has been used. The force constants corresponding to the tetrahedral and octahedral complexes are calculated by using the standard formulae given below,

$$K_1 = 7.62 \times M_1 \times v_1^2 \times 10^{-7}$$
 1.2

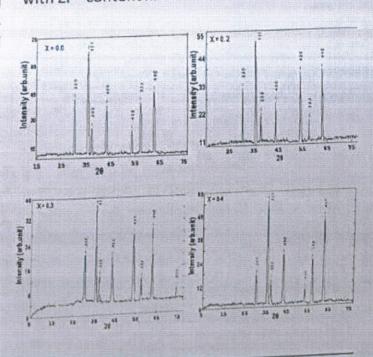
$$K_o = 10.62 \times \frac{M_2}{2} \times v_2^2 \times 10^{-7}$$
 1.3

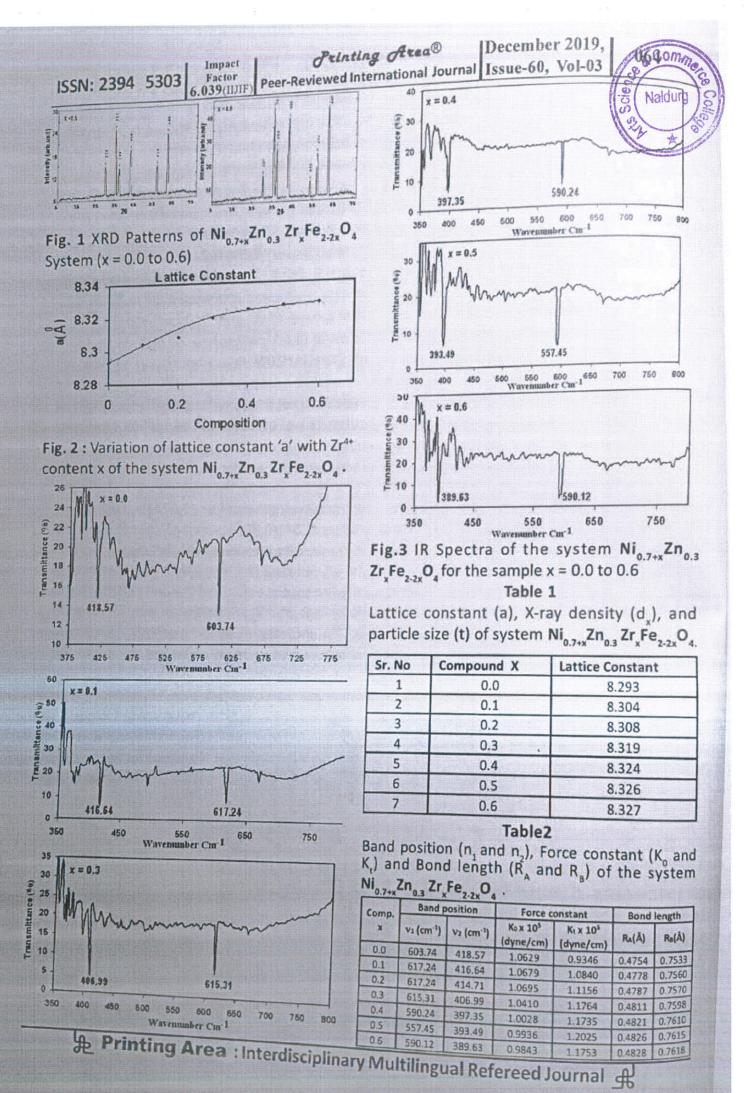
Where Ko is the force constant on octahedral site, K, is the force constant on tetrahedral site, M, is molecular weight of Asite, M₂ is molecular weight of B- site, I₁ is the corresponding center frequency on tetrahedral site, and iz is the corresponding center frequency on octahedral site. The molecular weight M₁ and M, for each sample is calculated from the cation

distribution. M, is molecular weight of A-site M, is molecular weight of B- site, i, is the corresponding center frequency on tetrahedral site, and i, is the corresponding center frequency on octahedral site. The molecular weight M1 and M2 for each sample is calculated from the cation distribution. The values of K, and K, are summarized in Table 1.2. The values of bond length R and R have also been calculated from IR data and some are presented in Table 1.2. The values of R, and R, are increases with increase in Zr4+ content.

III. CONCLUSION

The analysis of X-ray diffraction patterns indicates that the samples possess single phase cubic spinel structure. The lattice constant obtained from XRD data increases with the increase in Zr4+ content x. The linear increase in 'a' with x is due to the replacement of smaller ionic crystal radius of Fe3+ (0.67Å) by larger ionic radius of Zr4+ (0.80Å) obeying Vegard's Law. The X-ray density increases with Zr4+ concentration for all, because the increase in mass over takes the increase in volume of the unit cell. The bond lengths R_A and R_B are increased with Zr⁴⁺ content x. The increase in bond lengths R and R are attributed to increase in lattice parameter 'a' with Zr⁴+ content x.





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Geographical Aspects Of Maalgaon Watershed

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Introduction :

Geographers prefered water as their study to make up the geographic scene. They considered hydrologic cyde precipation as the frist phase. The georaphers recived sporadic attention to the land phases of the cycle.

The american study of hydrology taken under economic heads. The Geologist, meteorologists and engineers have been behaviour of water, treid to keep it during floods the work and study of hydrology taken plac in the last of th e19 th centtrory. The american taxbook on hydrology published two years later. The early neglected study of water by geographers resumed in the beginning of hte contrary. Running water was the touch stone to an understanding of land forms.

II: Location:

Beed district located in the central part of the marathwada region. Beed district lines between 18' 20' to 19' 27 north latitudes and 74' 57' to 76' 57 east longitude. This district has Majalgaon tahsil is the gar study area locatd to the north-east direction of Beed district. To the direction of east there is Parali tahsil and prabhani district to the direction of west theire is some part of georal tahsil and wadwani towards southern direction. Majalgaon is located in the centre of these Tahsil Godawari river is very important for Majalgaon taluka regarding crops and cultivation sindhphana river is so importanant because dam is on the river, sindhphana unite godawari at Manjrath

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