

Synthesis and Characterizations of Nickel Doped Zinc Oxide Nanocomposites by Solvothermal Method.

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Abstract

Nanocrystalline Nickel doped Zinc Oxide nanocomposites were successfully synthesized by Solvothermal method from Zinc acetate dihydrate and nickel nitrate hexahydrate using micro wave irradiation. The nanocomposite was annealed at 500⁰ C. Ni doped Zinc oxide nanocomposite was analysed through X- ray diffraction, transmission voltage microscopy, selected area diffraction, SEM, EDAX, Fourier transformed infrared spectroscopy and Photoluminescence spectroscopy. The particle size of the nanocomposite was estimated to be 49 nm. From SEM, the nanoparticles are in spherical shape. In FT-IR gives the Zn- O vibration gives its characteristic intense peak at about 500 cm⁻¹. The transmission peak was observed at 642 nm in the photoluminescence spectroscopy.

Key words: Solvothermal, Nanocomposite, nanoparticles, annealed, photoluminescence.

Introduction:

Admired for ZnO radioactivity was consumed in vast chemistry, electricity and radiance at immense conditions, ZnO has entered deep production and is one of the analytical architectures in present civilizations ^[1]. ZnO is surely an essential component in many modern construction techniques counting pigments, ointments, medicine, elastic, cluster, mechanical apparatus, stretching material, detergent, fibre, housing mat etc. The advance automation of ZnO frame was grainy and individual translucent. ZnO material will desire grow into developing useful in the neighboring subsequent ^[2]. The grains of rare earth metals were broadly considered as a result of their effective utilizations in the field of spintronics ^[3], auto electronic ^[4], sensor ^[5], radiation equipment ^[6], beam transmitting diodes ^[7] etc. The equity of the particular nanomaterials are transformed by virtue of sum internment appreciates exterior amount proportion ^[8]. Zinc oxide is a useful component ^[9]. The magnetic and more electric equity of Zinc oxide can be used as a sensor, converter, power alternator and light stimulant manufacture of hydrogen ^[10]. The more electric capacity of Zinc oxide was used in the different operations of audio flap vibrator, audio visible inflector etc ^[11]. Zinc oxide has absolutely, broad line difference of 3.3 eV and 60 meV activation efficiency ^[12].

Chemicals required:

The analytical grade reagents were used in this work. These were: Zinc acetate dihydrate, Nickel nitrate hexahydrate. Urea, Ethylene glycol and Acetone were acquired. Deionized water was used complete this procedure.

Manufacture of Nickel doped Zinc Oxide nanocomposite:

50 ml of Ethylene glycol was mixed with Zinc acetate dihydrate, nickel nitrate hexahydrate and Urea. The chemicals were stored in the heated klin. A great amount green precipitate was occurred. The composite was refreshed and cleaned with water. Finally acetone was used to remove the organic impurities. The composite was dehydrated as well as preserved. The composites were analysed by using the following approaches such as X- ray diffraction, transmission voltage microscopy, selected area diffraction, SEM, EDAX, Fourier transformed infrared spectroscopy and Photoluminescence spectroscopy.

Results and discussion:

I. Structural property:

1. XRD :

Fig (i) shows the XRD pattern of unannealed Nickel doped Zinc Oxide

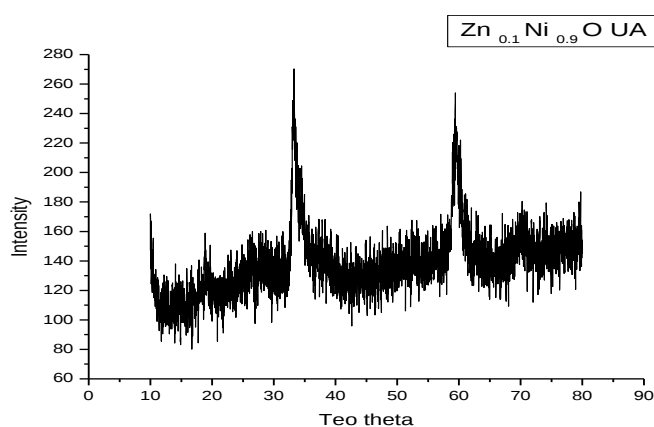


Fig (i) XRD pattern of unannealed Nickel doped Zinc Oxide

The crystals were amorphous in nature. Therefore the samples were annealed at 500⁰ C.

Fig (ii) & Fig (iii) shows the XRD spectrum of Nickel doped Zinc Oxide nanocomposite annealed at 500⁰ C.

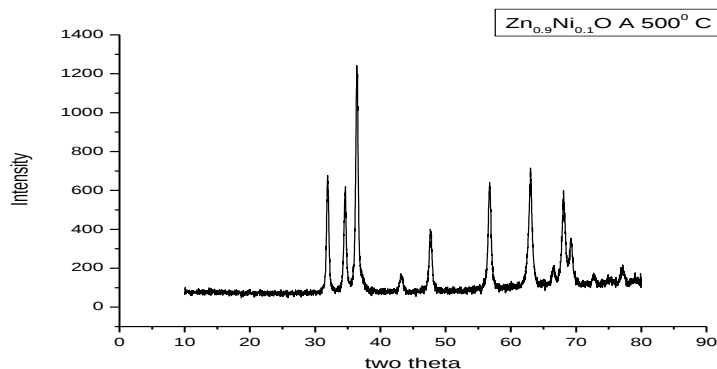


Fig (ii) XRD spectrum of Nickel doped Zinc Oxide

Fig (ii) provides a single development of ZnO with a hexagonal frame work. The lattice parameter values of ZnO are $a=b=3.2409$, $c= 5.1819$. XRD patterns reveal that the ZnO peaks were well matched to the standard diffraction of hexagonal Zinc oxide structure. (JCPDS file No.89 – 7103). The equate range of the samples were established to be 45 nm using Debye - Scherer formula^[13].

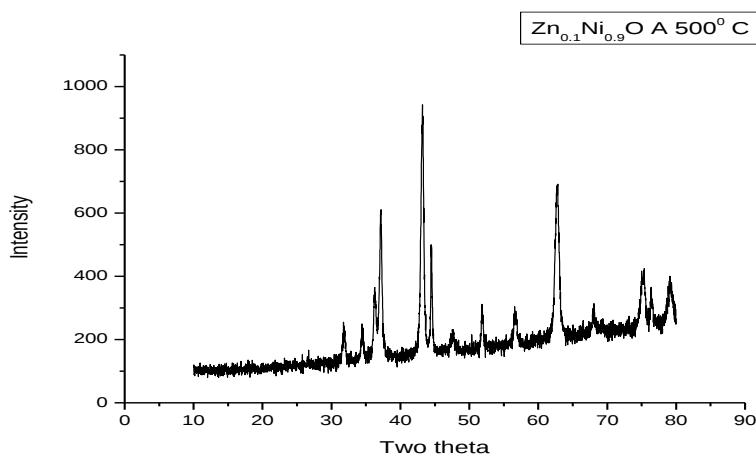


Fig (iii) XRD pattern of Nickel doped Zinc Oxide

Fig (iii) gives a secondary phase of NiO with a rhombohedral structure. In this case the Zn peaks along with corresponds to NiO. The magnitude and position of the lines were excellent arrangement with literature values (JCPDS file No.89 – 3080). The size of the nanocomposite was observed at 49 nm.

Table (i) & (ii) The lattice parameter values of Nickel doped Zinc Oxide nanocomposite annealed at 500⁰ C.

Peak Fitting											
lambda0 = 1.5406±0.0000 Å D:\GROUP-2\Ni11-1\data Data ...XRD D:\GROUP-2\Ni11-1.DAT											
#	hkl	d (Å)	2*theta (°)	Area	fwhm	diff	type	comment	offset	slope	quad
hexagonal a=3.2409±0.0063 c=5.1819±0.0022 c/a=1.5989±0.0007 v=47.135											
0	100	2.7874	32.085±0.011	9179±962	1.118±0.046	-0.0193	II		132966	-4295.62	-
1	002	2.5910	34.591±0.007	417±62	0.467±0.036	-0.0000	II		204	-5.18	-
2	101	2.4657	36.408±0.005	1021±110	0.493±0.027	-0.0022	II		1963	-56.59	-
3		2.0960	43.125±0.018	7±5	0.147±0.082	-	II		-3793	91.14	-
4	102	1.9048	47.707±0.029	535±359	0.732±0.203	0.0010	II		1051	-23.75	-
5	110	1.6210	56.744±0.007	549±83	0.576±0.046	0.0006	II		-1963	35.08	-
6	103	1.4736	63.029±0.007	683±76	0.682±0.039	0.0026	II		6624	-104.83	-
7	200	1.4039	66.553±0.018	59±18	0.476±0.102	0.0006	II		-436	8.41	-
8	112	1.3758	68.098±0.012	486±103	0.678±0.073	0.0019	II		-1458	22.65	-
9	201	1.3558	69.241±0.012	138±40	0.472±0.077	0.0013	II		3019	-41.24	-
10	202	1.2363	77.085±0.084	123±177	0.984±0.641	0.0023	II		-1372	19.17	-
no structure selected											
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Table (i) Lattice parameter value of Nickel doped Zinc Oxide

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no structure selected											
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no structure selected											
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Table (ii) Lattice parameter value of Nickel doped Zinc Oxide

2. TEM & SAED:

Fig (iv) was the TEM image of Nickel doped Zinc Oxide nanocomposite. The grain sizes of the samples were estimated at 48 – 50 nm. SAED appearance Fig (v) presents that the hexagonal and rhombohedral structure of as prepared Ni doped ZnO nanocomposite.^[14].

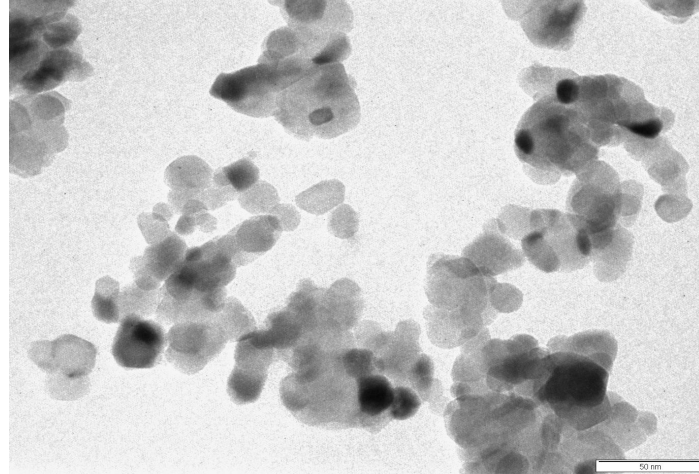


Fig (iv) TEM image

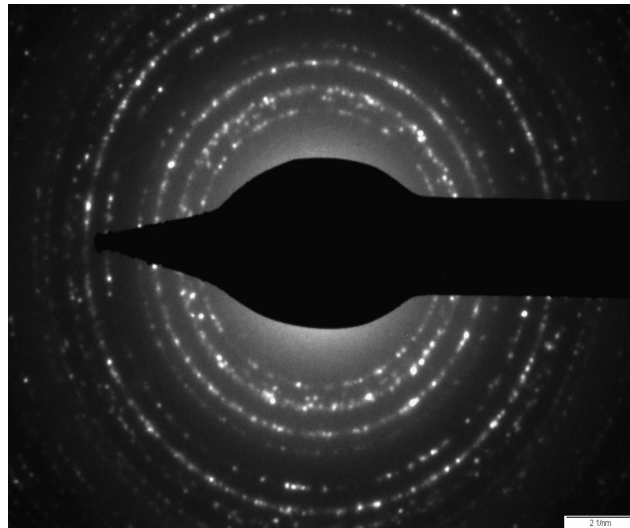


Fig (v) SAED pattern

3. SEM & EDAX:

Scanning electron microscope of the Nickel doped Zinc Oxide nanocomposite was shown in the following Fig (vi). It reveals that the Nickel doped Zinc Oxide particles forming the spherical structure. The energy dispersive x- ray analysis of Nickel doped Zinc Oxide nanoparticles were presented in Fig (vii). It is evident from the X –ray patterns , the dopants are found in the respective spectrum.

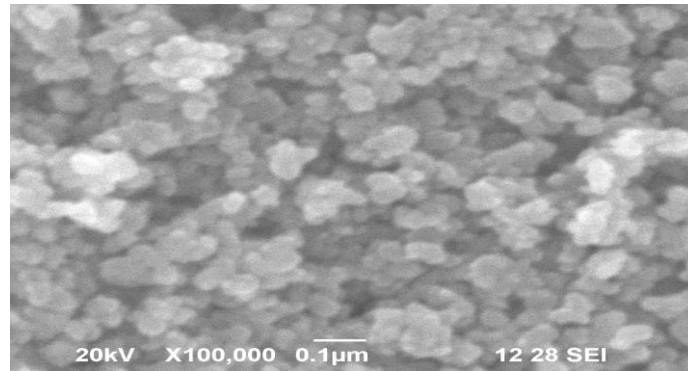


Fig (vi) SEM image

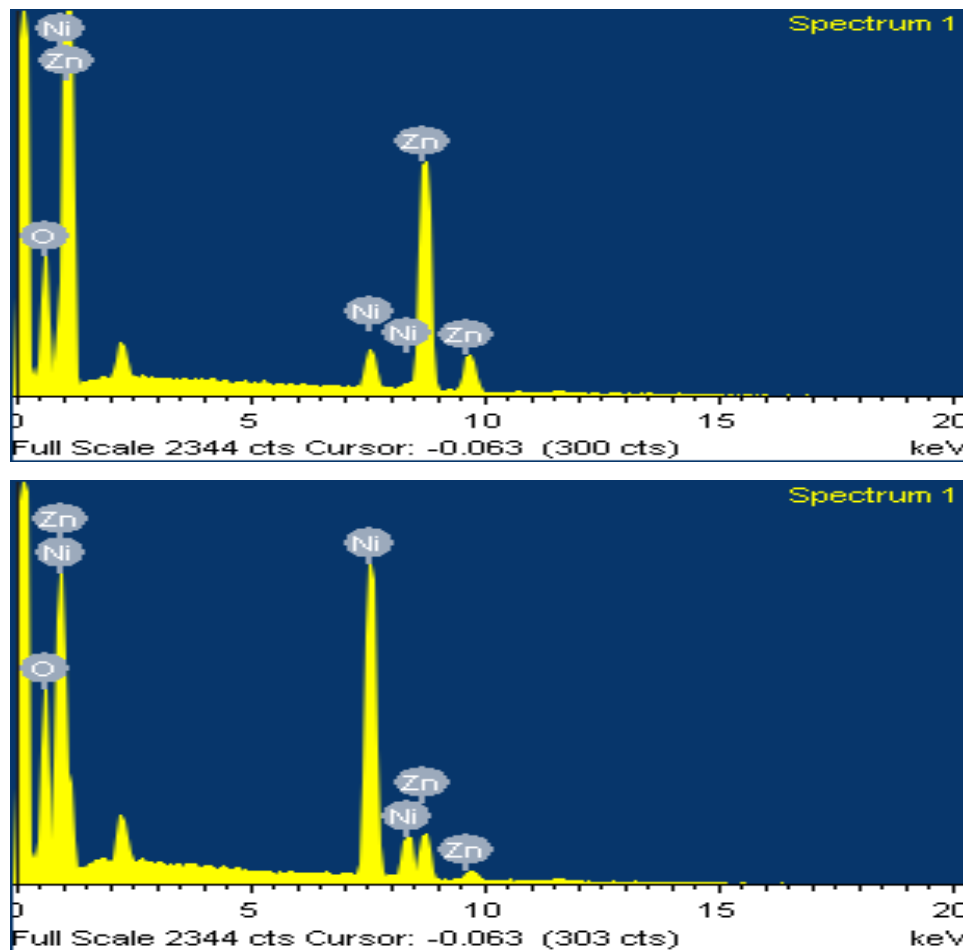


Fig (vii) EDAX spectrum of Ni doped ZnO nanocomposite

II. Optical property:

1. Fourier Transformed Infra Red Spectroscopy:

We have carried out the Fourier Transformed infra red spectra for the Nickel doped Zinc Oxide nanocomposite was shown in Fig (viii). In this figure, the higher energy region, there is a broad peak between $3000 - 3600 \text{ cm}^{-1}$. It might be due to OH stretching of H_2O . The presence of H_2O is once again confirmed by its bending mode at 1641 cm^{-1} . The presence of adsorbed EG is also confirmed by its CH_2

bending mode at 1377 cm^{-1} . The broad line has been recognized around at 500 cm^{-1} for the development of Zinc Oxide band.

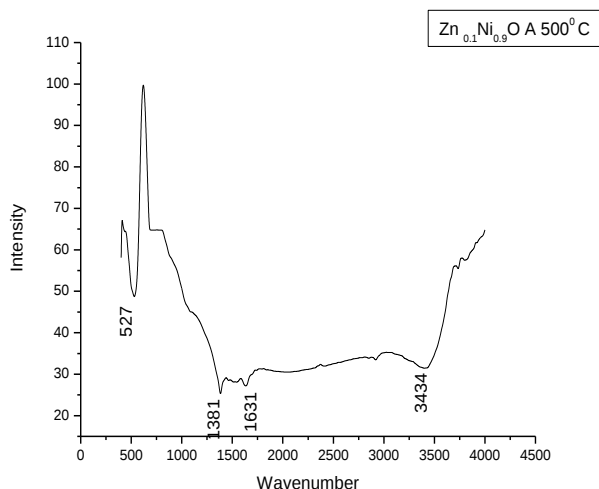


Fig (viii) FT – IR spectrum of Ni doped ZnO

2. Photoluminescence Spectroscopy:

Fig (ix) gives the PL spectrum of Nickel doped ZnO nanocompositie. A peak at 390 nm, it was the band edge excitonic luminescence of Nickel doped ZnO. This was due to the exciton recombination through an exciton – exciton collision process. The emission peak approximately observed at 642 nm in orange red region of Ni in ZnO.

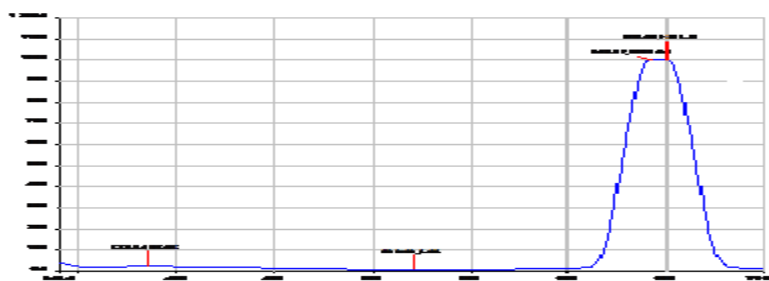


Fig (ix) Photoluminescence Spectrum of Ni doped ZnO

Conclusion:

Ni doped ZnO nano particles with single phase of Hexagonal structure of ZnO and secondary phase of Rhombohedral structure of NiO are synthesised successfully by Solvothermal method using micro wave irradiation. From XRD, the average size of nanoparticles were found to be 49 nm. Further the particle size was confirmed by TEM analysis. From TEM, the sample size was observed in the arrange of 50 nm. From SEM, the nanoparticles are in spherical shape. In FT-IR gives the NiO - ZnO vibration gives its characteristic intense peak at about 500 cm^{-1} . The peak at 390 nm, it was the band edge

through an exciton – exciton collision process. Nickel doped nanocomposites are further used in nano electronics, particularly in front of areas of flat panel displays and electron emitter devices.

References:

1. Drexler Eric kK (1986) Engines of Creation: The coming Era of Nanotechnology Doubleday.
2. Su SJ, Kuramoto N (2000) processable Poly aniline – titanium dioxide nanocomposites: effect of titanium dioxide on the conductivity Synth Met 114 : 147 – 153.
3. Meron T, Markovich G (2005) Ferromagnetism in colloidal Mn^{2+} doped ZnO nanocrystals. J phys Chem B109: 20232.
4. Liang S, Sheng H , Liu Y, Huo Z, Lu Y et al (2001) ZnO Schottky ultra violet photodetectors . J Cryst Growth 225: 110 – 113.
5. Shishiyanu ST, Shishiyanu TS, Lupan OL (2005) Sensing Charactersitics of of tin – doped Zno thin films as NO_2 gas sensor. Sens Actuators B 107: 379.
6. Huang MH, Mao S, Feick H, Yan H, Wu Y, et al (2001) Room temperature ultra violet nanowire nanolasers. Science 292 : 1897.
7. Saito N, Haneda H, Sekiguchi T, Ohasi N, Sakaguchi I, et al (2002) low – Temperature Fabrication of Light Emitting Zinc Oxide Micro patterns using Self Assembled Monolayers, Adv. Mater 14 : 418.
8. Ma Y, Ricciuti C, Miller T, Kadlowec J, Pearlman H (2008) Enhanced Catalytic Combustion Using sub micrometer and nano size Platinum Particles. Energy and Fuels 22 : 3695.
9. Segets S, Gradi J, Taylor RK, Vassilev V, Peukert W (2009) Analysis of Optical absorbance Spectra for the determination of ZnO nanoparticles Size distribution , Solubility and Surface energy. ACS Nano 3: 1703 – 1710.
10. Chaari M, Matoussi A (2012) Electrical conduction and dielectric Studies of ZnO Pellets. Phys B Condens matter 407: 3447.
11. Coroso AD, Posternak M, Resta R, Baldereschi A (1994) Ab initio study of piezoelectricity and Spontaneous Polarisation in ZnO. Physical Review B 50: 10715.
12. Wang ZL, (2004) Zinc Oxide nanostructures: growth, properties and applications. J Phys Condens Matter 16: R829 – R 858.
13. H.Klug, L.Alexander, X – ray diffraction procedurees, wiley, Newyork, 1962, p 125.
14. H.Wang, J.Zhang, J.J.Zhu, H.Y. (2002),Chen..*J.cryst.Growth*, 244, 88.