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Thermal, structural and magnetic properties of TeO₂-MgO-Na₂O-Nd₂O₃ glass system with NiO nanoparticles



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ABSTRACT

In this work, we have five samples of NiO Nanoparticles (NPs) embedded in tellurite glass system of composition Neodymium doped tellurite nanostructured glass having a composition of (72.5-x)TeO2-15MgO-10Na2O-2.5Nd₂O₃-(x)NiO, x = 0.5, 1.0, 1.5, 2.0, 2.5 mol% has successfully been completed by melt-quenching procedure. Structure of crystallization is determined using XRD technique verifies amorphous natures of produced glass. Differential thermal stability analyser (DTA-Parkin-Elmer) is utilized to illustrate the thermal stability of a sample at a temperature ranged from 300 to 600 °C. A Transmission Electron Microscope (TEM) is used to detect the presence of NiO NPs and High-Resolution Transmission Microscope (HRTEM) is utilizing to measure the dspacing of the NPs. The structure in glass network is identified by FT-IR and Raman spectrometer. The elemental trace in the glass sample is detected using the energy dispersive of X-ray (EDX) analysis. Meanwhile, magnetization measurements are carried out using Variable sample magnetometer (VSM). The transition temperature, T_g is found at 388 °C, the melting temperature T_m is ranged at 530-540 °C depending on composition while the thermal stability is around 117 °C. The structural and magnetic properties of the glass samples are observed to be delicate to NiO NPs concentration variations. The d-spacing at (111) plane having a size of 0.24 nm proves the presence of NiO NPs. The glass network contains of bending vibrations of Te-O-Te linkages centred at 448 cm⁻¹which is also the stretching band of NiO mode. It is also found that the glass structure is dominated with the existence of TeO3, TeO3+1 and TeO4 tribypiramidal. The glass system reveals paramagnetic behaviour by magnetic hysteresis at room temperature with Hc values were observed to be decreased from 610 ± 30.5 to 139 \pm 6.95 (Oe) and ratio of (Mr/Ms) from 0.153 \pm 0.008 to 0.071 \pm 0.004with the increase of NiO NPs concentration

1. Introduction

International progress in tellurite glasses modified with rare earth oxides (R.E.) or transition metal ion (T.M.) have been published caused by their interesting properties [1–5]. Tellurite glass is widely known due to better glass stability, high non-linear refractive index, highly corrosion resistance and unaffected to moisture for long periods compare to those of silicate, phosphate and fluoride glasses relatively. This implies that tellurite glasses are suitable for optical fibres, modulators, non-linear and laser applications [6–8]. Tellurium oxide is considered as alluring hosts for dynamic medium, considering their low phonon vitality and low liquefying temperature. So, adding a glass modifier enhances glass formation ability (GFA) through the breaking chain of the truth of the properties of the properti

applications [9,10]. However, the phosphate host materials are known to be lack of during ty, thus it need to be substituted with more suitable host materials. The combination of tellurite oxide and neodymium oxide offers novel and high optical efficiency in laser glass [11].

Recent technological applications have generated more attraction in the studies of NPs of magnetic material of the their properties which usually vary from bulk materials. They have important applications in magnetic data-storage media, in magnetic liquids (ferrofluids), i.e. stable suspensions of magnetic NPs, in magnetic beads that are applied in biotechnology, for contrast enhancement in magnetic resonance imaging (MRI), and for the target drug delivery [12]. In this regard, tellurite glasses embedded with magnetic metallic NPs system has also gained a massive interest. The incorporation of magnetic metal NPs in glass improve the electrical field around rare earth and produce quantum effect to increase the optical efficiency of the glass system

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[13]. Among magnetic metal NPs, NiO NPs are particularly attractive due to their beneficial magnetic and electronic properties. They can be applied in the gas sensor, lithium - ion battery, p - type semiconductors (cubic, 3.5 eV), alkaline battery cathode and electrochemical capacitors [14–16].It also exhibits anodic electrochromy, excellent durability, electrochemical stability, and high optical spin density. A recent study on NiO NPs has shown that NPs with an average diameter of 3–24 nm exhibit superparamagnetic along with magnetic hysteresis under a blocking temperature [17]. Other reported that small amount of NiO particles were known to be super - paramagnetic [18] while for pure NiO, it exhibits antiferromagnetic ordering with plane of opposite spin [19].

Although the behaviour of local environr [14] of NiO NPs has been well established, advanced investigation on the influence of NiO NPs embedded in tellurite glass on the structural characteristics and magnetic properties are still deficient. Thus, in this research, the thermal, structural and magnetic properties of (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO glasses are studied using XRD, TEM, HRTEM, EDX, DTA, FTIR and Raman spectroscopy techniques. All the results will be discussed in detail with respect to composition.

2. Experimental work

The glasses with nomina 19 pmposition of (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-xNiO with x = 0.5, 1.0, 1.5, 2.0, 2.5 mol% are made through melt quenching method. By mixing the oxides, batches of 20 g of well - mixed starting materials were prepared. Then, the well - mixed raw material is melted in an electric furnace at 900 °C in platinum crucible for half an hour. Af 12 hat, the melts were quenched on a preheated stainless-steel mould at 300 °C before cooling gradually down to the room temperature. The obtained glass samples are named as 0.5NiO, 1.0NiO, 1.5NiO, 2.0NiO and 2.5NiO for x = 0.5, 1.0, 1.5, 2.0, and 2.5 respectively. The X-rassiffraction (XRD) measurement is done on a relatively fine powder by CuKα radiation at 40 kV, 30 mA utilizing Siemens Diractrometer D5000 to identify the amorphous nature of glass. The differential thermal stability analyser (DTA-Parkin-Elmer) is utilized to in strate thermal stabilities of sample in the range of 300-600 °C. The energy dispersive X-ray spectrometer (model Swift ED3000 EDX) was utilized to perceive elemental cost of glass samples. The existence of NiO NPs is noticed utilizing Transmission Electron Microscope (TEM) (model Philips CM12 along with Docu Version 3.2 and High-Resolution Transmitted Electron Microscope (HRTEM)functioned at a voltage acceleration of 20 kV. All infrared spectrum of the glass is noted at the room temperature (25 °C) utilizing a Perkin-Elmer Spectrum on FTIRspectrometer over the ranged 4000-400 cm⁻¹by 10 scans419at 4 cm⁻¹ resolutions. The Raman spectra are measured through a confocal Horbia Jobin Yvon (HR800 UV) at the spectral ranged 2012000 cm⁻¹, with the laser power on the samples being 1000 mW. Magnetization measurements are carried out using Variable Sample Magnetometer (VSM).

3. Result and discussions

Table 1 should the formation, colour and transparency of produced glass samples (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, x = 0.5, 1.0, 1.5, 2.0, 2.5 mol%. All samples are transpart and slightly change in colour from green yellowish to light brown. Fig. 1 displays the disordered nature of the glass sample in the 2.5NiO glass spectra as confirmed by XRD measurement. The XRD pattern verifies the glass is in the amorphous states, as indicated by the absence of sharp peaks with a broad hump for 2θ around the region 20-35°. Fig. 2(a) demonstrates the TEM images for 2.5NiO glass. The selected area in Fig. 2(a) is acceptable to predict the size of NiO NPs. Some black non-spherical spots having various shapes and sizes are visibly distributed inside the glass network. The inset of the Fig. 2(a) demonstrates the size of NiO NPs is observed to be gaussian having an average diameter 4.9 nm. Fig. 2(b)

clearly shows the sample SAED pattern suggesting strong bright diffraction spots revealed single crystalline cubic nickel oxide formation [20–23]. The appearance of NiO NPs lattice spacing for sample in HR-TEM image is shown in Fig. 2(c). The estimated distance between crystal lattice spacing is about 0.24 nm corresponding to (111) crystal facet of NiO NPs which matched with JCPDS card no. 00–004-0835 [24] and 03–065-5745. Fig. 2(d) demonstrates an optical absorption spectrum of [18] bulk samples (\approx 2.0 mm thick) with seven absorption bands of; $(^2P_{1/2},\ ^2D_{5/2}),(^4G_{7/2}),(^4G_{5/2},\ ^2G_{7/2}),\ (^4F_{9/2}),(^4S_{3/2},\ ^4F_{7/2}),\ (^2H_{9/2},\ ^4F_{5/2})$ and $(^4F_{3/2})$ as explained in previous report [25]. It can be seen in the present glasses that optical absorption edge has no sharp peaks indicates their glassy nature [26]. It is observed that the cut-off wavelength is in the range of (367–370) nm which indicate the limiting absorption at UV range. This result is in agreement with the other tellurite glass system reported elsewhere [27].

Creation of NiO NPs is as cribbed to nucleation and growth and process, where potential for reducing redox system elements is demonstrated as follows:

$$Nd^{3+}/Nd^{2+} = 2.6 V$$

$$Nd^{3+}/Nd^{0} = -2.32 V$$

$$Te^{6+}/Te^{4+} = 1.02 \text{ V}$$

The reduction procedures involved areas follows:

$$Te^{4+} + 2Nd^{2+} = Te^{6+} + 2Nd^{2+} \& \Delta E^{0} = -6.22 \text{ V}$$

$$3\text{Te}^{4+} + 2\text{Nd}^{3+} = 3\text{Te}^{6+} + 2\text{Nd}^{0}\&\Delta E^{0} = -7.70 \text{ V}$$

$$NiO \rightleftharpoons Ni^{2+} + O^{2-}$$

where ΔE^0 is the total of reduction process. Furthermore, NiO does not have ΔE^0 due to the bonding between Ni and O cannot be broken. The existence of NiO NPs in a glass may be strengthened by HRTEM lattice spacing, TEM images and EDX result. The calculated optical spectrum of nanosized nickel SPR absorption was reported to be between 300 and 400 nm [28]. Besides, this result is consistent with other reports [29,30]. Fig. 3(a) illustrates the distribution of elements in 2.5NiO glass. It can seemly be observed that Te, Mg, Nd, and Ni elements are randomly distributed among each other in scale of 5 µm image. However, the blue colour for these NPs cannot be seen clearly because of too small particle size (≈ 4.9 nm) of NiO NPs and low composition content of NiO (2.5 mol%), in line with Fig. 1 that cannot be detected by XRD either. However, EDX spectrum reveals some weak peaks of Ni from spectra as shown in Fig. 3(b). Table 2 encapsulates the details of the data collected from EDX. It shows the experimental and calculated weight of elements for the 2.5NiO glass sample. From Table 2, it can be seen that some Ni elements are detected which confirmed that the black spots showed in Fig. 2(a) are indeed NiO NPs.

Fig. 4 shows the DTA thermogram for the present (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, x = 0.5, 1.0, 1.5, 2.0, 2.5 mol% and glass crystallization temperature (T_c), transition temperature (T_g) and melting temperature (T_m) have been tabulated in Table 3. It indicates one blunt endothermic hump pertaining to T_g at about 338 °C which due to arrangement of NiO in the glass system [31]. Another endothermic band is corresponded to T_m in the range 530–540 °C. The exothermic peak at about 455 °C is translated for T_c . The glass stability (ΔS) and the Hruby (T_g) are the ways to evaluate the glass thermal stability [32]. These equations are defined as follows [33]:

$$\Delta S = (T_c - T_g) \tag{1}$$

$$H_R = (T_c - T_g)/(T_m - T_c)$$
 (2)

The ΔS of 2.5% NiO discovered the maximum thermal stability around 117 °C which is considered to be stable, in good agreement with the preceding value of ΔS when it is > 100 °C [34]. While the value of $H_Ris1.52$ that can be defined as the denominator of growth rate and nucleation rate [32]. These values are also consistent with the one

Table 1 The nominal composition of (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, where $0 \le x \le 2.5$ mol% glasses.

Sample		Nominal composition (mol%)		(o)	Color	Samples figure	
No.	TeO ₂	MgO	Na ₂ O	Nd ₂ O ₃	NiO		
TMNN1	72.0	15.0	10.0	2.5	0.5	Transparent, yellow	2
TMNN2	71.5	15.0	10.0	2.5	1.0	Transparent, yellow	2
TMNN3	71.0	15.0	10.0	2.5	1.5	Transparent, light brown	Q
TMNN4	70.5	15.0	10.0	2.5	2.0	Transparent, brown	
TMNN5	70.0	15.0	10.0	2.5	2.5	Transparent, brown	0 00

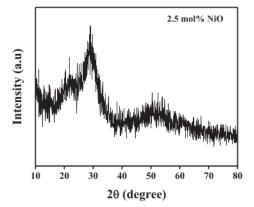


Fig. 1. XRD pattern of glass 2.5NiO.

reported by R. El-Mallawany [34].

Fig. 5 shows FTIR spectra of the present glass samples with 0.5, 1.0, 1.5, 2.0, 2.5 mol% of NiO NPs in a range of 400–2000 cm⁻¹. Table 4 abridges the illustrated peak at the 15 le spectral area and correlating band assignment. Any band centred at around 448 cm⁻¹ is allocated to combined bending vibration of the O-Te-O or Te-O-Telinkage [35]. This band also allocated to stretching NiO mode [36,37]. The broadening band in between 610 and 680 cm⁻¹ is suggested due to Te-O stretching vibration of TeO₄ trigonal bipyramidal unit with connecting oxygen [38]. With a gradual increase in concentration of NiO, the TeO₄ band shifted to lower number corresponding to a reduction in the number TeO₃ units [39]. Meanwhile, the other band centred at 702 cm⁻¹matches to a transitional coordination of tellurium atom between 4 and 3 [40]. The shoulder band appearance at 766 cm⁻¹ is attributed to TeO₃trigonal pyramidal (tp) units with non-bridging oxygen inside glass network.

As a complementary technique to the infrared spectra, Raman spectrum of present glass system with 0.5, 1.0, 1.5, 2.0, 2.5 mol% of NiO NPs is shown in Fig. 6. (a) Raman patterns consisting of broad peaks in a range: 550 to 950 cm⁻¹ and 300 to 550 cm⁻¹. Fig. 6. (b) shows the deconvolution of the Raman spectra of 2.5 mol% NiO NPs

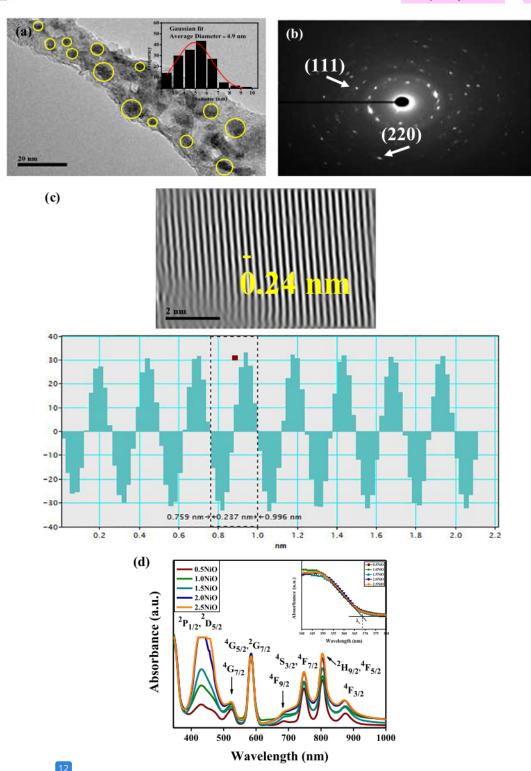


Fig. 2. (a) TEM images displaying the distribution of non-spherical NiO NPs (inset shows corresponding size distribution). (b) SAED pattern of 2.5NiO glass. (c) HR-TEM image and the lattice fringe profile of 2.5NiO glass. (d) UV absorption spectra of glass samples (inset illustrating cut off wavelength of glass samples).

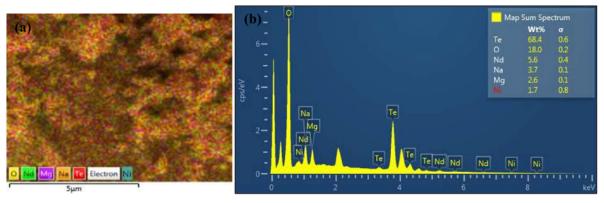


Fig. 3. (a) The distribution (b) EDX spectrum of elements Te, Mg, Na, Nd and Ni in 2.5NiO glass.

Table 2 Experimental and calculated weight of elements for the 2.5NiO glass sample.

Element	Experiment	cal	Calculated	
	Wt%	Atomic%	Wt%	Atomic%
Na	3.69	8.03	4.50	18.43
Mg	2.55	5.26	3.11	12.07
Ni	1.74	1.49	2.12	3.42
Te	68.40	26.85	83.45	61.63
Nd	5.59	1.94	6.82	4.45

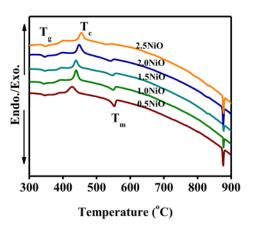


Fig. 4. DTA curve of glass samples.

Table 3 Glass transition temper 3 re (T_g) , crystallization temperature (T_c) and melting temperature (T_m) of $(72.5\text{-x})\text{TeO}_2\text{-15MgO-10Na}_2\text{O-}2.5\text{Nd}_2\text{O}_3\text{-}(x)\text{NiO}$, where $0 \le x \le 2.5$ mol%.

Sample no	Temperature (± 0.01 °C)	
	$T_{\rm g}$	$T_{\rm c}$	$T_{\rm m}$
TMNN1	334.46	427.56	552.72
TMNN2	330.83	439.47	549.44
TMNN3	333.47	438.63	547.68
TMNN4	334.06	448.19	541.79
TMNN5	337.86	454.91	531.78

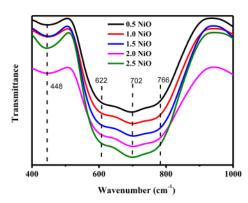


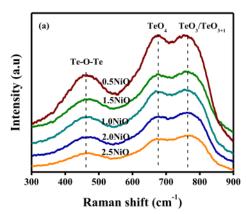
Fig. 5. NiO NPs content (mol%) dependent IR transmission bands of glass samples.

with base 10 is corrected and peaks are centred at 464, 689, and 789 cm⁻¹. The assignment of Raman band is summarized in the Table 5. In the frequency range of 300 to 550 cm⁻¹, the existence of bending vibrations of Te-O-Te and O-Te-O linkage is observed at about 464 cm -1 with O at a position alternatively equatorial and axial [41,42]. The strong band that appeared at peak around 689 cm⁻¹ is ascribed to stretching vibrational modes of TeO4 tetrahedral with bridging oxygen (BO) atoms. It has been proposed that this band determined to a coupled symmetric vibration along Te-O-Te axis in the TeO₃₊₁/TeO₄ unit and TeO₄/TeO₃ pair [43]. Meanwhile, it has been considered that band to be related measure of connectivity among TeO4, TeO3 and TeO3+1 species [44]. A Raman band centred at 789 cm⁻¹ can be assigned to the TeO₃or TeO₃₊₁ units [42,45,46].Increasing the NiO content of samples results in a shift to higher wave numbers from 752 cm^{-1} when x = 0.5 mol% to 789 cm^{-1} when x = 2.5 mol%. This is because an increase in the NiO content increases the quantity of non-bridging oxygen atom and transforms of TeO4 to TeO_3 and TeO_{3+1} structural unit in glass networks [44,47].

Fig. 7 displays the set of glass system hysteresis loop data obtained from magnetization M versus field H. It can be said that, the glass system at room temperature is an indication of paramagnetic behaviour. It is also in a similar trend with the hysteresis loop as reported previously [48]. The value of their remnant magnetization (Mr), saturation magnetization (Ms) and the coercivity field (Hc) can be found from the curve which are listed in Table 6. The obtained values of Ms. of bulk Ni (77.65 emu/g) [49]and Ni-Co oxide NPs [50] at room temperature are much higher than Ms. of glass system because of the implication of the non-magnetic surfactant coated [51]. It is might be due

Table 4 :IR band assignments of (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, where $0 \le x \le 2.5$ mol%glass samples with varying concentration of NiO NPs.

Glass					Band assignments
0.5NiO	1.0NiO	1.5NiO	2.0NiO	2.5NiO	
448	450	448	444	448	Bending vi 16 ons of Te-O-Te
622	618	616	614	614	Stretching vibrations of Te-O of TeO ₄
696-698	702	702-704	702	700–702	Stretching vibrations of Te-O of TeO_{3+1}
764	764	766	774	774	Stretching vibrations of Te-O of TeO ₃ units



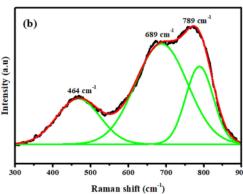


Fig. 6. (a) Raman spectra of TeO₂-MgO-Na₂O-Nd₂O₃-NiO, (b) the deconvolution of the Raman spectra of 2.5NiO.

Glass	Glass				Band assignments
0.5NiO	1.0NiO	1.5NiO	2.0NiO	2.5NiO	
457	471	469	469	464	Bending vibrations of Te-O-Te and O-Te-O
671	675	670	679	689	Stretching vibrations of TeO ₄
752	752	757	764	789	Stretching vibrations of TeO ₃ or TeO ₃₊₁

to disorder canting spin (Spin-glass-like)onto the exterior of NiO NPs [52]. From Fig. 8(a), it was found that Hc value decreases as NiO NPs increase, which can be attributed to the effects of porosity that work as a demagnetizing field generator. A higher field is required to push that domain wall by decreasing the porosity and thus decreasing Hc [53].According to Brown's relation [54] as per below,

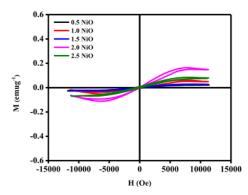


Fig. 7. The magnetic hysteresis loops (M-H curve) of NiO NPs embedded in tellurite glass.

Hcvarying inversely to Ms. which in accordance with current result and similar for $\rm NiFe_2O_4$ prepared by others [53].

$$H_c = 2K_1/\mu_s M_s \qquad (3)$$

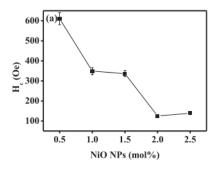
Therefore, a rising trend of both Ms. and Mr. are shown in Table 6 with an increased concentration of NiO NPs. Which clearly shows that changes NPs local field environment and their electron spins can lead to an increase in Ms. in the NiO NPs [55]. The comparison of the present values to other glass systems revealed higher values than phosphate glasses containing nickel NPs [49]. The modification of (Mr/Ms) as a function of NiO NPs concentration is presented in the Fig. 8(b). The (Mr/Ms) is an extent of how square a hysteresis loop is, which proposed a dominant role performed by an uniaxial magnetocrystalline anisotropy [56]. It is typically related with the bigger particles and the formation of domain - walls and high texture value [57]. The lower values of (Mr/Ms) in the range 0.01 to 0.1 indicates that the materials are in multidomain magnetic grains composed in the structure of the present tellurite glasses.

4. Conclusion

Transparent tellurite glass samples in the form (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, x = 0.5, 1.0, 1.5, 2.0, 2.5 mol% have been achieved in the amorphous matrix by using melt quenching technique. TEM observation shows the nucleation of NiO NPs has nonspherical shape with average size around 4.9 nm. Compositional analysis (EDX) discovered the existence of Ni, O,Te, Na, Mg and Nd in the prepared samples. Differential thermal analysis showed that T_g at 338 °C, T_m in the range 530–540 °C, exothermic peak T_c at about 455 °C, glass stability (Δ S) and the Hruby (H_R) are Δ S > 100 °C implies the glass system is regarded as stable and the value of H_R is 1.52. FTIR and Raman analysis have observed the involvement of all functional groups, and this study has confirmed that adding NiO content raises the quantity of non - bridging oxygen atoms. They revealed the presence of various groups such as bending vibrations of Te-O-Te, stretching vibration of TeO₄and stretching NiO mode in a title of glass. The magnetic

Table 6 Magnetic parameters of glass system of (72.5-x)TeO₂-15MgO-10Na₂O-2.5Nd₂O₃-(x)NiO, where $0 \le x \le 2.5$ mol%.

Sample No.	Ms(emu/g)	$Mr \times 10^{-3}$ (emu/g)	Hc(Oe)	Mr/Ms
0.5NiO	0.023 ± 0.001	3.51 ± 0.176	610 ± 30.5	0.153 ± 0.008
1.0NiO	0.063 ± 0.003	6.47 ± 0.324	349 ± 17.45	0.103 ± 0.005
1.5NiO	0.079 ± 0.001	1.88 ± 0.094	336 ± 16.8	0.024 ± 0.001
2.0NiO	0.164 ± 0.008	2.49 ± 0.125	125 ± 6.25	0.015 ± 0.001
2.5NiO	0.084 ± 0.004	6.00 ± 0.300	139 ± 6.95	0.071 ± 0.004



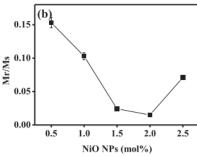


Fig. 8. (a) The variation of Hc as a function of NiO NPs concentration. (b) The variation of Mr./Ms. as a function of NiO NPs concentration.

measurements showed that the existence of NiO NPs in a glass pecimen is paramagnetic behaviour at room temperature (25 °C) and values were observed to be decreased from 610 \pm 30 1 to139 \pm 6.95 (Oe) and ratio of (Mr/Ms) from 0.153 \pm 0.008 to0.071 \pm 0.004with the increase NiO NPs concentration indicates that the materials are in multidomain magnetic grains composed in the glass structure.

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8 Declaration of Competing Interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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