Role of Praseodymium Addition in the Microstructure and Magnetic Properties of ZnCo Ferrite Nanopowders: Positive or Negative?

B. Shahbahrami\textsuperscript{a}, S. M. Rabiee\textsuperscript{a,}\textsuperscript{*}, R. Shidpoor\textsuperscript{b}, H. Salimi-Kenari\textsuperscript{b}

\textsuperscript{a}Department of Materials Engineering, Babol Noshirvani University of Technology, Babol, Iran
\textsuperscript{b}Faculty of Engineering & Technology, University of Mazandaran, Babolsar, Iran

**ABSTRACT**

In order to investigate the effect of praseodymium (Pr) addition on the microstructure and magnetic properties of cobalt-zinc ferrite nanoparticles (NPs), different values of Pr element (x=0.2, 0.4 and 0.6) were added to the initial composition (Co\textsubscript{0.8}Zn\textsubscript{0.2}Fe\textsubscript{2}O\textsubscript{4}) in the co-precipitation method, and the prepared precipitates calcined at 750 °C for 2 h. The synthesized powders were characterized by X-ray diffraction (XRD), field emission electron microscopy (FESEM), Fourier transform infrared spectroscopy (FTIR), and vibrating sample magnetometer (VSM). XRD pattern revealed the formation of a secondary phase of Pr-Fe oxide in addition to the ZnCo ferrite phase in the samples. FESEM images showed changes in the morphology and size of the particles by adding Pr to the composition. For specimen with x=0.2, the homogeneous spherical like particles with the size about less than 30 nm was formed. Whereas, for composition containing x=0.6 of Pr, a non-uniform powder with plate-like particles was obtained and NPs had a thickness of approximately less than 30 nm. VSM analysis indicated that by increasing the element Pr to the cobalt-zinc ferrite composition, especially for values higher than x=0.2, the powder become a completely non-magnetic material.

**Keywords:**
Coprecipitation
Cation Distribution
Microstructure Properties
Magnetic Properties

**NOMENCLATURE**

\begin{tabular}{ll}
XRD & X-ray diffraction \\
FESEM & Field emission scanning electron microscopy \\
VSM & Vibrating sample magnetometer \\
M\textsubscript{s} & Saturation magnetization \\
K & Anisotropic constant \\
H\textsubscript{c} & Coercive force \\
\mu\textsubscript{B} & Bohr magneton \\
Mr & Remanence magnetization \\
a & Lattice parameter (Å) \\
D & Crystallite size (Å) \\
\end{tabular}

**1. INTRODUCTION**

The magnetic properties of ferrites with spinel structure and their application depend on the composition and distribution of the cations at the tetrahedral (A) and octahedral (B) spaces [1-4]. Accordingly, many researchers have replaced various elements in the spinel structure and have studied changes in magnetic properties such as the saturation magnetism (M\textsubscript{s}), coercive force (H\textsubscript{c}), anisotropy constant (K), remanence magnetization (M\textsubscript{r}), and the magnetic moment (n\textsubscript{m}). Cobalt ferrite is an interesting material due to its high magnetic permeability, high coercive force, good saturation magnetization (M\textsubscript{s}) and usability in a wide range of applications such as telecommunications and medicine [2, 5]. Therefore, research is underway to study the effect of element substitution on cobalt ferrite materials. For example, Topkaya et al. [6] synthesized cobalt ferrite nanoparticles (NPs) substituted with zinc element (Zn\textsubscript{0.4}Co\textsubscript{0.6}Fe\textsubscript{2}O\textsubscript{4}) by hydrothermal method. The highest M\textsubscript{s} of 76.5 emu/g reported for ferrite containing x=0.2. In addition, they stated that Zn\textsuperscript{2+} ion is located in A site, forcing Fe\textsuperscript{3+} ions to move from the A position to the B space. Dalal et al. [7] also synthesized Ni\textsubscript{0.2}Zn\textsubscript{0.8}Co\textsubscript{0.2}Fe\textsubscript{2}O\textsubscript{4} NPs by co-precipitation method and obtained crystallites with a size of about 34.7 nm and

\*Corresponding Author Institutional Email: rabiee@nit.ac.ir (S. M. Rabiee)
high magnetization (84 emu/g) at room temperature. Ramakrishna et al. [8] synthesized Co_{0.5}Zn_{0.5}Cu_{0.13}Fe_{2}O_{4} NPs with crystallite diameters 82.41 nm via sol-gel method. They reported Ms, Hc, K and n0 of the NPs equal to 50.35 emu/g, 198.9 Oc, 1.7 x 10^4 erg/Oe and 2.9 μB, respectively.

In another study by Naik and, Salker [9], the effect of adding Dy and Gd elements on magnetic properties of the cobalt ferrite NPs at room temperature (300 K) and very low temperature (5 K) investigated. With analysis of their results, it was found that firstly, lowering the temperature increased the saturation magnetization and the coercive field. The reason for this is activation of the magnetic moments of Dy and Gd ions at temperatures below 40 K. By substituting Dy^{3+} and Gd^{3+} ions instead of Fe^{3+}, the magnetic moment of the B sites, saturation magnetization and the coercive field increases. In addition, it observed that the amount of M_{s} in Dy-doped cobalt ferrite NPs (70.29 emu/g) is higher than that of Gd-doped specimen (60.85 emu/g) at room temperature. The higher magnetic moment in Dy ions than Gd ions mentioned for this difference.

In the work of Xavier et al. [10], the effect of samarium on the structural and magnetic properties of cobalt ferrite NPs (CoFe_{2-x}Sm_{x}O_{4}) investigated. They stated that with increasing Sm^{3+} concentration to x=0.25, the crystallites size increases from 11.26 to 16.53 nm, which is due to the higher ionic radius of the samarium than the iron element. For this reason, the coercive force reduced from 1372.2 Oe to 814.73 Oe, which is due to the easier movement of magnetic domains.

Alves et al. [11] studied the effect of adding Y^{3+} ions on the structural and magnetic properties of cobalt ferrite NPs (CoY_{2-x}Fe_{2}O_{4}). By increasing the Y^{3+} ion values to 0.04, the grain size of the sample decreased from 35.32 to 16.05 nm. The saturation magnetization also decreased from 69 emu/g to 33 emu/g, while the coercive force increased from 1100 Oe to 1900 Oe.

In order to control particle size at high calcination temperatures and improve magnetic properties, Panda et al. [12] doped rare earth elements Pr or Gd in cobalt ferrite compositions and calcined them at different temperatures. They stated that the single-domain size of the pure cobalt ferrite materials is 70 nm. On the other hand, with increasing the calcination temperature to more than 500 °C, the size of the crystallites significantly increases, so the presence of rare earth elements in the ferrite structure prevents their growth. The crystallite sizes of the samples synthesized by them for the compounds CoGd_{1-x}Fe_{2}O_{4}, CoPr_{1-x}Fe_{2}O_{4}, CoGd_{1-x}Fe_{3}O_{4}, and CoPr_{1-x}Fe_{3}O_{4} at 900 °C were equal to 41.4, 33.2, 65.6 and 57.5 nm, respectively, while for pure cobalt ferrite it was 87.5 nm.

We know that the particle size plays a role in the relaxation time according to Equation (1), and the rapid switching of magnetic behavior occurs with smaller particles [13-15].

$$\tau_N = \tau_0 e^{(K/B \tau)}$$

(1)

where \(\tau_N\) is the Neel relaxation time, \(\tau_0\) is the exponential parameter \((\tau_0=10^{-2}.10^{-3} s^3)\), K is the anisotropy constant, V is the volume of NPs, K_B is Boltzmann constant, and T is the temperature (K_BT is thermal energy).

In the other hand, the amount of magnetic moment theoretically depends on the difference in magnetic moment between the tetrahedral and octahedral spaces according to Equation (2) [6, 16, 17]. Therefore, with the introduce of an even non-magnetic element into the ferrite composition and its replacement instead of one of the elements in A or B spaces, the initial order of ions is disturbed and may lead to a difference in the magnetic moment of the material.

$$nB(c.e.L.) = MB(x) - MA(x)$$

(2)

Therefore, in the present work, the element praseodymium is added to the zinc cobalt ferrite composition and its effect on the microstructure and magnetic properties is studied, which has not been investigated by others. In this composition, by doping Zn in cobalt ferrite structure, it is expected to increase the crystallite sizes and saturation magnetization, and reduce the coercive force. By adding Pr to the composition, this element is located in the octahedral spaces instead of iron element and acts like non-magnetic atoms at room temperature. However, it is used for this research for two reasons; first, due to the unpaired electrons in the 4f layer, it has a single ion anisotropy in the crystal lattice, which may be useful. Second, by changing the particle size and surface effects due to inhibiting the growth of particles and the crystallite diameters beyond the range of single-domain size (70 nm) may improve the microstructure and magnetic properties.

2. EXPERIMENTAL

Nanocrystalline powders (Co_{0.5}Zn_{0.5}Fe_{2-x}Pr_{x}O_{4}, x=0.0, 0.4 and 0.6) were prepared by co-precipitation method using raw materials of iron nitrate (Fe(NO_{3})_{3}.9H_{2}O) (Merck), cobalt nitrate (Co(NO_{3})_{2}.6H_{2}O) (Merck), zinc nitrate (Zn(NO_{3})_{2}.4H_{2}O) (Merck), and praseodymium nitrate (Pr(NO_{3})_{3}.6H_{2}O) (Merck). Using deionized water (DI), 0.5 M solutions were prepared from each of the starting materials. The solutions were stirred for 10 m on a magnetic stirrer (50 °C, 400 rpm). These solutions were mixed together. The pH of the mixture was raised to 11 using NaOH (Merck) and stirred again (5 h, 80 °C, 700 rpm). The resulting precipitates were dried (110 °C, 24 h), then calcined (750 ° C, 2 h). The phase formation and strain created in the particles were studied by XRD (D8 ADVANCE, BRUKER, Germany) technique. FESEM (Mira 3-XMU) was used to investigate the
morphology and size of the particles, and FTIR (Shimadzu Co., Japan) was performed in order to evaluate the chemical bonds in the synthesized specimens. Also magnetic properties of the prepared samples were identified using VSM (Danesh Pajooh Co., Iran) analysis. From now on, specimens with different Pr values are introduced as Pr0 to Pr6 according to Pr content (x).

3. RESULTS AND DISCUSSION

The XRD results of the synthesized samples at 750 °C for 2 h are shown in Figure 1. It should be noted that in order to compare and evaluate the effect of adding Pr to the composition, XRD analysis of the Pr free sample is also provided.

In the sample without Pr (Pr0) (Figure 1-a), the appearance of planes (220), (311), (222), (400), (422), (511), (440), (622) and (533) according to JCPDS card 00-022-1086 confirms the formation of single-phase ZnCo ferrite with cubic spinel structure. By adding Pr to the composition equal to x=0.2 (Pr2) (Figure 1-b), it can be seen that in addition to the previously appeared peaks, other peaks are formed at other angles, which according to JCPDS 01-074-1472 belong to the new phase of Pr-Fe oxide (PrFeO3). As the amount of praseodymium increased, the intensity of the new phase peaks also increased, and the height of the initial spectra in the Pr-free sample was significantly reduced, and even many peaks were removed (Figure 1-d). In fact, the predominant phase in these specimens is PrFeO3. This means that contrary to expectations, with the presence of praseodymium in the reaction system, no solid solution of CoZnPr ferrite is formed. This result contradicts the findings reported by Panda et al. [12] that synthesized cobalt ferrite substituted with Pr4+ (CoFe2+3Pr2O4) using citrate raw materials by sol-gel method and obtained CoPr0.6Fe1.4O4 and CoPr0.4Fe1.6O4. This is probably due to the higher ionic radius of praseodymium (1.13 Å) than iron (0.67 Å) [16], which makes it difficult to place as a substituent or interstitial element within the structure.

The Bertaut method was used to calculate the amount of phases which is based on the intensity of the scattered peaks and is summarized in Table 1. It is observed that in the composition with x=0.2, equivalent to 30.94 wt. % of the new phase (PrFeO3) is formed. This amount increased to 84.23 wt. % in the composition containing x=0.6, which means a reduction of the initial phase (Co0.6Zn0.4Fe2O4) to 15.77 wt.% in the powder.

The crystallite diameters and the strain created inside the crystalline particles were obtained by Williamson-Hall equation [18-20] as follows:

\[ B \cos(\theta) = \frac{0.9 \lambda}{D} + 4 \varepsilon \sin(\theta) \]  

where, D is the grain size (Å), λ is the wavelength of CuKα (1.54056 Å), B is the broadening line at half the peak height (rad.), ε is the lattice strain, and θ is the Bragg angle of the scattered peak.

The results of the Williamson-Hall equation for the synthesized samples are summarized in Table 2.

It is observed that the sample without Pr has a crystallite diameter of 47.81 nm and a compressive strain equal to 10^-4. When praseodymium introduce to the composition, the strain introduced in the ZnCo ferrite is tensile type and this strain increases with increasing amount of Pr element. Due to the increase in tensile strain, the diameter of its crystallites also decreases further. On the other hand, the strain created in the Pr-Fe oxide phase is compressive type. As a result, the diameter of its crystallites increases. The diagram of changes in the crystallite diameters in terms of the Pr value for each of the phases is given in Figure 2.

The lattice parameter of the synthesized powders was also obtained using extrapolate function \( \cos^2(\theta)/\sin(\theta) \) [21] (Table 3).

The chemical bonds in the synthesized specimens are shown in Figure 3. The vibrational modes appeared at 450 cm^-1 and 590 cm^-1 are related to M-O bonds in the B and A spaces of the ferrite phase, respectively. The stretching modes at 1650 cm^-1 and 3450 cm^-1 are due to hydroxyl groups which can help to establish the surface bonds of the particles. The frequency band around 2350 cm^-1 is attributed to C-H bond. Weak bands in the range 1250-1700 cm^-1 are also related to \(-\text{C}=\text{C}-\) bonds [22-25].

**TABLE 1.** The relative intensity of the spectra appearing (110) and the amount of phases formed in the synthesized powders

| Sample                  | Pr0 | Pr2 | Pr4 | Pr6 |
|-------------------------|-----|-----|-----|-----|
| Relative intensity (I110) of Co0.6Zn0.4Fe2O4   | 100 | 100 | 31.83 | 18.72 |
| Relative intensity (I110) of PrFeO3             | 0   | 44.81 | 100 | 100 |
| I110/ICuKα                                      | 0   | 0.45 | 3.14 | 5.34 |
| PrFeO3 (Wt. %)                                  | 0   | 30.94 | 75.86 | 84.23 |
| Co0.6Zn0.4Fe2O4 (Wt. %)                         | 100 | 69.06 | 24.14 | 15.77 |
The FESEM images of the synthesized samples are shown in Figure 4. It is seen that the morphology of the particles with the increase of praseodymium for content of x=0.4 and 0.6 is deviates from the spherical state and appears as a plate like. The non-uniformity in morphology and particle size is especially evident in Figure 4-c. The larger particles seems to be related to the Pr-Fe oxide phase, which appears mainly in the form of plates with the thickness about less than 30 nm and diameter approximately more than 100 nm. EDS analysis of samples and the weight percent of the elements are presented in Figure 5 and Table 4, respectively.

natural that by substituting Pr instead of iron in the composition and increasing its concentration, the presence of this element also increases and the concentration of Fe decreases. The concentrations of other elements (Co, Zn, O) are also changed by maintaining their atomic ratio in the composition. The extracted data are close to the stoichiometric state of the desired compounds.

Magnetic properties of the synthesized samples were investigated using VSM analysis (Figure 6) and the results are given in Table 5.
The composition without the element praseodymium (cobalt-zinc ferrite; Pr0) has excellent magnetic properties, so that its saturation magnetization and coercive force at 10^4 Oe are 100.34 emu/g and 150 Oe, respectively. By adding the element Pr to the composition, the magnetic properties are drastically reduced and the sample becomes a completely non-magnetic material. The question is, what is the reason for these drastic changes? It should be noted that if the powders were a single phase (as in the Pr0 sample), since this phase (ZnCo ferrite) is a magnetic material, changes in magnetic properties could be attributed to changes in microstructure such as crystallite diameter and particle size. In this case, due to the decrease in crystallite diameter (Table 2) and particle size (Figure 4), it was expected that the Ms would decrease and the Hc would decrease.

### Table 4. Weight percent of elements in synthesized samples containing various praseodymium

| Sample | O    | Fe   | Co   | Zn   | Pr  |
|--------|------|------|------|------|-----|
| Pr2    | 32.30| 31.09| 13.47| 8.81 | 14.32|
| Pr4    | 29.07| 24.78| 12.39| 8.04 | 25.72|
| Pr6    | 29.64| 18.73| 10.88| 7.58 | 33.16|

### Table 5. Magnetic properties obtained from the hysteresis loops in terms of different concentrations of the Pr element

| Magnetic properties | Pr0 | Pr2 | Pr4 | Pr6 |
|---------------------|-----|-----|-----|-----|
| M (emu/g) at (3000 Oe) | 91.54 | 28.22 | 4.58 | 0.43 |
| M (emu/g) at (4000 Oe) | 94.16 | 29.26 | 5.31 | 0.56 |
| M (emu/g) at (5500 Oe) | 96.48 | 30.37 | 6.23 | 0.76 |
| M (emu/g) at (7000 Oe) | 98.01 | 31.32 | 6.99 | 0.94 |
| M (emu/g) at (8500 Oe) | 99.18 | 32.03 | 7.67 | 1.12 |
| Ms (emu/g) at (10000 Oe) | 100.34 | 32.72 | 8.34 | 1.31 |
| Hc (Oe) | 150 | 0 | 0 | 0 |
| Mr (emu/g) | 12.16 | 0.07 | 0.03 | 0.01 |
| Mr /Ms | 0.121 | 0.002 | 0.004 | 0.008 |
increase. However, we see that the coercive forces are completely eliminated and are almost zero, and the saturation magnetization of the samples are very negligible, especially for composition containing Pr equal to $x=0.6$ (1.31 emu/g). Therefore, it is concluded that this change is due to the magnetic nature of the powder and its conversion to a non-magnetic material with the formation of PrFeO$_3$ phase, and not the microstructure. However, the microstructure of the samples, as shown in Figure 4, had non-uniform particles containing ZnCo ferrites and Pr-Fe oxides which made effect on the magnetic properties.

4. CONCLUSIONS

Contrary to the initial predictions of solid solution formation with the entry of the element praseodymium into the crystal lattice, in the present study this does not happen at least for Pr values higher than $x=0.2$ and a secondary phase of PrFeO$_3$ forms next to the primary ZnCo ferrite phase. By increasing the presence of Pr in the composition, especially at $x=0.6$, about 84% by weight of the PrFeO$_3$ phase is formed and a completely non-magnetic material with zero coercive force and almost negligible saturation magnetization (1.13 emu/g) is produced. It has no positive effect on improving magnetic properties. However, the formation of plate like particles with the thickness about less than 30 nm can be seen in the sample containing Pr with $x=0.6$. Due to the magnetic properties of the Pr-doped sample with $x=0.2$, it seems that if smaller amounts of this additive are used in the initial compositions, it is possible to create a solid solution of the initial ferrite and improve the microstructure because of particle size control.

5. ACKNOWLEDGEMENTS

The authors would like to acknowledge Department of Materials Engineering, Babol Noshirvani University of Technology.

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**Persian Abstract**

چکیده

به‌منظور بررسی اثر Pr بر روی خواص برخی از ترکیب فلزی (Me-Zn)PrO 
ارو در ترکیب اولیه (Me=Fe,Co) از اکسید Fe,Co,PrO 
می‌توان استفاده از ماده اولیه نیترات و با استفاده از فناوری اکسیداسیون میکرووکوئستی فنی (FTIR) و ترکیب نمونه ارتقاء (VSM) در فاصله 300-1000 nm شیشه برای تغییرات در شکل و اندازه ذرات سنتز شده و تأثیر این تغییرات بر مقاومت ارتعاشی نمونه را ارزیابی کرد. نتایج نشان داد که با افزودن Pr به ترکیب اولیه ترکیب فلزی تغییراتی را باعث کرده که در شکل و اندازه ذرات سنتز شده با افزودن Pr به ترکیب اولیه ترکیب فلزی تغییراتی را باعث کرده که در شکل و اندازه ذرات سنتز شده با افزودن Pr به ترکیب اولیه ترکیب فلزی تغییراتی را باعث کرده که در شکل و اندازه ذرات سنتز شده با افزودن Pr به ترکیب اولیه ترکیب فلزی تغییراتی را باعث کرده که در شکل و اندازه ذرات سنتز شده با افزودن Pr به ترکیب اولیه ترکیب فلزی T1021/11014.