

Short Communication

Optical, Thermal and Magnetic Properties of Strontium Ferrite Nanoparticles

Sigamani Saravanan^{1,*}, Tamilarasan Sivanandan² and Gopal Ramalingam³

¹Department of Nanotechnology, Swarnandhra College of Engineering & Technology, Narsapur-534 280, West Godavari, Andhra Pradesh, India

²Department of Physics, Indian Arts & Science College, Tiruvannamalai (TN), India

³Department of Nanoscience and Technology, Alagappa University, Karaikudi (TN), India

(*) Corresponding author: shasa86@gmail.com

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Abstract

This study endeavors to investigate the influence of calcination temperatures (650, 750 & 850°C) on the strontium ferrite (SrFe₂O₄) nanoparticles synthesized by the co-precipitation method. The prepared powder samples were characterized by various measurement techniques such as X-ray diffractometer (XRD), scanning electron microscopy (SEM), thermo gravimetric analysis (TGA), and vibrating sample magnetometer (VSM). Initially, the XRD patterns were confirmed the presence of spinel SrFe₂O₄ phases. Overall, the number of diffraction peaks increased due to the enhancement of calcination temperature. The SEM morphological features are shown the spherical-shaped nanoparticles with less agglomeration. Considerably, the agglomeration between the nanoparticles increased due to the higher calcination temperatures. However, the structural and morphological investigation was helpful and carried out for the TGA and VSM investigation. At 850°C calcination temperature, TGA revealed 5.8% of weight loss and VSM endorsed the magnetic properties such as high saturation magnetization (M_s), remanent magnetization (M_r) and coercivity (H_c) come out to be 37.26 emu/g, 19.788 emu/g and 6188.4 Oe, respectively.

Keywords: SrFe₂O₄, Nanoparticles, Calcination, Thermal, Coercivity.

1. INTRODUCTION

The strontium ferrite (SrFe₂O₄) materials have great fundamentals, an abundance of raw material, low manufacturing cost, and stable properties that are technologically applied in permanent material applications such as photocatalyst (biodiesel), optical, electronic and magnetic (regarding media) properties [1-4]. Also, the ferrites are classified into two different types like soft and hard magnets. The magnetic property of nanoscale materials has been depending on the pH, particle size, precursor, solvents, and the amount of interactions, calcination (or preparation) temperatures and preparation of methods. The various synthesis and deposition techniques including the co-precipitation, salt-melt, sonochemistry, micro emulsion,

hydrothermal, sol-gel, sol-gel auto-combustion, glass crystallization, the mechanical alloying methods and ultrasound-assisted synthesis methods have been successfully used to develop the efficient strontium ferrite nanostructures [5-14]. Compared to the above methods, the co-precipitation method is simple, promising and easy to synthesis ferrite nanoparticles. The solvent plays a pivotal role and is studied by various solvents in the preparation of ferrite nanostructures like acetonitrile, formamide, N, N-dimethyl formamide, water, ethylene glycol and glycerol, dimethylacetamide, aqueous NH₃, (NH₄)₂CO₃, NH₄Ac, NH₄HCO₃ and N-methyl acetamide [15]. The co-precipitation method is well-

established and used to synthesis important oxide materials due to its simple and good control of grain size. Ghahfarokhi et al. (2020) prepared 10-25 nm size of SrFe_2O_4 nanoparticles using sol-gel method using ethylene glycol as a surfactant. X-ray diffraction (XRD) pattern revealed the cubic spinel- SrFe_2O_4 structures and field emission scanning electron microscopy (FESEM) morphological image confirmed the reduced particle size by the addition of ethylene glycol surfactant. The ultraviolet-visible spectroscopy and vibrating sample magnetometer (VSM) results depict a wider energy gap and improved magnetic properties due to the increment of surfactant addition [16]. Zafar et al. (2018) synthesized SrFe_2O_4 nanoferrites and SrFe_2O_4 /ground eggshell nanocomposites using micro emulation and solvothermal methods. The prepared samples were utilized as an efficient adsorbent for anionic dye, eriochrome black T, and cationic dye methylene blue. The SrFe_2O_4 nanoferrite and SrFe_2O_4 /ground eggshell nano-composites powders were tuned and proved as efficient adsorbents [17]. Yasmin et al. (2018) investigated the structural and magnetic properties of Cr-doped SrFe_2O_4 clusters using the sol-gel auto-combustion method. XRD pattern reveals that crystallite size and lattice parameters were improved with the effect of Cr-substitution increment. SEM micrographs showed inhomogeneous grains with agglomeration and VSM studies confirmed the soft magnetic properties [18]. Zhang et al. (2016) investigated the SrFe_2O_4 /reduced graphene oxide (RGO) composite prepared using an ultrasonic-assisted sol-gel process and studied the photocatalytic performance. The experimental studies showed great applied potential in environmental purification due to its improved photocatalysis, easy separation and good reusability [19]. Ali Ghasemi (2012) prepared the SrFe_2O_4 nanoparticles and thin films (on a silicon wafer) using sol-gel and thermal oxidation methods. The

narrow size particle distribution with an average particle size of 50 nm was confirmed by transmission electron microscopy (TEM). Pardeshi and Pawar (2011) synthesized spinel-type SrFe_2O_4 catalyst using citrate gel method. They have investigated the optimization and effects of different environmental conditions on styrene conversion and product distributions. After the 18 hrs chemical reaction, the maximum benzaldehyde yield was received at 70°C using water solvent over 0.1 g of catalyst [20].

In this study, SrFe_2O_4 nanoparticles were synthesized via the chemical co-precipitation method. The influences of different calcination temperatures (650 , 750 , and 850°C) were investigated on the SrFe_2O_4 nanoparticles by their structural (XRD), morphological (SEM), and magnetic (VSM) properties. The required synthesis materials and preparation methods are discussed in section 2 and the results are discussed in section 3. Finally, summarize the work in section 4.

2. EXPERIMENTAL APPROACH

2.1. Materials

The analytical grade raw materials were accurately weighted for the preparation of SrFe_2O_4 nanoparticles. Initially, strontium nitrate ($\text{Sr}(\text{NO}_3)_2$) and ferric nitrate ($\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$) are the primary sources of the materials. Next, ethylene glycol, distilled water, and ethanol procured to carry out the experimental work.

2.2 Synthesis & Characterizations

The SrFe_2O_4 nanoparticles were synthesized by using the co-precipitation method as depicted in Figure 1. Strontium nitrate and ferric nitrate are used as the primary source materials. Strontium nitrate is used for strontium species and ferric nitrate is used for iron species. Additionally, ethylene glycol was used chemicals for the synthesis as a surfactant and solvent. Among various solvents (methanol, formamide, ethylene glycol,

dimethyl acetamide, glycerol etc.), mostly ethylene glycol (C₂H₆O₂) is one of the best due to their formation of inter- and intramolecular hydrogen bonds and self-assembly of surfactants [15]. The coprecipitation method is simple to synthesize SrFe₂O₄ nanoparticles. All starting precursors were high purity compounds. First, 0.846 g Sr(NO₃)₂ and 1.6 g of Fe(NO₃)₂·9H₂O with Sr:Fe, atomic ratio of 1:2 were dissolved in ethylene glycol by maintaining gentle heat and stirring. Using a magnetic stirrer, the solution stirred for 2 hrs at room

temperature. Finally, observed gelatinous precipitates were washed by several times using distilled water and ethanol. Further, the obtained precipitates were dried at room temperature. Thereafter, the dried powders were calcinated (muffle furnace) at 650, 750 and 850°C for 2 hrs. Ultimately, the prepared SrFe₂O₄ nanopowder was collected when the temperature cooled down to the room temperature. The following chemical reaction occurred during the synthesis process is given below,

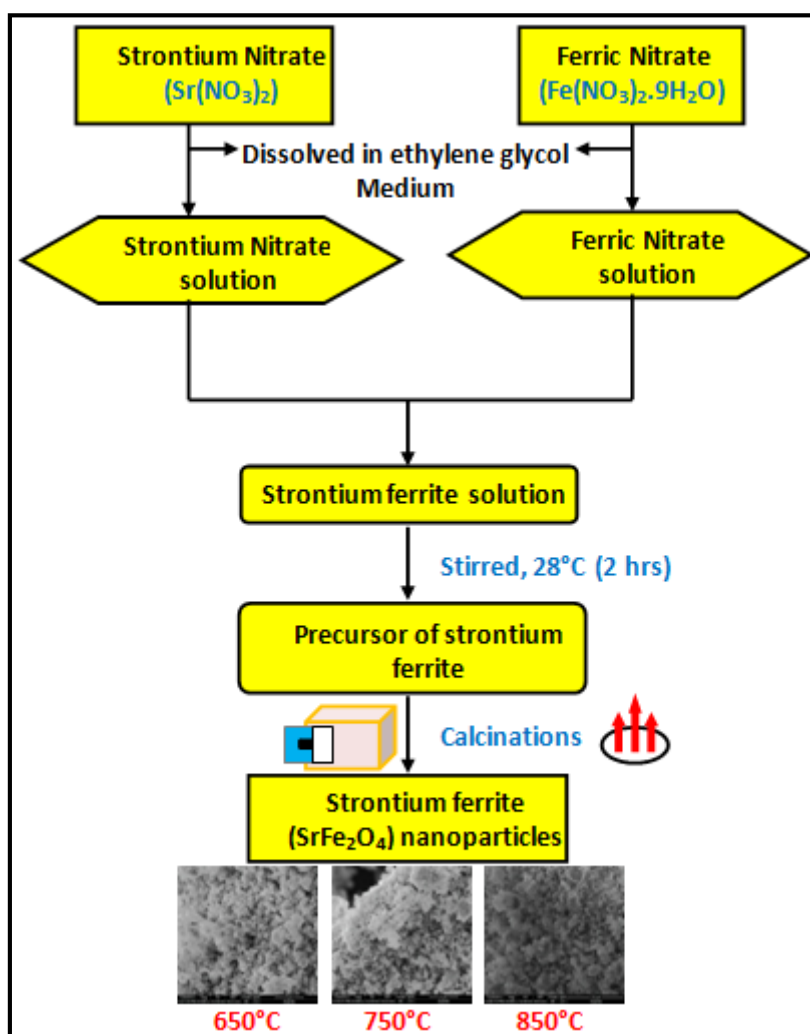
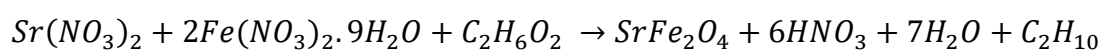


Figure 1. The procedural steps of SrFe₂O₄ nanoparticles with different calcination temperatures.

After the preparation of the samples, tested for their structural, functional, magnetic, and morphological properties of the SrFe₂O₄ nanoparticles by using an X-ray diffractometer (Rigaku Ultima III, Japan), Fourier transform infrared (Perkin Elmer-LS 45, USA) spectroscopy, vibrating sample magnetometer (Lakeshore, USA; Model 7404) and scanning electron microscopy (FBI Quanta).

3. RESULTS AND DISCUSSION

3.1. Results of X-ray Diffraction (XRD)

Figure 2 (a)-(c) shows the XRD patterns of prepared samples were calcinated at 650, 750, and 850°C. The sharp and relative intensities of all diffraction peaks of SrFe₂O₄ nanoparticles are well aligned with the (002), (311), (331), (420), (520), and (440) planes matching with the standard JCPDS card no. 00-01-1027. It is indicated that the diffraction peaks are

indexed by the formation of spinel SrFe₂O₄ structure [21-23]. With the influence of enhanced calcination temperatures, the intensity of diffraction peaks increased obviously and no other impurity peaks were presented. The average crystallite size can be calculated using the Scherrer's equation as follows,

$$D = \frac{K\lambda}{\beta \cos\theta}$$

Where 'D' is the crystalline size of the particles, 'K' is a constant (k=0.89), 'λ' is the wavelength of Cu Kα radiation, 'β' is the full-width half-maximum (FWHM) of the peak, and 'θ' is Bragg's angle. Further, the lattice constant 'a', 'b' & 'c' is calculated such as 8.03, 18.02 and 5.45 Å° were similar to the M-type strontium ferrite phase and evidenced the presence of materials [24]. Kahlenberg (2001) and Berthet (1992) reported the same crystallographic data of SrFe₂O₄ phases and compared them [25-26].

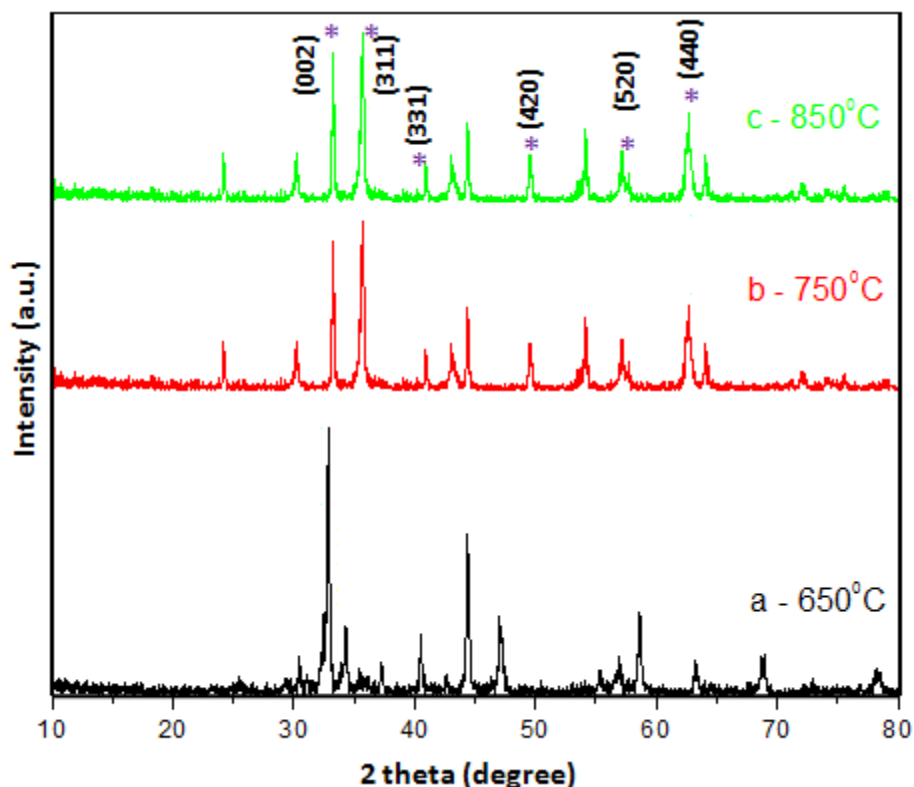


Figure 2. XRD spectrum of SrFe₂O₄ nanoparticles a) 650°C, b) 750°C and c) 850°C.

3.2. Results of Scanning Electron Microscopy (SEM)

Figure 3(a)-(c) shows the surface morphology of the various calcined

SrFe₂O₄ nanoparticles were explored by SEM. Figure 3(a)-(c) shows the spherical nanoparticles with the agglomerated nanoparticles due to magnetic properties among the particles. With the effect of calcination temperature, observed the same nature (size) of the particles with some agglomeration. Usually, the magnetic nanoparticles are agglomerated with the effect of dipole-dipole interaction between the nanoparticles. This interaction generates a negative effect on the various applications. The different morphological investigations employed by Zhang et al.

(2016) [19]. The agglomerates could be reduced by covering the particle surface with surfactants (non-magnetic/non-metallic compounds). Surfactant materials include hydrophilic head and hydrophobic tails [27-31]. Larger particles, as well as large pores, were located at the agglomeration junctions. Here, ethylene glycol acts as a solvent as well as a surfactant due to the size of the nanoparticles decreased. However, the nanoparticles size enhanced with water solvent based strontium ferrite nanoparticles as reported [32].

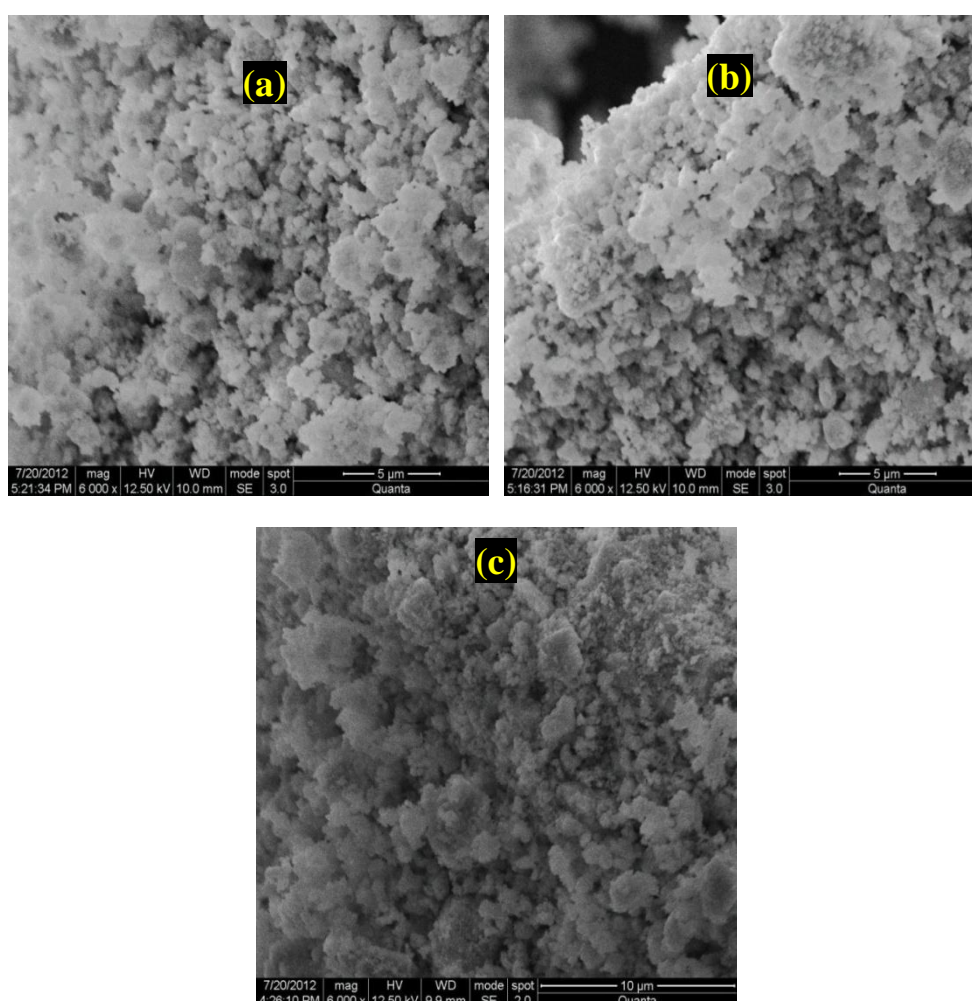


Figure 3. SEM images of SrFe₂O₄ nanoparticles 650°C (a), 750°C (b) and 850°C (c).

Natarajan and Wang (2000) reported that the aggregation (cluster) performance is due to the selection of surfactant, solvent dielectric constant, and less magnitude of

the hydrocarbon mixed solvent interfacial tension [15]. Sivanadhan (2021) reported highly agglomerated SrFe₂O₄ nanoparticles (size, 112-130 nm) with the effect of water

medium [32]. Among various solvents, ethylene glycol is a suitable choice for the preparation of ferrite nanoparticles.

3.3. Results of Thermogravimetric Analysis (TGA)

The weight loss (%) as a function of temperature using thermogravimetric analysis (TGA) is shown in Figure 4. It is the TGA analysis curve of the SrFe_2O_4 nanoparticles which is calcinated at 850°C . This curve shows four distinct steps of weight loss.

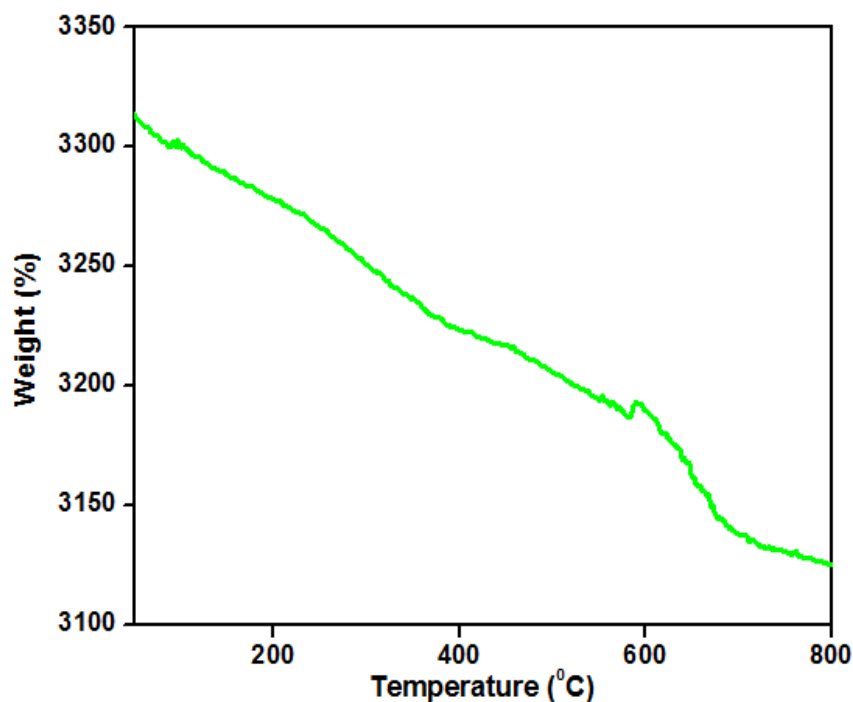


Figure 4. TGA curve of as-prepared SrFe_2O_4 nanoparticles (850°C).

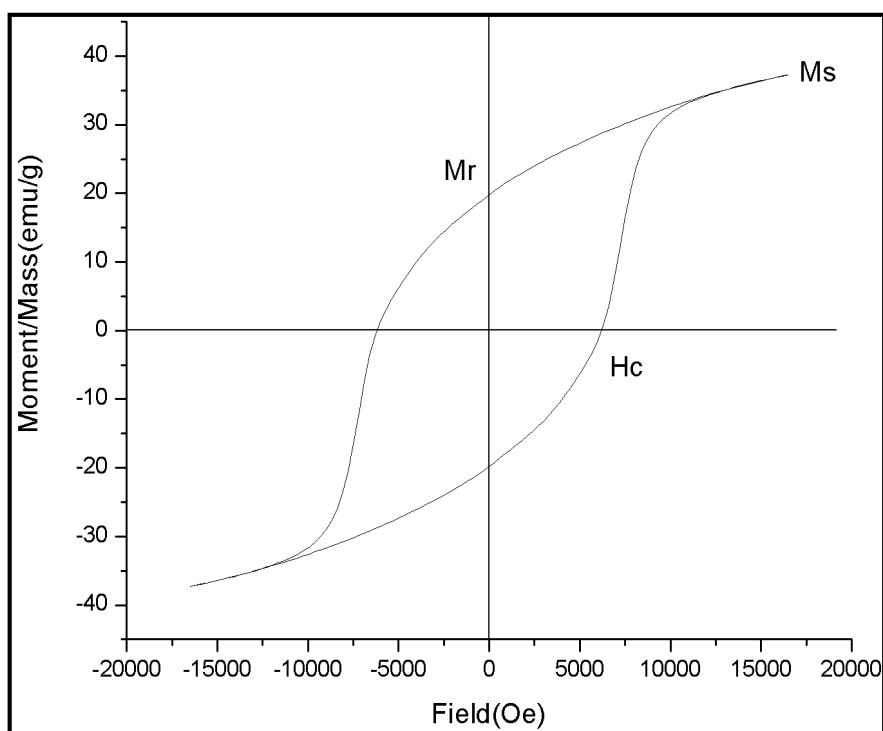


Figure 5. Hysteresis loop of SrFe_2O_4 nanoparticles at 850°C calcination temperature.

The first step weight loss is between 50 to 100°C due to adsorptions of water molecules and the second step the weight loss between 350 to 400°C which is due to hydroxyl ions (O-H) [33]. The weight loss of 550 to 580°C and 640 to 710°C due to the decomposition of nitrates and metal nitrate. Saleh et al. reported the formation of strontium oxide phases depending on the reaction temperature and the molar ratio among the reactants. The monoferrite (SrFe₂O₄) formation obtained from 592 to 765°C [34-35].

3.4. Results of Vibrating Sample Magnetometer (VSM)

The magnetic properties are saturation magnetization (M_s), remanent magnetization (M_r), and coercivity (H_c) increasing with the effect of calcination temperatures.

Table 1. The list of coercivity of strontium ferrite nanoparticles.

Temp. (°C)	Nanoparticles	Coercivity (Oe)	Solvent	Ref.
850	SrFe ₂ O ₄	5662	Water	[32]
1000	SrFe ₁₂ O ₁₉	4900	Water	[5]
800		5000		
700	SrFe ₂ O ₄	299	Water	[19]
950	SrFe ₁₂ O ₁₉	3500	Tween-80, DI Water	[33]
1100		2375		
700	SrFe ₂ O ₄	709	DI water	[21]
		453		
		440		
		15		
		137		

These magnetic properties are depending on the size of the nanoparticles, magneto-crystalline anisotropy, preparation/ calcination temperatures, the effects of tilting, the distribution of cations, and the degree crystallinity as reported by Ghahfarokhi

and Shobegar (2020) [21]. Consequently, the magnetocrystalline value is increasing with the effect of the highest calcination temperature [5]. Here, VSM studies revealed the hysteresis curve of SrFe₂O₄ nanoparticles prepared under the conditions of Fe³⁺/ Sr²⁺ mole ratio and calcination temperature which was calcinated at 850°C as shown in Figure 5. The improved coercivity (6188.4 Oe), remanent magnetization (19.788 emu/g), and the saturation magnetization (37.26 emu/g) reached a maximum as compared to water solvent [32]. Vogel and Evans (1979) investigated the SrFe₂O₄ composition and improved the complex chemical and physical interactions that could lead the superior magnetic properties [36]. The coercivity of different strontium ferrite, and calcination temperatures as compared (Table-1). Accordingly, coercivity, remanent magnetization and saturation magnetization of SrFe₂O₄ nanoparticles can be useful in various applications. The spinel strontium ferrites were found as highly active in catalytic, permanent magnetic, hard magnetic devices, and microwave applications [19, 37-39]. The magnetic properties of nanoparticles are varied by size which could play significant role in physics [40-44]. By comparing other methods [40], this co-precipitation method is a good candidate for the preparation of nano material structures because it is a relatively easy, low-cost and reproducibility.

4. CONCLUSION

In summary, the strontium ferrite (SrFe₂O₄) nanoparticles were synthesized by the co-precipitation method. The primary source materials are strontium nitrate and ferric nitrate. The prepared

samples were calcinated at three different temperatures. The spinel SrFe_2O_4 phase was confirmed with the XRD diffraction spectra. SEM morphological investigation revealed small spherical-shaped nanoparticles, and some agglomeration. The average particle size of 117, 24, and 19 nm corresponds to the calcination temperature of 650, 750, and 850°C. The reduced thermal properties were noticed with the effect of the

highest calcination temperature by the TGA. The VSM magnetic study indicated magnetic properties such as the H_c (6188.4 Oe), H_s (37.26), and M_r (19.78). Accordingly, we studied spinel SrFe_2O_4 nanoparticles that can be tested for various applications such as a photocatalyst.

CONFLICT OF INTEREST

The authors declare that they have no conflict of interest.

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