Review

Recent Advances in Synthesis and Applications of MFe$_2$O$_4$ (M = Co, Cu, Mn, Ni, Zn) Nanoparticles

Thomas Dippong $^1$, Erika Andrea Levei $^2$,* and Oana Cadar $^2$,*

$^1$ Faculty of Science, Technical University of Cluj-Napoca, 430122 Baia Mare, Romania; thomas.dippong@cunbm.utcluj.ro
$^2$ National Institute for Research and Development for Optoelectronics INOE 2000, Research Institute for Analytical Instrumentation Subsidiary, 400293 Cluj-Napoca, Romania; erika.levei@icia.ro
* Correspondence: oana.cadar@icia.ro

Abstract: In the last decade, research on the synthesis and characterization of nanosized ferrites has highly increased and a wide range of new applications for these materials have been identified. The ability to tailor the structure, chemical, optical, magnetic, and electrical properties of ferrites by selecting the synthesis parameters further enhanced their widespread use. The paper reviews the synthesis methods and applications of MFe$_2$O$_4$ (M = Co, Cu, Mn, Ni, Zn) nanoparticles, with emphasis on the advantages and disadvantages of each synthesis route and main applications. Along with the conventional methods like sol-gel, thermal decomposition, combustion, co-precipitation, hydrothermal, and solid-state synthesis, several unconventional methods, like sonochemical, microwave assisted combustion, spray pyrolysis, spray drying, laser pyrolysis, microemulsion, reverse micelle, and biosynthesis, are also presented. MFe$_2$O$_4$ (M = Co, Cu, Mn, Ni, Zn) nanosized ferrites present good magnetic (high coercivity, high anisotropy, high Curie temperature, moderate saturation magnetization), electrical (high electrical resistance, low eddy current losses), mechanical (significant mechanical hardness), and chemical (chemical stability, rich redox chemistry) properties that make them suitable for potential applications in the field of magnetic and dielectric materials, photoluminescence, catalysis, photocatalysis, water decontamination, pigments, corrosion protection, sensors, antimicrobial agents, and biomedicine.

Keywords: transition metal; ferrites; magnetic nanoparticles; synthesis; applications

1. Introduction

Nanocrystalline magnetic materials have attracted considerable interest due to their uniqueness and remarkable properties in various fields including physics, chemistry, biology, medicine, materials science, and engineering. Compared to their bulk counterparts, nanomaterials have particle size in the 1–100 nm range and a high surface to volume ratio that determines different or enhanced reactivity, thermal, mechanical, optical, electrical, and magnetic properties [1]. While in the case of bulk materials, the chemical composition is the main factor that determine their properties, in the case of nanomaterials, besides the chemical composition, the particle size and morphology determine most of their characteristics [1,2]. Moreover, these properties can be tuned based on the particle size and chemical composition [1,2].

Ferrites are important and interesting materials, from a practical as well as fundamental point of view. Among various ferrites, nanosized CoFe$_2$O$_4$, MnFe$_2$O$_4$, ZnFe$_2$O$_4$, NiFe$_2$O$_4$, and CuFe$_2$O$_4$ have attracted considerable attention due to their chemical and thermal stability and unique structural, optical, magnetic, electrical, and dielectric properties and wide potential technological applications in photoluminescence, photocatalysis, humidity-sensors, biosensors, catalysis, magnetic drug delivery, permanent magnets, magnetic refrigeration, magnetic liquids, microwave absorbers, water decontamination, ceramics pigment, corrosion protection, antimicrobial agents, and biomedicine (hyperthermia) [1,3–8].
CoFe$_2$O$_4$ has an inverse spinel structure with Co$^{2+}$ ions mainly placed in octahedral (B) sites and Fe$^{3+}$ ions almost equally distributed between tetrahedral (A) and octahedral (B) sites. It presents large coercivity ($H_C$), high magnetocrystalline anisotropy constant ($K$), high Curie temperature ($T_C$), moderate saturation magnetization ($M_S$), low remanent magnetization ($M_R$), excellent chemical and mechanical stability, rich redox chemistry, large magnetostrictive coefficient ($\lambda$), high electrical resistance, low eddy current losses, significant mechanical hardness, and low toxicity [7,9–13]. The synthesis of CoFe$_2$O$_4$ with different structures (i.e., nanoparticles, hollow mesoporous nanospheres, nanorods and three-dimensional ordered macroporous structure) have been reported [9,10,13]. Beside many advantages, CoFe$_2$O$_4$ owns poor electrical conductivity, poor cyclic stability, structural strain, and large volume expansion [9,10,14].

ZnFe$_2$O$_4$ has a normal spinel structure, where Zn$^{2+}$ ions preferably occupy the tetrahedral (A) sites and Fe$^{3+}$ ions the octahedral (B) sites [6,15,16]. The absence of Fe$^{3+}$ in the A sites results in weak antiferromagnetic exchange interactions within Fe$^{3+}$ in B sites, making ZnFe$_2$O$_4$ antiferromagnetic below 9 K [15–17]. The enhanced magnetization originates from the super-exchange interaction ascribed to the inversion of Fe$^{3+}$ and Zn$^{2+}$ ions in the tetrahedral (A) and octahedral (B) sites [18].

NiFe$_2$O$_4$ possesses an inverse spinel structure, where the tetrahedral (A) sites are occupied by Fe$^{3+}$ ions, while the octahedral (B) sites by Fe$^{3+}$ and Ni$^{2+}$ ions, the Fe$^{3+}$ ions being easily distributed between the A and B sites [19,20]. NiFe$_2$O$_4$ is one of the most versatile and technologically important soft ferrite materials due to its low electrical conductivity, high electrochemical stability, catalytic behavior, abundance in nature, low $K$ and $H_C$ values, high $M_S$, paramagnetic, superparamagnetic or ferrimagnetic behavior depending on the particle size and shape, low eddy current loss and conductivity, high electrical resistance, and electrochemical stability [4–6,11,21].

CuFe$_2$O$_4$ has an inverse spinel structure with 8 Cu$^{2+}$ ions on octahedral sites and 16 Fe$^{3+}$ ions equally distributed between the tetrahedral (A) and octahedral (B) sites [22,23]. The Cu$^{2+}$ ion displays the transition from tetragonal phase, technologically less important (at low temperature) to cubic phase, which is relatively more useful (at high temperature) [24,25]. Moreover, magnetic and electrical properties of CuFe$_2$O$_4$ vary significantly with the change of cation distribution [25]. CuFe$_2$O$_4$ exhibits ferrimagnetism, high thermal stability, high resistance to corrosion, excellent catalytic properties, and sufficient band gap, acting as an efficient photocatalyst [22–24,26,27].

Manganese ferrite (MnFe$_2$O$_4$) has a partially inverse spinel structure, fine structural, magnetic and electrical properties, with 20% of Mn$^{2+}$ ions located at octahedral (B) sites and 80% located at tetrahedral sites [3,8]. MnFe$_2$O$_4$ nanoparticles (NPs) has attracted attention in biomedicine due of its good biocompatibility, controllable size, high magnetization value, superparamagnetic nature and ability to be monitored by an external magnetic field. Moreover, MnFe$_2$O$_4$ is an inorganic heat-resistant, non-corrosive, environmentally friendly, non-toxic, high shock resistant [28], reusable adsorbent, often employed for adsorption and desorption processes [29] and magnetic drug targeting/delivery [3].

The surface coating of magnetic NPs is required in order to generate non-toxic, biocompatible and water dispersible NPs, along with drug targeting ability. The most used surface coating materials are poly(vinylalcohol), poly(N—isopropylacrylamide), polyethylene glycol (PEG), chitosan, Au, ZrO$_2$, and SiO$_2$ [3]. Of these, due to the excellent drug loading capacity, stability, non-toxicity, biocompatibility, and water dispersibility, the mesoporous SiO$_2$ is a multipurpose candidate for the development of nanocarriers, as it enhances the stabilization of ferrite NPs in water, improves the chemical stability, and minimizes the agglomeration of NPs, without influencing their magnetic and dielectric properties [30]. Non-magnetic SiO$_2$ can easily promote conjugation with many functional groups, thus allowing selective and specific coupling and labeling of biotargets. Moreover, SiO$_2$ coatings may change the surface properties of magnetic NPs and offer a chemically inert layer, which is particularly useful in biological systems [30,31].
By searching in the Web of Science Core Collection the keyword “M ferrite”, where M = cobalt, copper, nickel, manganese, and zinc, we observed that Co ferrite attracted the attention, with the number of publications on Co-ferrite exceeding by far those published on the other ferrites. Between the studied ferrite, Co-ferrite was the topic of 4276 papers, followed by Zn-ferrite (3073), Ni-ferrite (2432), Mn-ferrite (895), and Cu-ferrite (880). For every ferrite, the number of papers started to grow exponentially in the last 20 years (Figure 1), the highest increasing rate being observed for the Co-ferrite. The increasing interest in these ferrites may be attributed both to the development of new equipment that allowed the ferrites characterization and the increase of the demand for materials with special properties for a wide range of applications.

Figure 1. Papers having M-ferrite (M = Co, Cu, Ni, Mn, Zn) in the topic published in Web of Science Core Collection between 1975–June 2021.

In the last few years, the number of papers that review data on nanosized ferrites with a focus on different ferrites, different synthesis methods, or different applications significantly increased. The Web of Science Core Collection have indexed 58 publication that contains the words “ferrite” and “review” in their title of which 25 were published between 2019 and June 2021.

Zate et al. [32] reviews the mechanical, chemical, spray and electrospinning methods used for the synthesis of different ferrites with magnetic properties. Vedrtnam et al. [33] reviews the properties, classification, synthesis, and characterization of hexagonal and spinel ferrites with a focus on the main four synthesis routes (sol-gel, hydrothermal, co-precipitation and solid-state), magnetic properties, and characterization of the ferrites. Vinosha et al. [34] review the recent advances of synthesis, magnetic properties, and water treatment applications of cobalt ferrite, while Masunga et al. [35] reviews the recent advances in copper ferrite synthesis, magnetic properties and application in water treatment. Kumar et al. [36] review magnetic nano ferrites and their composites used in the treatment of pollutants from waste waters, emphasizing pros and cons of several synthetic pathways, the adsorption mechanism, and ferrite regeneration, while Kefeni and Mamba [37] review the photocatalytic application of spinel ferrite NPs in pollutant degradation with emphasis on the possible recovery and reuse of NPs. Kharisov et al. [38] review the use of cobalt, nickel, copper, and zinc ferrites and their doped derivates as catalysts in organic processes, while Dalawai et al. [39] overview the spinel-type ferrite thick films together with preparation strategies and sensors, microwave, magnetic, and advanced applications. Kefeni et al. [40] review the ferrite’s applications in electronic devices, such as sensors and biosensors, microwave devices, energy storage, electromagnetic interference shielding, and high-density recording media together with the advantages and drawbacks of most important ferrite NPs synthesis methods.

Our previous works reported the thermal, structural, morphological, and magnetic characterization of ferrites ($MFe_2O_4$, M = Co, Mn, Zn, Ni and Cu) and doped ferrites with different divalent transition metals, produced by sol-gel synthesis [41–46]. Furthermore,
since the physical, chemical, magnetic, electrical, and optical properties can be tailored by the dopant type and content, we also reviewed the potential applications of $\text{CoFe}_2\text{O}_4$ and divalent transition metal-doped cobalt ferrites ($\text{M}_x\text{Co}_{1-x}\text{Fe}_2\text{O}_4$, $\text{M} = \text{Zn}, \text{Cu}, \text{Mn}, \text{Ni}$, and Cd) [47]. In addition, this paper aims to deepen our previous studies and review a number of topics including the various synthesis methods of $\text{CoFe}_2\text{O}_4$, $\text{MnFe}_2\text{O}_4$, $\text{ZnFe}_2\text{O}_4$, $\text{NiFe}_2\text{O}_4$, and $\text{CuFe}_2\text{O}_4$ NPs together with their advantages and disadvantages, and the most important applications in conventional and modern technologies. The review not only summarizes the existing literature from theoretical and methodological points of view, but also synthetizes it from a new perspective, representing a significant and useful contribution to subsequent research.

2. Synthesis Methods

The chemical and physical properties of transition metal nanoferrites ($\text{MFe}_2\text{O}_4$; $\text{M} = \text{Co}, \text{Ni}, \text{Zn}, \text{Mn}, \text{Cu}$) are dependent on the synthesis method and conditions. Thus, the selection of an appropriate synthesis route plays a crucial role in tailoring the properties and obtaining high quality nanoferrites [1]. Nanosized ferrites can be synthesized by various techniques (Figure 2) and the possibility to obtain almost any solid solution of nanoferrites unlocks the way to tailor their properties for a wide range of applications [22].

![Figure 2. Classification of nanoferrite synthesis methods.](image)

There are different classification approaches of the ferrite synthesis methods: (i) physical, chemical, and biological, based on the type of processes that take place in the synthesis methods, (ii) dry and wet methods, based on the presence or absence of a solution, and (iii) conventional and non-conventional, based on their novelty. Despite several classification methods, it is difficult to univocally group the synthesis methods, as sometimes different processes are coupled to obtain NPs with specific characteristics.

Many efforts have been made to tailor the size, shape, particle size distribution, surface area, composition, structure, and properties of the ferrite NPs by employing different synthesis methods, or changing the synthesis parameters, such as the annealing temperature and duration, concentration of reactants, pH value, stirring speed, doping additives, etc. [48,49]. Numerous physical and chemical methods have been used for the synthesis of ferrites [1,19,25]. Some synthesis methods are high-energy consuming, complex procedures, demanding a high processing temperature and long reaction time to complete the crystallization, and the use of reduction agents with negative effects on the environment [6,13]. The importance of these factors is different in every method, making the achievement of reproducibility in desired
properties difficult [50]. Another issue that appears in the majority of the synthesis methods is the agglomeration of NPs after production, which limits the control of size, shape and function [51]. The wet-chemical synthesis has multiple advantages. However, single phase ferrite can be obtained only after annealing at high temperatures and is accompanied by particle growth, aggregation, and coarseness of NPs [14,20].

2.1. Dry synthesis Methods

2.1.1. Combustion Method

Combustion method is simple, fast, and inexpensive, as it does not involve intermediate decomposition or calcinations steps. This method exploits an exothermic, usually very rapid, and self-sustaining chemical reaction between metal salts and an organic fuel (glycine, urea, citric acid, sucrose, hydrazine, and polyvinyl alcohol) that act as a reducing agent [14,52]. The powder characteristics (i.e., crystallite size, surface area) are governed by the enthalpy or flame temperature produced during the combustion, which is dependent on the type of fuel and fuel-to-oxidant ratio. Beside the key role of fuel in the morphology of the NPs it also determines the phase formation [53]. An important aspect is that the heat necessary to sustain the chemical reaction is provided by the reaction itself and not by an external source [8]. Nitrates are the favorite metal precursors because of their high solubility in water and easy combustion following their mixing with a suitable fuel [7,54]. Ammonium nitrate is used as an extra oxidant in the combustion reaction, producing the expansion of the microstructures and eventually the increase of NPs surface area, without changing the proportion of the other participants [7]. The combustion method is a very popular method for the synthesis of ceramic materials, composites, and ferrimagnetic nanomaterials, due to its efficiency, short preparation time, use of relatively inexpensive precursors and simple equipment, good control of stoichiometry, desired particle size distribution, formation of high-purity products, and stabilization of metastable phases. The main disadvantages are high combustion temperature and low production yields [6,11,13,14,55]. The combustion method is a good choice for the preparation of high quality CoFe$_2$O$_4$, NiFe$_2$O$_4$ and MnFe$_2$O$_4$ NPs. By varying the nitrates to fuel ratio, the size, $M_S$ and $H_C$ values are tailored [20,56,57]. Nanocrystalline NiFe$_2$O$_4$ was successfully prepared by mixing of metal nitrates and citrate with the formation of a colloidal solution (sol), followed by the continuous heating of xerogel, the auto combustion process until the formation of a loose powder, and the annealing of the powder at 700 °C [21]. This method requires less time and produces pure and homogeneous NPs, without any type of waste product, but involves a high temperature [7,58,59]. The high purity of materials prepared by the combustion process is attributed to the removal of unwanted impurities as volatile species at high temperatures [6].

2.1.2. Solid State Synthesis

Solid state synthesis produces polycrystalline ferrite nanomaterials from solid reagents at high temperature [20]. This method assumes the grinding and mixing of metal nitrates or sulphates with NaOH or NaCl in an agate mortar or mills for short times. After the removal of NaCl by washing, the powders are dried at 80 °C for two hours and then annealed at 700 °C for two hours [60]. CoFe$_2$O$_4$ NPs have been synthesized by a low-temperature solid state process using various salts (sulphate, acetate, nitrate and chloride). By using chloride salts, the obtained nanomaterials have smaller particle size than by using other salts [61]. Furthermore, in comparison to other synthesis techniques, the low temperature solid state process requires neither complex apparatus, nor solvent or solution, and is a convenient, environment-friendly, low-cost, time-saving and low energy consumption process [62]. The CoFe$_2$O$_4$ NPs prepared using solid state technique display high $M_S$ values, as well as low $M_R$ and low $H_C$ values [63]. The major advantages are that the synthesis procedure is completed under atmospheric pressure, which is economic and feasible, as well as the use of non-expensive and toxic solvents and cost-effective raw materials [58]. Mechanical milling is a simple, low-cost, solid state and non-equilibrium process in which the final product have a very fine, typically nanocrystalline or amorphous structure. Generally, the final
product has a nanosized structure with enhanced properties and performance compared to the bulk material [15]. The main advantages of this method are simplicity, low cost and ability to produce large volumes [63]. Mechanical milling of appropriate metal oxides for 12 h and subsequent sintering at 600 °C for 2 h were applied to prepare ZnFe$_2$O$_4$ and NiFe$_2$O$_4$ ferrites [15]. CoFe$_2$O$_4$ NPs were obtained by ball milling of Fe$_2$O$_3$ and Co$_3$O$_4$ powders in stoichiometric amounts using high-energy vibratory mill for 8 h, followed by calcination at 900 °C for 12 h [64]. CoFe$_2$O$_4$ NPs of 10 nm size were also prepared by the precipitation of hydroxide/oxidhydroxide, followed by mechanical milling at lower speeds, and subsequent heat treatment for a short time. NaCl was added before milling to avoid agglomeration [62]. NiFe$_2$O$_4$ crystallites were prepared by high energy ball milling. The increase of milling time led to a significant increase in the NiFe$_2$O$_4$ formation and to a progressive decrease in its particle size and lattice parameter [20]. MnFe$_2$O$_4$ NPs (4–8 nm) were prepared by solid state ball-milling and calcinations (300–400 °C) of nitrate precursors and citric acid [65].

2.2. Wet Synthesis Method

2.2.1. Co-Precipitation Method

Co-precipitation method is one of the most widely used methods due to its high yield and simplicity in producing high purity ultrafine magnetic nanostructured ferrites [58,66]. The co-precipitation method is a very simple and cost-effective method that allows an easy control of particle size and composition, requires low temperature and leads to materials with high crystallinity, homogeneity and good textural properties [1,58]. The major drawbacks of co-precipitation methods are extensive agglomeration, poor crystallinity and particle size distribution, and the necessity of pH control [57,67].

In this case, homogeneous solutions are formed by dissolving inorganic salts (chloride, sulfate, nitrate) in water or solvents. After pH adjustment in the range of 7–12 under continuous stirring, the precipitate is collected by filtration or centrifugation, washed, and dried. The pH change rate causes the particles aggregation and crystal growth [58]. The most common way to synthesize nanostructured ferrites by chemical co-precipitation method is using Co$^{2+}$ and Fe$^{3+}$ salts in the presence of a strong base [51]. By adjusting the experimental parameters (i.e. reaction temperature and time, reagents feeding rate and concentration, pH, drying temperature, etc.), the size, shape, and magnetic properties of the nanostructures may be controlled [68].

CoFe$_2$O$_4$ NPs (2–47 nm) were synthesized from metal chloride salts using different concentration of aqueous NaOH and NH$_4$OH solution using a reaction time of 2 h and variable reaction temperature (20–100 °C). To achieve the precipitation reaction, the aqueous metal chloride solutions were slowly added into the preheated boiling aqueous alkaline solution [68]. The crystallite size depends on the reaction temperature, time, concentration of the base solution, and pH [68]. Another study reported the preparation of CoFe$_2$O$_4$ NPs (2–14 nm) by controlling the co-precipitation temperature of Co$^{2+}$ and Fe$^{3+}$ ions in alkaline solution from 20 to 80 °C [49].

The co-precipitation of CoFe$_2$O$_4$ nanocrystals by mixing Fe$^{2+}$, Co$^{2+}$ and NaOH in the presence of oxidizing agent such as KNO$_3$ to convert Fe$^{2+}$ into Fe$^{3+}$ was reported by Chia et al. [66] and Senapati et al. [69]. However, due to the coexistence of both Fe$^{2+}$ and Fe$^{3+}$ ions, besides CoFe$_2$O$_4$, secondary phase such as magnetite (Fe$_3$O$_4$) is also formed [66]. Moreover, the $H_C$ values of particles obtained by this procedure are not very high and depends on the amount of surfactant used, that favor a stable colloidal dispersion of the NPs, as well as on the annealing temperature [69].

A detailed investigation on the effect of precursors, solvents, precipitating agent, and heat treatments on the particle size, particle size distribution, morphology, and chemical composition of CoFe$_2$O$_4$ NPs synthesized at various liquid phase was reported by Prabhakaran et al. [68]. In some cases, in order to facilitate the solubilization of metal salts at low temperatures or to prevent oxidation and particle agglomeration, some surfactants such as oleic acid are added to the solution before precipitation [1].
The temperature may influence the size and stability of the NPs [50]. The particle size growth in the co-precipitation method is influenced by the difference between local and surface temperature, as the formation of spinel nano-ferrites is usually an exothermic reaction, with the latent heat being released at the surface [70]. At low temperatures particles tend to aggregate, the particle size increasing with the decrease of temperature. High purity and no aggregation of the NPs were reported at high temperatures due to directed crystallization [50].

Reverse co-precipitation is similar to conventional co-precipitation, except that instead of adding a precipitant into the precursor ions solution, the precursor ions are added into the precipitant solution. Thereby, the precipitant is in a supersaturated state, ensuring the complete precipitation. The resultant NPs have smaller particle-size than those obtained by traditional co-precipitation [71]. The synthesis of CoFe$_2$O$_4$ NPs by reverse co-precipitation was reported by Huixia et al. [58].

2.2.2. Sol-Gel Method

The sol-gel method is a low temperature process based on hydrolysis and condensation reactions of metal precursors (salts or alkoxides), leading to the formation of a three-dimensional inorganic network [72]. Sol results from the conversion of monomers into a colloidal solution, while gel is obtained after the solvent evaporation by joining together particles into a network [58]. The sol-gel method is a simple, low cost, and environmentally friendly method to prepare nanocomposites (NCs) as it allows good control of the microstructure, particle size, dispersion, structure, and chemical composition by carefully monitoring the preparation parameters [58,73,74]. The sol-gel method has been used to prepare very fine, highly dense, homogenous, and single-phase ferrite NPs. Compared to other conventional methods, the sol-gel displays a good stoichiometric control and also allows the production of ferrites at relatively low temperature [20]. The obtained nanomaterials may be formed either as films or as colloidal powders [58]. The main disadvantage is the limited efficiency and long duration of the synthesis process [32,48,58]. NiFe$_2$O$_4$ nanostructures with an average particle size of 27 nm were synthesized by sol-gel method using glycolic acid as a chelating agent [20]. In the case of CoFe$_2$O$_4$@SiO$_2$, the SiO$_2$ network protects the NPs and minimizes the surface roughness and spin disorder. The $H_C$ values at room temperature (RT) for CoFe$_2$O$_4$@SiO$_2$ were much higher than that of unembedded CoFe$_2$O$_4$ [73]. The most commonly used reagents are metal nitrates as metals source, 1,2-ethanediol, 1,3-propanediol or 1,4-butanediol as chelators, tetraethyl orthosilicate (TEOS) as matrix precursor, ethanol as solvent and HNO$_3$ [32]. By annealing, the NPs agglomerate and form larger particles. Thus, further research on the influence of thermal energy released during heat treatment on the undesired growth of particle size is needed [70].

The Pechini method is an alternative approach to sol-gel synthesis and consists in the complexing of cations with hydroxycarboxylic acids (usually citric acid or ethylenediaminetetraacetate) in an aqueous-organic medium. The obtained chelates are cross-linked and transformed into polymers using polyalcohol (ethylenediol, polyvinyl alcohol) through polyesterification. By heating, the viscous liquid is dried and forms a gel. By annealing the gel, the organic part is removed, resulting in reactive oxides, which further form the ferrites [75–77]. The microwave-assisted Pechini method combines the advantages of microwave and Pechini methods, such as low temperature and time of process, accurate control of stoichiometry, uniform mixing of various components on molecular level, low price of precursor and equipment, and capability of industrial scale-up [31,75]. However, the main drawback of this method is the long and energy-wasting thermal treatments in order to remove the large amounts of organic precursors [75]. The microwave-assisted sol-gel Pechini method is a faster, energy-saving procedure for obtaining single-phase nanopowders of high purity [75].

Combining the sol-gel and auto-combustion methods results in a simple and inexpensive preparation method for high purity, homogenous nanopowders at low annealing temperature [56,61]. The method consists in dissolving nitrate salts in water, addition of the organic complexing agent (citric acid), adjustment of the solution to pH 7, heating at
70 °C to form the sol, and then at 110 °C to remove the residual water and form the gel, and initiation of the autocatalytic self-combustion process of nitrate-fuel gel [78]. The citric acid and glucose are two of the most used fuels due to their strong complexing ability and low ignition temperature (200–250 °C) [78]. The synthesis of CoFe$_2$O$_4$ NPs in the size range of 11–40 nm by the sol-gel auto combustion method and annealing at different temperatures (800–1000 °C) was also reported [78].

The organic precursor method is similar to the Pechini method and involves the preparation of aqueous solution containing cations, chelation of cations using carboxylic acid, followed by heating of the solution until precursor formation and annealing of the precursors. The carboxylic acid acts both as complexing agent and as organic rich fuel. Magnetic CoFe$_2$O$_4$ NPs (38.0–92.6 nm) were produced by the organic precursor method using oxalic and tartaric acids as precursors [79].

2.2.3. Microwave-Assisted Combustion Method

The microwave-assisted combustion method applies microwave radiations to the reaction solution to synthesize NPs. Microwave radiation is absorbed and converted to thermal energy, the heat being generated inside a material, unlike the conventional heating methods where the heat is transferred from the outside. This heating allows for a considerable reduction of processing time and energy [13,54,80]. The main advantages of this method compared to other synthesis methods are simplicity, homogeneous nucleation, rapid reaction time, high production rates, environmentally friendly, excellent reproducibility, low energy cost, easy handling, and control of parameters. In the microwave-assisted combustion process, the reagents are mixed at the molecular level, due to the interaction of microwaves, offering an excellent control of stoichiometry, purity, homogeneity, and morphology [53,58,59]. However, the method is expensive mainly due to the expensive fuels such as urea, glycine, L-alanine, carbohydrazide, or citric acid used to promote and control the combustion process in accordance with the propellant chemistry principles, and inappropriate for scale up and reaction monitoring [13,53,80–82].

In the case of the microwave hydrothermal method, the necessary heat is generated by microwaves, which have the advantage of very fast heating to a certain depth, allowing the generation of homogenous nanomaterials with fine particle size distribution [52,53]. Zn, Fe, Mn, Cu, Ni, and Co nitrates were dissolved in deionized water, maintaining a pH equal to 9.4. The mixture was sealed with tetrafluorometoxil (TFM) and placed in a microwave oven at 160 °C for 30 min. The resulting wet mixture was dehydrated, followed by adding polyvinyl alcohol (PVA) as a binder and sintering for 30 min [58]. This method provides faster heating, is more economical, and produces very fine, uniform NPs [59].

2.2.4. Citrate Precursor Method

The citrate precursor method is a wet chemical process that involves the mixing of aqueous solutions of precursor salts (metal nitrates) with an aqueous solution of citric acid, followed by heating at about 80 °C and annealing at 700 °C [83,84]. During this process, the precursors are thermally decomposed into ferrite powders [83]. The advantage of this method is the high reactivity, low reaction time, low synthesis temperature, homogenous distributions of ions, and low cost over other chemical methods [83]. To reduce the particle agglomeration, the dispersion of NPs in SiO$_2$ matrix is a commonly used. The synthesis of CoFe$_2$O$_4$@SiO$_2$ NPs by citrate method using TEOS, ethanol, Fe and Co nitrates, citric acid, and ethylene glycol was successfully reported by Garcia [83] and Varma [62], with the latter reporting 50 nm size and $M_S$ values in the range of 4–25 emu/g.

2.2.5. Thermal Decomposition Method

The thermal decomposition method is based on the heating of reactants (metallic precursors) at different temperatures. To control the nucleation and growth of the NPs and consequently improve the physical properties of the material, such as crystallite size, porosity, and specific surface, surfactants (oleylamine, oleic acid) are added in the
decomposition step. The surfactants act as a protective envelope coating the NPs, limiting the coalescence, and leading to the improvement of the physical properties of the material, such as crystallite size, porosity, and specific surface. These parameters allow the control of the magnetic properties [85].

The method is relatively simple, low-cost, takes place at low reaction temperature, is environment friendly, produces highly monodispersed particles with a narrow size distribution, and does not produces byproducts. The main drawback is the removal of surfactants simultaneously with the particle size controlling [50, 86]. The use of thermal treatment to synthesize ferrite NPs is very limited, although this method requires reduced time as compared to other synthesis techniques [87]. The thermal decomposition method, reflux temperature, and time play an important role in controlling size distribution and aggregation of CoFe$_2$O$_4$ NPs, which will significantly influence the magnetic properties such as permeability and coercivity of the products. The reaction rate depends on the concentration of metal precursors, surfactants, reducing agents and temperature [88]. One example of thermal decomposition is the dissolution of metal nitrates in deionized water, followed by their dispersion in a solution of polyvinylpyrrolidone (PVP) at 70 °C, followed by a thermal treatment at 600 °C for four hours [87]. Asghar et al. [3] reported the synthesis of MnFe$_2$O$_4$ by dissolving Fe and Mn acetates (2:1, molar ratio) in dibenzyl ether, in the presence of 1,2-dodecanediol, oleic acid, and oleylamine, followed by a thermal treatment at 250 and 350 °C, respectively. Peddis et al. reported the synthesis of crystalline CoFe$_2$O$_4$ NPs coated by oleic acid and organized in a self-assembling arrangement with narrow size distribution (∼5 nm) by high thermal decomposition using Fe(III) acetylacetonate, Co(II) acetylacetonate, 1,2-hexadecanediol, oleylamine, oleic acid, and phenyl ether as solvents, heated to 200 °C for 30 min and to 265 °C for 30 min [89]. Following the thermal treatment method, in aqueous solution containing metal nitrates and polyvinyl pyrrolidone, followed by grinding and calcination at temperatures ranging between 723 and 873 K, MnFe$_2$O$_4$ NPs (12–22 nm) were produced [86].

2.2.6. Pyrolysis

Spray Pyrolysis Method

The spray pyrolysis method consists in converting the reagent mixture into aerosol droplets, solvent evaporation, solute condensation, and drying followed by thermolysis of the particles at high temperature [50]. This method allows the control of the particle formation environment by dividing the solution into droplets [90]. Generally, it is suitable for the synthesis of mixed metal ferrites as it ensures complete stoichiometry retention on the droplet scale and provides a highly homogeneous distribution of the components [90]. By controlling the type of thermolysis reaction, the type of precursors, the gaseous carrier, deposition time, and substrate temperature, this method allows the synthesis of a broad range of hollow or porous particles with potential applications in thermal insulation or catalyst support [90, 91]. Non-agglomerated particles smaller than 10 nm are achieved by using soluble inert additives (NaCl, H$_3$BO$_3$) to the reaction mixture. Furthermore, the inert components allow subsequent thermal treating of the obtained powder without significant particle size growth [89]. The method’s advantages include obtaining predictably sized, finely dispersed particles of variable shape and composition, high production rate, scalability, and reduced time [50, 89]. The disadvantages of the method are the high costs and possibility to obtain aggregated particles [50]. Zn$_x$Fe$_{3-x}$O$_4$ and CoFe$_2$O$_4$ thin films were produced by spray pyrolysis technique, by spraying aqueous solution of metal nitrates at a temperature of 300 °C. After deposition, film was naturally cooled [92]. CoFe$_2$O$_4$ thin film is prepared by the spray pyrolysis technique due to its easy handling, simple experimental setup, and cost effectiveness as large surfaces of thin film can be produced [92].

Laser Pyrolysis

Laser pyrolysis is another method for the synthesis of nanosized ferrites. The gaseous phase precursors are transported by a carrier gas introduced into a reaction chamber where a high-power laser beam (2400 W) generates elevated local temperatures, which trigger the
nucleation and growth of NPs that are further collected by a filter [93]. Liquid reactants are introduced into the reaction chamber reactants as vapors or microscale droplets [94]. This technique allows the production of high purity nanomaterials with small and narrow particle size distribution in one step. The method is versatile and flexible, and allows the control of operational parameters, such as laser power, type of gases, gas flow rate, and concentration of the precursors [94]. Moreover, it can be easily scaled-up to industrial production. The disadvantages of the method include the high cost of equipment used and the need for a precursor with absorption band at 10.6 µm, or addition of C2H4 or NH3, which leads to further contamination with carbon or nitrogen [93,94]. The synthesis of ZnFe2O4 nanopowders by laser pyrolysis using Fe and Zn nitrate solutions as precursors, ethylene as sensitizer gas and air as carrier gas was reported. The obtained nanopowders were tested as negative electrode materials for Li-ion batteries [94].

2.2.7. Hydrothermal Method

The hydrothermal method is used for the preparation of ferrite NPs at large scale owing to its high yields [58,59]. The method consists in the NPs formation by mixing the aqueous divalent metal (Ni, Co, Mn, Zn, Mn) acetate solutions with iron nitrate and a carbon-based nano template at alkaline pH and its dispersion around 200 °C in an autoclave. The obtained precipitate is washed, separated by centrifugation, and annealed around 500 °C to eliminate the carbon-based template [95]. In order to control the crystal structure, particle size, and morphology, surfactants are often employed in solution process [96]. Ethylenediamine and citric acid may also be used in the synthesis. This method requires neither sophisticated processing nor high processing temperature and allows the selection of the NPs properties by using different temperatures, pressure, reaction times, and nano templates [59,97].

Advantages of the hydrothermal method are good nucleation control, production of low particle size with narrow particle size distribution, controlled morphology at a high reaction rate and different temperature and pressure levels, high yield, excellent stoichiometry, and simplicity [67].

The production of magnetic nanosized CoFe2O4 by hydrothermal method without subsequent calcination processes was reported [98]. However, the influence of the synthesis time on the morphology and the particle size of CoFe2O4 NPs is still under discussion [96]. CoFe2O4 nanorods using cetyltrimethylammonium bromide (CTAB) as the surfactant were also synthesized by the hydrothermal method [99]. Porous anodic aluminum oxide is also used as template in the synthesis of one-dimensional CoFe2O4 nanostructures, however the costs of the synthesis are high while the yields are low [99]. The decrease of the reaction medium pH favors the CoFe2O4 crystallite size decrease. However, pure CoFe2O4 at pH < 10 was not obtained independently of the temperature and reaction time [100]. CoFe2O4 nanoplatelets and NPs were also prepared by hydrothermal reaction from FeCl3, cobalt dodecyl sulfate and NaOH aqueous solution, in strictly controlled synthesis environment [101]. CoFe2O4 NPs were also synthesized by the supercritical hydrothermal method at different temperatures of 25–390 °C for two hours [58,59,68,97].

NiFe2O4 nanocrystals were synthesized from FeCl3·6H2O and NiCl2·6H2O, using cetyltrimethylammonium bromide (CTAB) as the surfactant and NH3 as hydrolyzing agents [101]. Moreover, ZnFe2O4 were synthesized using metallic Zn sheet and FeCl2 as reactants in ammonia solutions [101]. Hydrothermal synthesis of MFe2O4 (M = Ni, Co, Mn, Zn) using metal acetylacetonate and aloe vera extracts provided high-yield nanosized ferrite with well crystalline structure using high calcination temperature [101].

2.2.8. Solvothermal Method

The solvothermal method has emerged as a promising technique to synthesize monodisperse spherical magnetic microspheres with high surface area, magnetic saturation and good dispersion in liquid media [102]. CoFe2O4 magnetic NPs in the size range 2–15 nm were prepared using a non-aqueous solvothermal method by dissolving Co(III) acetylacetonate and Fe(III) acetylacetonate in acetonaphone, followed by solvothermal treatment between
120 and 200 °C for 22 h in a Parr Acid Digestion Bomb autoclave [103]. The solvothermal method was successfully applied to synthetize CoFe$_2$O$_4$, NiFe$_2$O$_4$, MnFe$_2$O$_4$, and ZnFe$_2$O$_4$ using sodium or ammonium acetate, polyethylene glycol, urea, and oleylamine at around 200 °C [102]. Size-controlled NiFe$_2$O$_4$ NPs were successfully synthesized via a simple solvothermal method using ethylene glycol as solvent and sodium acetate as electrostatic stabilization, by adjusting the experimental parameters (time, initial concentration of reactants, amount of protective reagents, and the type of acetates) [20]. The main advantages of the method are high performance in biological applications by obtaining magnetic NPs with smooth surfaces, narrow size distributions, large surface areas, and high magnetic saturation in order to provide maximum signal and good dispersion in liquid media [49].

2.2.9. Microemulsion Method

The microemulsion method consists in the mixing of microemulsions containing reactants, the formation of micro droplets, followed by the trapping of fine aqueous micro droplets inside surfactant molecule assemblies. This procedure results in a locking up effect that limits the growth dynamics, particle nucleation, and agglomeration during the nanoparticle formation. The major advantage is the obtaining of monodispersed NPs with various morphologies, while among the major disadvantages can be listed the low efficiency and difficulty to scale up [69]. CoFe$_2$O$_4$ NPs lower than 50 nm, with high $H_C$ and $M_S$ were synthetized by a simple micro-emulsion synthesis route using water-in-oil emulsions consisting of water, cetyltrimethyl ammonium bromide as surfactant, n-butanol as co-surfactant, and n-octane as oil phase [104]. CoFe$_2$O$_4$ NPs of 4–25 nm were prepared using normal micelle microemulsion methods [105].

2.2.10. Reverse Micelle Method

The reverse micelle method is based on the formation of water-in-oil emulsions in which the water to surfactant ratio controls the size of water pools within which aqueous chemical syntheses take place [106]. The method consists in mixing the metal salt precursors and the precipitating agent and the formation of microemulsion droplets, which act as a nano-reactor, in which nanometric size precipitates are formed following the coalescence and droplet collision of reactants [107]. This method allows an excellent control of the particle size, size distribution, chemical stoichiometry, and cation occupancy and is suitable for room temperature reactions such as the precipitation of oxide NPs [106].

Metallic chlorides and NaOH were dissolved by ultrasonication in two different microemulsion systems prepared by mixing water, sodium dodecyl sulfate, 1-butanol and n-hexane. The two microemulsions were mixed together till metallic hydroxides were precipitated, the obtained precipitates were filtered, washed, dried and annealed at 400 °C [107]. The nature of surfactant, water to surfactant ratio and pH value of solution strongly influences the particle size of ferrites, which in turn affects their properties [101]. For the synthesis of CoFe$_2$O$_4$ a surfactant system formed by mixing sodium dioctylsulfosuccinate with a 2,2,4-trimethylpentane oil phase was used [106]. CoFe$_2$O$_4$ magnetic NPs were synthesized by reverse micelle methods using a micro emulsion consisting of three independent phases, (petroleum oil, pyridine and water) as a template to control the size of the NPs [108]. MnFe$_2$O$_4$ NPs with particle size of approximately 8–25 nm were prepared using reverse micelle microemulsion methods [105]. Using the reverse micelle microemulsion method, a wide range of spinel ferrite nanoparticle cores can easily be coated with a silica shell [105]. Such a method increases the potential for development of tunable magnetic silica for magnetoelectronic and biomedical applications [105].

2.2.11. Sonochemical Method

The sonochemical method is considered one of the most promising procedures to obtain ferrite NPs. The sonochemistry arises due to acoustic cavitation phenomenon consisting in the formation, growth and collapse of bubbles in liquid medium. The extreme reaction conditions (i.e., high temperature (5000 K), pressure (20 MPa) and cooling rate
(1010 K/s)) lead to numerous unique properties of the produced particles [109]. Highly crystalline, monodisperse CoFe$_2$O$_4$ NPs with uniform spherical shape and high $M_S$ values were synthetized in a one-step, surfactantless, sonochemical process. Additionally, the synthesis time was low (about 70 min) and no subsequent annealing was necessary [109]. The advantages of this approach over the conventional methods include simplicity, low cost, safety, environment friendly, uniform size distributions, high surface area, fast reaction time, and good phase purity [109]. Disadvantages comprise a very small concentration of prepared NPs and particle agglomeration [50].

2.2.12. Spray Drying Method

The spray drying method is used for the large-scale production of ZnFe$_2$O$_4$ and MnFe$_2$O$_4$. The method is based on the spraying of the ferrite slurry droplets into a hot air vertical evaporating tube. The main drawbacks are the inflation defects visible as large voids inside the granule and the development of capillary stress due to the rapid water evaporation. This stress may lead to undesirable diffusion and segregation phenomena, which reduce the compositional and morphological homogeneity of NPs [110]. As the defects may resist annealing, to preserve the requested particle size and avoid the occurrence of defects, the spray droplets may be frozen using liquid nitrogen and freeze-dried. During water sublimation, there are no undesirable capillary stresses that generate hard unbreakable aggregates and result in voids that lead to a rigid porous product of loose and non-agglomerated particles [110].

2.2.13. Polyol Method

The polyol method consists in the reduction of metallic oxide or metallic complexes by a high boiling point solvent that acts as a solvent as well as a reducing agent of the metallic ions under reflux conditions. The method allows controlling particle growth and preventing the particles agglomeration by the suspension of precursors into liquid polyol (ethylene glycol, diethylene glycol and triethylene glycol), followed by thermal treatment up to boiling point of polyol. The major advantage of this method is the obtaining of uniform size soft and hard magnetic NPs, while the main disadvantages are the high temperature and long time [111,112]. The polyol method has been used for synthesis of many oxides and NCs materials [73]. To produce CoFe$_2$O$_4$, FeCl$_3$·6H$_2$O and CoCl$_2$·4H$_2$O were individually dissolved in glycol, mixed together under continuous stirring. To the homogenous solution, water and sodium acetate were added, and the pH value of the solution was adjusted (8, 10, and 12) by adding ammonia solution. Under reflux conditions, the mixture was heated to the boiling temperature of glycol, while the formed NPs were separated by centrifugation [111]. The synthesis of NiFe$_2$O$_4$ NCs starting from tetraethoxysilane without, or with modifiers as formamide, citric acid, and PVA were also reported [4].

2.2.14. Biosynthetic Method

The biosynthetic method uses plant extracts as a simple and viable alternative to chemicals during the synthesis process. The use of microorganisms, enzymes, and plant extracts has been proposed as a possible eco-friendly alternative to chemicals of physical synthesis. Furthermore, the biosynthetic route is very simple and provides high-yield, crystalline nanomaterials with adequate properties [13]. Aloe vera (Aloe barbadensis mill.) plant-extract may be used as a bio-reducing agent in the preparation of ferrites, providing a simple, efficient and green alternative route for the synthesis of nanomaterials. A possible explanation for this could be the presence of long chain polysaccharides in the Aloe vera plant extract that allows the homogeneous distribution of ferrite [13]. The major advantages consist in the selectivity and precision of NPs formation, cost-effectiveness, as well as the use of an eco-friendly reducing agent and non-hazardous gelling agent for stabilizing the nanostructures. Disadvantages are the difficult size and properties controlling, high temperature required to convert the precursors in crystalline materials and the formation of polydisperse, surface capped or unpurified NPs, in addition to poor reproducibility [13,50].
The use of Ni and Fe nitrates and freshly extracted egg white (ovalbumin) in an aqueous medium, results in the formation of NiFe$_2$O$_4$ NPs with ferrimagnetic behavior, $M_S$ values in the range of 26.4–42.5 emu/g for the applied field of 10 kOe, at room temperature [20].

In conclusion, although different synthetic routes have been adjusted, the key challenge in the synthesis field remains the obtaining of size- and phase-controlled synthesis, with good reproducibility. However, reproducibility is difficult to achieve in the synthesis methods where partial mixing of reactants and undesired reactions take place, affecting the properties of NPs. Moreover, the heat and mechanical treatments can alter crystal structure, affecting the photocatalytic activity. The combustion method is simple and quick, but the rate of organic fuel to nitrates can highly affect the size and magnetic properties. The solid-state method at higher temperatures produced micron-sized CoFe$_2$O$_4$ particles, while the citrate precursor method at lower temperatures produced NPs. To overcome these challenges in an ideal, controlled reproducible synthesis, the following steps should be achieved: (i) direct active and complete mixing of reactants, (ii) automation, and (iii) enabled reaction parameters controlled precisely. Each synthesis method has its own pros and cons, and selection depends on many factors, but amongst all methods of synthesis, the sol-gel and chemical co-precipitation techniques stand to be favorable routes to synthesize homogeneous, highly pure, and narrow size distribution nanoferites.

3. Applications

The most frequent applications of MFe$_2$O$_4$ (M = Co, Cu, Mn, Ni, Zn) are presented in Figure 3. Considering the cation type and synthesis route, the structure of the ferrite and its particle size and shape changes and thus also its properties, which further lead to different applications.

![Figure 3. Application of nanosized ferrites.](image-url)

3.1. Magnetic Applications

The dependence of the magnetization on the grain size is due to the variations of exchange interaction between tetrahedral and octahedral sites and [20,113]. For their application in high-density magnetic recording, the magnetic particles should have nanosize to prevent the exchange interaction between adjacent grains, and thus reducing the media noise in the materials. The particles must also possess high $H_C$ values to obtain high storage density [98]. The magnetic properties ($M_R$, $M_S$, and $H_C$) of spinel ferrites depend upon the composition, particle size, crystal structure, and cationic distribution between octahedral and tetrahedral sites. Moreover, they may exhibit antiferromagnetic, ferrimagnetic and paramagnetic behavior [101,114].
The increase of $H_C$ values may also occur due to a combination of spin disorder, spin canting effect, and improved surface barrier potential in surface layers [87,115]. In the case of magnetic NPs, the presence of a large number of atoms at the surface due to large surface-to-volume ratio results in interesting and superior properties in comparison to corresponding bulk materials. Moreover, the low number of coordination surface atoms relative to interior atoms of the nanoparticle leads various surface effects such as spin canting, spin disorder and existence of a magnetically dead layer [115].

High coercivity materials are known as hard materials, while low coercivity materials are known as soft materials [113]. The soft materials are used for inductor cores, transformers, and microwave devices, while hard materials are used for permanent magnets. Generally, the $H_C$ value is low in soft ferrites and the magnetization can be tailored for cutting-edge applications in electronic engineering such as microwave components, high-frequency inductors and transformer cores [58]. Furthermore, due to the high $H_C$ values of hard ferrites, it is not so easy to magnetize, and hence it is used as enduring magnets with applications in washing machine, refrigerator, communication systems, microwave absorbing systems, loudspeaker, TV, switch-mode power supplies, DC-DC converters, and high-frequency applications [58].

CoFe$_2$O$_4$ is a ferrimagnetic ceramic with excellent properties, such as high $K$, $H_C$, and $T_C$ values, moderate $M_S$ values, and large magnetostrictive coefficient value [42,116]. The high $M_S$ and $M_R$ values can be explained on the basis of Neel’s theory and cation distribution at tetrahedral (A) and octahedral (B) sites [11,25,117]. Generally, in single-phase samples the magnetic characteristics are highly influenced by the material microstructure such as shape and size of crystals, residual stress and crystal defects [51]. Moreover, the synthesis temperature plays a key role in controlling particle size of CoFe$_2$O$_4$ with a significantly influence on its magnetic properties [66]. As expected, $M_S$ values increase with the CoFe$_2$O$_4$ content and temperature, due to the increase of the crystallinity degree as well as to the average size of the magnetic CoFe$_2$O$_4$ NPs [14,42,99]. Due to the high values of $K$ and high value of spin-orbit coupling constant of Co$^{2+}$ ions, CoFe$_2$O$_4$ displays the highest magnetostriction (i.e., up to 200 ppm) among oxide-based materials. The increment of $H_C$ value is mainly attributed to the increasing particle size as a consequence of the coarsening of nanocrystals and diminished crystal defects at high temperature [98]. In the case of CoFe$_2$O$_4$ embedded in SiO$_2$ matrix (CoFe$_2$O$_4$@SiO$_2$), the reduction of $M_S$ and $M_R$ values is attributed to the reduced amount of magnetic material per gram of nanocomposite [105]. Moreover, the increase of annealing temperature leads to the decrease of $K$ and $H_C$ values [74]. In contrast, the $H_C$ value of CoFe$_2$O$_4$@SiO$_2$ NPs does not show any change after coating, while the $H_C$ value of MnFe$_2$O$_4$ NPs decreases by 10% after coating [105]. In the case of CoFe$_2$O$_4$ obtained by the citrate gel method, higher $M_S$ values were obtained for samples synthesized at high temperatures than the samples prepared at low temperatures, suggesting that the $M_S$ value rises with increase in particle size, while the $M_R$ decrease with the increase of particle size [62]. Peixoto et al. [43] produced CoFe$_2$O$_4$ NPs embedded in a SiO$_2$ matrix via sol–gel method and evaluated the magnetic properties at 5 K and 100–200 K, revealing a superparamagnetic behavior.

The $M_S$ value of 27.09 emu/g found for CuFe$_2$O$_4$ obtained by solid state chemistry (particle size of 36 nm) was similar to the $M_S$ value of 33 emu/g for CuFe$_2$O$_4$ NPs of ~32 nm obtained by other synthesis method. However, the $H_C$ of 112 Oe found for CuFe$_2$O$_4$ obtained by solid state chemistry (particle size 36 nm) is greater compared to other syntheses, as a consequence of the presence of some impurities (CuO). Further, an $M_R$ value of 4 emu/g was reported for CuFe$_2$O$_4$ of particle size ~8 nm synthesized by the sol-gel method [113]. The $H_C$ of CoFe$_2$O$_4$ NPs show a maximum $H_C$ value at a particle size of about 25 nm, while in the case of CuFe$_2$O$_4$ NPs obtained by sol–gel, co-precipitation, solid-state reaction, thermal decomposition, and solution auto-combustion methods, the $H_C$ decreases linearly with the decreasing size, indicating that the particles are in single-domain area and the surface spins dominate the magnetization decreases with size reduction [57]. The $H_C$ and
The increase of $M_R$ and $M_S$ values of MnFe$_2$O$_4$ can be ascribed to the metal cations distribution at octahedral and tetrahedral sites of crystal lattice structure, meaning to the tendency of Mn$^{2+}$ ions to be positioned at octahedral (B-site) [118]. MnFe$_2$O$_4$ NPs and MnFe$_2$O$_4$@SiO$_2$ nanocomposite exhibit superparamagnetic behavior attributed to the ultra-small size of MnFe$_2$O$_4$ NPs and are widely used for drug delivery applications. The low $M_S$ value of MnFe$_2$O$_4$@SiO$_2$ nanocomposite is due to the presence of non-magnetic SiO$_2$ matrix [3]. The $H_C$ of MnFe$_2$O$_4$@SiO$_2$ NPs slightly decreases compared to that of native magnetic NPs. No similar behavior was observed for CoFe$_2$O$_4$@SiO$_2$ probably due to the larger contribution of the surface anisotropy to the total anisotropy of MnFe$_2$O$_4$ NPs [105]. The $M_S$ value of 52.4 emu/g obtained in the MnFe$_2$O$_4$ (diameter of ~15.9 nm) is lower than values 67.0 emu/g for synthesized MnFe$_2$O$_4$ crystallites using a triethanolamine-assisted route under mild conditions (diameter of ~1 μm), but higher than the value of ~48.6 emu/g for the polymer-pyrolysis route MnFe$_2$O$_4$ NPs (diameters of ~9 nm) [101].

ZnFe$_2$O$_4$ displays antiferromagnetic behavior when the temperature is below the Néel temperature and diamagnetic, superparamagnetic, or ferrimagnetic behavior when particle sizes are at nanometer scale [119]. The superparamagnetic behavior is due to the increased disorder of magnetic moments orientation in the various sites when the ratio surface/volume increases. The $M_S$ value of ZnFe$_2$O$_4$ strongly depends on various factors, such as the synthesis route and its conditions, type of the precursors, and annealing treatments [63]. The $M_S$ value of 7.06 emu/g for ZnFe$_2$O$_4$ (diameter of ~17.9 nm) is lower than values of 54.6 emu/g for hydrothermal-synthesized ZnFe$_2$O$_4$ ultrafine particles (crystallite size of ~300 nm) [101]. However, the paramagnetic behavior of ZnFe$_2$O$_4$ was remarked upon in bulk [46] and NPs, due to the increase of an atypical Fe$^{3+}$ cation distribution in the tetrahedral coordination sites upon nanosizing [47,101].

In the crystal structure of NiFe$_2$O$_4$, the Ni$^{2+}$ occupies the octahedral site with Fe$^{3+}$ cations distributed equally at tetrahedral and octahedral sites, while their antiparallel spins produce a net magnetic moment of $2\mu_B$ owing to Ni$^{2+}$ ions at octahedral site [87]. Due to high crystallinity and uniform morphology, NiFe$_2$O$_4$ ferrites show high $M_S$ and low $H_C$ values [119]. The magnetic properties of NiFe$_2$O$_4$ NPs are attributed to the cumulative effect of various factors, such as super-exchange interaction, magneto crystalline anisotropy, canting effect, and dipolar interactions on the NP’s surface. Besides, the variation of $H_C$ value with particle size can be explained by the domain structure, critical size and anisotropy of the crystal [20]. The large NiFe$_2$O$_4$ is due to the scattering in direction of anisotropy field due to inhomogeneous broadening, at high temperature, it has the tendency to make magnetic moment isotropic causes the decrease of $H_C$ value [21]. The $M_S$ value of 31.9 emu/g found for NiFe$_2$O$_4$ (particle size of ~8.2 nm) is close to the values of 34.5 emu/g for NiFe$_2$O$_4$ NPs (crystallite size of ~68 nm) obtained by egg-white solution route, but lower than the theoretical $M_S$ of 50 emu/g calculated using Neel’s sublattice theory and the reported value of 56 emu/g for the bulk sample [96]. The $M_S$ value of 55.3 emu/g obtained in the CoFe$_2$O$_4$ (diameter of ~8.5 nm) is lower than values of 80 emu/g for bulk CoFe$_2$O$_4$ and ~65 emu/g for the CoFe$_2$O$_4$ NPs with crystallite size of ~40 nm synthesized by aerosol route, and higher than the values of 30 emu/g for hydrothermal-synthesized CoFe$_2$O$_4$ NPs (diameter of ~30 nm).

The synthesis of nanocrystalline spinel ferrite of type MFe$_2$O$_4$ (e.g., $M = $ Mn, Co, Ni, Cu, Zn) has great relevance to modern technological applications in several industrial and biological fields, including magnetic recording media and magnetic fluids for the storage and/or retrieval of information, magnetic resonance imaging enhancement, and magnetically guided drug delivery [101]. Furthermore, the magnetic NPs showed promising results in the field of medicine and healthcare treatment owing to their biocompatibility, low toxicity, and ability to be handled by the application of a magnetic field [50].

Table 1 presents the main magnetic parameters ($M_S$, $M_R$, $H_C$, $K$), particle diameter, and synthesis method of the studied ferrites.
Table 1. Average crystallites size ($D$), saturation magnetization ($M_S$), remanent magnetisation ($M_R$), coercivity ($H_C$) and anisotropy constant ($K$) of CoFe$_2$O$_4$, CuFe$_2$O$_4$, MnFe$_2$O$_4$, NiFe$_2$O$_4$, ZnFe$_2$O$_4$, produced by different synthesis methods; RT-room temperature.

| Ferrite          | Synthesis Method          | Temp. | $D_{XRD}$ (nm) | $M_S$ (emu/g) | $M_R$ (emu/g) | $H_C$ (kOe) | $K \cdot 10^3$ (erg/cm$^3$) | Ref.  |
|------------------|--------------------------|-------|----------------|---------------|---------------|-------------|-----------------------------|-------|
| CoFe$_2$O$_4$    | sol-gel                  | RT    | 28.0           | 26.3          | 11.9          | 1.440       | 2.378                       | [46]  |
|                  | sol-gel auto-combustion  | RT    | 36.0           | 89.0          | 20.0          | 0.650       | -                          | [11]  |
|                  | sol-gel auto-combustion  | RT    | 52.0           | 65.6          | 71.9          | 1.117       | -                          | [78]  |
|                  | solid-state              | RT    | 80.0           | 77.2          | -             | 0.525       | -                          | [120] |
|                  | co-precipitation         | RT    | 17.3           | 78.6          | 15.8          | 0.778       | -                          | [121] |
|                  | co-precipitation         | RT    | 37.0           | 72.6          | 34.5          | 1.060       | -                          | [117] |
|                  | co-precipitation         | RT    | 11.70          | 58.4          | 12.45         | 0.286       | -                          | [108] |
|                  | combustion               | RT    | 52.0           | 52.6          | 20.8          | 1.274       | -                          | [122] |
|                  | hydrothermal             | RT    | 17.3           | 63.4          | 23.1          | 0.831       | 0.450                      | [123] |
|                  | hydrothermal             | RT    | 8.5            | 55.3          | -             | 0.074       | -                          | [101] |
|                  | hydrothermal             | RT    | 16.3           | 58.3          | -             | 1.029       | -                          | [121] |
|                  | normal micelles          | RT    | 5.58           | 12.6          | 0.17          | 0.0237      | -                          | [108] |
|                  | reverse micelles         | RT    | 7.62           | 29.4          | 0.84          | 0.0252      | -                          | [108] |
|                  | reverse micelle          | RT    | 6.00           | 34.4          | 8.50          | 0.500       | -                          | [124] |
|                  | polyl                    | RT    | 14.0           | 57.0          | 21.6          | 1.123       | -                          | [125] |
|                  | thermal decomposition    | RT    | 13.0           | 68.0          | -             | 0.560       | -                          | [88]  |
|                  | thermal decomposition    | RT    | 82.8           | 48.38         | 11.72         | 0.64        | -                          | [126] |
| CuFe$_2$O$_4$    | sol-gel                  | RT    | 60.0           | 14.5          | 3.03          | 0.018       | 0.163                      | [46]  |
|                  | sol-gel auto-combustion  | RT    | 8.00           | 16.5          | 4.00          | 0.450       | -                          | [113] |
|                  | solid-state              | RT    | 38.0           | 27.1          | 1.41          | 0.112       | -                          | [113] |
|                  | co-precipitation         | RT    | 31.0           | 22.9          | 1.12          | 0.114       | -                          | [113] |
|                  | co-precipitation         | RT    | 52.0           | 28.1          | 8.52          | 0.189       | -                          | [117] |
|                  | combustion               | RT    | 22.3           | 14.0          | 0.17          | 0.006       | -                          | [118] |
|                  | auto-combustion          | RT    | 22.0           | 18.9          | 1.51          | 0.140       | -                          | [113] |
|                  | thermal decomposition    | RT    | 53.0           | 29.0          | 0.91          | 0.102       | -                          | [113] |
|                  | thermal decomposition    | RT    | 10−250         | 0.90          | -             | 0.600       | -                          | [128] |
|                  | sonochemical             | RT    | 40.0           | 21.5          | 12.6          | 0.235       | -                          | [24]  |
| Ferrite   | Synthesis Method   | Temp.   | $D_{\text{XRD}}$ (nm) | $M_s$ (emu/g) | $M_r$ (emu/g) | $H_c$ (kOe) | $K \cdot 10^3$ (erg/cm$^3$) | Ref.    |
|----------|--------------------|---------|-----------------------|--------------|--------------|-------------|-----------------------------|---------|
| MnFe$_2$O$_4$ | sol-gel           | RT      | 49.0                  | 28.5         | 14.8         | 0.119       | 3.392                       | [46]    |
|          | co-precipitation   | RT      | 6.10                  | 13.5         | 1.24         | 0.046       | -                           | [129]   |
|          | hydrothermal       | RT      | 22.0                  | 65.5         | 3.86         | 0.054       | 0.018                       | [130]   |
|          | hydrothermal       | RT      | 15.9                  | 52.4         | -            | 43.9        | -                           | [101]   |
|          | hydrothermal       | RT      | 25.4                  | 67.3         | -            | 0.173       | -                           | [124]   |
|          | polyl              | RT      | 7.10                  | 51.9         | -            | -            | -                           | [130]   |
|          | sonochemical       | RT      | 25.5                  | 59.4         | 3.41         | 0.024       | 1.53                        | [131]   |
|          | microwave combustion| RT      | 27.9                  | 61.0         | 11.4         | 0.064       | -                           | [53]    |
|          | thermal decomposition| RT      | 61.0                  | 56.0         | 10.0         | 0.08        | -                           | [132]   |
| NiFe$_2$O$_4$ | sol-gel           | RT      | 23.0                  | 20.1         | 6.82         | 0.061       | 0.770                       | [46]    |
|          | sol-gel auto-combustion | RT      | 70.0                  | 37.3         | -            | 0.321       | -                           | [133]   |
|          | co-precipitation   | RT      | 17.3                  | 43.9         | 16.6         | 0.051       | -                           | [121]   |
|          | combustion         | RT      | 25.0                  | 30.2         | 4.00         | 0.159       | -                           | [122]   |
|          | microwave combustion| RT      | 18.5                  | 37.9         | 2.57         | 0.016       | -                           | [134]   |
|          | hydrothermal       | RT      | 8.20                  | 31.9         | -            | 0.007       | -                           | [101]   |
|          | thermal decomposition| RT      | 25.0                  | 36.5         | 10.6         | 0.263       | -                           | [87]    |
|          | thermal decomposition| RT      | 79.0                  | 43.60        | 17.63        | 0.645       | -                           | [135]   |
|          | thermal decomposition| RT      | 10.7                  | 36.8         | -            | 0.007       | -                           | [136]   |
| ZnFe$_2$O$_4$ | sol-gel           | RT      | 49.0                  | 10.8         | 1.67         | 0.015       | 0.102                       | [46]    |
|          | sol-gel auto-combustion | RT      | 54.0                  | 5.31         | 0.08         | 0.113       | -                           | [78]    |
|          | solid-state        | RT      | 80.0                  | 77.27        | -            | 0.525       | -                           | [120]   |
|          | co-precipitation   | RT      | 4.80                  | 11.9         | 0.01         | 0.004       | -                           | [129]   |
|          | microwave combustion| RT      | 37.5                  | 2.60         | 0.01         | 0.007       | -                           | [53]    |
|          | microwave combustion| RT      | 21.1                  | 3.85         | 0.51         | 0.010       | -                           | [134]   |
|          | hydrothermal       | RT      | 15.9                  | 52.4         | -            | 0.044       | -                           | [101]   |
|          | solvothermal       | RT      | 80.0                  | 77.0         | -            | 0.090       | -                           | [102]   |
|          | reverse micelle    | RT      | 8.30                  | 4.90         | 0.01         | 0.010       | -                           | [107]   |
|          | thermal decomposition| RT      | 30.0                  | 12.8         | -            | -            | -                           | [137]   |
The magnetostrictive materials are solids in which the application of a magnetic field results in strong strain and deformation as a consequence of the strong coupling of magnetic moments with crystal lattice [138]. The large magnetostriction in a wide range of temperature is very useful when a mechanical coupling is required in multiferroic composite systems [139]. The electronic and optical properties of magnetostrictive solid materials are more sensitive to magnetic field comparing to solids with weak magnetostriction [138]. Magnetostrictive smart materials are generally used to design sensors, actuators, sonar transducers, motors, etc. In this regard, there is an increasing interest on metal oxide based magnetostrictive materials such as CoFe$_2$O$_4$ due to various practical advantages over alloy based magnetostrictive materials, i.e., the high magnetostriction and strain-field derivative with low saturation field of CoFe$_2$O$_4$ results in potential for sonar detector and force sensor applications and vibration components in high frequency ultrasonic transducer [140–142].

The magnetostriction value of CoFe$_2$O$_4$ prepared using micron-size powders without magnetic annealing decreases from 130 ppm to 200 ppm [142]. The magnetic annealing may increase the magnetostriction and strain derivative values due to the uniaxial anisotropic distribution of magnetic domains, while various magnetic or non-magnetic ions could decrease the magnetostriction and increase the strain derivative values [142]. The application of a magnetic field to magnetostrictive ferrimagnetic CoFe$_2$O$_4$ leads to strong strain and deformation of the crystal lattice and to change of spectrum-strain-magneto-optics, while high magnetoreflection is remarked on in the case of great magnetostriction [138].

Beside the excellent soft magnetic properties, Fe-based soft magnetic composites have unique properties such as high electric resistivity and low eddy loss, making them good electrical insulation coatings without any notable decrease in magnetic properties [143]. $H_C$ value confirms the magnetic properties of hard magnetic or soft magnetic materials. CoFe$_2$O$_4$ obtained using co-precipitation method and low heat treatment have $M_S$ of 21.74 emu/g, $M_R$ of 2.37 emu/g, and $H_C$ of 556.57 Oe [144]. The in-situ oxidation is an effective and novel method for manufacturing soft magnetic composites with low core loss for applications in medium and high frequency fields. The nano-ZnFe$_2$O$_4$ layer effectively improves the magnetic properties of soft magnetic composites [143]. Soft magnetic spinel ferrite properties probably occur due to the interactions within metal oxide and particular vacancy oxygen ions in their cubic spinel structure [144].

3.2. Photoluminescent Applications

Room temperature photoluminiscence is one of the important properties of mixed spinels as CoFe$_2$O$_4$, NiFe$_2$O$_4$, and Zn$_2$Fe$_2$O$_4$ nanostructures. The photoluminescence spectrum offers information regarding the surface oxygen vacancies and defects, as well as the efficiency of the charge carrier trapping, immigration and transfer [17]. The broad visible band emission was ascribed to the charge transfer among Fe$^{3+}$ at octahedral sites, M$^{2+}$ (M = Co, Ni and Zn) at both tetrahedral and octahedral sites, and its surrounding O$^{2-}$ ions [1]. The blue emission peak at 460 nm is attributed to the Fe$^{3+}$ transition in the ferrite sites, while the main peak at 418 nm resulted from the free electrons trapped at the oxygen vacancies. The intensity of every peak diminished with the increasing Ni content, as a consequence of the increasing band gap which reduces the electron-hole recombination ratio [145]. The violet emissions resulted from the radiating defects associated to the interface traps within the grain boundaries. The decrease in the luminescence intensity of Co$^{3+}$ from CoFe$_2$O$_4$ with the increase of doping fraction suggests a lower electron-hole recombination ratio [81]. Photoluminescence measurements showed a value around 2.13 eV for the energy gap of nanocrystalline ZnFe$_2$O$_4$ [63].

3.3. Catalytic Applications

Spinel ferrites have been extensively used as heterogeneous catalysts as they can be simply recovered from the reaction mixture by filtration or in the presence of an external magnetic field and recycled numerous times, making the process more economically and environmentally feasible [53]. Heterogeneous catalytic nanomaterials play a central role
in the selective protection of functional groups in order to be economically valuable and environmentally friendly. The catalytic property of MFe₂O₄ (M = Cu, Ni, Co, Zn) spinel revealed that CoFe₂O₄ catalyst displayed the best performance using benzaldehyde with 63% conversion and 93% selectivity, while CuFe₂O₄ catalyst using H₂O₂ oxidant showed 57.3% conversion and 89.5% selectivity [53]. The particle size, surface area, morphology, and chemical composition considerable influence the catalytic activity of materials [134]. In addition, the catalytic property of nanocrystalline spinel ferrites depends on the distribution of cations among the tetrahedral (A) and octahedral (B) sites. As catalyst, CoFe₂O₄ displays the best performance compared to other ferrites [13]. The catalytic activity of CoFe₂O₄ was successfully reported for the synthesis of arylidene barbituric acid derivatives. The advantages of this method were short reaction time, high yields, simple and economic work-up procedure, high turnover frequency, chemoselectivity, environmental sustainability, and green conditions [38]. Large-size magnetic CoFe₂O₄ NPs were successfully used as catalyst for the oxidation of various alkenes in the presence of tert-butylhydroperoxide. The catalyst from the medium was easily separated using an external magnet and the catalyst was recycled several times without important loss of activity. CoFe₂O₄ exhibited high catalytic activity and selectivity in methanol decomposition to CO and H₂ [38]. Moreover, non-nanosized CoFe₂O₄ displayed catalytic applications, such as the conversion of CO to CO₂. In the case of the catalytic oxidation of benzyl alcohol to produce a mixture of benzaldehyde, benzoic acid, and benzyl benzoate, the CoFe₂O₄ NPs have been proven to be highly selective [13]. CoFe₂O₄ NPs can be also used as catalyst for Knoevenagel condensation reactions between aromatic aldehydes and ethylcyanoacetate, in mild conditions. The catalyst could be easily recovered from the reaction mixture using an external magnet and can be re-used four times, without a significant loss of activity [69].

NiFe₂O₄ is an effective catalyst in several industrial processes. Its catalytic activity related to O₂ yield and evolution rate under ambient reaction conditions, is comparable to that of Ir, Ru or Co-based materials [119]. Due to its importance for both industry and domestic safety, the catalytic combustion of CH₄ has attracted considerable interest, because of its higher energy conversion efficiency and low emissions of environmental pollutants. Therefore, the development of a high activity catalyst with sensitivity for CH₄ detection is of great interest [19]. In this regard, the catalytic performance of NiFe₂O₄ for CH₄ combustion, but also for alkylation and oxidation reactions has been reported [19]. NiFe₂O₄ was successfully used also as a catalyst in photocatalytic water oxidation using [Ru(bpy)₃]²⁺ as a photosensitizer and S₂O₈²⁻ as a sacrificial oxidant [119].

CuFe₂O₄ has been used in industrial processes as a catalyst in both organic and inorganic reactions [146]. The main advantages of CuFe₂O₄ NPs are simple work-up and low-cost procedure, mild reaction conditions, reusable catalyst, high yield, short reaction times, no isomerization during the reaction [38]. The use of CuFe₂O₄ NPs in organic catalysis, such as the reaction of substituted aromatic aldehydes, ethyl acetooacetate, and ammonium acetate, at room temperature was also reported. In all cases, the nano-catalyst can be easily recovered and re-used [9,38]. CuFe₂O₄ NPs were also employed as reusable heterogeneous initiator in the synthesis of 1,4-dihydropyridines, of α-amino nitrites and conversion of CO to CO₂ [38,146]. The successful use of CuFe₂O₄ NPs as catalyst for the ligand free N-arylation of N-heterocycles and for the cross-coupling of aryl halides with diphenyl diselenide to produce diaryl selenides was also reported [147,148]. The CuFe₂O₄ NPs also proved to be efficient catalysts for a simple, one pot, green, and efficient synthesis method for substituted benzoazoles by redox condensation [149].

ZnFe₂O₄ spinel acts as a wide band gap semiconductor with application in hydrogen generation by light-activated water splitting and in photocatalytic removal of harmful organic molecules [85]. Non-nanosized ZnFe₂O₄ have been used as catalysts in the oxidative conversion of methane, oxidative coupling of methane, and in methanol decomposition to CO and H₂O [38]. Additionally, the coupling between the catalyst and magnetic material is a new approach to enhance the catalytic performance of a catalyst [150].
3.4. Photocatalytic Applications

Photocatalysts are important materials that support the use of solar energy in oxidation and reduction processes, with applications in various areas, such as the removal of water and air contaminants, odor control, bacterial inactivation, water splitting to produce hydrogen, the inactivation of cancer cells, etc. [151]. Nowadays, the usage of photocatalysis is the preferred treatment methods for dyes removal, as alongside irradiation of light on a semiconductor, the produced electron-hole pairs are used also for the oxidation and reduction process [81]. The degradation of dyes occurs due to the formation of active radicals during the photocatalytic reaction [22,145].

Only few materials are capable of both photo-oxidation and photo-reduction, namely to complete the decomposition of harmful organic compounds and, concomitantly, to absorb visible light efficiently. The low crystallite size of ferrites leads to a large surface area and more reaction sites, increasing the photocatalytic activity [151]. The type of photocatalyst, crystallinity, size of NPs, accessibility of the active surface to pollutants, and diffusion resistance of organic pollutants are the most important characteristics that enhance the photocatalytic properties. Thus, small particle sizes with high crystallinity are of great interest due to their higher specific surface area and active sites that favor the photocatalytic activity [37]. Due to their spinel crystal structure and a band gap capable to absorb visible light, ferrites are suitable as photocatalysts for the degradation of a wide range of contaminants [151].

CoFe$_2$O$_4$ was used as photocatalyst for the degradation of organic dyes (i.e., methylene blue (MB) and rhodamine B (RhB)), its chemical stability and the narrow bandgap (1.1–2.3 eV) making it active under visible light [81,145]. CuFe$_2$O$_4$ photocatalyst was found to be more effective than ZnFe$_2$O$_4$ and NiFe$_2$O$_4$ in decomposing hazardous dye compounds, probably due to lower crystallite size, large surface area, small band gap and occurrence of pores that trap oxygen molecules and produce large number of oxidizing species and OH$^-$ radicals. The photocatalyst was recovered using a magnet and reused in four consecutive cycles under visible light [22]. CuFe$_2$O$_4$ “oversized” nanostructures displayed improved photocatalytic activity in the conversion of benzene under Xe lamp irradiation. Moreover, CuFe$_2$O$_4$ powders exhibited a good catalytic efficiency (~60%) at 200 °C and pH = 12 due to the high surface area (~120 m$^2$/g) [38]. The photocatalytic ozonation of dyes with CuFe$_2$O$_4$ NPs prepared by co-precipitation method was reported to successfully decolorize and degrade textile dyes (Reactive Red 198 and 120) without using high pressure of oxygen or heating [152].

The photocatalytic activity of ZnFe$_2$O$_4$ NPs relies on the surface properties and defects. ZnFe$_2$O$_4$ was tested for solar energy conversion and photochemical hydrogen production due to its relatively narrow band gap energy (1.9 eV), excellent visible-light response, good photochemical stability and low-cost [17]. The use of ZnFe$_2$O$_4$ NPs for the photocatalytic degradation of 4-chlorophenol and the dependence of the degradation efficiency on their particle size and morphology was also reported [134]. ZnFe$_2$O$_4$ was found to be an appropriate visible-light-driven catalyst, which completely degraded the RhB dye in short time, due to the small particle size and narrow size distribution [107]. The sol–gel/precipitation hybrid synthesis method of magnetic NiFe$_2$O$_4$@TiO$_2$ and its use as photocatalyst for the production of hydrogen from aqueous systems was reported by Kim et al. [153].

3.5. Water Decontamination

In the last years, the industrial wastewater management is one of the main challenges in developed countries. Wastewaters resulted from the textile industry contain many non-biodegradable organic dyes mixed with different contaminants in a variety of ranges. Untreated effluents negatively affect not only humans and animals, but also the flora and fauna [81]. RhB is a synthetic, highly toxic, water soluble, organic dye, widely used as colorant in many industries and frequently found in wastewaters. Various techniques, such as ozonation, electrochemical method, and Fenton process, have been employed for the treatment of RhB containing water. The use of CoFe$_2$O$_4$ for wastewater treatment is
based on its high adsorption capacity, magnetic properties that allow the NPs separation using an external magnetic field, and on its low energy band gap that enables the photocatalytic degradation of RhB under visible light irradiation, [37,81]. Moreover, CuFe$_2$O$_4$ is considered a prospective material for the water decontamination by photoelectrochemical processes. In combination with a suitable adsorbent, CuFe$_2$O$_4$ can be applied as a cost-effective adsorbent for removal of MB from water and wastewater [146]. The atrazine degradation by a three-dimensional electrochemical process using CuFe$_2$O$_4$ NPs as electrodes and catalyst for the activation of persulfate for were also reported [149]. Further, ZnFe$_2$O$_4$ can be effectively used as magnetically recyclable material for the removal of chemical contaminants (i.e., dyes), as well as biological contaminants from water/industrial wastewater treatment [154].

3.6. Coloring

Ceramic pigments are metal transition oxides with high thermal and chemical stability, high tinting strength when dispersed and fired with glazes or ceramic matrices, high refractive index, acid and alkali resistance, and low abrasive strength. The color of each pigment is obtained by adding chromophore agents (usually transition metals) in an inert oxide matrix [155].

CoFe$_2$O$_4$ is a black pigment widely used in the ceramic industry. The color performance depends on the coating crystallization degree, a larger number of crystals in the glass resulting in a brighter color [156]. The ZnFe$_2$O$_4$ spinels are thermally stable, insoluble, and resistant to aggressive media, exhibit a high covering power and a reasonable cost. ZnFe$_2$O$_4$ spinel pigments improve the mechanical strength of the binder by a chemical reaction producing cationic soaps, resulting in low solubility and a tendency towards saponification in contact with a corrosive environment [157]. The color of ZnFe$_2$O$_4$ depends on the annealing temperatures and particle size as it causes a reduction of the total reflecting surface of the powder. At high annealing temperatures, the system suffers a notable modification due to the disappearance of defects (as oxygen vacancies), leading to less distorted tetrahedral and octahedral sites, and consequently better-defined colors [158]. The possibility to select the desired cations in the spinel lattice (Zn$^{2+}$ in A sites and Fe$^{3+}$ in B sites for ZnFe$_2$O$_4$) can be used to change or to create new properties of the pigment dispersed in an organic binder [157].

3.7. Corrosion Protection

Metal corrosion is caused by an electrochemical process, generally promoted by Cl$^-$ ions, which dissolve Fe oxides [159]. The corrosion of a metallic substrate may be constrained by the transformation of ferrous (Fe$^{2+}$) ions to ferric (Fe$^{3+}$) ions, which act as inorganic corrosion inhibitors in the electrochemical process, leading to the formation of a passivation layer, which will overcome the cathodic and/or anodic reaction [157].

Organic coatings act as barrier or mechanical protection against corrosive environments [157]. Nanocontainers can release corrosion inhibitors and subsequently protect the metal substrate from corrosion when incorporated into an organic coating [160]. The CoFe$_2$O$_4$@SiO$_2$ nanopigment enhances the corrosion protection performance of the coating better than CoFe$_2$O$_4$, by its appropriate dispersion in the coating and by filing and blocking the free cavities and all electrolyte pathways in the coating. Besides, it releases inhibitive species (Co$^{2+}$ cations) in the scratched area and restricts corrosion inhibition at the coating/metal interface. The incorporation of CoFe$_2$O$_4$@SiO$_2$ nanopigment into epoxy coatings results in a noticeable improvement of the coating corrosion protection [160]. Furthermore, the corrosion protection of pigments, such as diatomite, talc, wollastonite, and kaolin, was considerably improved using the surface treatment with a layer of ZnFe$_2$O$_4$ on the particle surface. Low amounts (16–20 wt.%) of ZnFe$_2$O$_4$ used for the surface treatment also enhance the corrosion protection [157].
3.8. Sensors

Sensors are devices used for signaling changes that appear in a specific material in a certain environment [40]. Sensors based on ferrite NPs are highly sensitive, have low detection limits, and a high signal to noise ratio [40]. One of the most frequent uses of sensors is for detecting changes of humidity. The monitoring of humidity is frequently used in industrial and domestic environments in order to assure human comfort and the appropriate storage of various goods, as well as the optimal functioning condition for industrial process and various instruments [161]. Generally, the humidity-sensing phenomenon is based on the surface effect of water vapor–solid interaction. The ceramic type humidity sensors based on metal oxides displayed superior advantages comparing to polymer films in terms of physical stability mechanical strength, thermal capability, and resistance to chemical attack, making them promising materials for electrochemical humidity sensor applications. By water adsorption, the electrical properties (i.e., capacitance, resistance or electrolytic conduction) of the ceramic surfaces change. In this regard, by increasing the humidity, the conductivity increases, resulting in a higher dielectric constant [145]. A ceramic thick film humidity sensor of MnZn ferrite based on interdigitated electrodes has been reported by Arshaka et al. [162].

The humidity sensing efficiency of a material depends on its microstructural characteristics that are related to the synthesis method [161]. In this sense, the higher sensitivity of ZnFe$_2$O$_4$ can be attributed to the small grain size, high surface area accessible for adsorption of water vapor and is correlated with the low barrier height. Nanosized CoFe$_2$O$_4$, CuFe$_2$O$_4$ and NiFe$_2$O$_4$ have also been demonstrated to be good sensors for detecting oxidizing gases such as chlorine [60]. Owing to its high $H_C$ values and stability, the sensors and actuators made by CoFe$_2$O$_4$ are more durable and of extensive use [7]. MagnetostRICTive CoFe$_2$O$_4$ composites attracted interest for the development of magnetoelastic sensors based on high magnetostriiction, response to applied stress, chemical inertness and low cost [163]. Furthermore, due to the bifunctional nature in terms of stress sensing and actuation or contraction under the influence of a magnetic field, CoFe$_2$O$_4$ has been classified as a smart material with various technological applications among position sensors [120].

Fiber optic biosensors offer great advantages over conventional sensors. Due to its technological advantages (high stability, possibility to be easily removed from the reaction medium, and reuse), the NPs immobilized in glucose oxidase are of great significance for the development of fiber optic glucose oxidase sensors, widely used in food technology, fermentation products, and glucose biosensor [164]. Besides, the immobilization of glucose oxidase sensor on functionalized CoFe$_2$O$_4$@SiO$_2$ NPs via cross-linking with glutaraldehyde can be an effective way to produce the ideal immobilized enzyme [164].

A carbon paste electrode containing ZnFe$_2$O$_4$ NPs was proven to have high sensitivity and good reproducibility in detecting trace levels of 5-fluorourcile in drug samples [165].

3.9. Dielectric Applications

Generally, the dielectric structure involves well conducting grains, separated by low conductivity grain boundaries [166,167]. The dielectric properties of spinel ferrites are determined by structural homogeneity, cation distribution, particle size, density, and porosity [167]. These properties are highly dependent on the synthesis method and thermal treatment parameters, such as temperature, duration, or heating and cooling rate [74].

In case of CoFe$_2$O$_4$, the small Co$^{2+}$ ion occupying the tetrahedral site, reduces the lattice constant, facilitates the electron hopping between Fe$^{3+}$ and Fe$^{2+}$ ions and acts as a source of charge carrier enhancing the dielectric behavior of CoFe$_2$O$_4$ [8]. At low frequencies, the polarization in CoFe$_2$O$_4$ is determined by the hopping of charge carriers, that accumulates and produces polarization upon reaching the grain boundaries, leading to high dielectric constants. In contrast, at the higher frequencies range, the hopping of charge carriers is not able to follow the alternating current induced field, and are thus incompletely polarized, resulting low dielectric constants [166,167]. The dielectric constant decreases by increasing the grain size, which further decreases the grain boundary between the
small grains [118]. The dielectric constant is high at lower frequencies, and decreases with increase of frequency [167]. This decrease becomes constant beyond a certain frequency as only the dielectric polarization contributes to dielectric constant [25]. The dielectric constant gradually also increases with increasing temperature due to dielectric polarization [168]. At low temperature, the charge carriers are unable to orient along the applied electric field direction, therefore the polarization is weak and the dielectric constant is very low [168]. However, as temperature increases, a high number of charge carriers liberate and contribute to polarization, that further increases the dielectric constant [168]. The charge carriers exchange between different valence states of elements present in nanosized ferrites highly influences the polarization [166]. High dielectric constants at high frequencies are attributed to the occurrence of space charge polarization due to inhomogeneous dielectric structure of grain size and impurities [169]. Gopalan et al. [170] reported lower dielectric constant values of CoFe$_2$O$_4$ NPs prepared by the sol-gel method compared to bulk CoFe$_2$O$_4$. The CoFe$_2$O$_4$@SiO$_2$ NCs with controlled magnetic and dielectric properties are promising candidates for biological and high frequency applications [30]. CuFe$_2$O$_4$ is a potential candidate for microwave devices due to the adequate magnetic and dielectric properties at high frequency [27].

NiFe$_2$O$_4$ has also a dielectric structure with grains and grain boundaries of different conducting properties [171]. The exchange electron between Fe$^{2+}$ and Fe$^{3+}$ ions and the hole that transfer between Ni$^{3+}$ and Ni$^{2+}$ ions assure the electrical conduction and dielectric polarization [171]. By increasing the frequencies, the electron/hole exchange frequency will not be able to follow the applied electric field, resulting in lower polarization [171].

3.10. Antimicrobial Applications

Iron plays a key role in microbial pathogenesis, many organisms using Fe sequestration as a defense against infection [172]. The Fe(II)- and Fe(III)-resistant microorganisms showed unexpected resistance to a range of antibiotics such as ampicillin, chloramphenicol, rifampicin, sulfanilamide and tetracycline. Ferrite NPs exhibit good antibacterial properties against *Bacillus cereus*, *Escherichia coli*, *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Serratia marcescens*, and *Candida albicans*. The strain of *Escherichia coli* was capable of evolving resistance to excess Fe mainly through changes in the uptake of Fe [172].

CoFe$_2$O$_4$ NPs are well-known candidates for biomedical applications due to the antimicrobial activity against many human pathogens [55]. The high surface-to-volume ratios and nanoscale size of CoFe$_2$O$_4$ NPs improves their reaction with the pathogenic microbes [173]. CoFe$_2$O$_4$@SiO$_2$@Ag nanocomposite and its combination with Streptomycin exhibited significant antibacterial activity against both Gram-positive and Gram-negative bacteria. The deposition of Ag NPs on CoFe$_2$O$_4$@SiO$_2$ can prevent agglomeration. Moreover, it can be easily recovered from solution after disinfection, by applying of an external magnetic field due to the presence of magnetic core in the composite [174].

CoFe$_2$O$_4$ adheres to the membranes of microorganisms, thus increasing the lag stage of the bacterial growth period, spreading the production time of microorganisms and increasing the bacterial cell division. The antimicrobial activity of CoFe$_2$O$_4$ is due to its produced effective oxides (i.e., superoxide (O$_2^-$) and hydrogen peroxide (H$_2$O$_2$)) [173]. The ferrite NPs antibacterial effect on unicellular fungi is dependent on the diffusion flow of an effective oxide within the bacteria cell surface [173].

The antimicrobial activity of CoFe$_2$O$_4$ NPs was studied against multidrug resistant clinical pathogens (*Staphylococcus aureus*, *Escherichia coli*, *Candida parapsilosis* and *Candida albicans*) by assessing the colony forming units [175,176]. Furthermore, CoFe$_2$O$_4$ NPs functionalized with oleine (CoFe$_2$O$_4@$Ole) and lysine (CoFe$_2$O$_4@$Lys) demonstrated high efficiency against all tested microorganisms most probably due to (i) CoFe$_2$O$_4@$Lys NPs can easily interact electrostatically with negatively charged *S. aureus* and *E. coli* cell walls, causing destruction of the cytoplasm; (ii) positively charged CoFe$_2$O$_4@$Lys NPs and negatively charged CoFe$_2$O$_4@$Ole NPs by can affect bacteria cell homoeostasis. The high antimicro-
bial efficiency of CoFe$_2$O$_4$@Ole NPs could be due to their higher stability compared to CoFe$_2$O$_4$@Lys NPs or to a negative curvature wrapping of anionic membranes [175].

The ZnFe$_2$O$_4$ nanomaterials also display good antibacterial activity against Lactobacillus, Bacillus cereus (Gram-positive), Escherichia coli, Aeromonas hydrophila and Vibrio harveyi (Gram-negative) bacterial strains [145,146,155]. The low crystallite size and surface area play a critical role in the antimicrobial activity against tested pathogenic microbes. The ZnFe$_2$O$_4$ NPs had no activity against Escherichia coli and was only active against Staphylococcus aureus and Pseudomonas aeruginosa [173].

NiFe$_2$O$_4$ had no activity against E. coli, but was active against S. aureus and P. aeruginosa [176]. In addition, the antibacterial effect of NiFe$_2$O$_4$@carbon nanocomposite on the degradation of P. aeruginosa bacteria was rapid and sensitive [95].

### 3.11. Biomedical Applications

The biomedical applications require non-agglomerated and stable aqueous dispersion of magnetic NPs with high $M_S$ values and good biocompatibility [177]. The magnetic NPs can be highly effective for magnetic fluid hyperthermia, drug release and thermal excitation of metabolic pathways within a single cell [178]. In this regard, the MnFe$_2$O$_4$ NPs have attracted considerable attention in biomedicine due their easy synthesis process, controllable size, high magnetization value, superparamagnetic nature, ability to be monitored by external magnetic field, surface manipulation capability, and greater biocompatibility [3,179]. Moreover, the surface modification of MnFe$_2$O$_4$ NPs with mesoporous SiO$_2$ or its incorporation into mesoporous SiO$_2$ nanosphere could play an important role in the stabilization of NPs in water, in the enhancement of its biocompatibility, and in reducing the agglomeration and degradation of MnFe$_2$O$_4$, respectively [3].

The physical characteristics of CoFe$_2$O$_4$, especially high stability, led to extensive attention regarding potential biomedical applications [173]. Generally, CoFe$_2$O$_4$ has been used for drug delivery, imaging factor and therapy of brain tumors [173]. At high concentrations of CoFe$_2$O$_4$ NPs, metal ions are released and some of them enter the cells leading to cytotoxicity, according to the Trojan horse mechanism. A possible explanation could be the prevention of cell transcription and protein synthesis and subsequent altered cellular function [31]. The biocompatible SiO$_2$ coating could remove or reduce the adverse effects. By coating CoFe$_2$O$_4$ NPs, their adhesion decreased due to the negative charge of SiO$_2$ matrix, and they displayed superparamagnetic behavior with a low $M_S$ value and higher compatibility, which made them suitable for various medical applications (drug delivery, specifically cancer cells, magnetic resonance imaging contrast agent for cancer diagnosis) [31,180].

Hyperthermia is a noninvasive treatment procedure for oncological pathologies in which both healthy and carcinoma cells of a living tissue are exposed to necrosis, by prolonged overheating (>43 $^\circ$C) [181]. The existing tumor heating methods (hot water, microwave, infrared ray, ultraviolet ray, etc.) are ineffective for deep-seated cancers, but the magnetic induction hyperthermia, i.e., via magnetic NPs, may be effective in these types of tumor due to the selective heating and destruction of tumor tissue, with minimum collateral damage [178,182]. Magnetic hyperthermia is an emerging adjuvant therapy for malign tumors, where the appropriate magnetic NPs are located under a magnetic field and the heat released may remove the cancerous tissue, at an optimum temperature of 41–46 $^\circ$C [15,123]. Ferrimagnetic materials own hysteretic properties under time-varying magnetic field, which lead to magnetically induced heating [183]. Moreover, the magnetic NPs have low toxicity, are biocompatible and well-tolerated by the living organisms, leading to a simple and fast analysis of specific absorption rate [177]. Some studies reported an effective technique of inserting ferrimagnetic NPs in a tumor region based on the magnetophoresis, a phenomenon caused by a magnetic field gradient on magnetically induced magnetic moment of particle [180,183]. Moreover, magnetophoresis is used in numerous commercial and industrial processes for the separation of magnetic NPs dangling in fluids [183].
However, the main challenge of hyperthermia is to lower the damage to nearby normal tissues. In this regard, NPs must be concentrated at the tumor site rather than in the tumor surroundings [50]. The uncoated magnetic NiFe$_2$O$_4$ NPs have the potential to be used in cell differentiable hyperthermia agents due to their high value of cell survival rate (85%) and non-cytotoxicity under different pH levels (pH = 7 normal cell and pH = 6 tumor cell) [15]. Further, ZnFe$_2$O$_4$ is a good candidate for hyperthermia due to its low toxicity [15,123]. The nanosized ferrite spinel’s enzyme-like activities include peroxidase, oxidase, and catalase, and their applications as enzyme mimetics in biosensing, molecular detection, cancer therapy, and drug delivery were recently reviewed [184].

The hyperthermia measurements CoFe$_2$O$_4$ and MnFe$_2$O$_4$ NPs were performed under various conditions, in order to normalize the obtained results by removing their dependence on magnetic field frequency and amplitude. MnFe$_2$O$_4$ NPs obtained in gelatin medium at 175 °C display the best reported heating efficiency, with values comparable to those of commercial magnetic NPs [185]. CoFe$_2$O$_4$ NPs with narrow size distribution and small particle size (~10 nm), and hence small hydrodynamic diameter, are successfully used to produce high magnetic hyperthermia within short duration [181]. Moreover, the magnetic CoFe$_2$O$_4$@MnFe$_2$O$_4$ NPs have proven to be superior to the individual NPs for producing effective heating in hyperthermia [178].

ZnFe$_2$O$_4$ type bioactive glass ceramics possess both the magnetic properties that generate adequate heat and the ability to bond to natural tissues (via hydroxyapatite layer) [182]. Consequently, this type of materials can be used for the hyperthermia treatment of cancer, but also as a substitute for a cancerous/ damaged bone. Such magnetic heat generation materials depend on various factors such as the structure and magnetic characteristics of the material, strength and frequency of alternating magnetic field, quantity of implant, etc. [182].

The synthesis of CoFe$_2$O$_4$@DMSA/DOX NPs to improve the efficiency of magnetic CoFe$_2$O$_4$ in therapeutic applications, by functionalizing its surface with meso-2,3-dimercaptosuccinic acid, followed by conjugation with the anticancer agent, doxorubicin was also reported [186]. As expected, the experimental results demonstrate that the combined thermal and chemotherapy effect induced by CoFe$_2$O$_4$@DMSA/DOX NPs displays an excellent antitumor efficacy by mitochondrial membrane disruption [186]. The well dispersed spherical magnetic ZnFe$_2$O$_4$ NPs exhibit a better heat efficiency compared to the aggregated polyhedral magnetic ZnFe$_2$O$_4$ NPs, being able to produce a threshold hyperthermia temperature of 42–45 °C in a short time, a key feature for magnetic hyperthermia applications [187].

The cytotoxicity of NPs considered as potential drug delivery systems must be taken into account, especially since it may vary with the type of cell line, exposure dosage, exposure time, particle size and aggregation tendency [188]. The in vitro estimation of cytotoxicity using different cancer cell lines (HeLa, cervical cancer cell and PC-3, prostate cancer cell) revealed good biocompatibility of NiFe$_2$O$_4$ NPs synthesized by co-precipitation and subsequent thermal annealing and less toxic effect to normal cell L929 [188].

The superparamagnetic response and biocompatibility of magnetic NPs make them potential candidates as contrast agents in magnetic resonance imaging (MRI) and tracer agents in magnetic particle imaging (MPI). Due to its soft magnetic behavior, NiFe$_2$O$_4$ is used as a contrast agent in MRI [189]. NiFe$_2$O$_4$@PAA composites (PAA - polyacrylic acid) could be considered as potential tracer agents for MPI, outperforming the commercial, commonly used tracer agents due to the minimum relaxation time of 3.10 µs and high resolution of 7.75 mT, [189]. Besides its use in photothermal and sonodynamic therapy of melanoma, MnFe$_2$O$_4$/C nanocomposite may also be used as a novel theranostic agent in nanomedicine [190]. In this regard, the intratumoral injection of MnFe$_2$O$_4$/C may induce deep tumor tissue necrosis, while the imaging studies endorse MnFe$_2$O$_4$/C as a contrast agent for MRI with a dose-dependent decrease in the signal intensity [189,190]. However, comprehensive studies on the in vitro and in vivo biodegradability and pharma-
cokinetics of MnFe$_2$O$_4$/C nanocomposite are necessary for additional investigations and evaluation [189,190].

4. Conclusions

Nanosized CoFe$_2$O$_4$, MnFe$_2$O$_4$, ZnFe$_2$O$_4$, NiFe$_2$O$_4$, and CuFe$_2$O$_4$ NPs have received impressive attention in the last decade due to their special properties, such as chemical, mechanical, and thermal stability, large coercivity, high anisotropy constant and Curie temperature, moderate saturation magnetization, high electrical resistance, and low eddy current loss. Amongst all the reviewed synthesis methods, the sol-gel and chemical co-precipitation techniques stand as superlative routes for synthesizing fine, homogenous, nanostructured ferrites. However, several unconventional methods allow the cost-efficient preparation of high-quality NPs. Although bulk ferrites remain a key group of magnetic materials, due to their advantageous properties, the investigated nanostructured ferrites are used in many areas, including material sciences, engineering, physics, chemistry, biology, and medicine. Notably, these magnetic ferrite NPs have displayed outstanding results in the field of medicine because of their low toxicity, biocompatibility, and ability to be easily manipulated using a magnetic field.

Author Contributions: T.D., E.A.L. and O.C. conceived and designed the work; T.D., E.A.L. and O.C. wrote the manuscript. All authors have read and agreed to the published version of the manuscript.

Funding: This work was supported by a grant of the Romanian National Authority for Scientific Research CNCS-UEFISCDI, project number PN-III-P2-2.1-PED-2019-3664.

Data Availability Statement: The data presented in this study are available on request from the corresponding author.

Conflicts of Interest: The authors declare no conflict of interest. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results.

References

1. Chand, P.; Vaish, S.; Kumar, P. Structural, optical and dielectric properties of transition metal (MFe$_2$O$_4$; M = Co, Ni and Zn) nanoparticles. Phys. B Condens Matter. 2017, 524, 53–63. [CrossRef]
2. Jeevanandam, J.; Barhoum, A.; Chan, Y.S.; Dufresne, A.; Danquah, M.K. Review on nanoparticles and nanostructured materials: History, sources, toxicity and regulations. Beilstein. J. Nanotechnol. 2018, 9, 1050–1074. [CrossRef]
3. Asghar, K.; Qasim, M.; Das, D. Preparation and characterization of mesoporous magnetic MnFe$_2$O$_4$m-SiO$_2$ nanocomposite for drug delivery application. Mater. Today Proc. 2020, 28, 87–93. [CrossRef]
4. Sivakumar, P.; Ramesh, R.; Ramanand, A.; Ponnumas, Y.; Muthamizhchelvan, C. Synthesis and characterization of NiFe$_2$O$_4$ nanoparticles and nanorods. J. Alloy Comp. 2013, 563, 6–11. [CrossRef]
5. Ozçelik, B.; Ozçelik, S.; Amaveda, H.; Santos, H.; Borrell, C.J.; Saez-Puche, R.; de la Fuente, G.F.; Angurel, L.A. High speed processing of NiFe$_2$O$_4$ spinel using laser furnace. J. Materiomics 2020, 6, 661–670. [CrossRef]
6. Džunuzović, A.S.; Ilić, N.I.; Vijatović Petrović, M.M.V.; Bobić, J.D.; Stojadinović, B.; Dohčević-Mitrović, Z.; Stojanović, B.D. Structure and properties of Ni-Zn ferrite obtained by auto-combustion method. J. Magn. Magn. Mater. 2018, 374, 245–251. [CrossRef]
7. Kaur, H.; Singh, A.; Kumar, A.; Ahlawat, D.S. Structural, thermal and magnetic investigations of cobalt ferrite doped with Zn$^{2+}$ and Cd$^{2+}$ synthesized by auto combustion method. J. Magn. Magn. Mater. 2019, 474, 505–511. [CrossRef]
8. Naik, A.B.; Naik, P.P.; Hasolkar, S.S.; Naik, D. Structural, magnetic and electrical properties along with antifungal activity & adsorption ability of cobalt doped manganese ferrite nanoparticles synthesized using combustion route. Ceram. Int. 2020, 46, 21046–21055.
9. Xiong, Q.Q.; Tu, J.P.; Shi, S.J.; Liu, X.Y.; Wang, X.L.; Gu, C.D. Ascorbic acid-assisted synthesis of cobalt ferrite (CoFe$_2$O$_4$) hierarchical flower-like microspheres with enhanced lithium storage properties. J. Power Sources 2014, 256, 153–159. [CrossRef]
10. Li, X.; Sun, Y.; Zong, Y.; Wei, Y.; Liu, X.; Li, X.; Peng, Y.; Zheng, X. Size-effect induced cation redistribution on the magnetic properties of well-dispersed CoFe$_2$O$_4$ nanocrystals. J. Alloy Comp. 2020, 841, 155710. [CrossRef]
11. Torkian, S.; Ghasemi, A.; Razavi, R.S. Cation distribution and magnetic analysis of wideband microwave absorptive Co$_x$Ni$_{1-x}$Fe$_2$O$_4$ ferrites. Ceram. Int. 2017, 43, 6987–6995. [CrossRef]
12. Mahala, C.; Sharma, M.D.; Basu, M. 2D nanostructures of CoFe$_2$O$_4$ and NiFe$_2$O$_4$: Efficient oxygen evolution catalyst. Electrochim. Acta 2018, 273, 462–473. [CrossRef]
13. Manikandan, A.; Sridhar, R.; Arul, S.A.; Ramakrishna, S. A simple aloe vera plant-extracted microwave and conventional combustion synthesis: Morphological, optical, magnetic and catalytic properties of CoFe$_2$O$_4$ nanostructures. J. Molec. Struct. 2014, 1076, 188–200. [CrossRef]

14. Salunkhe, A.B.; Khot, V.M.; Phadatate, M.R.; Thorat, N.D.; Joshi, R.S.; Yadav, H.M.; Pawar, S.H. Low temperature combustion synthesis and magnetostuctural properties of Co-Mn nanoferrites. J. Magn. Magn. Mater. 2014, 352, 91–98. [CrossRef]

15. Ghayour, H.; Abdellahie, M.; Ozada, N.; Jabbrzare, S.; Khandan, A. Hyperthermia application of zinc doped nickel ferrite nanoparticles. J. Phys. Chem. Solids 2017, 111, 464–472. [CrossRef]

16. Kremenovic, A.; Antić, B.; Vulić, P.; Blanuša, J.; Tomić, A. ZnFe$_2$O$_4$ antiferromagnetic structure redetermination. J. Magn. Magn. Mater. 2017, 426, 264–266. [CrossRef]

17. Manikandan, A.; Durka, M.; Arul, S.A. Magnetically recyclable spinel Mn$_3$Zn$_{1-x}$Fe$_2$O$_4$ (0.0 ≤ x ≤ 0.5). Adv. Sci. Eng. Med. 2015, 7, 33–46. [CrossRef]

18. Ge, Y.-C.; Wang, Z.-L.; Yi, M.-Z.; Ran, L.-P. Fabrication and magnetic transformation from paramagnetic to ferrimagnetic of ZnFe$_2$O$_4$ hollow spheres. Trans. Nonferrous Met. Soc. China 2019, 29, 1503–1509.

19. Feng, S.; Yang, W.; Wang, Z. Synthesis of porous NiFe$_2$O$_4$ microparticles and its catalytic properties for methane combustion. Mater. Sci. Eng. B 2011, 176, 1509–1512. [CrossRef]

20. Alarifi, A.; Deraz, N.M.; Shaban, S. Structural, morphological and magnetic properties of NiFe$_2$O$_4$ nano-particles. J. Alloy Comp. 2009, 486, 501–506. [CrossRef]

21. Shamugavel, T.; Raj, S.G.; Rajarajan, G.; Kumar, G.R. Tailoring the structural and magnetic properties of NiFe$_2$O$_4$ nanoparticles synthesized by a sonochemical method. Mater. Today Proc. 2020, 33, 30–45. [CrossRef]

22. Iqbal, M.J.; Yaqub, N.; Sepiol, B.; Ismail, B. A study of the influence of crystallite size on the electrical and magnetic properties of NiFe$_2$O$_4$. Solid State Sci. 2013, 111, 1837–1842. [CrossRef]

23. Shetty, K.; Renuka, L.; Nagaswarupa, H.P.; Prashantha, S.C.; Ashwini, K. A comparative study on CuFe$_2$O$_4$, ZnFe$_2$O$_4$ and NiFe$_2$O$_4$: Morphology, impedance and photocatalytic studies. Mater. Today Proc. 2017, 4, 11806–11815. [CrossRef]

24. Qibl, M.J.; Yaqub, N.; Sepiol, B.; Ismail, B. A study of the influence of crystallite size on the electrical and magnetic properties of CuFe$_2$O$_4$. Mater. Res. Bull. 2011, 46, 1837–1842. [CrossRef]

25. Dar, M.A.; Varshney, D. Effect of d-block element Co$^{2+}$ substitution on structural, Mössbauer and dielectric properties of spinel copper ferrite. J. Magn. Magn. Mater. 2014, 436, 101–112. [CrossRef]

26. Mohanty, D.; Mallick, P.; Biswall, S.K.; Behera, B.; Mohapatra, R.K.; Behera, A.; Satpathy, S.K. Investigation of structural, dielectric and electrical properties of ZnFe$_2$O$_4$. Mater Today Proc. 2020, 33, 4971–4975. [CrossRef]

27. Mohanty, D.; Satpathy, S.K.; Behera, B.; Mohapatra, R.K. Dielectric and frequency dependent transport properties in magnesium doped CuFe$_2$O$_4$ composite. Mater Today Proc. 2020, 33, 5226–5231. [CrossRef]

28. Sivakumar, A.; Dhas, S.S.J.; Dhas, S.A.M.B. Assessment of crystallographic and magnetic phase stabilities on MnFe$_2$O$_4$ nanocrystalline materials at shocked conditions. Solid State Sci. 2020, 107, 106340. [CrossRef]

29. Junlabhut, P.; Nuthongkum, P.; Pecharap, W. Influences of calcination temperature on structural properties of MnFe$_2$O$_4$ nanodispersers synthesized by co-precipitation method for reusable absorbent materials. Mater Today Proc. 2018, 5, 13857–13864. [CrossRef]

30. Nadeem, K.; Raut, S.D.; Sangale, S.; Kadam, A.S. Ferrite nanofibers: Synthesis methods. In Nanomaterials 2021, 11, 1560.

31. Vedrttnam, A.; Kalauni, K.; Dubey, S.; Kumar, A. A comprehensive study on structure, properties, synthesis and characterization of ferrites. AIMS Mat. Sci. 2020, 7, 800–835. [CrossRef]

32. Vinosha, P.A.; Manikandan, A.; Preetha, A.C.; Dinesh, A.; Srimani, Y.; Almessiere, M.A.; Baykal, A.; Xavier, B.; Nirmala, F.G. Review on recent advances of synthesis, magnetic properties, and water treatment applications of cobalt ferrite nanoparticles and nanocomposites. J. Supercond. Nov. Magn. 2021, 34, 995–1018. [CrossRef]

33. Masunga, N.; Mmelesi, O.K.; Kefeni, K.K.; Mamba, B.B. Recent advances in copper ferrite nanoparticles and nanocomposites synthesis, magnetic properties and application in water treatment: Review. J. Environ. Chem. Eng. 2019, 7, 103179. [CrossRef]

34. Kumar, M.; Doanjhi, S.J.; Singh, J.; Monir, K.; Singh, H. Review on magnetic nano ferrites and their composites as an alternative in waste water treatment: Synthesis, modifications and applications. Environ. Sci. Water Res. Technol. 2020, 6, 491–514. [CrossRef]

35. Kefeni, K.K.; Mamba, B.B. Photocatalytic application of spinel ferrite nanoparticles and nanocomposites in wastewater treatment: Review. Sustain. Mater. Tech. 2020, 23, e00140. [CrossRef]

36. Kharisov, B.I.; Rasika Dias, H.V.; Kharissova, O.V. Mini-review: Ferrite nanoparticles in the catalysis. Arab. J. Chem. 2019, 12, 1234–1246. [CrossRef]
39. Dalawai, S.P.; Kumar, S.; Al Saad Aly, M.; Khan, M.Z.H.; Xing, R.; Vasambekar, P.N.; Liu, S. A review of spinel-type of ferrite thick film technology: Fabrication and application. J. Mat. Sci. Mat. Electr. 2019, 30, 7752–7779.

40. Kefeni, O.; Dippong, T.; Deac, I.G. Formation, structure and magnetic properties of Ni$_{0.5}$Co$_{0.5}$Fe$_2$O$_4$ nanocomposites by sol-gel method using different polyols. J. Magn. Magn. Mater. 2020, 498, 166168. [CrossRef]

41. Dippong, T.; Levei, E.A.; Deac, I.G. Microstructure, porosity and magnetic properties of Zn$_{0.5}$Co$_{0.5}$Fe$_2$O$_4$/SiO$_2$ nanocomposites prepared by sol-gel method using different polyols. J. Magn. Magn. Mater. 2020, 498, 166168. [CrossRef]

42. Dippong, T.; Levei, E.A.; Deac, I.G.; Neag, E.; Cadar, O. Influence of Cu$^{2+}$, Ni$^{2+}$, and Zn$^{2+}$ ions doping on the structure, morphology, and magnetic properties of Co-ferrite embedded in SiO$_2$ matrix obtained by an innovative sol-gel route. Nanomaterials 2020, 10, 580. [CrossRef]

43. Dippong, T.; Levei, E.A.; Deac, I.G.; Goga, F.; Cadar, O. Investigation of structural and magnetic properties of Ni$_{x}$Zn$_{1-x}$Fe$_2$O$_4$/SiO$_2$ (0 ≤ x ≤ 1) spinel-based nanocomposites. J. Anal. Appl. Pyrol. 2019, 144, 104713. [CrossRef]

44. Hashim, M.; Alimuddin; Kumar, S.; Koo, B.H.; Shirsath, S.E.; Mohammed, E.M.; Shah, J.; Kotnala, R.K.; Choi, H.K.; Chung, H.; et al. Structural, electrical and magnetic properties of Co–Cu ferrite nanoparticles. J. Alloy. Comp. 2012, 518, 11–18. [CrossRef]

45. Houshiar, M.; Zebhi, F.; Razi, Z.J.; Alidoust, A.; Askari, Z. Synthesis of cobalt ferrite (CoFe$_2$O$_4$) nanoparticles using solid state, citrate precursor and polymerized complex methods: A comparative study. J. Alloys Comp. 2019, 474, 56–62. [CrossRef]

46. Ajinka, N.; Yu, X.; Kaithal, P.; Luo, H.; Somani, P.; Ramakrishna, S. Magnetic iron oxide nanoparticle (IONP) synthesis to applications: Present and Future. Materials 2020, 13, 4644. [CrossRef]

47. Amirabadizadeh, A.; Salighe, Z.; Sarhaddi, R.; Lotfollahi, Z. Synthesis of ferrofluids based on cobalt ferrite nanoparticles: Influence of reaction time on structural, morphological and magnetic properties. J. Magn. Mater. 2017, 434, 78–85. [CrossRef]

48. Yousefi, M.H.; Manouchehri, S.; Arab, A.; Mozaffari, M.; Amighian, J. Preparation of cobalt–zinc ferrite (Co$_{0.8}$Zn$_{0.2}$Fe$_2$O$_4$) nanopowder via combustion method and investigation of its magnetic properties. Mat. Res. Bull. 2010, 45, 1792–1795. [CrossRef]

49. Baykal, A.; Kasapoglu, N.; Koseoğlu, Y.; Başaran, A.C.; Kavas, H.; Toprak, M.S. Microwave-induced combustion synthesis and characterization of Ni$_x$Co$_{1-x}$Fe$_2$O$_4$ Nanocrystals (x = 0.0, 0.4, 0.6, 0.8, 1.0). Cent. Eur. J. Chem. 2008, 6, 125–130. [CrossRef]

50. Naik, M.M.; Naik, H.S.B.; Nagaraju, G.; Vinuth, M.; Vinu, K.; Viswanath, R. Green synthesis of zinc doped cobalt ferrite nanoparticles: Structural, optical, photocatalytic and antibacterial studies. Nano Struct. Nano Object. 2019, 19, 100322. [CrossRef]

51. Bakhshi, H.; Vahdati, N.; Sedghi, A.; Mozharivskyj, Y. Comparison of the effect of nickel and cobalt cations addition on the structural and magnetic properties of manganese-zinc ferrite nanoparticles. J. Magn. Magn. Mater. 2020, 474, 56–62. [CrossRef]

52. Houshier, M.; Zebhi, F.; Razi, Z.J.; Alioudst, A.; Askari, Z. Synthesis of cobalt ferrite (CoFe$_2$O$_4$) nanoparticles using combustion, coprecipitation, and precipitation methods: A comparison study of size, structural and magnetic properties. J. Magn. Mater. 2014, 397, 43–48. [CrossRef]

53. Visnosa, P.A.; Minikandan, A.; Cecilia, A.S.J.; Dinesh, A.; Nirmala, G.F.; Preetha, A.C.; Simlani, Y.; Almessiere, M.A.; Baycal, A.; Xavier, B. Review on recent advances of zinc substituted cobalt ferrite nanoparticles: Synthesis characterization and diverse applications. Ceram. Int. 2021, 47, 10512–10535. [CrossRef]

54. Thakur, A.; Chahar, D.; Taneja, S.; Bhalla, N.; Thakur, A. A review on MnZn ferrites: Synthesis, characterization and applications. Ceram. Int. 2016, 42, 15374–15386. [CrossRef]

55. Jeseentharani, V.; George, M.; Jeyaraj, B.; Dayalan, A.; Nagaraj, K.S. Synthesis of metal ferrite (MFe$_2$O$_4$, M = Co, Cu, Mg, Ni, Zn) nanoparticles as humidity sensor materials. J. Exp. Nanosci. 2013, 8, 358–370. [CrossRef]

56. Varma, P.C.R.; Manna, R.S.; Benerjee, D.; Varma, M.R.; Suresh, K.G.; Nigam, A.K. Magnetic properties of CoFe$_2$O$_4$ synthesized by solid state, citrate precursor and polymerized complex methods: A comparative study. J. Alloys Comp. 2008, 453, 298–303. [CrossRef]

57. Dalawai, S.P.; Kumar, S.; Al Saad Aly, M.; Khan, M.Z.H.; Xing, R.; Vasambekar, P.N.; Liu, S. A review of spinel-type of ferrite thick film technology: Fabrication and application. J. Mat. Sci. Mat. Electr. 2019, 30, 7752–7779.
64. Atif, M.; Turtelli, R.S.; Grössinger, R.; Siddique, M.; Nadeem, M. Effect of Mn substitution on the cation distribution and temperature dependence of magnetic anisotropy constant in Co$_{1-x}$Mn$_x$Fe$_2$O$_4$ (0.0 ≤ x ≤ 0.4) ferrites. *Ceram. Int.* 2014, 40, 471–478. [CrossRef]

65. Aslibeiki, B.; Kameli, P.; Salamati, H.; Eshraghi, M.; Tahmasebi, T. Superspin glass state in MnFe$_2$O$_4$ nanoparticles. *J. Magn. Magn. Mater.* 2010, 322, 2929–2934. [CrossRef]

66. Chia, C.H.; Zakaria, S.; Yousif, M.; Goh, S.C.; Hawa, C.Y.; Ahmad, S.; Huang, N.M.; Lim, H.M. Size and crystallinity-dependent magnetic properties of CoFe$_2$O$_4$ nanocrystals. *Ceram. Int.* 2010, 36, 605–609. [CrossRef]

67. Zahraei, M.; Monshi, A.; del Puerto Morales, M.; Shahbazi-Ghroueli, D.; Amirnasr, M.; Behdadfar, M. Hydrothermal synthesis of fine stabilized superparamagnetic nanoparticles of Zn$_{2+}$ substituted manganese ferrite. *J. Magn. Mater.* 2015, 393, 429–436. [CrossRef]

68. Prabhakaran, T.; Mangalaraja, R.V.; Denardin, J.C.; Jimenez, J.A. The effect of reaction temperature on the structural and magnetic properties of nano CoFe$_2$O$_4$. *Ceram. Int.* 2017, 43, 5599–5606. [CrossRef]

69. Senapati, K.K.; Borgohain, C.; Phukan, P. Synthesis of highly stable CoFe$_2$O$_4$ nanoparticles and their use as magnetically separable catalyst for Knoevenagel reaction in aqueous medium. *J. Mol. Catal. A Chem.* 2011, 339, 24–31. [CrossRef]

70. Singh, A.; Pathak, S.; Kumar, P.; Sharma, P.; Rathi, A.; Basheer, G.A.; Maurya, K.K.; Plant, R.P. Tuning the magnetocrystalline anisotropy and spin dynamics in Co$_x$Zn$_{1-x}$Fe$_2$O$_4$ (0 ≤ x ≤ 1) nanoferrites. *J. Magn. Magn. Mater.* 2020, 493, 165737. [CrossRef]

71. Huixia, F.; Baiyi, C.; Deyi, Z.; Jianqiang, Z.; Lin, T. Preparation and characterization of the cobalt ferrite nano-particles by reverse coprecipitation. *J. Magn. Mater.* 2014, 388, 68–72. [CrossRef]

72. Sajja, M.; Oubaha, M.; Hasanauzzaman, M.; Olabi, A.G. Development of cobalt ferrite nanoparticles prepared by the sol–gel process. *Ceram. Int.* 2014, 40, 1147–1154. [CrossRef]

73. Wang, L.; Mu, M.; Liu, Y.; Li, J.; Liu, M.; Lin, H. The structure, magnetic properties and cation distribution of Co$_{1-x}$Mg$_x$Fe$_2$O$_4$/SiO$_2$ nanocomposites synthesized by sol–gel method. *Ceram. Int.* 2015, 41, 4176–4181. [CrossRef]

74. Zhang, R.; Sun, L.; Wang, Z.; Hao, W.; Cao, E.; Zhang, Y. Dielectric and magnetic properties of CoFe$_2$O$_4$ prepared by sol-gel auto-combustion method. *Mat. Res. Bull.* 2018, 98, 133–138. [CrossRef]

75. Barison, S.; Fabrizio, M.; Fasolini, S.; Montagner, F.; Mortalò, C. A microwave-assisted sol–gel Pechini method for the synthesis of BaCe$_{0.65}$Zr$_{0.35}$O$_{2−δ}$ powders. *Mater. Res. Bull.* 2010, 45, 1171–1176. [CrossRef]

76. Sharma, S.; Kumar Verma, M.; Sharma, N.D.; Choudhary, N.; Singh, S.; Singh, D. Rare-earth doped Ni–Co ferrites synthesized by coprecipitation. *Mater. Res. Bull.* 2017, 89, 477–486. [CrossRef]

77. Dimesso, L. Pechini Processes: An Alternate Approach of the Sol–Gel Method, Preparation, Properties, and Applications. In *Handbook of Sol-Gel Science and Technology*; Klein, L., Aparicio, M., Jitiani, A., Eds.; Springer International Publishing: Berlin/Heidelberg, Germany, 2016; pp. 1–22.

78. Raut, A.V.; Barkule, R.S.; Shengule, D.R.; Jadhav, K.M. Synthesis, structural investigation and magnetic properties of Zn$_{2+}$ substituted cobalt ferrite nanoparticles prepared by the sol–gel auto-combustion technique. *J. Magn. Magn. Mater.* 2014, 358, 87–92. [CrossRef]

79. Mohamed, R.M.; Rashad, M.M.; Haraz, F.A.; Sigmund, W. Structure and magnetic properties of nanocrystalline cobalt ferrite powders synthesized using organic acid precursors. *J. Magn. Magn. Mater.* 2010, 322, 2058–2064. [CrossRef]

80. Barathiraja, C.; Mnikandan, A.; Mohideen Uduman, A.M.; Jayasree, S.; Arul Antony, S. Magnetically recyclable spinel Mn$_{0.65}$Ni$_{0.35}$Fe$_2$O$_4$ nanocrystals. *Ceram. Int.* 2017, 43, 1250–1259. [CrossRef]

81. Sundararajan, M.; Sailaja, V.; Kennedy, L.J.; Vijaya, J.J. Photocatalytic degradation of rhodamine B under visible light using nanostructured zinc doped cobalt ferrite catalysts: Kinetics and mechanism. *Ceram. Int.* 2017, 43, 540–548. [CrossRef]

82. Sundararajan, M.; Kennedy, L.J.; Aruldoss, U.; Pasha, S.K.; Vijaya, J.J.; Dunn, S. Microwave combustion synthesis of zinc substituted nanocrystalline cobalt ferrite: Structural and magnetic studies. *Mat. Sci. Semicond. Proc.* 2015, 40, 1–10. [CrossRef]

83. Singh, R.K.; Rai, B.C.; Prasad, K. Synthesis and Characterization of Copper Substituted Cobalt Ferrite Nanoparticles. *Int. J. Adv. Mater. Sci.* 2012, 3, 71–76.

84. Garcia Cerda, L.A.; Montemayor, S.M. Synthesis of CoFe$_2$O$_4$ nanoparticles embedded in a silica matrix by the citrate precursor technique. *J. Magn. Magn. Mater.* 2005, 294, e43–e46. [CrossRef]

85. El-Foulani, A.-H.; Aamouche, A.; Mohseni, F.; Amoral, J.S.; Tobald, D.M.; Pullar, R.C. Effect of surfactants on the optical and magnetic properties of cobalt zinc ferrite Co$_{0.5}$Zn$_{0.5}$Fe$_2$O$_4$. *J. Alloys Comp.* 2019, 774, 1250–1259. [CrossRef]

86. Naseri, M.G.; Bin Saion, E.; Ahangar, H.A.; Hashim, M.; Shaari, A.H. Synthesis and characterization of manganese ferrite nanoparticles by thermal treatment method. *J. Magn. Magn. Mater.* 2011, 323, 1745–1749. [CrossRef]

87. Chakradhary, V.K.; Ansaria, A.; Akhtar, M.J. Design, synthesis, and testing of high coercivity cobalt doped nickel ferrite nanoparticles for magnetic applications. *J. Magn. Magn. Mater.* 2019, 469, 674–680. [CrossRef]

88. Soundararajan, D.; Kim, K.H. Synthesis of CoFe$_2$O$_4$ magnetic nanoparticles by thermal decomposition. *J. Magn.* 2014, 19, 5–9. [CrossRef]

89. Peddis, D.; Cannas, C.; Musinu, A.; Ardu, A.; Orru, F.; Fiorani, D.; Laureti, S.; Rinaldi, D.; Muscas, G.; Concas, G.; et al. Beyond the effect of particle size: Influence of CoFe$_2$O$_4$ nanoparticle arrangements on magnetic properties. *Chem. Mater.* 2013, 25, 2005–2013. [CrossRef]
Improvement of the properties of MnZn ferrite power cores through spray pyrolysis method. *Solid State Sci.* 2015, **39**, 69–73. [CrossRef]

Saritas, S.; Sakar, B.C.; Turgut, E.; Kundakci, M.; Yildirim, M. Cobalt metal doped magnesium ferrite and zinc ferrite thin films grown by Spray Pyrolysis. *Mater. Today Proc.* 2021. [CrossRef]

Jundale, A.V.; Chorage, G.Y.; Yadav, A.A. Structural, morphological and optical properties of CoFe$_2$O$_4$ thin film by spray pyrolysis technique. *Mater. Today Proc.* 2021, **43**, 2678–2681. [CrossRef]

Kotsikau, D.; Ivanovskaya, M.; Pankov, V.; Fedotova, Y. Structure and magnetic properties of manganese-zinc ferrites prepared by spray pyrolysis method. *Solid State Sci.* 2015, **39**, 69–73. [CrossRef]

Morrison, S.A.; Cahill, C.L.; Carpenter, E.E.; Calvin, S.; Swaminathan, R.; McHenry, M.; Harris, V.G. Magnetic and structural properties of Mn and Co spinel ferrite-silica nanoparticles with tunable saturation magnetization and coercivity cobalt ferrite nanocrystals without subsequent calcination. *Mat. Chim. Phys.* 2017, **41**, 9236–9243. [CrossRef]

Sharifi, I.; Shokrollahi, H.; Doroodmand, M.M.; Safi, R. Magnetic and structural studies on CoFe$_2$O$_4$ nanoparticles. *Ceram. Int.* 2020, **53**, 14021–14027. [CrossRef]

Jia, Z.; Ren, D.; Zhu, R. Synthesis, characterization and magnetic properties of CoFe$_2$O$_4$ nanorods. *Mater. Lett.* 2012, **66**, 128–131. [CrossRef]

Halalai, M.; Mirkazemi, S.M.; Alamolhoda, S. The effect of poly vinyl alcohol (PVA) surfactant on phase formation and magnetic properties of hydrothermally synthesized CoFe$_2$O$_4$ nanoparticles. *J. Magn. Magn. Mater.* 2016, **419**, 363–367. [CrossRef]

Zhao, D.; Wu, X.; Guan, H.; Han, E. Study on supercritical hydrothermal synthesis of CoFe$_2$O$_4$ nanoparticles. *J. Superfluids* 2007, **42**, 226–233. [CrossRef]

Goh, S.C.; Chia, C.H.; Zakaria, S.; Yusoff, M.; Haw, C.Y.; Ahmad, S.; Huang, N.M.; Lim, H.N. Hydrothermal preparation of high saturation magnetization and coercivity cobalt ferrite nanocrystals without subsequent calcination. *Chem. Phys.* 2010, **120**, 31–35. [CrossRef]

Jia, Z.; Ren, D.; Zhu, R. Synthesis, characterization and magnetic properties of CoFe$_2$O$_4$ nanorods. *Mater. Lett.* 2012, **66**, 128–131. [CrossRef]

Pilai, V.; Shah, D.O. Synthesis of high-coercivity cobalt ferrite particles using water-in-oil microemulsions. *J. Magn. Magn. Mater.* 1996, **163**, 243–248. [CrossRef]

Vazquez-Vazquez, C.; Lopez-Quintero, M.A.; Bujan-Nunez, M.C.; Rivas, J. Finite size and surface effects on the magnetic properties of cobalt ferrite nanoparticles. *J. Nanopart Res.* 2011, **13**, 1663–1676. [CrossRef]

Singh, C.; Jauhar, S.; Kumar, S. Cation vacancy and magnetic properties of ZnFe$_2$O$_4$ microspheres. *Phys. B Condens Matter.* 2020, **583**, 412015. [CrossRef]

Abbas, M.; Rao, P.B.; Nazrul Islam, M.; Woo Kim, K.W.; Naga, S.M.; Takahashi, M.; Kim, C. Size-controlled high magnetization CoFe$_2$O$_4$ nanospheres and nanocubes using rapid one-pot sonochemical technique. *Ceram. Int.* 2014, **40**, 3269–3276. [CrossRef]

Singh, C.; Jauhar, S.; Kumar, S.; Singh, J.; Singhal, S. Synthesis of zinc substituted cobalt ferrites via reverse micelle technique involving in situ template formation: A study on their structural, magnetic, optical and catalytic properties. *Mat. Chem. Phys.* 2015, **156**, 188–197. [CrossRef]

Shahid, I.; Shokrollahi, H.; Doroodmand, M.M.; Safi, R. Magnetic and structural studies on CoFe$_2$O$_4$ nanoparticles synthesized by co-precipitation, normal micelles and reverse micelles methods. *J. Magn. Magn. Mater.* 2012, **324**, 1854–1861. [CrossRef]

Abbas, M.; Rao, P.B.; Nazrul Islam, M.; Woo Kim, K.W.; Naga, S.M.; Takahashi, M.; Kim, C. Size-controlled high magnetization CoFe$_2$O$_4$ nanospheres and nanocubes using rapid one-pot sonochemical technique. *Ceram. Int.* 2014, **40**, 3269–3276. [CrossRef]

Kogias, G.; Tsakaloudi, V.; Van der Valk, P.; Zaspalis, V. Improvement of the properties of MnZn ferrite power cores through improvements on the microstructure of the compacts. *J. Magn. Magn. Mater.* 2012, **324**, 235–241. [CrossRef]

Shanmugam, S.; Subramanian, B. Evolution of phase pure magnetic cobalt ferrite nanoparticles by varying the synthesis conditions of polyol method. *Mater. Sci. Eng. B* 2020, **252**, 114451. [CrossRef]

Ibrahim, A.M.; Abd El-Latif, M.M.M.; Mahmoud, M.M. Synthesis and characterization of nano-sized cobalt ferrite prepared via polyol method using conventional and microwave heating techniques. *J. Alloy. Comp.* 2010, **506**, 201–204. [CrossRef]

Selima, S.S.; Khairy, M.; Mousa, M.A. Comparative studies on the impact of synthesis methods on structural, optical, magnetic and catalytic properties of CuFe$_2$O$_4$. *Ceram. Int.* 2019, **45**, 6335–6540. [CrossRef]

Shakil, M.; Inayat, U.; Arshad, M.I.; Nabi, G.; Khalid, N.R.; Tarig, N.H.; Shahd, A.; Iqbal, M.Z. Influence of zinc and cadmium co-doping on optical and magnetic properties of cobalt ferrites. *Ceram. Int.* 2020, **46**, 7767–7773. [CrossRef]

Margarbandhu, M.; Sendhilnathan, S.; Senthikumar, S.; Gajalakshmi, D. Investigation of structural, morphological, magnetic properties and biomedical applications of Cu$_2$+ substituted uncoated cobalt ferrite nanoparticles. *Braz. Arch. Biol. Technol.* 2016, **59**, e16161046. [CrossRef]
116. Chen, Z.-H.; Sun, Y.-P.; Kang, Z.-T.; Chen, D. Preparation of Zn$_x$Co$_{1-x}$Fe$_2$O$_4$ nanoparticles by microwave-assisted ball milling. *Ceram. Int.* 2014, 40, 14687–14692. [CrossRef]

117. Balavijayalakshmi, J.; Suriyanarayanan, N.; Jayaprakash, R. Influence of copper on the magnetic properties of cobalt ferrite nanoparticles. *Mater. Lett.* 2012, 81, 52–54.

118. Jabbar, R.; Sabeeh, S.H.; Hameed, A.M. Structural, dielectric and magnetic properties of Mn+2 doped cobalt ferrite nanoparticles. *J. Magn. Magn. Mater.* 2020, 494, 165726. [CrossRef]

119. Majid, F.; Rauf, J.; Ata, S.; Bibi, I.; Ibrahim, S.M.; Ali, A.; Iqbal, M. Synthesis and characterization of NiFe$_2$O$_4$ ferrite: Sol–gel and hydrothermal synthesis route effect on magnetic, structural and dielectric characteristics. *Mater. Chem. Phys.* 2021, 258, 123888. [CrossRef]

120. Bhame, S.D.; Joy, P.A. Enhanced strain sensitivity in magnetostrictive spinel ferrite Co$_{1-x}$Zn$_x$Fe$_2$O$_4$. *J. Magn. Magn. Mater.* 2018, 447, 150–154. [CrossRef]

121. Ati, A.A.; Othaman, Z.; Samavati, A. Influence of cobalt on structural and magnetic properties of nickel ferrite nanoparticles. *J. Mol. Struct.* 2013, 1052, 177–182. [CrossRef]

122. Ortiz-Quinñonez, J.-L.; Pal, U.; Villanueva, M.S. Structural, magnetic, and catalytic evaluation of spinel Co, Ni, and Co–Ni ferrite nanoparticles fabricated by low-temperature solution combustion process. *ACS Omega* 2018, 3, 14986–15001. [CrossRef]

123. Nasrin, S.; Chowdhury, F.U.Z.; Hoque, S.M. Study of hydrodynamic size distribution and hyperthermia temperature of chitosan encapsulated zinc-substituted manganese nano ferrites suspension. *Phys. B. Cond. Mater.* 2019, 561, 54–63. [CrossRef]

124. Reddy, M.P.; Zhou, X.; Yann, A.; Du, S.; Huang, Q.; Mohamed, A.M.A. Low temperature hydrothermal synthesis, structural investigation and functional properties of Co$_{1-x}$Mn$_x$Fe$_2$O$_4$ (0 ≤ x ≤1) nanoferrites. *Superlattices Microstruct.* 2015, 81, 233–242. [CrossRef]

125. Abdallah, H.K.I.; Moyo, T.; Msoni, J.Z. The effect of annealing temperature on the magnetic properties of Mn$_x$Co$_{1-x}$Fe$_2$O$_4$ ferrites nanoparticles. *J. Supercond. Nov. Magn.* 2012, 25, 2625–2630. [CrossRef]

126. Tomar, D.; Jeevanandam, P. Synthesis of cobalt ferrite nanoparticles with different morphologies via thermal decomposition approach and studies on their magnetic properties. *J. Alloys Comp.* 2020, 843, 155815. [CrossRef]

127. Rathod, S.M.; Chavan, A.R.; Jadhav, S.S.; Bato, K.M.; Haidi, M.; Raslan, E.M. Ag+ ion substituted CuFe$_2$O$_4$ nanoparticles: Analysis of structural and magnetic behavior. *Chem. Phys. Lett.* 2021, 765, 138308. [CrossRef]

128. Manikandanath, N.T.; Rimal, Isaac, R.S.; Sanjith, S.; Ramesh Kumar, P.; Anooj, E.S.; Vallinayagam, S. Superb catalytic activity of as-grown synthesized copper ferrite’s thermal decomposition of ammonium perchlorate. *J. Molec. Struct.* 2021, 1234, 130161. [CrossRef]

129. Cahyana, A.H.; Liandi, A.R.; Yulizar, Y.; Romdoni, Y.; Wendi, T.P. Green synthesis of CuFe$_2$O$_4$ nanoparticles mediated by Morus alba L. leaf extract: Crystal structure, grain morphology, particle size, magnetic and catalytic properties in Mannich reaction. *Ceram. Int.* 2021. [CrossRef]

130. Sabale, S.R. Studies on catalytic activity of MnFe$_2$O$_4$ and CoFe$_2$O$_4$MNP as mediators in hemoglobin based biosensor. *Mater. Today Proc.* 2020, 23, 139–146. [CrossRef]

131. Yadav, R.S.; Kuritka, I.; Vilakova, J.; Jamatia, T.; Machovsky, M.; Skoda, D.; Urbánek, P.; Masaf, M.; Urbánek, M.; Kalina, L.; et al. Impact of sonochemical synthesis condition on the structural and physical properties of MnFe$_2$O$_4$ spinel ferrite nanoparticles. *Ultrason. Sonochem.* 2020, 61, 104839. [CrossRef]

132. Golsefidi, M.A.; Abbasi, F.; Abroudi, M.; Abroudi, M.; Abbasi, Z.; Gatei, A. Facile synthesis of MnFe$_2$O$_4$ nanoparticles and investigation of various reducing and capping agents on their size and morphology. *J. Mater. Sci. Mater. Electron.* 2017, 28, 1378–1385. [CrossRef]

133. Mozaffari, M.S.; Amighian, J.; Darsheshdar, J.A.E. Magnetic and structural studies of nickel-substituted cobalt ferrite nanoparticles, synthesized by the sol–gel method. *J. Magn. Magn. Mater.* 2018, 350, 19–22. [CrossRef]

134. Padmapiya, G.; Manikandan, A.; Krishnasamy, V.; Jaganathan, S.K.; Arul Antony, S. Spinel Ni$_x$Zn$_{1-x}$Fe$_2$O$_4$ (0.0 ≤ x ≤1.0) nano-photocatalysts: Synthesis, characterization and photocatalytic degradation of methylene blue dye. *J. Mol. Struct.* 2016, 1119, 37–39. [CrossRef]

135. Karpova, T.; Vassiliev, V.; Vladimirova, E.; Osotov, V.; Ronkin, M.; Nosov, A. Synthesis of ultradisperse NiFe$_2$O$_4$ spinel by thermal decomposition of citrate precursors and its magnetic properties. *Ceram. Int.* 2012, 38, 373–379. [CrossRef]

136. Kiran Babu, L.; Rami Reddy, Y.V. A novel thermal decomposition approach for the synthesis and properties of superparamagnetic nanocrystalline NiFe$_2$O$_4$ and its antibacterial, electrocatalytic properties. *J. Supercond. Novel Magn.* 2020, 33, 1013–1021. [CrossRef]

137. Popkov, V.I.; Tolstoy, V.P.; Semenov, V.G. Synthesis of phase-pure superparamagnetic nanoparticles of ZnFe$_2$O$_4$ via thermal decomposition of zinc-iron layered double hydroxysulphate. *J. Alloys Comp.* 2020, 813, 152179. [CrossRef]

138. Sukhorukov, Y.P.; Telegin, A.V.; Bebenin, N.G.; Nosov, A.P.; Bessonov, V.D.; Buchkevich, A.A. Strain-magneto-optics of a magnetostrictive ferrimagnetic Co$_2$Fe$_3$O$_4$. *Solid State Commun.* 2017, 263, 27–30. [CrossRef]

139. Salazar-Kuri, U.; Estevez, J.O.; Silva-González, N.R.; Pal, U. Large magnetostriction in chemically fabricated CoFe$_2$O$_4$ nanoparticles and its temperature dependence. *J. Magn. Magn. Mater.* 2018, 460, 141–145. [CrossRef]

140. Anantharamaiah, P.N.; Joy, P.A. Large enhancement in the magnetostriction parameters of the composite of CoFe$_2$O$_4$ and CoFe$_2$O$_4$MNP. *Mater. Lett.* 2019, 236, 303–306. [CrossRef]

141. Wang, J.; Li, J.; Li, X.; Bao, X.; Gao, X. High magnetostriction with low saturation field in highly textured CoFe$_2$O$_4$ by magnetic field alignment. *J. Magn. Magn. Mater.* 2018, 462, 53–57. [CrossRef]
170. Gopalakrishnan, E.V.; Joy, P.A.; Al-Omari, I.A.; Sakthi Kumar, D. On the structural, magnetic and electrical properties of sol-gel derived nanosized cobalt ferrite. J. Alloys Comp. 2009, 485, 711–717. [CrossRef]

171. Anupama, M.K.; Rudraswamy, B.; Dhananjaya, N. Investigation on impedance response and dielectric relaxation of Ni-Zn ferrites prepared by self-combustion technique. J. Alloy. Comp. 2017, 706, 554–561. [CrossRef]

172. Ashour, A.H.; El-Batal, A.I.; Abdel Maksoud, M.I.A.; El-Sayed, G.S.; Labib, S.; Abdeltwab, E.; El-Okry, M.M. Antimicrobial activity of metal-substituted cobalt ferrite nanoparticles synthesized by sol-gel technique. Particuology 2018, 40, 141–151. [CrossRef]

173. Ashour, A.H.; El-Batal, A.I.; Abdel Maksoud, M.I.A.; El-Sayyad, G.S.; Labib, S.; Abdeltwab, E.; El-Okr, M.M. Antimicrobial activity of monodisperse CoFeO4 nanoparticles. J. Alloys Comp. 2017, 706, 554–561. [CrossRef]

174. Anupama, M.K.; Rudraswamy, B.; Dhananjaya, N. Investigation on impedance response and dielectric relaxation of Ni-Zn ferrites prepared by self-combustion technique. J. Alloy. Comp. 2017, 706, 554–561. [CrossRef]

175. Gharineh, S.; Mehrkhah, M.; Shaker, A.; Motamedi, H. Preparation and anti-bacterial activity of CoFe2O4/SiO2/Ag composite impregnated with streptomycin. Chem. Eng. J. 2015, 259, 34–42. [CrossRef]

176. Ishaq, K.; Saka, A.A.; Kamardeen, A.O.; Ahmed, A.; Alhassan, M.H.; Abdullahi, H. Characterization and antibacterial activity of nickel ferrite doped-alumina nanoparticles. J. Magn. Magn. Mater. 2020, 510, 166143. [CrossRef]

177. Shah, S.A.; Hashmi, M.U.; Alam, S.; Shamim, A. Magnetic and bioactivity evaluation of ferrimagnetic ZnFe2O4 nanoparticles. J. Molec. Magn. Mater. 2010, 322, 375–381. [CrossRef]

178. Suleman, M.; Riaz, S. In silico study of hyperthermia treatment of liver cancer using core-shell CoFe2O4@MnFe2O4 magnetic nanoparticles. J. Magn. Magn. Mater. 2020, 498, 166143. [CrossRef]

179. Issa, B.; Obaidat, I.M.; Albiss, B.A.; Haik, Y. Nanoparticles: Surface effects and properties related to biomedicine applications. Int. J. Mol. Sci. 2013, 14, 21266–21305. [CrossRef] [PubMed]

180. Kurugan, E.; Gas, P. Magnetophoretic placement of ferromagnetic nanoparticles in RF hyperthermia. In Proceedings of the 2017 Progress in Applied Electrical Engineering (PAEE), Koscielisko, Poland, 25–30 June 2017; pp. 1–4. [CrossRef] [PubMed]

181. Manohar, A.; Geleta, D.D.; Krishnamoorthi, C.; Lee, J. Synthesis, characterization and magnetic hyperthermia properties of nearly monodisperse CoFe2O4 nanoparticles. Ceram. Int. 2020, 46, 28035–28041. [CrossRef]

182. Shah, S.A.; Hashmi, M.U.; Alam, S.; Shamim, A. Magnetic and bioactivity evaluation of ferrimagnetic ZnFe2O4 containing glass ceramics for the hyperthermia treatment of cancer. J. Magn. Magn. Mater. 2010, 322, 375–381. [CrossRef]

183. Kurugan, E.; Gas, P. Methods of calculation the magnetic forces acting on particles in magnetic fluids. In Proceedings of the 2015 Selected Problems of Electrical Engineering and Electronics (WZEE), Kielce, Poland, 17–19 September 2015; pp. 1–6.

184. Kurgan, E.; Gas, P. Magnetophoretic placement of ferromagnetic nanoparticles in RF hyperthermia. In Proceedings of the 2017 Progress in Applied Electrical Engineering (PAEE), Koscielisko, Poland, 25–30 June 2017; pp. 1–4. [CrossRef] [PubMed]

185. Ishaq, K.; Saka, A.A.; Kamardeen, A.O.; Ahmed, A.; Alhassan, M.H.; Abdullahi, H. Characterization and antibacterial activity of nickel ferrite doped-alumina nanoparticles. J. Magn. Magn. Mater. 2020, 46, 28035–28041. [CrossRef]

186. Shah, S.A.; Hashmi, M.U.; Alam, S.; Shamim, A. Magnetic and bioactivity evaluation of ferrimagnetic ZnFe2O4 nanoparticles. J. Molec. Magn. Mater. 2010, 322, 375–381. [CrossRef]

187. Gharineh, S.; Mehrkhah, M.; Shaker, A.; Motamedi, H. Preparation and antibacterial activity of CoFe2O4/SiO2/Ag composite impregnated with streptomycin. Chem. Eng. J. 2015, 259, 34–42. [CrossRef]

188. Ishaq, K.; Saka, A.A.; Kamardeen, A.O.; Ahmed, A.; Alhassan, M.H.; Abdullahi, H. Characterization and antibacterial activity of nickel ferrite doped-alumina nanoparticles. J. Magn. Magn. Mater. 2020, 46, 28035–28041. [CrossRef]

189. Gorgizadeh, M.; Behzadpour, N.; Salehi, F.; Daneshvar, F.; Dehdari Vais, R.; Nazari-Vanani, R.; Azarpira, N.; Lotfi, M.; Sattarhamd, N. A MnFe2O4/C nanocomposite as a novel theranostic agent in MRI, sonodynamic therapy and photothermal therapy of a melanoma cancer model. J. Alloys Comp. 2020, 816, 152597. [CrossRef]