Synthesis of cobalt ferrite (CoFe₂O₄) by combustion with different concentrations of glycine

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Abstract. Among the wide range of applications of Cobalt Ferrite (CoFe₂O₄), its usage for the manufacturing of modern electronic devices such as solar panels, capacitors and batteries has been studied lately. Hence, solar-powered vehicles, which strongly rely their performance on energy efficient electronics, is a sector that could particularly benefit from enhanced applications of such material. In this work, the synthesis of Cobalt Ferrite was studied by means of Solution Combustion Synthesis (SCS) using iron nitrate nonahydrate and cobalt nitrate hexahydrate as precursors, and glycine as fuel. The nitrates were dissolved in distilled water and placed under stirring and heating for 5 minutes; when the temperature reached 60°C, glycine was added; after complete homogenization, the solution was placed in an electric oven (400°C) until complete combustion (approximately 15 minutes). Three syntheses were carried out using different concentrations of fuel: lean, stoichiometric and rich. The product crystallite size and composition were investigated in order to determine the influence of the fuel concentration on the structure of the produced cobalt ferrite.

1. Introduction

Ferrite is a ceramic material produced from burning large proportions of iron (III) oxide (Fe₂O₃) blended with small concentrations of one or more metallic elements, such as cobalt, manganese, nickel, magnesium, and zinc [1]. Ferrites usually present a spinel structure with the formula AB₂O₄, where A and B represent various metal cations, including iron (Fe) [2].

Many studies on ferrites nanoparticles including CoFe₂O₄ spinel have been recently presented due to their distinct magnetic and electronic properties, which open up numerous possibilities including cancer treatment [3], biosensors [4], sensors [5], memory shape alloys [6] and water purification [7]. In addition, ferrite cobalt has been emerging as a promising material for the development of modern electronic devices, such as capacitors, sensors or high capacity batteries [8–10]. Thus this material can be particularly useful in applications related to the sustainable mobility in general and to the solar vehicles in particular. Optimizing the synthesis and usage of such material, for instance, has become a topic of interest for the design of solar-powered electric vehicles, given that power management and energy efficiency are the key aspects of such mobility branch [11-13].
Solution combustion synthesis (SCS) has been increasingly applied in the production of oxide catalysts due to the possibility of producing low-cost, highly pure and homogeneous nanostructured powders. SCS mainly consists of combining the reactants in an aqueous medium using a complexing agent (usually glycine and urea), as well as oxidizing agents (usually metal nitrates), to oxidize the fuel [14,15]. The mixture is heated to 150°C and 500°C to carry out self-ignition in a rapid combustion reaction that can reach more than 1700°C [16]. A solid product that is typically well dispersed and crystalline is formed at the end of the process [16].

The concentration ratio between the complexing agent and oxidizing agent is directly related to the final properties of the powder produced [15]. Thus, this study aims to formulate a methodology to use SCS to produce, at a low cost, homogeneous CoFe₂O₄ catalyst nanoparticles. For this purpose, we studied the influence of different concentrations of fuel on the product composition and morphology. With this scope, we used an experimental approach that, coupling diffractometry and microscopy, permitted relevant and unexpected results in similar situations [17-19].

2. Materials and methods

2.1 Synthesis of cobalt ferrite

CoFe₂O₄ nanostructured particles were produced by SCS with various concentrations of fuel. Co(NO₃)₂.6H₂O and Fe(NO₃)₃.9H₂O were employed as precursors and oxidizers, while glycine (C₂H₅NO₂) is used as fuel. All of the reactants were obtained from Merck. The fuel stoichiometric, rich and lean ratios were established by the chemistry of propellants method [20] and applied according to Equations 1, 2 and 3, respectively.

\[2\text{Fe(NO}_3\text{)}_3.9\text{H}_2\text{O} + \text{Co(NO}_3\text{)}_2.6\text{H}_2\text{O} + 4.5\text{C}_2\text{H}_5\text{NO}_2 + 0.25\text{O}_2 \rightarrow \text{CoFe}_2\text{O}_4 + 9\text{CO}_2 + 35.25\text{H}_2\text{O} + 6.25\text{N}_2\] (1)

\[2\text{Fe(NO}_3\text{)}_3.9\text{H}_2\text{O} + \text{Co(NO}_3\text{)}_2.6\text{H}_2\text{O} + 5.5\text{C}_2\text{H}_5\text{NO}_2 + 4.75\text{O}_2 \rightarrow \text{CoFe}_2\text{O}_4 + 11\text{CO}_2 + 37.75\text{H}_2\text{O} + 6.75\text{N}_2\] (2)

\[2\text{Fe(NO}_3\text{)}_3.9\text{H}_2\text{O} + \text{Co(NO}_3\text{)}_2.6\text{H}_2\text{O} + 3.5\text{C}_2\text{H}_5\text{NO}_2 \rightarrow \text{CoFe}_2\text{O}_4 + 7\text{CO}_2 + 32.75\text{H}_2\text{O} + 5.75\text{N}_2 + 4.25\text{O}_2\] (3)

The nitrates were individually dissolved in water and then mixed. The solution was stirred and heated for approximately 5 min. When reaching 60°C, the fuel that had previously been dissolved in water was added to the solution. The solution was maintained under stirring and temperature for a few minutes to promote perfect homogenization. The solution was placed in an electric muffle furnace preheated to 400°C until complete combustion (around 15 min).

2.2 Chemical and structural characterization

The crystallinity of the samples was evaluated by X-ray diffraction (XRD) using a PHILIPS diffractometer (model X'Pert MPD) at 40 kV and 40 mA using a Cu anode. The powder crystallite sizes were calculated using Scherrer's equation 4:

\[D = \frac{K\lambda}{\beta \cos \theta}\] (4)

where D represents the crystallite size, K is a constant that depends on the particle shape (we assume the particles to be spherical, meaning K=0.94), λ is the wavelength of the electromagnetic radiation used (1.5406 Å, a value related to the main characteristic radiation emitted by copper), θ is Bragg’s angle, and β is the contribution of the crystallite size to the full width at half maximum (FWHM) of the corresponding diffraction peak in radians.

The morphology of the CoFe₂O₄ was characterized by scanning electron microscopy (SEM) using a JEOL microscope (JSM 6060) with a maximum operating voltage of 30 kV and a nominal resolution of 3.5 nm. The applied voltage was 10 to 20 k.
3. Results and Discussions
The diffractograms in Figure 1 show the characteristic crystal peaks corresponding to the cobalt ferrite phase for all samples according to JCPDS 73-1960. The sample lean presented a high degree of purity and showed only the COFe₂O₄ phase. The stoichiometric and rich samples showed the formation of a second phase of cobalt oxide (JCPDS 73-1701).

![Figure 1. Diffractograms of the CoFe₂O₄ catalysts for different fuel concentrations.](image)

SCS synthesis with elevated concentration of fuel leads to fast ignition and intense burn during reaction. With the excess of fuel, oxygen from the atmosphere is required in order to complete the reaction. This oxygen excess may cause the formation of secondary oxide phases besides the primary cobalt ferrite phase [21]. The crystallite sizes calculated by Scherrer’s formula showed for the different concentrations: 23.58 nm for lean (the smallest size among all samples), 31.14 nm for stoichiometric, and 33.16 nm for Rich. These results indicate that powder has a nanostructured crystallite. Figure 2 shows SEM micrographs of the CoFe₂O₄ samples just after combustion with no grinding process involved.

![Figure 2. SEM images of samples: a) Lean, b) Stoich, c) Rich.](image)

Figure 2 it is possible to observe the production of porous structures for all the compositions. The formation of the spongy features is attributed to the evolution of a large amount of gas during combustion [22].
The amount of oxygen involved in the reaction directly affects the velocity of the process and the product composition as discussed above. Glycine has a small carbonic chain, and the presence of N and OH in the structure provides an intense and fast high-temperature combustion during ignition, which releases a high volume of gases [21]. This release of gases leads to the production of more porous structures as can be seen when comparing the fuel-lean sample (Figure 2a) with the rich (Figure 2c) and stoichiometric sample (Figure 2b). It is possible to observe the increasing of structural porosity with the increasing of fuel concentration.

4. Conclusion
The synthetic route presented in this paper shown to be effective for production of nanostructured CoFe$_2$O$_4$. In this study was possible to conclude that in the SCS of CoFe$_2$O$_4$ higher concentrations of glycine lead to the formation of secondary phases of cobalt oxide. The fuel-lean sample was the only one that produced a nanometer pure phase of CoFe$_2$O$_4$. This sample is promising for applications in the electronics field, such as on solar-powered electric vehicles, since pure particles and good crystallinity are crucial in this area.

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