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Synthesis, Characterization, and Sensor Applications of Spinel ZnCo$_2$O$_4$ Nanoparticles

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Abstract: Spinel ZnCo$_2$O$_4$ nanoparticles were synthesized by means of the microwave-assisted colloidal method. A solution containing ethanol, Co-nitrate, Zn-nitrate, and dodecylamine was stirred for 24 h and evaporated by a microwave oven. The resulting solid material was dried at 200 °C and subsequently calcined at 500 °C for 5 h. The samples were characterized by scanning electron microscopy (SEM), transmission electron microscopy (TEM), X-ray diffraction (XRD), and Raman spectroscopy, confirming the formation of spinel ZnCo$_2$O$_4$ nanoparticles with average sizes between 49 and 75 nm. It was found that the average particle size decreased when the dodecylamine concentration increased. Pellets containing ZnCo$_2$O$_4$ nanoparticles were fabricated and tested as sensors in carbon monoxide (CO) and propane (C$_3$H$_8$) gases at different concentrations and temperatures. Sensor performance tests revealed an extremely high response to 300 ppm of CO at an operating temperature of 200 °C.

Keywords: spinel; faceted nanoparticle; cobaltite; sensors

1. Introduction

For a long time, pollutant gases from industry and internal combustion engines have been responsible for many human health issues, and to a large extent, for global climate change. Facing this, several groups have focused on the research and development of new sensor materials for the detection and monitoring of polluting gases such as CO and C$_3$H$_8$. In particular, sensor materials made from semiconductor oxides have been a valuable choice because they possess good chemical stability,
low price, and an easy integration into electronic circuits. However, it is still necessary to improve sensor parameters such as selectivity, sensitivity, and operating temperature.

Zinc cobaltite (ZnCo$_2$O$_4$) is a p-type semiconductor material with a spinel-type structure. This material has attracted the attention of several research groups due to its potential applications as an electrode for Li-ion batteries [1–5], as a catalyst [6–9], and in supercapacitors [10–13]. In the gas sensors field, sensor devices based on ZnCo$_2$O$_4$ nanoparticles have displayed an excellent sensitivity to liquefied petroleum gas [14–17], H$_2$S [18], ethanol [19,20], acetone [21], and formaldehyde [22,23], probably due to their high surface area. Additionally, ZnCo$_2$O$_4$ has also been used as Cl$_2$, NO$_2$, CO$_2$, H$_2$, NH$_3$, CH$_3$COOH, SO$_2$, CO, C$_3$H$_8$, ethylene, xylene, toluene, and methanol gas sensors [15,16,18–22]. Specifically, when ZnCo$_2$O$_4$ nanoparticles were exposed to CO and C$_3$H$_8$ concentrations, a poor sensitivity was exhibited at working temperatures in the range of 175 to 350 °C [15,16,19,21]. In contrast, sensors based on nanowire-assembled hierarchical ZnCo$_2$O$_4$ microstructures showed a good sensitivity towards CO and C$_3$H$_8$ at an operating temperature of 300 °C [22]. From these studies, it is fair to assume that the gas sensing properties depend on the shape and size of the nanoparticles [24]. Therefore, developing a ZnCo$_2$O$_4$-based gas sensor, with a high sensitivity and a low operating temperature, is of great interest to many people. Several synthesis methods of ZnCo$_2$O$_4$ nanoparticles have been reported, such as combustion [1], hydrothermal [3,12], thermal decomposition [14], co-precipitation/digestion [15], water-in-oil (W/O) microemulsion [18], sol-gel [25], and surfactant-mediated [26] methods. In recent years, the microwave-assisted colloidal method has been an efficient and low-cost synthesis process for obtaining micro and nanostructures of oxide materials [27–29]. In fact, microwave is a simple technique, which plays an important role in colloidal synthesis since it provides a rapid evaporation of the precursor solvent and a short reaction time [30–33].

To the best of our knowledge, there are not reports on the synthesis and use of faceted ZnCo$_2$O$_4$ nanoparticles for gas sensing applications, such as the ones presented in this investigation, but in recent works, sensors based on faceted SnO$_2$ and ZnSn(OH)$_x$ nanoparticles exhibited high response towards toluene and ethanol, respectively [34,35] and ZnCo$_2$O$_4$ nanoparticles are synthesized via a microwave-assisted colloidal method [27,36]. The experimental method and the different steps to synthesize ZnCo$_2$O$_4$ nanoparticles are described in detail. In the following, the results of SEM, XRD, Raman, and TEM characterizations are shown. In addition, sensitivity results for sensors based on ZnCo$_2$O$_4$ nanoparticles are thoroughly discussed.

2. Materials and Methods

Zinc nitrate hydrate (Zn(NO$_3$)$_2$$\cdot$6H$_2$O, Sigma Aldrich 99.99%), cobalt(II) nitrate hexahydrate (Co(NO$_3$)$_2$$\cdot$6H$_2$O, Sigma Aldrich 98%), dodecylamine (C$_{12}$H$_{25}$N, Sigma Aldrich 98%), and ethanol (C$_2$H$_6$O, Golden Bell 98%) were used as reagents. 5 mmol (0.947 g) of Zn(NO$_3$)$_2$$\cdot$6H$_2$O, 10 mmol (2.91 g) of Co(NO$_3$)$_2$$\cdot$6H$_2$O, and 5.4 mmol (1 g) of C$_{12}$H$_{25}$N were separately dissolved in 5 mL of ethanol. Additional syntheses were made with 10.8 mmol (2 g) and 16.2 mmol (3 g) of dodecylamine in order to obtain additional nanoparticle sizes. After 20 min of vigorous stirring on magnetic dishes at room temperature, the cobalt nitrate solution was added dropwise to the dodecylamine solution and kept stirring for 1 h. The zinc nitrate solution was then slowly added to the cobalt and dodecylamine mixture yielding a wine-color solution (the solutions with 10.8 and 16.2 mmol of dodecylamine were greenish-blue) with pH = 2 and a final volume of approximately 16 mL. This final solution was covered with a watch glass to avoid contamination and kept under stirring for 24 h, losing ~2 mL by evaporation. The solution (~14 mL) was evaporated afterwards using a microwave oven (General Electric JES769WK) at a low power (~140 W). The microwave radiation was applied over the solution for periods of 1 min in order to avoid splashing. The total time of evaporation was 3 h. The resulting solid was dried in air with a muffle (Novatech) at 200 °C. Finally, the calcination of the powder was done at 500 °C for 5 h in
an alumina crucible with a cover at a heating rate of 100 °C/h. The sample was kept in the furnace for
cooling at room temperature. The samples made with 5.4, 10.8, and 16.2 mmol of dodecylamine were
labeled as A, B, and C, respectively.

The calcined samples were characterized by SEM using a FEI-Helios Nanolab 600 system operated
at 20 kV. XRD characterizations were performed with a PANalytical Empyrean system with CuKα and
λ = 1.546 Å for phase identification. XRD patterns were obtained at room temperature in the range
2θ = 10–70° with steps of 0.02°, lasting 30 s for each step. The ZnCo2O4 crystallite size was calculated
by Scherrer’s equation [37] using the plane (311) at 2θ = 36.8°:

\[
\text{Cristallite size} = \frac{0.89\lambda}{\beta \cos \theta} \frac{180°}{\pi}
\]  

(1)

where λ is the X-ray wavelength and β is the full width at half maximum (FWHM). Raman spectroscopy
characterization was performed using a Thermo Scientific DXR confocal Raman microscope with
a 633 nm excitation source. The Raman spectra were measured from 150 to 800 cm⁻¹ at room
temperature, using an exposure time of 60 s and a Laser power of 5 mW. TEM, high-resolution
transmission electron microscopy (HRTEM), energy-dispersive X-ray (EDS) and high-angle annular
dark-field/scanning transmission electron microscopy (HAADF-STEM) were performed by means of
a FEI Tecnai-F30 system operated at 300 kV. The gas sensing measurements were carried out on pellets
of ZnCo2O4 with a thickness of 0.5 mm and a diameter of 12 mm. The pellets were prepared with
0.350 g of ZnCo2O4 powders employing a manual pressure machine (Simple Ital Equip) at 20 tons for
120 min. A TM20 Leybold detector was used to control the gas concentration and the partial pressure.
The sensing response was investigated by measuring the electric resistance using a digital-multimeter
(Keithley). A schematic diagram of the gas sensing measurement system is shown in Figure 1.

The sensing response, or sensitivity, was defined as \( S = R_a / R_g \) for the reducing gas, where \( R_a \) and
\( R_g \) refer to the resistances measured in air and gas, respectively [21,38,39].

\[
\beta = 36.8° \quad \theta = 10–70° \quad \text{with steps of 0.02°, lasting 30 s for each step.}
\]

3. Results and Discussion

SEM images of samples A, B, and C are shown in Figure 2. Sample A exhibited a high concentration
of nanoparticles with irregular shape and sizes in the range of 50–110 nm (see Figure 2a,d). Agglomerates
of nanoparticles were also observed (Figure 2d). SEM images of sample B revealed a large amount
of nanoparticles with diameters in the range from 40–85 nm (see Figure 2b,e). Sample C contained

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**Figure 1.** Experimental setup for the gas sensitivity measurements of ZnCo2O4 nanoparticles.
nanoparticles with diameters ranging from 25–70 nm (Figure 2c,d,f). The particle-size distribution of the three ZnCo$_2$O$_4$ samples can be seen in Figure 3. From SEM, it is clear that the dodecylamine concentration plays a key role in the morphologies and the particle sizes [40]: an increment of the dodecylamine concentration produced a large amount of nanoparticles and a decreasing particle size, which suggests that the dodecylamine inhibits the growth of the ZnCo$_2$O$_4$ nanoparticles.

**Figure 2.** SEM images of ZnCo$_2$O$_4$ nanoparticles: (a,d) sample A; (b,e) sample B; and (c,f) sample C.

**Figure 3.** Particle size distribution for the ZnCo$_2$O$_4$ samples: (a) A; (b) B; and (c) C.
The formation of the ZnCo$_2$O$_4$ nanoparticles follows the principles of nucleation and growth established by LaMer and Dinegar [41]. In our synthesis, the nucleation process could occur when the zinc nitrate solution was added to the cobalt and dodecylamine solution [30], and the particle growth continued developing by diffusion of the nuclei during agitation of the final solution [42].

As previously discussed, dodecylamine plays an important role in the microstructure of the ZnCo$_2$O$_4$ particles. In this research, it seems that the ethanol and the dodecylamine are working synergically to reduce cations to metals because ethanol alone is not capable of doing so (i.e., in order to produce ultrafine metal nanoparticles), but just mixed with strong reducing agents such as microwave radiation, other chemical compounds, etc. [43–45]. We have observed that the dodecylamine concentration is the rate-determining factor; therefore, we propose the following reaction mechanism.

Dodecylamine is a primary amine with properties of a weak base because of the nitrogen’s unshared electron pair. This kind of amine has nucleophilic behavior and can be used as reducing and surfactant agents. The reaction of dodecylamine with Co$^{2+}$ cations could occur in two ways: (i) the attraction of the unshared electron pair to the nucleus of the cations in a nucleophilic reaction; (ii) the formation of an electrostatic bond between the nitrogen of the dodecylamine and the electrons of the outer shell of Co$^{2+}$, because of the strong electronegativity of the nitrogen (≈ 3.04). The result of these interactions is the formation of the dodecylamine-Co$^{2+}$ complex [46]:

$$\text{Co}^{2+} + \text{NH}_2-(\text{CH}_2)_{11}\text{NH}_3 \rightarrow \text{Co}^{2+} \cdots \cdots \text{NH}_2-(\text{CH}_2)_{11}\text{NH}_3$$ (2)

In this reaction, dodecylamine is working as the surfactant; therefore, nitrogen’s electrons could participate as reducing agents, reducing the cationic Co$^{2+}$ to metallic nanoparticles of Co$^0$.

$$\text{Co}^{2+} + 2e^- \rightarrow \text{Co}^0$$ (3)

For 5.4 mmol of dodecylamine, the molar ratio of dodecylamine: Co$^{2+}$ was around 1:2, which suggests that half of the Co$^{2+}$ cations are not participating in the formation of complexes and should be reduced in some way by the solvent. When the dodecylamine-ethanol-Co$^{2+}$ solution is mixed with the ethanol-Zn$^{2+}$ solution, and once the Co$^{2+}$ and Zn$^{2+}$ have been reduced, Zn could attract two atoms of Co to form ZnCo$_2$, as Zn is slightly more electronegative than Co (1.90 vs. 1.65, respectively). Since 5.4 mmol of dodecylamine are not enough to complex all of the Co$^{2+}$ cations, there are several available cations to be reduced by the solvent, and as a result the ZnCo$_2$ grains are the larger ones. For 10.8 mmol of dodecylamine, most of the Co$^{2+}$ should be forming a complex, and the growing of ZnCo$_2$ grains is restricted through the slow release-reduction process of Co$^{2+}$. For 16.2 mmol of dodecylamine, the Co$^{2+}$ cations could be completely complexed; therefore, the excess of dodecylamine should form complexes even with Zn$^{2+}$, restricting even more the growing of ZnCo$_2$ grains. Summarizing, the role of dodecylamine is as a surfactant and as a reducing agent; however, it seems that the function of surfactant, through electrostatic interactions, is the dominant one, reducing the particle size and modifying the particles’ microstructure as the dodecylamine concentration increases. With heat treatment, the dodecylamine was finally removed from the ZnCo$_2$O$_4$.

Figure 4 depicts the XRD patterns of the samples. In them, the typical peaks corresponding to the cubic ZnCo$_2$O$_4$ spinel-structure were identified. The well-defined narrow peaks are an indication of the good crystallinity of the ZnCo$_2$O$_4$ samples. Table 1 shows the crystallite size of every sample. From these results, an increase in crystallite size was produced when the dodecylamine concentration was increased. A slight increase in the FWHM was also observed, indicating a particle size reduction as was confirmed by SEM.
The formation of the oxide ZnCo$_2$O$_4$ was also confirmed by Raman characterization. According to the group theory for oxides with a spinel structure, five active Raman bands were expected: $A_{1g} + E_g + 3F_{2g}$ [47]. Figure 5 shows the Raman spectrum indicating the main vibrational bands produced by the ZnCo$_2$O$_4$ spinel crystal structure. The bands labeled by $A_{1g}$ symmetry species. These results are consistent with those reported for ZnCo$_2$O$_4$ spinel structures [48]. Interestingly, the band located at 204 cm$^{-1}$ is a vibrational mode that could be attributed to a Co$_3$O$_4$ spinel structure [49]. The formation of Co$_3$O$_4$ is likely due to the cation disorder (substitution of Zn$^{2+}$ by Co$^{3+}$) in the spinel structure of the zinc cobaltite. Notice that the Co$_3$O$_4$ possesses a spinel structure similar to that of ZnCo$_2$O$_4$. Therefore, the XRD patterns between Co$_3$O$_4$ and ZnCo$_2$O$_4$ are almost indistinguishable.

![Figure 4](image_url)

**Figure 4.** XRD patterns of ZnCo$_2$O$_4$ samples synthesized with (A) 5.4; (B) 10.8; and (C) 16.2 mmol of dodecylamine. The peak signals for the ZnCo$_2$O$_4$ structure (JCPDF 23-1390) are labeled with black-filled circles.

![Table 1](table_url)

| Samples | FWHM  | Crystallite Size (nm) |
|---------|-------|-----------------------|
| A       | 0.338 | 24.75                 |
| B       | 0.351 | 23.84                 |
| C       | 0.421 | 19.92                 |

**Table 1.** Crystallite size of the ZnCo$_2$O$_4$ samples.

![Figure 5](image_url)

**Figure 5.** Raman spectra of the samples A, B, and C.
Figure 6 shows TEM images of the samples. The presence of faceted nanoparticles with a pockmarked structure was clearly identified (Figure 6a,c,e). The estimated average particle size was approximately 75, 61, and 49 nm for samples A, B, and C, respectively; these measurements are consistent with the SEM analysis. The standard deviation was \( \pm 12.64 \), \( \pm 8.78 \), and \( \pm 8.36 \) nm, respectively. Figure 6b,d,f shows HRTEM images of the selected area marked with a black rectangle. These images confirmed the presence of faceted ZnCo\(_2\)O\(_4\) nanoparticles as was done by the XRD patterns and the Raman characterization. Fringe spacings of 0.467 and 0.286 nm are clearly observed, which are attributed to the planes (111) and (220) of the ZnCo\(_2\)O\(_4\) spinel structure, respectively.

To investigate the nanoparticle composition, an EDS line scan was performed on sample A (see Figure 7). Figure 7a shows a HAADF-STEM image of the ZnCo\(_2\)O\(_4\) nanoparticles. The image confirms the presence of faceted nanoparticles with a pockmarked structure, which is consistent with the TEM images. In the EDS line scan, zinc, cobalt, and oxygen are observed across the linear mapping, confirming the presence of the expected elements, as seen in Figure 7b. In the central region \( p_2 \), a decrease of the element composition is observed in comparison to point \( p_1 \), which can be due to the irregular surface of the nanoparticle (pockmarked zone). Similar EDS elemental line scan spectra were obtained for samples B and C. Figure 7c shows an EDS microanalysis on \( p_2 \), where the individual elements Zn, Co, and O can easily be seen. The atomic ratio of Zn:Co is approximately 1:1.96, which is consistent with the composition of ZnCo\(_2\)O\(_4\). This EDS spectrum is in agreement with the literature [3,5].
To investigate the nanoparticle composition, an EDS line scan was performed on sample A (see Figure 7). Figure 7a shows a HAADF-STEM image of the ZnCo2O4 nanoparticles. The image confirms the presence of faceted nanoparticles with a pockmarked structure, which is consistent with the TEM images. In the EDS line scan, zinc, cobalt, and oxygen are observed across the linear mapping, confirming the presence of the expected elements, as seen in Figure 7b. In the central region p2, a decrease of the element composition is observed in comparison to point p1, which can be due to the irregular surface of the nanoparticle (pockmarked zone). Similar EDS elemental line scan spectra were obtained for samples B and C. Figure 7c shows an EDS microanalysis on p2, where the individual elements Zn, Co, and O can easily be seen. The atomic ratio of Zn:Co is approximately 1:1.96, which is consistent with the composition of ZnCo2O4. This EDS spectrum is in agreement with the literature [3,5].

Figure 7. (a) HAADF-STEM image; (b) elemental line scan; and (c) EDS microanalysis of sample A.

To investigate the sensing properties of the ZnCo2O4 oxide, pellets of the material were made and tested in CO and C3H8 atmospheres. Figure 8 shows the oxide’s response vs. CO concentration of sensors made from samples A, B, and C. As shown in Figure 8a,c, no response variation was measured for A and C at 100 °C. On the contrary, the sensor made from B exhibited response values of 1, 1.03, 1.21, 1.48, 2.50, and 5.68 for CO concentrations of 1, 5, 50, 100, 200, and 300 ppm, respectively (Figure 8b). At 200 and 300 °C, the response of the three sensors increased with an increase of the CO concentration. In the whole concentration range (1–300 ppm CO), the sensor made from C exhibited a high response at 200 °C, better than the sensors based on A and B. At this temperature, the response values of the sensor based on C were 2.56, 2.66, 3.18, 1274.29, 2622.22, and 2950 for CO concentrations of 1, 5, 50, 100, 200, and 300 ppm, respectively. The sensors made from A and B comparatively also showed a good response to 300 ppm of CO (305.07 and 19.37, respectively) at 300 °C. From these results, it is clear that the ZnCo2O4 sensors are highly sensitive to concentrations of carbon monoxide and working temperatures. As expected, the material’s gas response increased due to the raising of the gas concentration and operation temperature. The raise of the response is associated to an increased oxygen desorption at high temperatures. Some authors report that the response of a semiconductor material depends on the adsorption of several oxygen species as a function of temperature [50,51]. The mechanism to explain the interaction between the CO molecules and a semiconductor oxide like the one used in this work is based on the accumulation layer’s modulation due to the chemisorption of oxygen [33,52]. Therefore, in the tests at temperatures above 100 °C, the oxygen species O− and O2− (ionic form) that adsorb chemically on the sensor are more reactive than other oxygen species that adsorb at temperatures below 100 °C (like O2−) [29,36,37]. It means that below 100 °C, the thermal energy is not enough to produce the desorption reactions of the oxygen, and therefore, an electrical
response does not occur regardless of the gas concentration. By contrast, above 100 °C (in this case, 200 and 300 °C), the formation of oxygen species occurs causing an increase in the gas-solid interaction in the presence of CO [33,52,53].

Figure 8. Gas response of ZnCo$_2$O$_4$ sensors vs. CO concentration at different operating temperatures: (a) sample A; (b) sample B; and (c) sample C.

The response of the ZnCo$_2$O$_4$ sensors in propane atmospheres at different operating temperatures is shown in Figure 9. Such as in the case of CO, the response rose with the increasing of temperature and propane concentration. However, at temperatures below 100 °C, no changes were observed in the response. At 100 °C, a sensing response value of ~1 was calculated for the sensors A and B in the range of 1–300 ppm of C$_3$H$_8$ (Figure 9a,c). At the same temperature, the sensor B registered values of 1, 1.03, 1.13, 1.49, 1.76, and 2.62 for C$_3$H$_8$ concentrations of 1, 5, 50, 100, 200, and 300 ppm, respectively (as seen in Figure 9b). Again, the sensor based on sample C also exhibited a higher response than those of A and B at a working temperature of 200 °C. The response values for this sensor C were 1, 1.03, 1.18, 1.56, 1.94, and 8.99 at C$_3$H$_8$ concentrations of 1, 5, 50, 100, 200, and 300 ppm, respectively. The sensors based on A and B also exhibited good response to 300 ppm of C$_3$H$_8$: 5.89 at 200 °C, and 8 at 300 °C, respectively. Additionally, the three ZnCo$_2$O$_4$ sensors showed a decrease in gas response when the test gases were removed from the vacuum chamber.
C₃H₈: 5.89 at 200 °C, and 8 at 300 °C, respectively. Additionally, the three ZnCo₂O₄ sensors showed a decrease in gas response when the test gases were removed from the vacuum chamber.

**Figure 9.** Response of the ZnCo₂O₄ sensors as a function of C₃H₈ concentration at different working temperatures: (a) sample A; (b) sample B; and (c) sample C.

As discussed above, the gas detection ability of a material such as the one used in this work depends on the microstructure obtained during the synthesis process [28,29,33,36]. If the particle size is nanometric, the response of the material is substantially improved [54]. It has been established that by reducing the particle size of the materials, their performance (i.e., their sensitivity) to detect different concentrations of gases is boosted [36,55–57], like in our case. Again, the most accepted mechanism to explain the response of the ZnCo₂O₄ is based on changes in the electrical resistance (or conductance) due to the adsorption and desorption of oxygen species on the surface [29,54,58–60]. Depending on the semiconductor type, the concentration of surface charge carriers can increase or decrease [59,61]. This is because during the chemical adsorption of oxygen molecules, a hole accumulation layer (space charge layer) is generated [52], provoking a chemical reaction between the gas and the surface of the ZnCo₂O₄ and resulting in changes in the electrical resistance of the material (i.e., a high sensitivity is recorded) [38,57]. Additionally, the ZnCo₂O₄ response is strongly related to the crystallite size (D), which is less than the thickness of the space charge layer, Lₛ, defined as [29,54,55,59]:

\[
L_s = L_D \sqrt{\frac{e V_s^2}{kT}}
\]

where \(L_D\) is the Debye length, \(e\) is the electron charge, \(V_s\) is the surface potential, \(k\) is the Boltzmann constant, and \(T\) is the absolute temperature. Generally speaking, \(L_s\) has a value between 1 and 100 nm [56]. Therefore, the conductivity mechanism is associated with the crystallite size and the space charge layer [29,54,56,62]: if \(D >> 2L_s\), the conductivity is limited by the Schottky barrier at the particle
Comparing the efficiency of our ZnCo$_2$O$_4$ nanoparticles (a maximum sensitivity of ~2950 and ~8.99 in 300 ppm of CO and C$_2$H$_8$, respectively, at 200 °C both) with similar semiconductor oxides, we found in our case greater sensitivity, stability, and efficiency to detect CO and C$_2$H$_8$ with sensing measurements. A.G.-B. and V.M.R.-B. performed the Raman experiments.

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4. Conclusions

In this work, we successfully synthesized ZnCo$_2$O$_4$ faceted nanoparticles with a size between 49 and 75 nm by means of a simple, economical, and efficient route: the microwave-assisted colloidal method using dodecylamine as a surfactant agent and a calcination temperature of 500 °C. Sensors prepared with these nanoparticles exhibited an excellent response (~2950 with sample C) at a relatively low operating temperature (200 °C) for the detection of CO, and they were capable of detecting up to 300 ppm of C$_2$H$_8$ at 200 °C, which is comparatively (with similar oxides) a very good performance. Hence, ZnCo$_2$O$_4$ is a promising material for applications as a gas sensor, especially in the detection of CO and C$_2$H$_8$.

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