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Ammonia gas sensors based on undoped and Ca-doped ZnO nanoparticles

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Due to its large use in different industrial sectors, high toxicity, and corrosion, the demand for sensing techniques towards ammonia gas has become urgent. In this study we report on the sensing performances of a conductometric sensor for NH₃ gas based on Ca-doped ZnO nanoparticles with different calcium concentrations (0, 1, and 3 at%) synthesized using the sol–gel process under supercritical dry conditions of ethanol. All samples were characterized using X-ray diffraction (XRD), scanning electron microscopy (SEM), transmission electron microscopy (TEM), and Fourier-transform infrared (FTIR) spectroscopy. Pure and Ca-doped ZnO are polycrystalline and well crystallized in the hexagonal wurtzite structure. TEM images revealed that pure ZnO is composed of spherical particles with dimensions in the nanometer range. Larger particles were observed after the incorporation of Ca ions. The average crystallite size, estimated by the Williamson–Hall method, was 43, 80, and 96 nm for pure, Ca-1 at% and Ca-3 at%, respectively. Furthermore, FTIR spectroscopy was used to prove the formation of ZnO and the incorporation of calcium ions in the Ca-doped ZnO samples. The gas sensing performances towards ammonia gas clearly ameliorated after the addition of Ca ions in the ZnO structure. The gas response to NH₃, R_0/R_g , of the 1% Ca-doped ZnO sensor reached a value of 33 for 4000 ppm of ammonia at $T = 300$ °C with good selectivity compared to other gases such as CO, CO₂, and NO₂. The response and recovery times were 5 s and 221 s, respectively. The reported good sensing performances indicate the potential application of Ca-doped ZnO as a sensor material for ammonia detection.

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1. Introduction

Ammonia gas (NH₃) is often used in the production of textiles, polymers, refrigerant systems, fertilizers, insecticides, explosives, and food-based products.^{1–3} It is a colorless poisonous gas with a strong odor that has a threshold limit of 25 ppm. The long-term inhalation of NH₃ can be fatal^{4,5} since it can harm the eyes, skin, and the human respiratory system. In one instance, the American National Institute for Occupational Safety and Health specified that the immediately harmful to life or health concentration (IDLH) should be no more than 300 ppm. The Occupational Safety and Health Administration (OSHA) has established a threshold limit of 50 ppm for the workplace. So, to

safeguard our clean environment and regularly monitor the leakage of this gas, it is crucial to create a gas sensor to detect ammonia gas.⁶

Since metal oxide materials offer great qualities, such as a high sensing response, long-term stability, and ease of synthesis, they may generally be employed for gas sensing detections.⁷ These types of materials adsorb oxygen molecules, which causes the oxygen to become ionized by abstracting an electron for the metal oxides conduction band, resulting in the production of a surface depletion layer, and consequently increasing resistance. SnO₂, TiO₂, ZnO,^{8,9} WO₃, and Fe₂O₃ (ref. 10) are a few examples of several metal oxide-based gas sensors with diverse morphologies. ZnO nanostructures, which are among these metal oxides, have been widely employed as gas sensors because of their superior sensing response, strong selectivity, low cost, simple fabrication, outstanding thermal and chemical stability, and non-toxicity. In addition, ZnO material exhibited n-type conductivity having a resistivity in the range of 10^{−4} to 10¹² Ω cm,¹¹ broad bandgap (3.3 eV) at ambient temperature, high exciton binding energy (60 meV), strong electron mobility, great thermal stability, and non-toxicity.¹²

During the previous several decades, the number of industries using ammonia gas sensors has increased largely. In fact,

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there has been a significant increase in the interest in developing NH_3 gas sensors, which should present key characteristics including good sensitivity, selectivity, stability, and reversibility. They also need to be safe, affordable, small, lightweight, and able to operate in a wide range of temperatures.^{13,14} For example, Kumar *et al.* synthesized MnO_2 nanofibers and tested them toward 100 ppm of NH_3 gas at ambient temperature. The fabricated sensors showed a response of 20% and response and recovery times of 200 and 100 s, respectively.¹⁵ Besides, the ammonia gas sensor based on SnO_2 nanostructures fabricated by Beniwal and his team presented a response of 92% toward 500 ppm at ambient temperature and response/recovery times of 29/49 s.¹⁶ To develop ammonia gas sensor-based ZnO material with higher performances, numerous ways are utilized such as (i) applying an electrostatic field, (ii) using ultraviolet radiations in sensing operation, and (iii) doping materials using suitable metals.^{17–23}

In this study, we selected calcium as a dopant of ZnO to improve the sensing properties of ammonia gas. Calcium is cheap and non-toxic to human health contrary to other elements. It is also the third most common metal in nature, after iron and aluminum. The incorporation of Ca^{2+} ions promotes the decrease of the near-band edge emission and the increase of oxygen vacancy defects, which could improve the sensing characteristics. A sol-gel method was used to fabricate ZnO and ZnO:Ca nanopowders. After performing the structural and morphological characterizations, the synthesized powders were tested for NH_3 sensing. The sensing mechanism was probed and discussed. The results and information acquired allowed the development of a simple conductometric platform for ammonia gas detection with good sensitivity and selectivity.

2. Experiments section

2.1. Synthesis of ZnO:Ca nanopowders

A sol-gel method was used in the preparation of metal oxide nanomaterials because of the cheaper price, simple setup, and reduced energy use associated with this method. In this context, ZnO nanomaterials doped with Ca and with varying Ca contents (0, 1, and 3 at%) were synthesized supercritically under dry ethanol circumstances (critical temperature = 243 °C; critical pressure = 63.6 bars). 16 g of zinc salt $[\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}]$ and the calcium salt $[\text{CaCl}_2 \cdot 6\text{H}_2\text{O}]$ were dissolved in methanol solvent, as suggested by El Mir and his team.^{24,25} The drying process was performed in a stainless-steel autoclave. The obtained ZnO, ZnO:Ca_{1%} (C1ZO), and ZnO:Ca_{3%} (C3ZO) nanopowders were then heat-treated for two hours at 400 °C.

2.2. Samples characterization

XRD measurements. An X-ray diffractometer (AXS D8 Advance; BRUKER, Billerica, MA, USA), working at 40 kV voltage and 40 mA with copper $K\alpha$ irradiation of wavelength of 0.154056 nm in the 2θ range from 20 to 80 2-theta degree, was used to study the nanopowder crystallographic structure. After being ground into a fine powder in a mortar, a small portion was placed in the sample holder to perform measurements.

Scanning electron microscopy (SEM). The surface morphology of the samples was observed using a scanning electron microscope (JEOL 5600LV, Tokyo, Japan). Samples of nanoparticles were coated with 10 nm Pt (to stop the creation of electrostatic charges) and then attached to an SEM holder using carbon tape. SEM was powered by a 15 kV acceleration voltage.

Transmission electron microscopy (TEM). Using a Tecnai G20-S-Twin transmission electron microscope (FEI, Eindhoven, The Netherlands) with a 200 kV accelerating voltage, the morphology of the nanopowders was investigated. A small quantity of nanopowdered samples was dissolved in ethanol using an ultrasonic bath (Branson CPX5800H-E, USA) for a few minutes. The resulting solution was then poured onto a Cu grid covered with carbon and allowed to dry at room temperature in a clean room.

Fourier transform infra-red (FTIR) spectroscopy. A PerkinElmer 100 Spectrum FTIR spectrophotometer was used to investigate the functional groups in the different samples. After cleaning up all apparatus with acetone to avoid any contamination, a small portion of the sample was placed in an agate mortar; then a KBr was added. The mixture was ground until a fine powder was obtained. Then, the sample was placed in the FTIR compartment for the measurement.

2.3. Sensing tests

Using alumina substrates (6 mm × 3 mm) and platinum interdigitated electrodes, thick films of ZnO and ZnO:Ca powders dispersed in water (1–10 m thick) were printed to fabricate the devices for the sensing tests. On the underside of the electrodes, there was a Pt heater. A synthetic dry air total stream of 100 sccm was used for the electrical experiments, which were conducted in the RT to 400 °C temperature range. The four-point method was used to obtain the resistance data. The gas sensing performances were evaluated using a flow-through method. A dual-channel power supplier instrument, Agilent E3632A, was utilized to bias the device's built-in heater while a multimeter data collecting unit, Agilent 34970A, was used to collect the data. In order to execute NH_3 detection, pulses of this gas from certified bottles were injected into the 5 cc test chamber, at 100 sccm. Mass flow controllers and dry synthetic air (20% O_2 in nitrogen) as the diluting carrier, were used to adjust the desired gas concentration from 0.025 to 0.4% (250 to 4000 ppm). The gas response, S , is given as $S = [R_0/R_g]$, where R_0 is the electrical resistance in synthetic air at the baseline and R_g is the electrical resistance of the sensor at various NH_3 concentrations.

3. Results and discussions

3.1. Structural and morphological properties

The XRD patterns of the ZnO, ZnO:Ca_{1%}, and ZnO:Ca_{3%} samples are displayed in Fig. 1A. The observed diffraction peaks are indexed to the hexagonal wurtzite structure of ZnO (space group: $P6_3mc$ (186)) in agreement with (COD card no. 96-230-0451). The 2θ peaks appeared at $2\theta \approx 32.07^\circ$, 34.72° , 36.55° , 47.85° , 56.9° , 63.17° , 66.67° , 68.24° , and 69.4° corresponding to



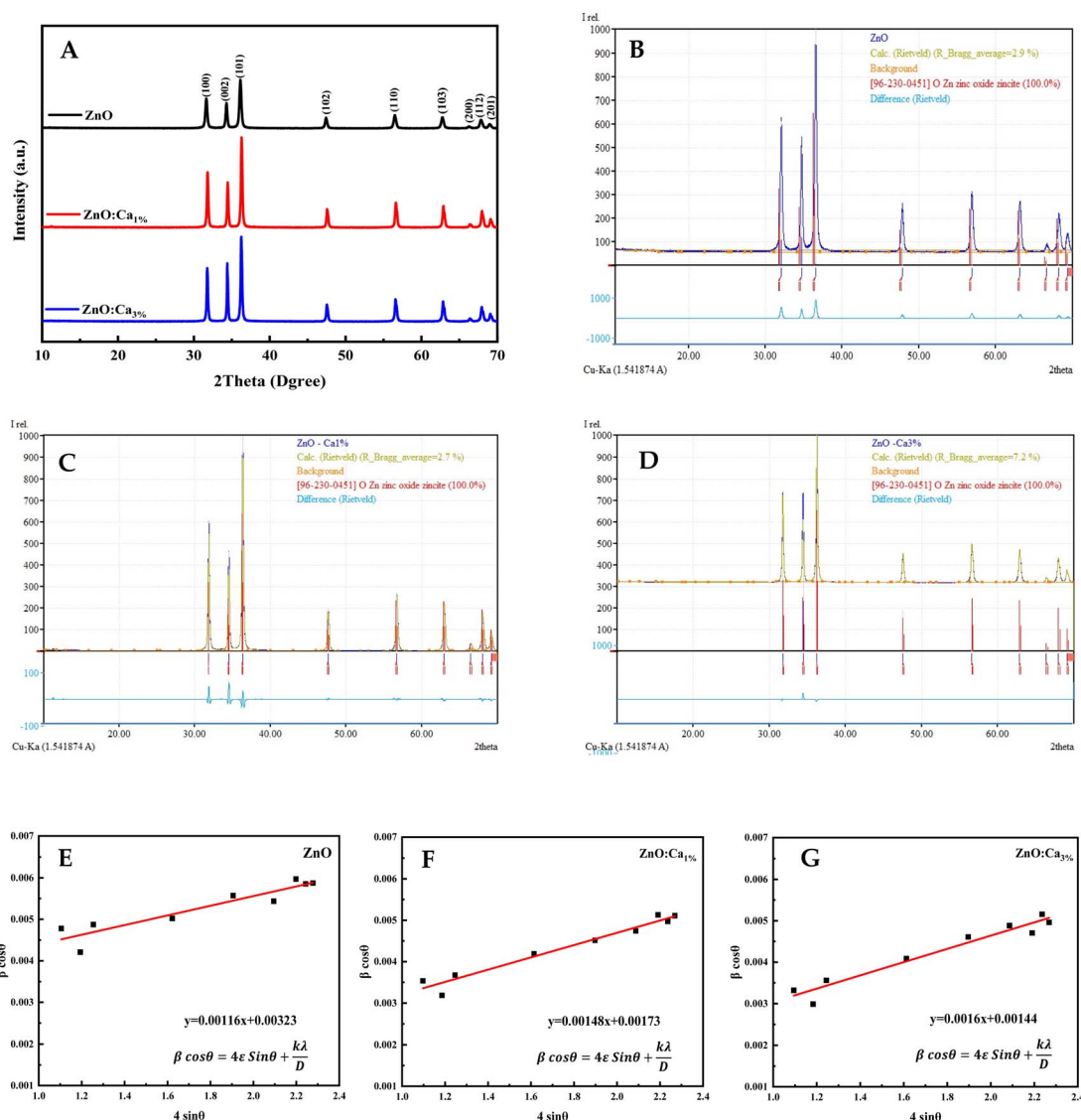


Fig. 1 (A) X-ray diffraction of ZnO and ZnO:Ca NPs. (B–D) Rietveld refinement plots of ZnO, ZnO:Ca_{1%}, and ZnO:Ca_{3%} (the blue curve represents the observed data, and the olive green color is the calculated pattern; the light blue curve shows the difference between the calculated and the observed intensities). (E–G) Williamson–Hall plots of ZnO, ZnO:Ca_{1%}, and ZnO:Ca_{3%}.

the crystal faces (100), (022), (101), (102), (110), (103), (200), (112) and (201), respectively.^{26,27} No extra peaks were observed due to the presence of any secondary phases.

The Rietveld refinement analysis shown in Fig. 1B and C was performed using the Match! software to obtain accurate structural parameters. The refined lattice parameters and other crystallographic data are summarized in Table 1. The lattice parameters “a” and “c” showed slight reductions upon calcium doping while the density increased, suggesting the successful incorporation of Ca into the ZnO lattice.

The crystallite sizes of all samples were determined using Scherrer's equation and Williamson–Hall's relation. Scherrer equation relates the crystallite size (D_s) to the peak broadening (β), wavelength (λ), and Bragg angle (θ) of the X-ray radiation as follows:²⁸

$$D_s = \frac{k\lambda}{\beta \cos \theta}$$

where K is the Scherrer constant (assumed as 0.9 for spherical particles).

Williamson–Hall's relation incorporates both the crystallite size (D_{WH}) and microstrain (ϵ) effects on the peak broadening:²⁹

$$\beta \cos \theta = 4\epsilon \sin \theta + \frac{k\lambda}{D_{WH}}$$

The calculated crystallite sizes using Scherrer's equation and Williamson–Hall's relation, along with the microstrain values, are also presented in Table 1. The comparison of the crystallite sizes obtained from Scherrer's equation and Williamson–Hall's relation revealed interesting trends. For all the samples, both



Table 1 The average crystallite sizes and parameters deduced from

Parameter	Samples		
	ZnO	ZnO:Ca ₁ %	ZnO:Ca ₃ %
<i>D_s</i> (nm)	26.55	32.84	33.78
<i>D_{WH}</i> (nm)	42.93	80.15	96.29
Microstrain ϵ	0.00116	0.00148	0.0016
Space group	<i>P6₃mc</i> (186)	<i>P6₃mc</i> (186)	<i>P6₃mc</i> (186)
Crystal system	Hexagonal	Hexagonal	Hexagonal
Density (g cm ⁻³)	5.671	5.678	5.679
Lattice parameters	<i>a</i> (Å)	3.2508	3.2495
	<i>c</i> (Å)	5.2078	5.2058

methods indicate an increase in the crystallite size with calcium doping. The ZnO:Ca₃% sample exhibited the largest crystallite size, followed by ZnO:Ca₁% and ZnO. Comparing the two methods, the Scherrer equation generally yields smaller crystallite sizes than the Williamson–Hall's relation. This discrepancy can be attributed to the inclusion of the microstrain effects in the latter. The presence of microstrain can arise from lattice defects, dislocations, or strain induced by the doping process.²⁹ The larger crystallite sizes obtained using Williamson–Hall's plot (Fig. 1E–G) indicate the presence of microstrain in the samples. Moreover, the microstrain also increased with an increase in calcium content, indicating the generation of more defects due to doping.

From FTIR, we investigated the functional groups in the undoped and Ca-doped samples. The spectra were recorded in the range of 500–4000 cm⁻¹. FTIR spectra of all samples, shown in Fig. 2 exhibit several characteristic absorption bands.

For undoped ZnO, the broad peak around 3100–3700 cm⁻¹ corresponds to the stretching vibration of O–H bonds in the hydroxyl groups (–OH) present on the surface of ZnO nanoparticles or adsorbed moisture.³⁰ The peaks around 1420 and 1638 cm⁻¹ are due to the production of carbonate species^{31,32} and the specifics related to asymmetric and symmetric COO⁻ vibrations. In addition, the sharp peaks at 650 cm⁻¹ and 870 cm⁻¹ correspond to Zn–O stretching vibrations, which is a characteristic feature of ZnO.³³ When calcium is doped into the ZnO

lattice, changes in the intensity of the O–H stretching peak around 3200–3700 cm⁻¹ were noted. The increase in the intensity suggests an increase in the concentration of O–H groups, likely due to the dopant calcium ions, which promote the adsorption of water. Further, whereas no intensity changes of the peaks in the fingerprint region (500–1000 cm⁻¹) for ZnO and ZnO:Ca₁% was observed; with ZnO:Ca₃%, there is an increase of the peak intensity in this region, due to the higher amount of calcium in the ZnO:Ca₃% sample. In any case, the presence of this impurity was not detectable by FTIR because the Ca–O bond overlaps with the Zn–O spectral bands in that region.¹⁸

Fig. 3A–C shows the surface morphology images of the samples recorded through SEM analysis. All the synthesized

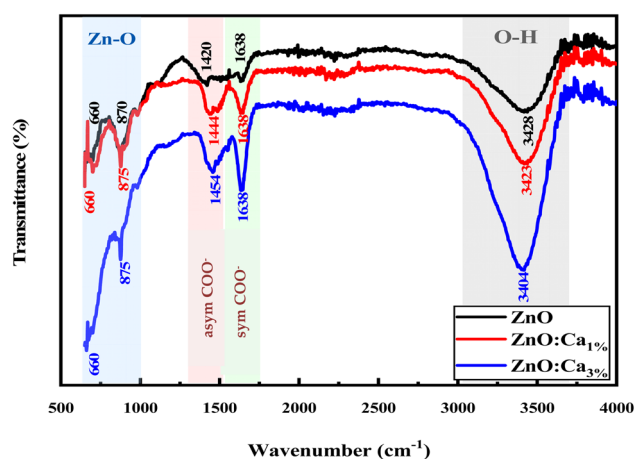
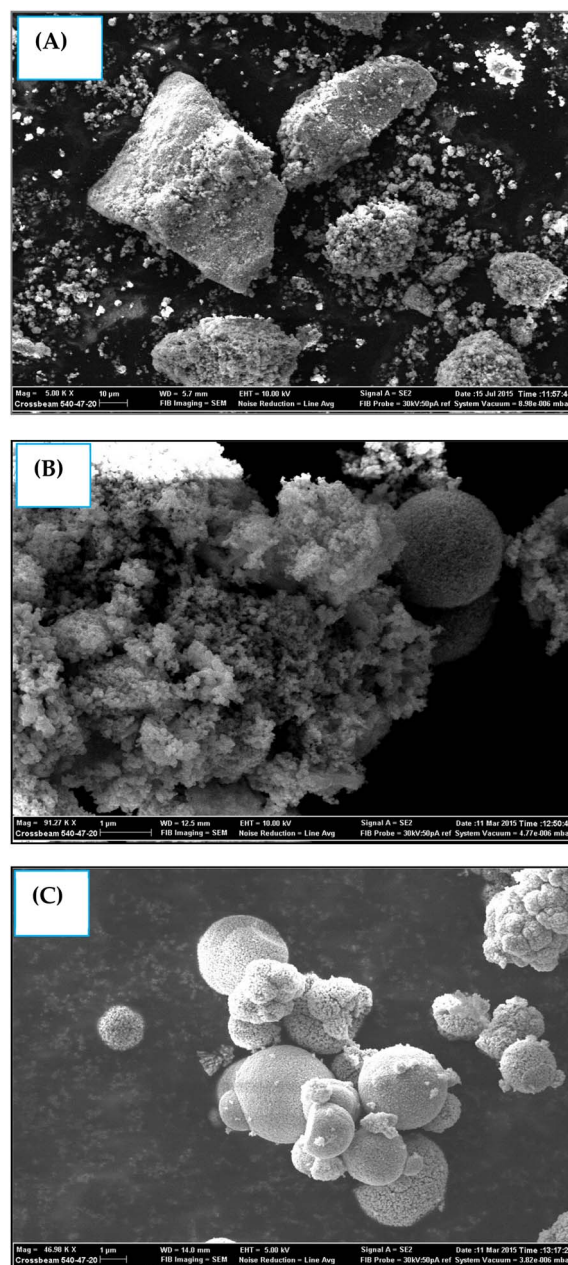


Fig. 2 FTIR spectra of ZnO and ZnO:Ca NPs.

Fig. 3 SEM photographs of (A) pure ZnO, (B) ZnO:Ca₁%, and (C) ZnO:Ca₃% nanoparticles.

samples appeared to be composed of small nanoparticles that were highly agglomerated.

Elemental analysis of CZO (1 at%) was carried out using color mapping analysis and EDX. The outcomes shown in Fig. 4 confirmed the presence of main elements, *i.e.*, Zn, O, and Ca in the sample with good uniform distribution. No meaningful presence of any other contaminating species was recorded.

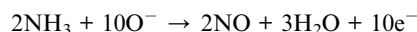
Fig. 5(A) and (B) shows TEM and (C) and (D) HRTEM images of ZnO and ZnO:Ca_{3%} samples. The images clearly show that the doped and undoped NPs have asymmetric spherical-like shapes and clearly visible lattice fringes.

The particle size distribution was determined from the TEM image using an image analysis software by measuring the diameter of individual nanoparticles and creating a histogram showing the frequency of nanoparticles at each diameter size as shown in Fig. 5(E) and (F). The size distribution illustrated that the mean particle sizes of ZnO and ZnO:Ca_{3%} are 37.62 nm, and 114.15 nm respectively. This confirmed the increase in particle size with calcium doping, supporting the XRD findings.

3.2. Sensing performances

Dynamic responses of different samples are reported in Fig. 6. The operating temperature was 300 °C and the samples were tested for various concentrations of NH₃ gas. In all the samples,

the resistance was decreased after the exposure of the gas, and then it returned to the starting value indicating that ZnO material kept its n-type nature with NH₃ injection. This was normal behavior because ammonia is a reducing gas and Ca dopant did not alter the n-type nature of ZnO. The same results were found by Radhi Devi and the team when the strontium-doped ZnO sensor was tested toward various concentrations of NH₃ gas.³⁴ Kathwate *et al.* also noticed that the resistance of the Al-doped ZnO films decreased after ammonia injection.³⁵ In contrast, Nakarungsee's team noticed that the resistance increased when Cr-doped ZnO sensors were tested toward NH₃ gas.³⁶ The decrease of material resistance could be explained using the traditional theory of resistance change³⁷ following the contact with a reducing molecule on the surface of the n-type semiconductors, which may be used to understand the observed drop in resistance. The resistance decrease was a result of gas molecules and adsorbed oxygen species interactions. The adsorbed oxygen species and reducing gas molecules will react on the surface of the material as described by the reaction below:



The resistance drop phenomenon after the NH₃ (reducing gas) injection was observed in ZnO and Ca-doped sensors

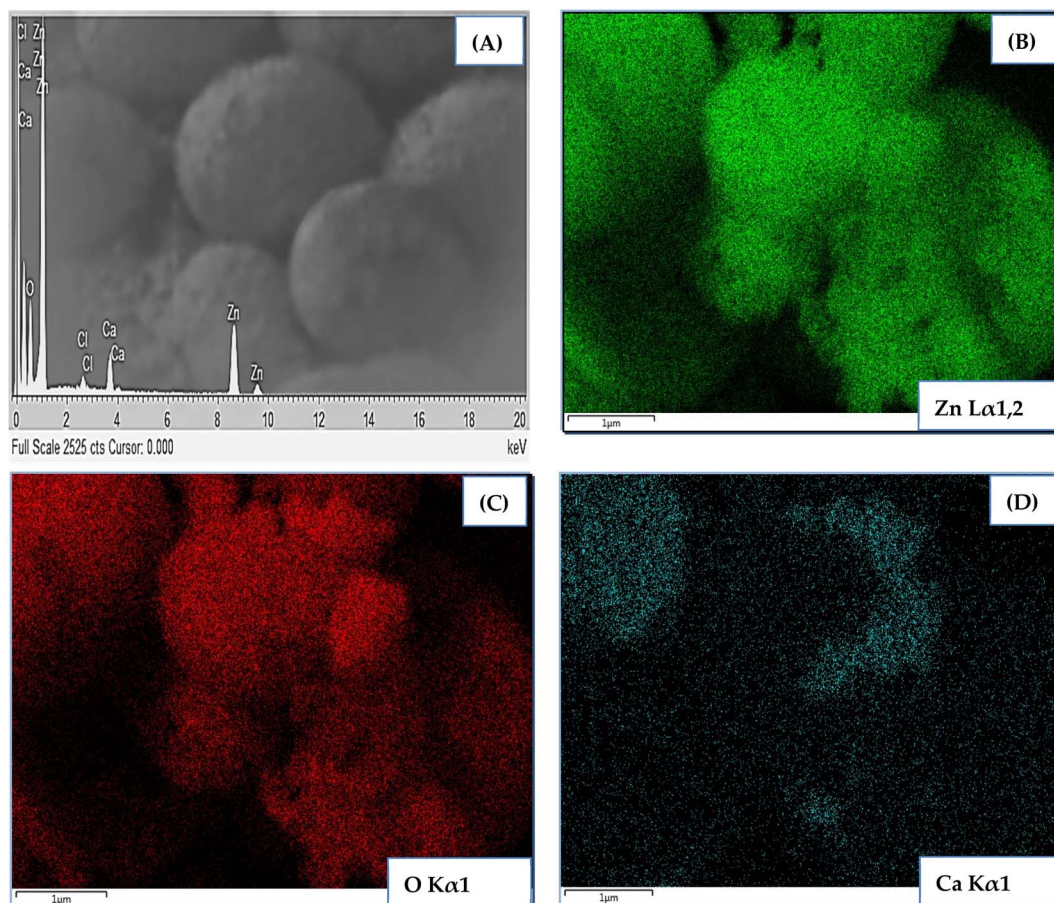


Fig. 4 EDS spectrum and elemental color mapping of 1 at% CZO.

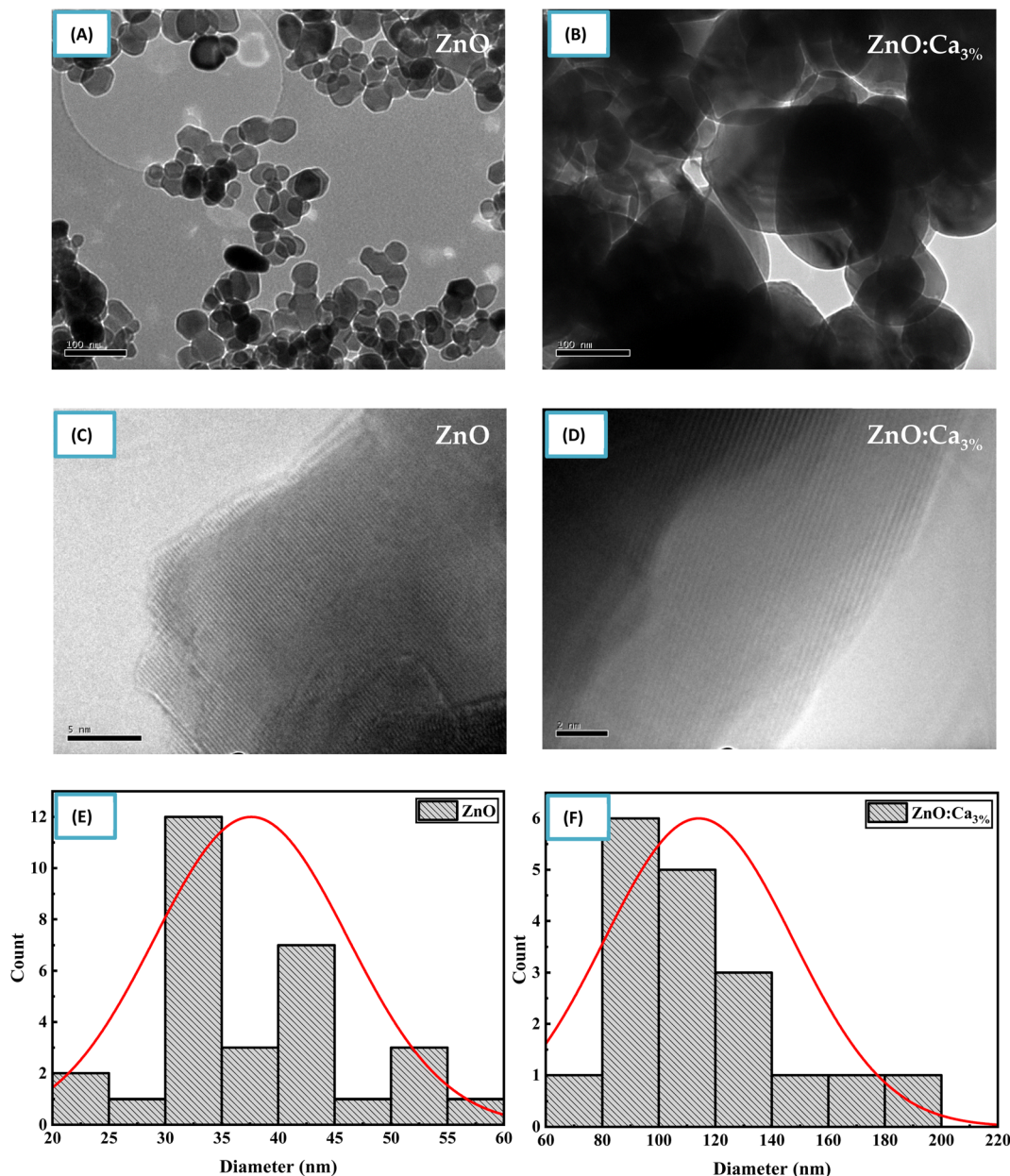


Fig. 5 (A–D) TEM and HRTEM images of ZnO and ZnO:Ca nanoparticles. (E and F) Particle size distribution of ZnO and ZnO:Ca nanoparticles.

suggesting that the incorporation of calcium ions did not alter the n-type nature of the ZnO material.

For pure ZnO sensors, the baseline resistance takes a value of 0.32 MΩ. A small change in resistance is noticed with NH₃ concentrations increase, as shown in Fig. 6(A). In Fig. 5(C) and 6(B), Ca-doped sensors exhibited a bigger baseline resistance (60 MΩ for C1ZO and 22 MΩ for C3ZO) in addition to a remarkable change in the resistance after the gas injection for both doped samples. The baseline resistance of doped samples was higher than that of the undoped ones. The same behavior was shown by Dhahri *et al.* in their research;³⁸ Misra's team³⁹ has also observed an increase in the baseline resistance in Ca-doped material. This is probably due to a drop in the concentration of carriers and/or due to the creation of a carbonate layer

after the calcium incorporation in the ZnO network, which changes the semiconductor surface.⁴⁰

As displayed in Fig. 6(D) the response, calculated as, $S = \frac{R_{0,41}}{R_g}$ reached a maximum value of 33 for the C1ZO sensor toward 0.4% NH₃ concentration at $T = 300^\circ\text{C}$, and it was reduced 10 times for the C3ZO sample. This response reduction for the C3ZO sensor can be explained as follows: at higher doping levels, calcium ions may not substitute zinc ions but may be going into the interstitial sites inside the ZnO or on the surface. In addition, as mentioned above ZnO:Ca_{1%} exhibited a particle size smaller than that of ZnO:Ca_{3%}. Smaller crystallite size means higher surface area and more sensitive sites. In addition, on the ZnO:Ca_{3%} sample, there is an excess of Ca that covers the



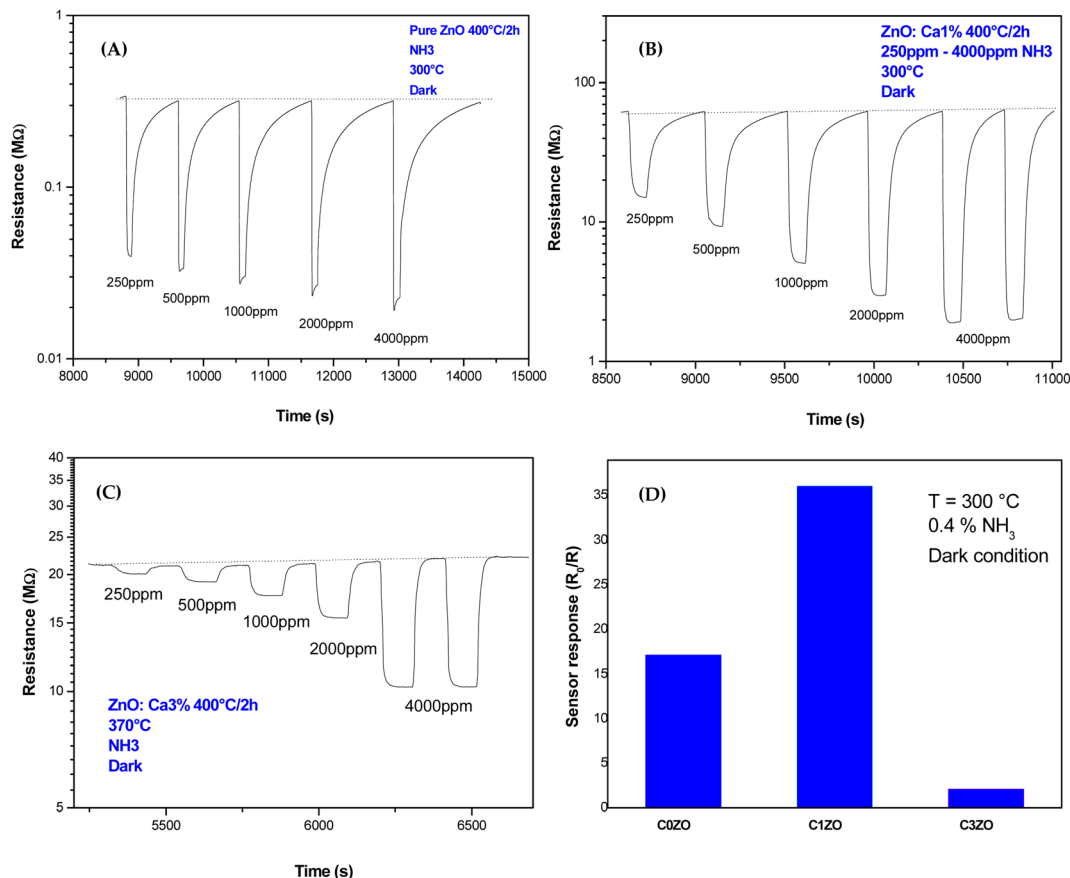


Fig. 6 Dynamic response of (A) pure ZnO, (B) ZnO:Ca_{1%}, and (C) ZnO:Ca_{3%} nanoparticles toward various concentrations of NH₃ gas. (D) Response of different samples toward 0.4% of NH₃ at $T = 300$ °C.

zinc oxide surface and decreases the interaction sites, consequently leading to a reduction of gas response.

Fig. 7 shows the response of the ZnO:Ca_{1%} material toward different ammonia concentrations at various temperatures. It is clearly seen that at low NH₃ concentrations, the gas response increases slowly upon reducing the temperature. After

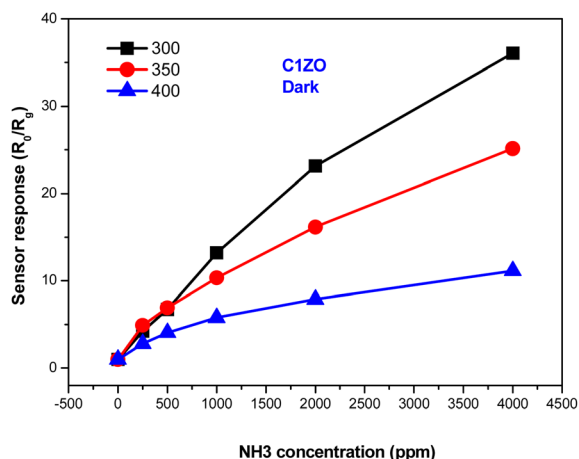


Fig. 7 Variation of the ZnO:Ca_{1%} gas response towards different ammonia concentrations at different temperatures.

increasing gas concentrations, the response enhances largely and reaches higher values at $T = 300$ °C. We can say that the greatest response toward ammonia was obtained at lower temperatures ($T = 300$ °C).

The gas response of the undoped and ZnO:Ca_{1%} sensors toward different ammonia concentrations at $T = 300$ °C is presented in Fig. 8. At low gas concentrations, the undoped sensor exhibits a higher response compared to the doped one. Increasing the gas concentrations, the C1ZO sensor response enhances and becomes better than that of the pure sensor until reaches a maximum value of $S = 33$. This behavior can be explained in terms of electronic defects such as oxygen vacancies (V_O) and zinc interstitials (Zn_i), which are created after the addition of Ca ions in the ZnO network, as cited by Saeedi and his colleagues.⁴² Zn_i and V_O acted as donor defects, leading to an increase in the concentration of free electrons.⁴³ These free electrons take part in the interaction between the sensing layer and ammonia molecules by reducing C1ZO sensing layer resistance and then the response increases. Besides, ammonia adsorption is favored on zinc active centers. However, increasing the concentration of ammonia, leads to the saturation of these active centers, while the presence of Ca ions can maintain the surface cleaner, favoring the desorption of ammonia from the Ca–ZnO active sites. This was confirmed by the very large difference in the recovery time, which is very

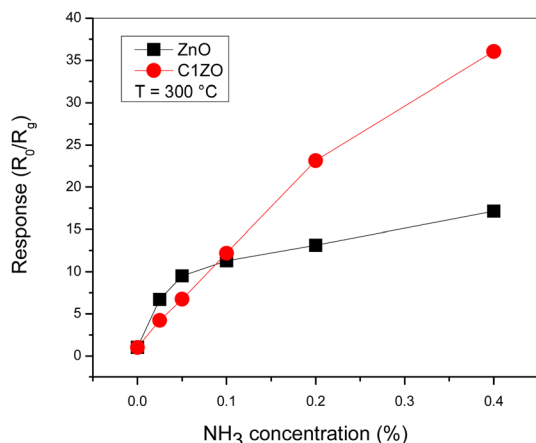


Fig. 8 Comparison between pure ZnO and ZnO:Ca_{1%} sensors response toward different NH₃ concentrations at $T = 300\text{ }^{\circ}\text{C}$.

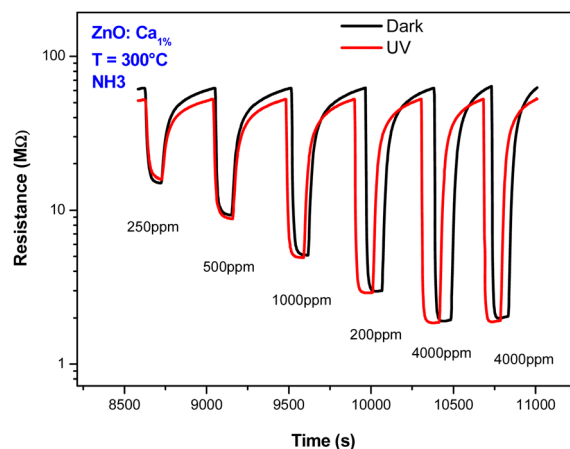


Fig. 10 Dynamic response behaviors of ZnO:Ca_{1%} sensor toward NH₃ concentrations with and without UV light illumination.

long for the pure ZnO and becomes faster increasing the Ca loading.

The response/recovery times of different sensors are displayed in Fig. 9. The response/recovery times are (6 s, 718 s), (5 s, 221 s), and (18 s, 37 s) for pure ZnO, C1ZO, and C3ZO samples, respectively. All the samples exhibited quick response times, but slow recovery times, especially ZnO and C1ZO. This outcome showed that there was a considerable quantity of ammonia adsorbing on the sensing layer's surface, leading to the hypothesis that the rate at which ammonia or its intermediary species desorb from the surface is slower than the rate at which ammonia is adsorbed.⁴⁴ The recovery time of the C3ZO sensor seems to be faster.

Also, the effect of UV light was investigated. Fig. 10 presents the dynamic response of the C1ZO sensor toward different concentrations of ammonia at $T = 300\text{ }^{\circ}\text{C}$ under dark conditions and UV radiation. Under UV light illumination, the baseline resistance of the sensing layer decreased a bit. The drop in the resistance of metal oxides by UV light is due to photoelectron activation. In fact, a decrease in the indium oxide resistance model suggested by Wagner's team highlighted the diffusion of oxygen species in and out of the In₂O₃ lattice after radiation with 350 nm.⁴⁵ The sensor response is also little

influenced by the presence of the ultra-violet light. This behavior can be explained by considering that operating with UV at high temperatures generally does not lead to an improvement of the sensing properties. The positive influence of ultra-violet illumination on electrical and gas sensing properties is instead well recognized at low operating temperatures.

To show the high selectivity of the suggested sensors, especially C1ZO one, toward ammonia gas, all sensors were tested toward several oxidizing and reducing gases such as CO, CO₂, and NO₂, as shown in Fig. 11. In the beginning, the addition of 1 at% of calcium well improved the sensor response toward NH₃. More calcium concentration (3 at%) in the ZnO network leads to poorer response compared to undoped and C1ZO sensors. The particle shape was excluded from the reasons for the gas sensor's response amelioration because all the samples maintained the same shape even after the incorporation of Ca²⁺ ions.

Testing sensors toward other gases (CO, CO₂, and NO₂) had a negligible effect on the gas response in comparison with ammonia indicating the high selectivity of the fabricated

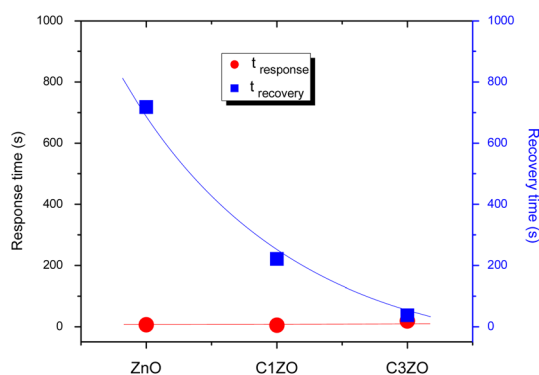


Fig. 9 Response/recovery times of various sensors.

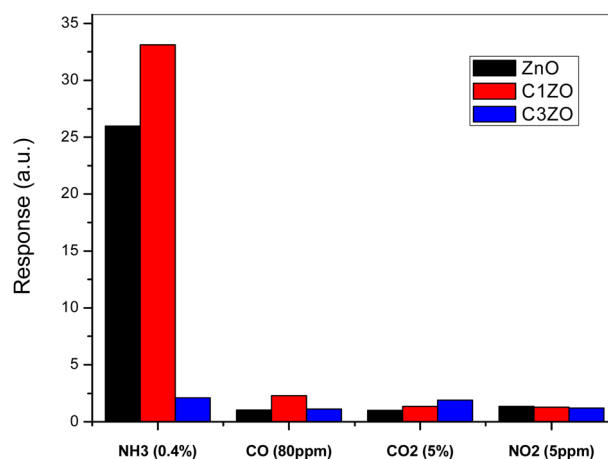


Fig. 11 Selectivity of the studied sensors when tested toward a mixture of gases (NH₃, CO, CO₂, and NO₂).



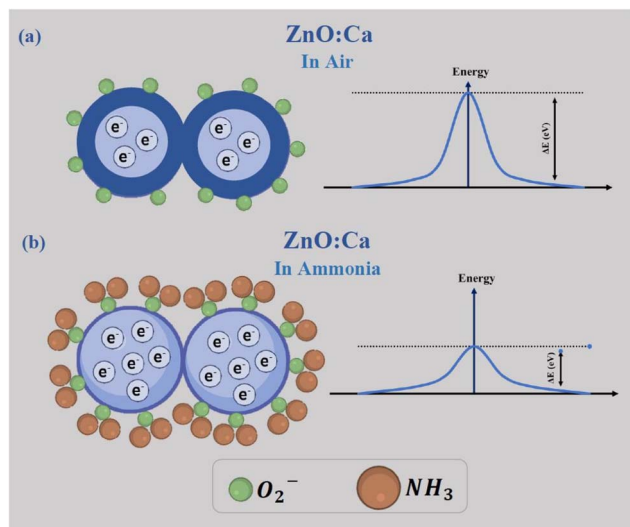


Fig. 12 The sensing mechanism of ZnO:Ca₁% nanopowders (a) in air and (b) in ammonia atmosphere.

sensors for ammonia. Chaudhary and his team attributed this high selectivity to the smaller kinetic diameter (0.26 nm) and low ionization energy (10.18 eV) of ammonia.⁴⁶ With its small kinetic diameter, NH₃ can easily enter pores of the sensing layer and this leads to an enhancement of the gas response toward NH₃.⁴⁶

3.3. NH₃ sensing mechanism

The chemical processes that occur between the sensing material and the target gas are at the basis of the ammonia gas sensing mechanism on metal oxide-based conductometric sensors.⁴⁷ As observed in Fig. 12, two main steps describe these chemical processes: the first step is the exposure of the ZnO:Ca layer to air (baseline condition) when the adsorption of oxygen molecules occurs on the sensing surface. Adsorbed oxygen interacts with free electrons from the conduction band of the Ca-doped ZnO sensing material, leading to the creation of ionized oxygen species (O²⁻, O⁻, O₂⁻), and the creation of a depletion layer on the surface and a potential barrier at the grain boundaries, establishing the baseline resistance (Fig. 12a).

In the second step, when NH₃ is injected, the gas molecules are adsorbed on the surface and react with the adsorbed oxygen species, liberating free electrons. This process involves a decrease of the barrier potential (Fig. 12b), so the free electrons can flow easily, and consequently, a noticeable drop in the resistance was obtained in response to NH₃ exposure, according to our experimental observations.

4. Conclusions

ZnO and ZnO:Ca thick films were screen printed on Pt-interdigitated alumina substrates and tested toward various concentrations of ammonia gas from 0.025–0.4%. The synthesized nanopowders were characterized by XRD, FTIR, TEM, and SEM. The samples were crystalline and showed the hexagonal

wurtzite structure. The sizes of the typical crystallite, as determined by the Williamson–Hall method, increased with the addition of Ca. The gas sensing performances were clearly improved with the ZnO sensor doped with 1% of Ca. The developed sensor also displayed high NH₃ selectivity with relation to other gases such as CO, CO₂, and NO₂, and fast response, thus demonstrating the possible use of ZnO:Ca for ammonia gas detection.

Data availability

The data used to support the findings of this study are available from the corresponding author upon request.

Author contributions

Mokhtar Hjiri: writing and editing; investigation; Saja Algessair: conceptualization; R. Dhahri: formal analysis; Hassan B Albargi: formal analysis; Nabil Ben Mansour: data curation; Amine Aymen Assadi: conceptualization; Giovanni Neri: supervision.

Conflicts of interest

The authors declare no conflict of interest.

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