Physicochemical conditions for ZnO films deposited by microwave chemical bath deposition

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Physicochemical analysis was carried out to obtain the species distribution diagrams (SDDs) for the deposition of ZnO films as a function of OH⁻ ion concentration ([OH⁻]) in the reaction solution. The study of SDDs predicts nucleation and ZnO film growth by means of the dominant species at a given pH value. To confirm this, a series of experiments were made varying the [OH⁻] in the reaction solution and keeping the others parameters constant. Structured zinc oxide (ZnO) films were obtained on glass substrates by microwave chemical bath deposition (MWCBD). Structural, optical and morphological ZnO film properties were investigated as a function of [OH⁻]. X-Ray diffraction technique (XRD) measurements show multiple diffraction peaks, indicating the polycrystalline nature of ZnO films. Scanning Electron Microscopy (SEM) images of ZnO structures showed morphological changes with the variation of [OH⁻]. The stoichiometry of the structures changed as the [OH⁻] was varied in solution. From Raman spectra, it was observed that the [OH⁻] of the reaction mixture strongly affects the crystal quality of ZnO structures. A reaction pathway for the synthesis of ZnO structures based on our results is proposed. Experimental results are consistent with the physical–chemical analysis.

1. Introduction

Due to technological and electronic advances, the world faces the challenge of creating materials with versatile properties. Those materials must fulfill some requirements such as having inert properties, a wide band gap, high electron mobility as well as a great resistance to field rupture. Zinc oxide (ZnO) is a semiconductor material, which has a band gap of 3.37 eV, an exciton binding energy of 60 meV at room temperature and a melting point of 1975 °C. Those properties are important in a semiconductor for the future needs. Besides, in recent years, ZnO has become one of the most studied materials due to its very interesting properties in optoelectronic devices application such as; room temperature lasers, light emitting diodes, ultraviolet (UV) detectors, field-emission displays, photonic crystals, solar cells and sensing in the nano-size range. The control of its properties is primarily in the context of novel applications as highly efficient gas sensing material and photocatalysis. For this reason, a better understanding of the synthesis of microstructures with different morphologies is important and necessary to achieve objectives and applications on materials for the modern world. A profitable, simple and easily scalable method to synthesize the material is the microwave chemical bath deposition (MWCBD). This technique allows us the production of multifunctional structures to a larger scale. In addition, it is possible to have control of the material properties by only changing the synthesis conditions. Moreover, in recent years, the process of microwave assisted heating has been used as an attractive alternative for the synthesis of nanoscale materials, since it is a fast and effective method that allows us to increase the kinetic reaction on one or two magnitude orders. On the other hand, to control the physicochemical properties of a final product synthesized by chemical bath deposition process, knowledge of the species generated during reaction is required. Therefore, an analysis on solution conditions needs to be done. This analysis could allow us to construct species distribution diagrams (SDDs) at the chemical equilibrium. SDDs have proven to be a successful tool to find the optimal films deposition conditions by chemical bath.

In this work, a methodology to synthesize structured ZnO films by the MWCBD technique based on a theoretical physicochemical analysis is presented. SDDs were constructed by using different OH⁻ ions concentration ([OH⁻]) in the analytical model. The [OH⁻] used in the experiments and the [OH⁻] used in the SDDs are related with the structural, optical and morphological properties of the deposited ZnO films. The ZnO deposition is carried out by means of a controlled urea
precipitation for the formation of the chelate \([\text{Zn(urea)}]^{2+}\), which provides the dissolution reaction of \(\text{Zn}^{2+}\) ions in a controlled way.\(^{19}\) It was found that adjusting pH in the initial solution, the grain size, morphology, stoichiometry \(\text{O}/\text{Zn}\) (% at) ratio and structural properties of the \(\text{ZnO}\) films can be controlled. A possible reaction pathway based on the SDDs and the experimental results is presented.

2. Physical–chemical analysis

2.1 Solubility and species distribution diagrams

Different authors have proposed \(\text{Zn(OH)}_2\) as a precursor of \(\text{ZnO}\).\(^{21,23}\) the \(\text{Zn(OH)}_2\) solubility product is about \(10^{-18}\) (mol \(\text{L}^{-1}\)). Thermodynamically, \(\text{ZnO}\) can be chemically deposited only if the \([\text{Zn}^{2+}]/[\text{OH}^-]\) ratio overcomes the solubility product, as it was reported the decomposition.\(^{26-28}\) The urea decomposition reaction consists of a series of steps, where the time plays an important role. The films deposition in MWCBD technique takes a short time period in contrast with the urea decomposition reaction, this ensure that the aqueous solution is not contaminated with the products of the urea decomposition during the films deposition.

In the analytical model \([\text{OH}^-]\) was adjusted from \(1.77 \times 10^{-5}\) to \(1.38 \times 10^{-3}\) at \(80\,^\circ\text{C}\) to construct the SDDs as shown in Fig. 1. When the \([\text{OH}^-]\) increases in the solution, the \([\text{NH}_4^+]\) increases too due to the ammonium hydroxide (\(\text{NH}_4\text{OH}\)) used, then in the analytical model the \([\text{NH}_4\text{OH}]\) is varied and the \([\text{OH}^-]\) is adjusted. Species or reaction products formed, for each \([\text{OH}^-]\) in the SDDs are listed in Table 1. We can also observe from Fig. 1 that due to reaction conditions, the number of nucleation centers \([\text{Zn(OH)}_2]^2\) and growth units \([\text{Zn(OH)}_3]^3\) and \([\text{Zn(OH)}_4]^4\) are also all different for each \([\text{OH}^-]\) in the solution. The \([\text{OH}^-]\) was calculated for each pH value, which are also listed in Table 1. Fig. 1a shows the SDDs for a \([\text{OH}^-]\) of \(1.77 \times 10^{-5}\), where we can see chemical species with different charges which form at a specific \([\text{OH}^-]\); species with positive charge are below of 9 pH value, the neutral specie \(\text{Zn(OH)}_2^2\) is between 8 to 11 pH values and species with negative charge are above of 9 pH value. It can be appreciated almost the same pattern for greater \(\text{OH}^-\) ions concentrations (Fig. 1b) \(7.58 \times 10^{-5}\), (Fig. 1c) \(2.45 \times 10^{-4}\), (Fig. 1d) \(8.70 \times 10^{-3}\) and (Fig. 1e) \(1.38 \times 10^{-3}\). As an example; in (Fig. 1e) species with positive charge are below of 10 pH value, the neutral specie \(\text{Zn(OH)}_2^2\) is between 9 to 11 pH values and species with negative charge are above of 9.5 pH value. Increasing \([\text{OH}^-]\) also increase the \([\text{NH}_4^+]\) (\(\text{NH}_4\text{OH}\)) concentration, resulting in an increment of the relative concentration of the species with positive charge \(\text{Zn(NH}_3)_3^{4+}\) and \(\text{Zn(NH}_3)_4^{5+}\) with a decrement on \(\text{Zn(OH)}^1\) and neutral \(\text{Zn(OH)}^0\) species. From these observations in the SDDS the neutral specie \(\text{Zn(OH)}^0\) is between 8.5 to 11 pH values and its relative concentration increase when the \([\text{OH}^-]\) increase, species with positive charge are below of 10 pH values and species with negative charge are above of 9 pH value, overlapping between them in 9 and 10 pH values, for all \([\text{OH}^-]\) studied in this work.

Solubility diagrams (SDS) can be constructed too, as well as SDDS were constructed. Thermodynamically, \(\text{ZnO}\) can be chemically deposited only if the \([\text{Zn}^{2+}]/[\text{OH}^-]\) ratio in the solution overcomes the \(\text{Zn(OH)}_2\) solubility product, as it was said before. MWCBD technique increase the kinetic reaction,
thus the previous thermodynamic condition could or could not influence the films deposition, in another way, chemical species that become dominant in the solution as a function of the pH value must influence the films deposition. The species with positive charge \( \text{Zn(NH}_3\text{)}_{2+}, \text{Zn(NH}_3\text{)}_{4}^{2+} \) rarely would participate in the ZnO growth because they do not contain oxygen, even though, the \( \text{Zn(OH)}^{1-} \) specie has positive charge and may participate. Besides, \( \text{Zn(OH)}_{2}^{0} \) (nucleation center), \( \text{Zn(OH)}_{3}^{-} \) and \( \text{Zn(OH)}_{4}^{2-} \) (growth units) species surely would participate in the ZnO growth because they contain oxygen. Thus, from the SDDs, chemical species that would participate in the ZnO growth are above of 9 pH values, and the species that would not participate are below of 10 pH values. Therefore, \( [\text{OH}^{-}] \) that produce pH values above 10 could be appropriate for ZnO growth by MWCBD process. When increase the \( [\text{OH}^{-}] \), the pH increase and the species \( \text{Zn(OH)}_{3}^{-} \) and \( \text{Zn(OH)}_{4}^{2-} \) (growth units) become dominant, thus ZnO growth is promoted.

3. Materials and methods

In order to investigate the physicochemical conditions and predictions described by the SDDs, initial solutions were prepared at room temperature by dissolving zinc nitrate (0.1 M) and urea (0.1 M) into deionized (DI) water (resistivity 18.2 MΩ) under magnetic stirring, keeping the molar ratio in 1 : 1. Different amounts of aqueous ammonia (\( \text{NH}_4\text{OH} \)) were added to the solutions in order to adjust the initial \( [\text{OH}^{-}] \) to: (a) \( 1.77 \times 10^{-5} \), (b) \( 7.58 \times 10^{-5} \), (c) \( 2.45 \times 10^{-4} \), (d) \( 8.70 \times 10^{-4} \) and (e) \( 1.38 \times 10^{-3} \) values obtained from the pH measurement in each solution. Three well-cleaned glass substrates were immersed vertically into the solutions and introduced simultaneously into a commercial microwave oven (Sharp, model R-408J, frequency 2.45 GHz) and were thereby four minutes in continuous microwave irradiation at full power (1200 W) until the reaction solution temperature reached 80 °C (±2 °C), after that, the microwave irradiation process was then continued for another 20 minutes at low microwave power (keeping a constant temperature of 80 °C). Finally, the substrates were taken out of the reaction beaker and rinsed with DI water several times under vigorous stirring and dried at room temperature in air.

For the purpose to observe the ZnO films properties, X-ray diffraction (XRD) was used to investigate the structures of the ZnO films. Morphological characterization of the ZnO films was carried out in a JEOL (JSM-5400LV) Scanning Electron Microscope (SEM) equipped with OXFORD analytical Energy Dispersive X-ray Spectroscopy (EDS) system. The Raman measurements were performed at room temperature using a LabRAM HR-Olympus Micro Raman spectrometer, equipped...

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### Table 1

| \( [\text{OH}^{-}] \) ions concentration | pH value | Predominate species for each pH value |
|----------------------------------------|---------|-----------------------------------|
| \( 1.77 \times 10^{-5} \)              | 9.25    | \( [\text{Zn(OH)}]^{+} \approx [\text{Zn(OH)}]_{2}^{+} \approx [\text{Zn(OH)}]_{3}^{-} \) |
| \( 7.58 \times 10^{-5} \)              | 9.88    | \( [\text{Zn(OH)}]_{3}^{0} > [\text{Zn(OH)}]_{2}^{+} > [\text{Zn(OH)}]^{+} \) |
| \( 2.45 \times 10^{-4} \)              | 10.39   | \( [\text{Zn(OH)}]_{3}^{0} > [\text{Zn(OH)}]_{2}^{+} \) |
| \( 8.70 \times 10^{-4} \)              | 10.94   | \( [\text{Zn(OH)}]_{3}^{0} > [\text{Zn(OH)}]_{2}^{+} \approx [\text{Zn(OH)}]^{0} \) |
| \( 1.38 \times 10^{-3} \)              | 11.14   | \( [\text{Zn(OH)}]_{3}^{0} > [\text{Zn(OH)}]_{2}^{+} \approx [\text{Zn(OH)}]_{3}^{-} \) |
with a He–Ne laser with an excitation of 633 nm. The film thickness was obtained on a profilometer Veeco (Bruker); model DEKTAK 150 with a Stylus of 12 μm.

4. Results and discussions

Fig. 2 shows the XRD patterns from ZnO films grown on glass substrates by using different [OH−] in the solution. The diffractograms revealed several well-defined diffraction peaks, all matching with the standard peak positions (JCPDS Card No. 36-1451) of ZnO in hexagonal wurtzite phase. Is interesting to note that for pH values of 9–10, the relative intensity of the diffraction peaks is small (in the SDDS there are species with positive and negative charge in those pH values), and as the pH increase (increasing [OH−]), from 9.25 to 11.14, the relative intensity of the diffraction peaks increases too, indicating a better crystal ZnO growth and confirming the predictions mentioned by the SDDS. On the other hand, for the 11.14 pH, the relative intensity of the diffraction peaks corresponding to planes (100) and (101) diminish and the (002) increase, this means that the crystal growth is promoted only along plane (100) and (101) for the different pH, those values are listed in Table 2.

The interplanar distance d was evaluated with the Bragg’s law: nλ = 2d sin θ (n = 1, λ = 1.54184 Å) for each crystallographic plane. The data in Table 2 were used to determine the structural parameters for the ZnO films in hexagonal wurtzite phase listed in Table 3. The lattice parameters a and c were calculated using the formula: \( \frac{1}{d^2} = \frac{4(h^2 + k^2 + l^2)}{a^2c^2} \), where h, k, and l are the Miller indices. The Zn–O bond length L was evaluated using the equation: \( L = \sqrt{\frac{a^2}{3} + \frac{1}{2}u^2} \), where a and c are the lattice parameters, u is the positional parameter that is a measure of the amount by which each atom is displaced with respect to the next along the ‘c’ axis and is obtained by the expression: \( u = \frac{a^2}{3c^2} + 0.25 \). The grain size D was determined for each crystallographic plane using Scherrer’s equation from full width at half maximum peak position:

\[ \beta = \frac{0.94λ}{D \cos θ} \]

where λ = 1.54184 Å and β = FWHM in Table 2. Finally, the strain (ε) in the films was calculated using the relation:

\[ \beta = \frac{λ}{D \cos θ} - ε \tan θ \]

All equation to determine the structural parameters in this work of the ZnO films in hexagonal wurtzite phase has been reported and are widely referenced. It can be observed in Table 2, that the interplanar distance d seems stretching and shrinking for all the diffraction planes when the pH increase in the solution. The same behavior can be noted for the lattice parameters a, c and the Zn–O bond length L in Table 3 when the pH increase. This behavior can be due to the total zinc or oxygen atomic in the ZnO lattice of the films, there is more zinc or oxygen when the pH increase, the ZnO unit cell volume stretching until a pH value about 10.39, increasing the pH it shrinks and stabilizes.

On the other hand, in Table 3 can observe that the grain size diminishes for the (100) plane until a pH value about 10.94 and for the (002) plane the grain size increases until a pH value about 10.39. The strain for its part increases for the (100) plane until a pH value about 10.94, increasing the pH it diminishes, for the (002) plane the strain increases and diminishes on every step that the pH increases. The ZnO wurtzite structure is composed of two interpenetrating hexagonal close packet (hcp) sublattices, where in the (100) crystallographic plane have equal proportion of zinc and oxygen atoms surrounded by each other, and in the (002) diffraction plane there is only oxygen or zinc atoms. Experimentally was reported that the (002) crystallographic plane is strongly dependent upon the O2 gas in sputtered ZnO films. The formation and ZnO growth is promoted when increasing the [NH4OH] in the solution allowing more zinc and oxygen atomic in the ZnO unit cell until a pH value about 10.94. The XRD results confirms the SDDS predictions, otherwise these results may indicate a change in the morphology of the ZnO films surface.

Fig. 3 shows SEM images of ZnO structures grown on glass substrates by MWCBD technique using different [OH−]: \( 1.77 \times 10^{-1}, 7.58 \times 10^{-5}, 2.45 \times 10^{-4}, 8.70 \times 10^{-4}, 1.38 \times 10^{-3} \) and the following pH values: 9.25, 9.88, 10.39, 10.94 and 11.14 that were measured respectively. Fig. 3a and b show the morphology of the ZnO films grown at 9.25 and 9.88 pH values. It is evident that in both cases the ZnO structures appear in
nucleation stage (some clusters can be appreciated), the samples mainly consist of nanorods assembled into a branched morphology. The nucleation stage is by the presence of ZnOH\(^+\) and Zn(OH)\(_2\) (nucleation center) in the SDDs. From Fig. 3, it also can be seen that by increasing the pH above 9.88 (Fig. 3c–d) these structures become larger; increasing [OH\(^-\)]/C\(_0\) promotes the ZnO film growth and the species Zn(OH)\(_3\)/C\(_0\) and Zn(OH)\(_4\)\(^{2-}\)/C\(_0\) are dominant above 10 pH value according the SDDs, thus these species promote the ZnO film growth and the others with positive charge do not. The increased shape ratio may be due to the Zn(OH)\(_2\) clusters in the substrate surface that act partly as nucleation centers, besides the Zn(OH)\(_3\)\(^-\) and Zn(OH)\(_4\)\(^{2-}\) ions

| pH  | Peak position 2\(\theta\) (degree) | FWHM peak position (degree) | d (Å)       |
|-----|----------------------------------|-----------------------------|-------------|
|     | (100) (002) (101)                | (100) (002) (101)           | (100) (002) (101) |
| 9.25| 31.72 34.36 36.21                | 0.2255 0.2153 0.2671        | 2.821 2.611 2.481 |
| 9.88| 31.72 34.36 36.18                | 0.2302 0.2228 0.2553        | 2.821 2.611 2.483 |
| 10.39| 31.64 34.32 36.15               | 0.2398 0.1932 0.2548        | 2.828 2.613 2.485 |
| 10.94| 31.81 34.44 36.27               | 0.2816 0.2269 0.2754        | 2.813 2.604 2.477 |
| 11.14| 31.82 34.48 36.29               | 0.2405 0.2181 0.2639        | 2.812 2.601 2.475 |

Table 3 Structural parameters of the ZnO films for different pH values

| pH  | Lattice parameters | Zn-O bond length | Grain size | Strain |
|-----|-------------------|-----------------|------------|--------|
|     | \(a\) (Å) \(c/\alpha\) | \(c/\alpha\) (Å) | \(D\) (nm) | \(\epsilon\) \(\times 10^{-4}\) |
| 9.25| 3.257 5.219 1.602 | 1.983           | 38.28     | 32.71  | 8.84 7.76 9.11 |
| 9.88| 3.257 5.219 1.602 | 1.983           | 37.51     | 34.22  | 9.03 8.03 8.71 |
| 10.39| 3.265 5.226 1.601 | 1.987           | 35.99     | 34.28  | 9.43 6.97 8.69 |
| 10.94| 3.248 5.208 1.603 | 1.977           | 30.66     | 31.73  | 11.01 8.16 9.37 |
| 11.14| 3.247 5.202 1.602 | 1.976           | 35.90     | 33.11  | 9.39 7.83 8.97 |

Fig. 3 SEM images of ZnO structures grown by MWCBD technique using the [OH\(^-\)]; (a) 1.77 \(\times\) 10\(^{-5}\), (b) 7.58 \(\times\) 10\(^{-5}\), (c) 2.45 \(\times\) 10\(^{-5}\), (d) 8.70 \(\times\) 10\(^{-4}\) and (e) 1.38 \(\times\) 10\(^{-3}\).
are responsible for the ZnO growth due to their oxygen excess. On the other hand, from Fig. 3e, it can also be observed that when the pH reaches 11.14 some of the ZnO structures become smaller again. This behavior is due to the total formation of zincate ions such as Zn(OH)$_3^{2−}$ and Zn(OH)$_4^{2−}$ when NH$_4$OH is added in excess.\textsuperscript{25-28} Once again, the predictions from the SDDs are confirmed with the morphological results in this work.

ZnO film thicknesses obtained by profilometry were: 563, 992, 2065, 4619 and 5798 (nm) as [OH$−$] was adjusted for 1.77 × 10$^{−5}$, 7.58 × 10$^{−5}$, 2.45 × 10$^{−4}$, 8.70 × 10$^{−4}$ and 1.38 × 10$^{−3}$ respectively, which correspond to initials pH values of 9.25 to 11.14. Regarding these thicknesses we can say that the substrate does not influence the compositional results obtained by EDS. Increasing [OH$−$] the ZnO film thickness increases as well, confirming the prediction made in the SDDs.

Fig. 4 shows the O/Zn (% at) ratio obtained by EDS analysis as a function of the [NH$_4$OH] used in the MWCBD. As it can be observed from this figure, the zinc stoichiometry rate and oxygen content into the films change as the [OH$−$] changes as well. At the beginning with the first two [OH$−$] the zinc atomic percent is low, until the solution reaches a pH value of 10.39 due to the [OH$−$] used. This is due to greater oxygen content in the solution when NH$_4$OH is added and/or by the species with positive charge that do not participate in the ZnO growth but they contain zinc. Then, when the pH increases above 10.39 the zinc atomic percent almost remains constant; this could mean that the ZnO growth process is stable on this pH value. It was experimentally observed that the solution becomes transparent at pH = 10.42, that should be caused by the ZnO amphoteric character, avoiding homogeneous nucleation; therefore, the thickness of the films increase as the pH increases according to profilometry measurements. This could indicate us that Zn(OH)$_3^{−}$ and Zn(OH)$_4^{2−}$ chemical species may be favorable for the ZnO grow as it was previous discussed. The O/Zn (% at) ratios for the MWCBD ZnO films obtained by EDS analysis are consistent with the XRD results as the [NH$_4$OH] increases in the reaction solutions.

Fig. 5 displays the Raman spectra from the ZnO structures in the 90–800 cm$^{−1}$ spectral range. On all the spectra at least seven Raman peaks located at around 102, 204, 335, 385, 439, and 583 cm$^{−1}$ can be observed. The peaks located at 102, 385, 415, 439 and 583 cm$^{−1}$ correspond to the $E_{2L}$, $A_1(TO)$, $E_1(TO)$, $E_{2H}$ and $E_1(LO)$ optic vibration modes of wurtzite ZnO, respectively.\textsuperscript{29} From the plots on the spectra (Fig. 5) it can be observed that the [OH$−$] modification in the reaction mixture strongly affects the intensity and width of the Raman peaks. The effects are most prominent for the $E_{2L}$ and $E_{2H}$ modes, located at around 102 and 439 cm$^{−1}$, the decreased intensity and observed broadening is a strong indication that the crystal quality is impoverished for the two lowest [OH$−$] used. This effect can be attributed to the presence of Zn(OH)$_3^{−}$, Zn(NH$_3$)$_4^{2+}$ and Zn(NH$_3$)$_3{\cdot}^{−}$ that do not promote the ZnO growth, in the SDDs (pH 9–10) by the two lowest OH$^{−}$ ions concentration. On the other hand, the zincate ions Zn(OH)$_3^{−}$ and Zn(OH)$_4^{2−}$ may affect the ZnO crystal quality inducing oxygen defects on the lattice at a pH about 11. The formations of these zincate ions linked to the (100) plane of the ZnO nanorods, when NH$_4$OH is added in excess, are responsible of the nanostructures crystallinity degradation when the pH value is increased above of 11.0. Finally, the peak that appeared around 583 cm$^{−1}$ can be ascribed to the $E_1$(LO) mode, associated with defects such as oxygen vacancy and interstitial zinc. Raman spectra clearly demonstrates that all ZnO structured films grown for various pH values have the hexagonal wurtzite structure and those with...
good crystal quality are for the [NH₄OH] that produced an initial pH between 10 to 11.

4.1 ZnO chemical reaction pathway
According the results, the SDDs and the main soluble species containing Zn²⁺ and OH⁻ ions, we can propose a possible chemical pathway to grow ZnO by MWCBD, which can be:

\[
\text{Zn(OH)}_2 \rightarrow \text{ZnO} + \text{H}_2\text{O} \quad (12)
\]

\[
\text{Zn(OH)}_3^- \rightarrow \text{ZnO} + \text{H}_2\text{O} + [\text{OH}^-] \quad (13)
\]

\[
\text{Zn(OH)}_4^{2-} \rightarrow \text{ZnO} + \text{H}_2\text{O} + 2[\text{OH}^-] \quad (14)
\]

Species with zero charge Zn(OH)₂ eqn (12) are responsible for ZnO nucleation while the species Zn(OH)₃⁻ and Zn(OH)₄²⁻ with negative charge are the responsible for ZnO growth. \{Zn(OH)₂\} form clusters on the substrate (start of precipitation heterogeneous) eqn (12). Following the pathway growth, the species \{Zn(OH)₃⁻\} and \{Zn(OH)₄²⁻\} have the availability to reconfigure the OH⁻ ions concentration in the solution, increasing the concentration of free ions in solution as eqn (13) and (14) describe. The solution is supersaturated again and the free ions have a tendency to condense on the surface of the formed particles. On the other hand, the substrate surface plays a role in the ZnO films heterogeneous precipitation, the MWCBD technique increases the kinetics of the reaction, so the catalytic activity of the substrate surface diminishes. This process is carried out due to the species responsible for the nucleation and growth coexists for a pH > 10 values in the SDDs.

5. Conclusions
A physicochemical analysis was carried out to investigate the conditions of ZnO deposited by microwave chemical bath deposition. The reaction solution was made with zinc nitrate, urea and different [NH₄OH]. SDDS was constructed as pH function in order to study and find the main chemical species in the solution at a specific [OH⁻] that affect the nucleation and ZnO growth. Structured ZnO films were grown on a glass surface, the effect of [NH₄OH] variation on the precursor solution is related on the structural, optical and morphological properties of ZnO films. The intensity of the (002) peak in the XRD diffraction patterns of ZnO films is favored when the [NH₄OH] increases up to 11-pH value and another diffraction planes are disfavored, below this pH value, more atomic oxygen enter in the ZnO unit cell. The SEM images of ZnO structures grown on glass substrates show that the shape ratio is affected by Zn(OH)₂ clusters and the formation of zinicate ions such as Zn(OH)₆⁻ and Zn(OH)₄²⁻ that act as nucleation center and grow units respectively during heterogeneous precipitation. Therefore, it is thought that the crystal quality degradation could be caused by these ions. The intensity and width of Raman peaks were an indicative that a change in the ZnO films occurs with the variation of [NH₄OH] in the precursor solution. Raman spectra clearly demonstrated that all ZnO structured films grown for different [NH₄OH] have the hexagonal wurtzite structure. Moreover, the samples grown with pH value above 11.0 presents a decrease of crystallinity. A MWCBD ZnO growth reaction pathway is proposed according to the results and the chemical species of the SDDS. According to SDDS and characterization results, ZnO films can grow from pH values of 9 to 11. The best chemical conditions to deposit ZnO films by MWCBD was found in an initial pH range from 10 to 11, for depositing ZnO films with good properties in this investigation.

Conflicts of interest
The authors declare no conflict of interest.

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