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Synthesis and Characterization of 1-D Cerium Oxide to Study Their Humidity Sensing Application

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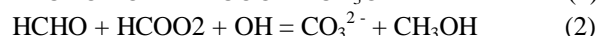
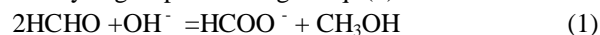
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Abstract: The formaldehyde-assisted hydrothermal system is used to synthesize CeO₂ nanorods by using precursor like cerium chloride. Concentration of precursors and hydrothermal temperature affects the morphologies of these one-dimensional (1D) CeO₂ Nanomaterial's. This is achieved to control the reaction degree of Cannizzaro disproportionation tuned by Na/Ce molar ratio. The XRD, FESEM, FTIR and TEM these characterization techniques have been used to characterize the crystal structure and morphology of the as-synthesized CeO₂. Humidity sensing characteristics such as hysteresis, repeatability, impedance-relative humidity (RH) characteristics, response and recovery behavior, and stability of CeO₂ nanostructure have been investigated in detail for given precursor. The sensor showed rapid response and recovery, prominent stability, high humidity sensitivity, good repeatability and narrow hysteresis loop i.e. suitable candidate for the fabrication of the humidity sensors.

Keywords: Nanorods, cerium chloride, humidity sensor, XRD, FESEM

I. INTRODUCTION

One-dimensional (1D) nanostructured materials such as nanowires, nanotubes, nanorods and nanobelts offer opportunities for fundamental research concerning the influence of size and dimensionality of a material on its physical and chemical properties.[1–3]. Cerium oxide (CeO₂) is a technologically important rare earth material because of its wide range applications as oxygen sensors [10], polishing agents [7], fuel cells [9], catalysts [8], and UV blockers [11]. Indirect synthetic pathway of 1D CeO₂ Nanorods results in increasing complexities and difficulties in the design and synthesis of their desired morphologies [12]. Formaldehyde, may used as an organic chemical, because of its rich abilities in synthetic reactions. [12–15] the Cannizzaro disproportionation reaction occurs immediately, when formaldehyde solution is mixed with strong alkali under the heated conditions in presence of the aldehyde group according to eqⁿ (1) and the formate is produced simultaneously.[12]



According to the cross disproportionation reaction, the aldehyde groups presented at formate can further react with formaldehyde to produce carbonate under superfluous alkali conditions (eqⁿ (2)). We can convert, formaldehyde into formate or/and carbonate by controlling the reaction conditions. The formate and carbonate are the great precursors for the synthesis of 1D CeO₂ Nanorods.[16–18] we can synthesize 1D precursors by controlling the reaction degree of the Cannizzaro disproportionation reaction. We can also state that formaldehyde solution is the most efficient solvent to synthesize 1D CeO₂ precursors among all attempted solvents such as acetaldehyde, isopropyl alcohol, ethanol, glyoxal, ethylene glycol, acetone,[15] etc

II. EXPERIMENTAL

A. Preparation of 1D CeO₂ nanorods

1.0 g of CeCl₃ was dissolved in 100 ml formalin solution at room temperature and then 2.1 gm of potassium hydroxide was slowly added to the solution and the whole solution was placed on magnetic stirrer for uniform mixing [5]. Then the solution was kept into a 200 ml Teflon-lined stainless autoclave and heated at 120^oC temperature for 20 hrs. The solution then cooled to room temperature, the precipitates were filtered, washed with double distilled water in several time and dried at 60^oC for 12 hrs [7]. The dried powders were calcinated to 400^oC for 2 hrs. The light yellow powder was obtained.

B. Preparation of Sensors Electrode

In present work the sensor consisting of CeO₂ nanorods layer coated on the top of an interdigitated electrode (IDE) was fabricated [10–13]. The IDE consists of five pairs of Cu tracks screen printed onto epoxy glass substrate (25 mm x 20 mm)[4]. The IDE-epoxy glass substrates were cleaned by an ultrasonic treatment in acetone and then rinsed with double-distilled water and dried in vacuum[17]. The powder of CeO₂ nanorods was mixed with double-distilled water in a weight ratio of 100: 25 to form a paste. The electrodes were used to evaluate the humidity sensing characteristics [14].

III. RESULTS AND DISCUSSION

A. Characterization of CeO₂ nanorods.

The XRD pattern of the CeO₂ nanorods as-synthesized by cerium chloride after calcinated at 400°C is depicted in Fig.1(a). All the diffraction peaks in the XRD pattern of CeO₂ are exactly matched with JCPDS file (JCPDS No.: 00-057-0401), indicating the formation of face centered cubic CeO₂ nanorods. No impurities were present and pure CeO₂ nanorods are obtained[16-18]. The average crystallite size of the CeO₂ nanorods from XRD graph was found to be in the range of 10-17 nm.

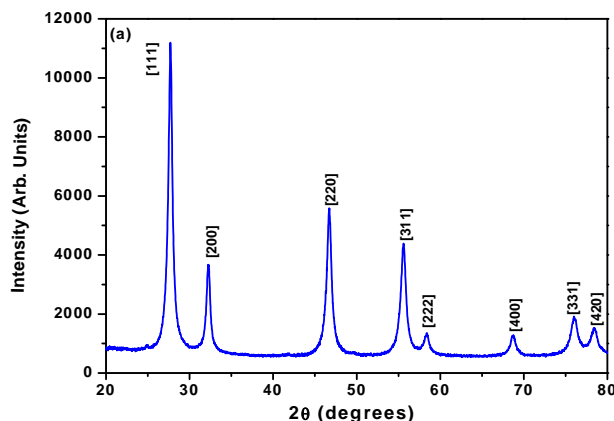


Fig.1 : XRD of CeO₂ nanorods

The FTIR spectrum of the CeO₂ nanorod synthesized by cerium chloride [Fig.2(a) and Fig.2(b)] shows the bands around 451 and 853 cm⁻¹ corresponding to a stretching vibrations characteristic of Ce-O. The bands around 1064 and 1327 cm⁻¹ shows to the characteristic vibrations of CeO₂. The stretching vibrational mode of the O-H group bonded to the Ce atom i.e. Ce-OH is given by band at ~ 3410 cm⁻¹. There is no distinction between FTIR of commercial and our obtained CeO₂ powders [18].

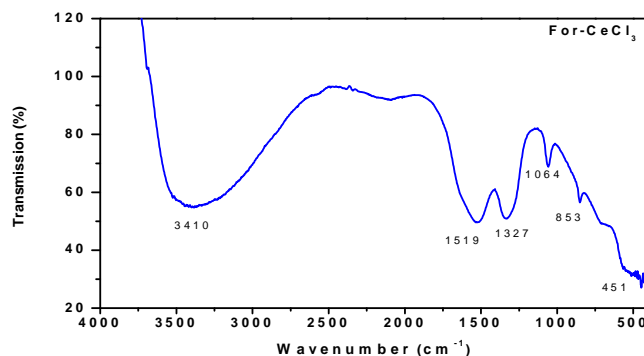


Fig.2 : FTIR of CeO₂

The field emission scanning electron microscopy (FESEM) and transmission electron microscopy (TEM) were used to characterize the morphology and size of the CeO₂ nanorods [1-3]. The FESEM images of as-synthesized CeO₂ product by formaldehyde-assisted hydrothermal system using cerium chloride, indicate that the 1D Nanomaterial's of CeO₂ nanorods with typically 300-500 nm in thickness, 8 -12 μm in length.

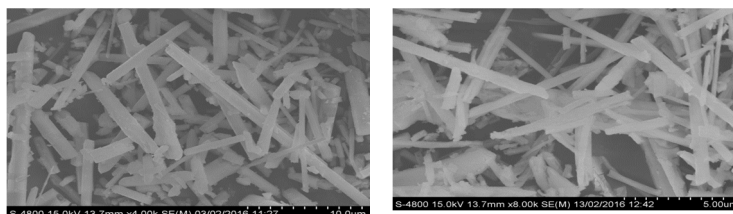


Fig.3 : (a) FESEM images of as-synthesized CeO₂ nanorods precursor using cerium chloride and (b) CeO₂ nanorods after calcinations

The TEM images shows CeO₂ nanorods structure with 494.92 to 1057.29 nm in thickness and 1065.02 to 4736.14 nm in length.

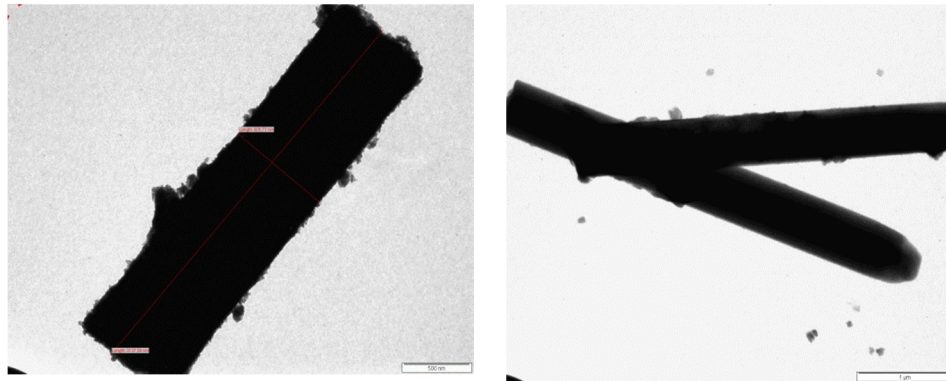


Fig.4 : (a) TEM images of as-synthesized CeO₂ nanorods precursor using cerium chloride (b) CeO₂ nanorods after calcinations

B. Humidity Sensing Measurements.

Different RH levels were generated by the different saturated salt solutions in air tight closed glass bottles at room temperature [3]. The six different standard saturated aqueous salt solutions of LiCl (11±0.30 %RH), MgCl₂ (33±0.14 %RH), K₂CO₃ (43±0.20 %RH), NaCl (75±0.15 %RH), KCl (85±0.24 %RH) and K₂SO₄ (97±0.16 %RH) were used to act as humidity source [18]. The sensing element was placed successively into the bottles with different RH levels at room temperature and the impedance of the sensor was measured as a function of RH at 27 oC (± 1 oC). The frequency range was varied between 60 Hz to 1 kHz for 1V applied voltage [14]. A humidity probe was also placed into the bottles along with the sensing element to monitor the RH during the measurement [4]. The response and recovery times were measured by switching the sensors back and forth between two closed bottles with RH values of 11% and 97% RH respectively [12]. The hysteresis was measured by switching the sensor between the closed bottles with 11%, 33%, 43%, 75%, 85% and 97% RH and then transferred back.

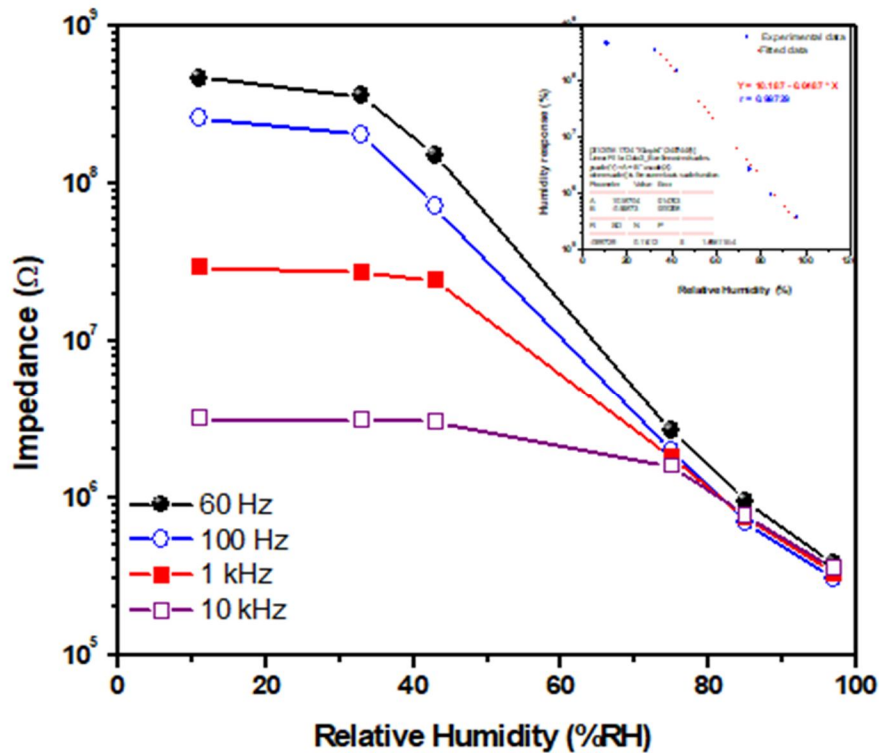


Fig.5 : Variations in impedance of CeO₂ nanorods based humidity sensor with change in RH (%) measured at various frequencies and 1 V. Inset depicts linear fit to the Impedance-RH curve measured at 60 Hz to 10 kHz and 1 V. calcinated 3000C

The linear frequency response was observed at relatively low measurement frequency i.e. at 60 Hz [10]. The impedance of the sensor changes by three orders of magnitude from 3.3×10^5 to $4.7 \times 10^8 \Omega$ as RH increases from 11% to 97%.

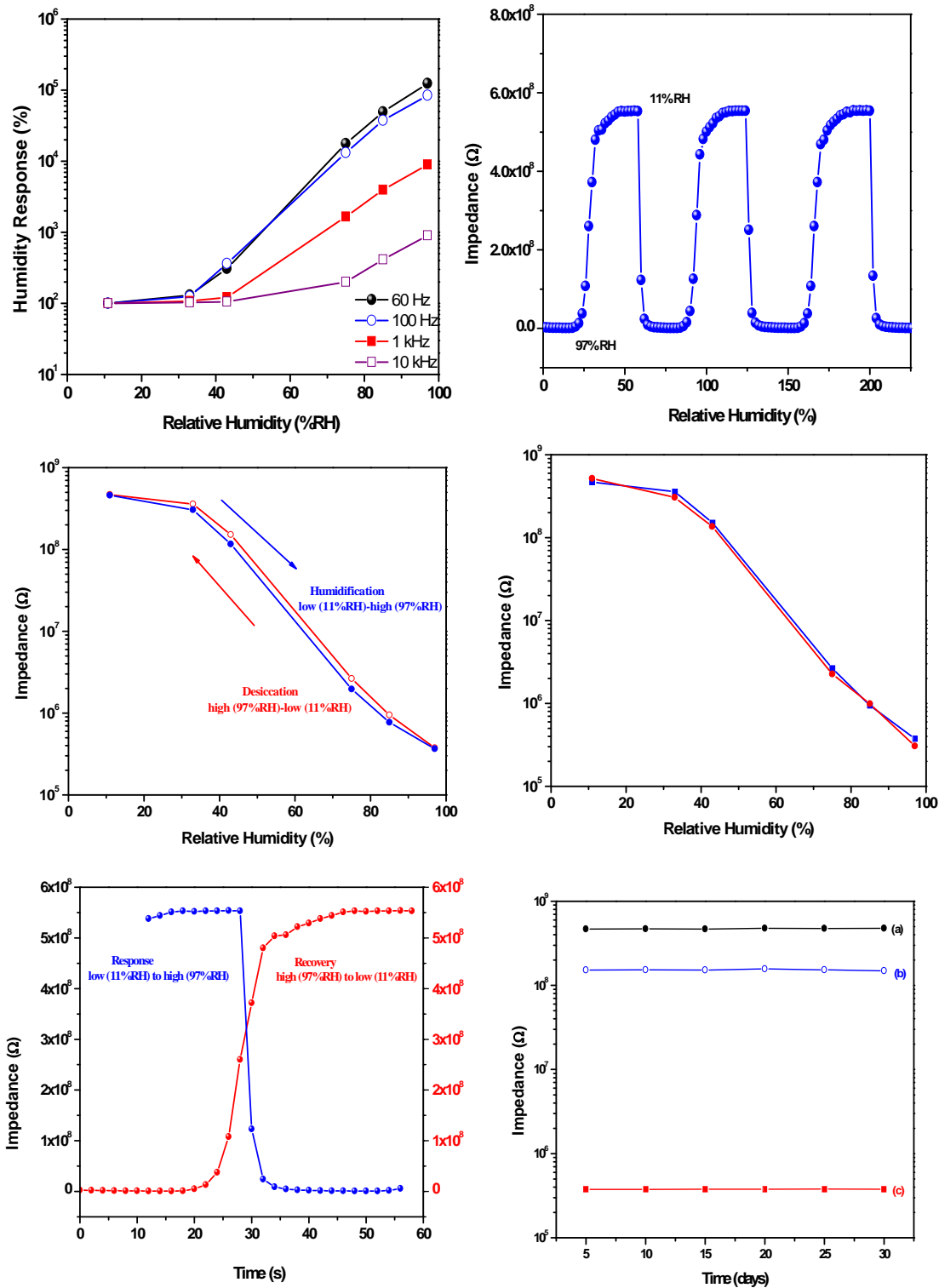


Fig.6 : (a) and (b) Dependence of response on RH and frequency for CeO₂ nanorods based sensor, (c) and (d) Humidity hysteresis and (e) Dynamic response of the sensor when exposed to six high (90% RH) – low (10% RH) – high (90% RH) cycles. The inset shows the response and recovery characteristics, (f) Stability of the sensor measured at 11, 43 and 97 % RH.

The dynamic response of the CeO₂ nanoparticles based sensor to rapid variations in the RH values of 11% and 97% is shown in Fig. 6(a). The sensor was switched back and forth between two closed bottles with RH values of 11% and 97%, respectively [17]. The response time (humidification from 11% RH to 97% RH) was 2-3 s and the recovery time (desiccation from 97% RH to 11% RH) was 9-10 s, respectively (see inset in Fig.6 (e)), demonstrating that the present sensor rapidly responds to the ambient RH [10]. The maximum absolute value of humidity hysteresis error γ_H is found to be ~1% in the range of 11-97% RH indicating a good reliability of the sensor in fig 6(c). The stability is an important parameter of humidity-sensing properties [9]. In the period of 30 days the sensor was tested repeatedly once in five days under fixed humidity levels (11%, 43% and 97% RH). The impedance variation (Fig.6(f)) is less than 2% at each humidity region for one month at 60 Hz, which shows that the impedance of the sensor fluctuates slightly with time and the data show good consistency [17].

IV. CONCLUSIONS

CeO₂ nanorods were successfully synthesized at low cost by using a simple hydrothermal method. The XRD results reveal the formation of face centered cubic phase of CeO₂ nanorods with good crystallinity and the crystalline size ranging in 10 -16 nm. The TEM images shows CeO₂ nanorods structure with 494.92 to 1057.29 nm in thickness and 1065.02 to 4736.14 nm in length. The FTIR bands around 1052 and 1385 cm⁻¹ show to the characteristic vibrations of CeO₂. FESEM shows CeO₂ Nanorods of 300-500 nm in thickness, 8 -12 μ m in length. The CeO₂ Nanorods exhibit excellent humidity sensing characteristics such as higher response, fast response time (~ 2-3 s), rapid recovery (~ 9-10 s), hysteresis within 1.0%, excellent reproducibility and broad range of operation (11-97% RH). It was demonstrated that the CeO₂ Nanorods can be used as reusable sensing material for the fabrication of humidity sensors.

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