

Research Article

Effect of Various Reducing Agents on Synthesis of ZnO Nanoparticles

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ABSTRACT:

Background: Nanotechnology is the science that deals with matter at the scale of 1 billion of a meter and is also the study of manipulating matter at the atomic and molecular scale. In this study ZnO nanoparticles are prepared by using sol gel method. This method is used by many researchers because of its ease of use, easy availability of apparatus, low cost and it is easy to perform.

Purpose: The aim of this research is to study the size of ZnO nanoparticles using different reducing agents e.g., NaOH or NH_3OH or Triton X-100. The used chemicals are highly pure with AR grade.

Methods: The precipitates were formed and annealed at temperature approx. $1000^{\circ}C$. The precipitate was collected and characterized by X-ray diffraction (XRD) to calculate the crystalline size of the nanoparticles.

Results: Nanosized ZnO powder was successfully synthesized and XRD analysis revealed that crystalline size is nearly independent of different reducing agents at $1000^{\circ}C$. The sharp peaks of ZnO confirm the crystallinity in nature and also confirms that synthesized nanopowder was free from impurities.

Conclusions: The results obtained show that sol-gel method can produce good quality ZnO nanoparticles by using any reducing agents.

KEYWORDS: Nanoparticles, Precursor, Reducing agents, XRD, Sol-Gel

1. Introduction

Nanotechnology is the science that deals with matter at the scale of 1 billionth of a meter and is also the study of manipulating matter at the atomic and molecular scale. A nanoparticle is the most fundamental component in the fabrication of a nanostructure and is far smaller than the world of everyday objects that are described by Newton's laws of motion. Nanoparticles are the particles in the range from 1nm-100nm [1] - [2]. These nanoparticles are useful in other fields like electronics, biological, medical etc as they show different properties with different materials [3]. Till date quantum dots, Integrated circuits are the best examples of nanoparticles use. Different methods like physical, chemical, biological are used for construction of nano-material [4] - [7]. Among these, sol-gel method is used because of low cost, easy availability of tools and many more [8] - [12]. Much research showed different

results and properties of nanoparticles which is useful in different applications and motivating others to invent such nanoparticles which could be a breakthrough [13]- [18]. Among various nanoparticles, ZnO nanoparticles are promising candidates for various applications, such as nanogenerators [19], gas sensors [20], biosensors [21], solar cells [22], varistors [23], and photodetectors [24] and photocatalysts [25]. We report the synthesis of ZnO nanoparticles using different reducing agents in chemical method and the characterization of ZnO nanoparticles using X-ray diffraction.

2. Method & Materials

2.1. Synthesis

Zinc oxide nanostructure was synthesized by using sol-gel method [8] - [12]. To prepare a sol, precursor 0.2M of zinc acetate dihydrate and reducing agent (0.2 M of sodium hydroxide or 0.2 M Ammonia Solution

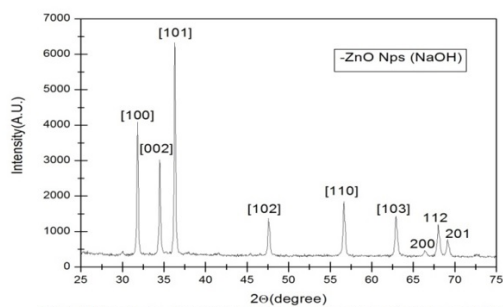


Fig.1 XRD pattern of ZnO powder prepared using reducing agent NaOH

Figure 1: XRD pattern of ZnO powder prepared using reducing agent NaOH.

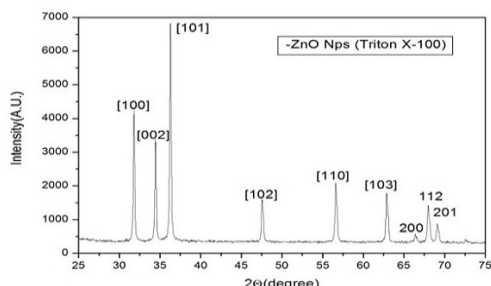


Fig.2 XRD pattern of ZnO powder prepared using reducing agent Triton X-100

Figure 2: XRD pattern of ZnO powder prepared using reducing agent Triton X-100.

(NH_3OH) or 5ml Triton X-100 [13]) were weighted using a weighing balance. Then 50ml of distilled water was measured by measuring cylinder. After that, 0.2 M of zinc acetate dehydrate was dissolved in 50ml of distilled water and reducing agent was dissolved in 50ml of distilled water. The solution was stirred with constant stirring for about 15 minutes each. After well mixed, reducing agent was poured to the solution containing zinc acetate with constant stirring by magnetic stirrer for about 15 minutes. Then burette was filled with 100ml of PVA (Poly Vinyl Alcohol) and titrate drop wise to the solution containing both reducing agent and zinc acetate [14]. After the reaction, white precipitate was formed. Dried precipitates was calcined at about $1000^{\circ}C$ for 24 hours in muffle furnace and sample was prepared.

3. Results and Discussion

3.1. Based on $ZnO + NaOH$

Based on the experimental work that has been done, there are series of chemical reaction that takes place. The complete reactions is in between transformation

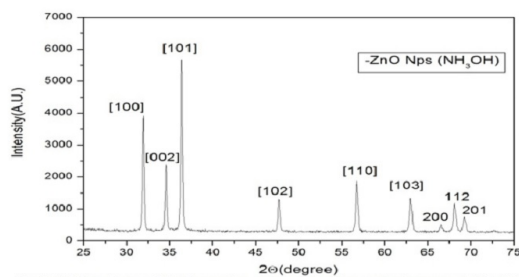
Fig.3 XRD pattern of ZnO powder prepared using reducing agent NH_3OH

Figure 3: XRD pattern of ZnO powder prepared using reducing agent NH_3OH .

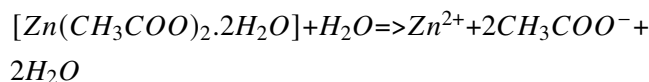
Table 1: Calculation of average size (D) of ZnO nanoparticles.

Angle ($2\theta^{\circ}$)	(θ°)	β° [FWHM]	D(nm)
36.31	18.155	0.2155	38.36

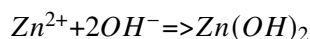
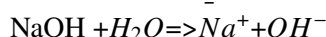
of zinc acetate dehydrate into zinc oxide nanoparticles. The addition of sodium hydroxide leading to the formation of zinc hydroxide intermediates in the form of Sol. PVA as a stabilizing agent ensure uniform dispersion of nanoparticles. After Gel formation and calcination, the resulting product comprises stable and dispersed zinc oxide nanoparticles [12].

3.2. Following reaction take place:

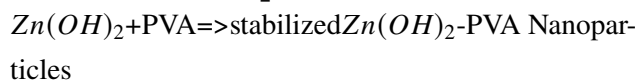
Precursor Dissolution:



Formation of Zinc Hydroxide Sol:



Polyvinyl Alcohol Addition: Gel Formation



Calcination:



These steps convert zinc hydroxide to zinc oxide nanoparticles at higher temperature. XRD analysis was carried using X-ray diffractometer with $Cu - K_{\alpha}$ radiation in the range $10 - 80^{\circ}$. Fig.1 Illustrate XRD spectrum of ZnO nanoparticles, synthesized using zinc acetate as precursor and $NaOH$ as reducing agent. The sharp peaks of ZnO indicate the crystallinity in nature and also confirm the synthesized nanopowder was free

Table 2: Calculation of average size (D) of ZnO nanoparticles.

Angle ($2\theta^0$)	(θ^0)	β^0 [FWHM]	D(nm)
36.28	18.14	0.2264	36.51

from impurities. The ZnO nanoparticles diameter was calculated for more intense peak corresponding to (101) plane using Debye-Scherrer formula [26], in Table.1.

$D = \frac{0.89\lambda}{\beta \cos\theta}$; Where λ is wave length of X-Ray (1.541\AA), β is FWHM (full width at half maximum) in radians, θ is the diffraction angle and D is particle diameter size.

3.3. Based on ZnO+Triton X-100

Following reaction take place:

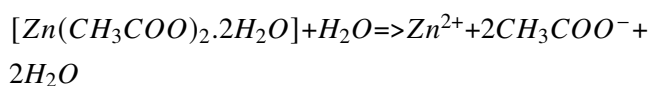


Fig.2 Illustrate XRD spectrum of ZnO nanoparticles, synthesized using zinc acetate as precursor and Triton X – 100 as reducing agent. Here Triton X – 100 used as reducing agent leads to prevent agglomeration of nanoparticles and control their growth. The sharp peaks of ZnO indicate the crystallinity in nature and also confirm the synthesized nanopowder was free from impurities. The ZnO nanoparticles diameter was calculated for more intense peak corresponding to (101) plane using Debye-Scherrer formula [26] include in Table.2

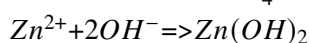
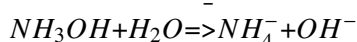
3.4. Based on ZnO + NH₃OH:

Following reaction take place:

Precursor Dissolution:



Formation of Zinc Hydroxide Sol:



Polyvinyl Alcohol Addition: Gel Formation



Calcination:

Table 3: Calculation of average size (D) of ZnO nanoparticles.

Angle ($2\theta^0$)	(θ^0)	β^0 [FWHM]	D(nm)
36.4	18.2	0.24153	34.26

Stabilized Zn(OH)₂- PVA Nanoparticles => ZnO+ Byproducts

Fig.3 Illustrate XRD spectrum of ZnO nanoparticles, synthesized using zinc acetate as precursor and as NH₃OH as reducing agent. The sharp peaks of ZnO confirm the crystallinity in nature and also confirm that synthesized nanopowder was free from impurities. The ZnO nanoparticles diameter was calculated for more intense peak corresponding to (101) plane using Debye- Scherrer formula [26] included in Table.3

4. Conclusion

Using different reducing agents, we have synthesized fine powder of ZnO nanoparticles. Also, XRD data produce the well-defined peaks of ZnO nanoparticles which confirm the crystallinity of nanoparticles. We have found that the average size of ZnO nanoparticles is independent of reducing agents at 1000^oC temperature. So, we may use any of the reducing agent to produce good quality of ZnO nanoparticles.

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Authorship contribution

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Conflict of Interest

Author declare there is no conflict of interest. The manuscript's contents have been reviewed and approved by author, and there are no competing financial interests to disclose.

Declaration

It is an original data and has neither been sent elsewhere nor published anywhere.

Similarity Index

I hereby confirm that there is no similarity index in abstract and conclusion while overall is less than 10% where individual source contribution is 2% or less than it.

References

- [1] C. N. R. Rao, A. Müller, A. K. Cheetham (eds), WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim, ISBN 3-527-30686-2, p.741.
- [2] A. K. Bandyopadhyay, New Age International, New Delhi (2008).
- [3] S. A. Akhoun, S. Rubab, M. A. Shah, Int Nano Lett 5: 9–13.(2015).
- [4] G. Amin, PhD thesis, Linköping University, Sweden (2012).
- [5] L. Changsong, Li Zhiwen, Z. Qifeng, material science, 22, 4, p. 603, (2007).
- [6] G. Kenanakis, D. Vernardou, E. Koudoumas, N. Katsarakis, Journal of Crystal Growth, 311, p.4799, (2009).
- [7] M. Gupta, et al., Bull. Mater. Sci., Vol. 32, 1, p.23, (2009).
- [8] H. Zhang, et al., Nanotechnology, 14, 423–426, (2003).
- [9] S. Fujihara, A. Suzuki, & T. Kimura, J. Appl. Phys., 94, p.2411, (2003) .
- [10] M. J. Alam & D. C. Cameron , J. Vac. Sci. Technol. A., 19, p.1642, (2001).
- [11] M. Toyoda, J. Watanabe & T. Matsumiya, J. Sol-Gel Sci. Technol., 1/2, 93, (1999).
- [12] D. A. Osman, M. A. Mustafa, Publishes by American Institute of Science, Vol. 1, N0. 4, p. 248, (2015)
- [13] T.V. Kolekar, H.M. Yadav, S.S. Bandgar and P.Y. Deshmukh, Indian Streams Research journal, ISSSN 2230 – 7850, Vol.1, Issue-1, (2011).
- [14] A. R. Bari, M. D. Shinde, V. Deo & L. A. Patil, Indian Journal of Pure & Applied Physics Vol. 47, p. 24, (2009).
- [15] J.N. Hasnidawani, H.N. Azlina, H.Norita, N.N. Bonnia, S. Ratim and E.S. Ali, Procedia Chemistry 19, p.211, (2016).
- [16] D. Bokov, A. T. Jalil, S. Chutradit, (2021).
- [17] O.W. Prez-Lopes, A.C. Faria, N.R.Marcilio, J.M.C. Bueno, Materials Reserch Bulletin, (2005).
- [18] L. Koudelka, J. Horak, P. Jariabka, Journal Materials Science, Vol. 29, No.6, p.1497, (2004).
- [19] P. X. Gao, Y. Ding, W. Mai, W. L. Hughes, C. Lao, and Z. L. Wang, Science, Vol. 309, No. 5741, p.1700, (2005).
- [20] X. L. Cheng, H. Zhao, L. H. Huo, S. Gao, and J. G. Zhao, Sensors and Actuators B, Vol. 102, No. 2, p. 248, (2004).
- [21] E. Topoglidis, A. E. G. Cass, B. O'Regan, and J. R. Durrant, Journal of Electroanalytical Chemistry, Vol. 517, No. 1-2, p. 20, (2001).
- [22] Y. Hames, Z. Alpaslan, A. Ko"semen, S. E. San, and Y. Yerli, Solar Energy, Vol. 84, No. 3, p. 426, (2010).
- [23] W. Jun, X. Changsheng, B. Zikui, Z. Bailin, H. Kaijin, and W. Run, Materials Science and Engineering B, Vol. 95, No. 2, p. 157, (2002).
- [24] P. Sharma, K. Sreenivas, and K. V. Rao, Journal of Applied Physics, Vol. 93, No. 7, p. 3963, (2003).
- [25] P. V. Kamat, R. Huehn, and R. Nicolaescu, Journal of Physical Chemistry B, Vol.106, No. 4, p. 788, (2002).
- [26] T. M. Al-Saadi, N. A. Bakr, N. A. Hameed, International Journal of Engineering and Technical Research, 2, 4 (2014).

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