



Synthesis of Transparent and Fluorescent ZnO Nanostructure for DSSC Application



Taehoon Lim^{1,2}, Eric Pichinte^{1,3}, Alfredo A. Martinez-Morales^{1*}

¹ Southern California Research Initiative for Solar Energy
College of Engineering Center for Environmental Research and Technology

² Department of Materials Science & Engineering
University of California, Riverside, California 92521

³ Engineering Department
Los Angeles Trade Technical College, Los Angeles, California 90015
(alfmart@ece.ucr.edu)



Abstract

The demand for clean energy has increased significantly in the past decade, and by a series of legislative measurements and public opinion, the development of renewable energy has been put at the forefront of power generation. In response, solar cells have been researched extensively because solar energy is one of the most promising clean and renewable energy sources. Dye-sensitized solar cells (DSSCs) are a certain type of next-generation solar cells, with wide applications, including building integrated photovoltaic (BIPV) technology and flexible solar cells.

In this poster, we demonstrate the synthesis of 1-dimensional semiconducting zinc oxide (ZnO) nanostructures for their future application as a photoelectrode in DSSCs. By adjusting synthesis conditions, ZnO with diverse dimensions have been successfully produced. Fluorescence emission characteristics have been found on certain dimensioned ZnO nanostructures. The fluorescence could be used as a secondary light source, with the potential to improve the photovoltaic characteristics of DSSCs.

Introduction

Dye-Sensitized Solar Cells

- Low cost and simple process
- High efficiency
- Aesthetically pleasant
- Suitable for BIPV
- TCO | TiO₂ : dye | electrolyte | TCO structure

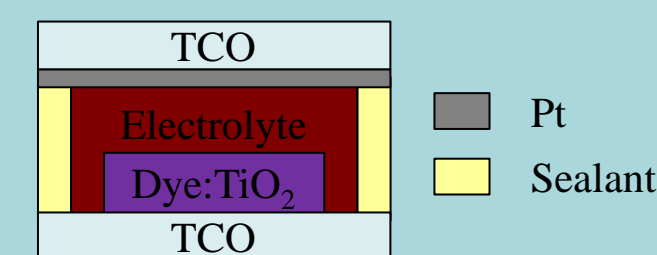


Figure 1. Structure of DSSCs

Zinc Oxide (ZnO)

- Wide-bandgap semiconductor
- Transparent, high electron mobility
- Hexagonal wurtzite
- Piezoelectricity, pyroelectricity characteristics

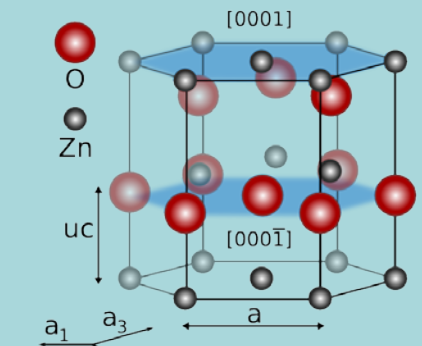


Figure 2. Crystal structure of wurtzite ZnO

Chemical vapor deposition (CVD)

- High-purity, high-performance solid product
- Continuous process for hybrid system
- Flexible conditioning
- Cost-effective
- Good step coverage

Fluorescence

- Electron excited by absorbing light
- Excited electron relaxes with light emission
- Longer wavelength emission than excitation

Experimental

CVD synthesis system

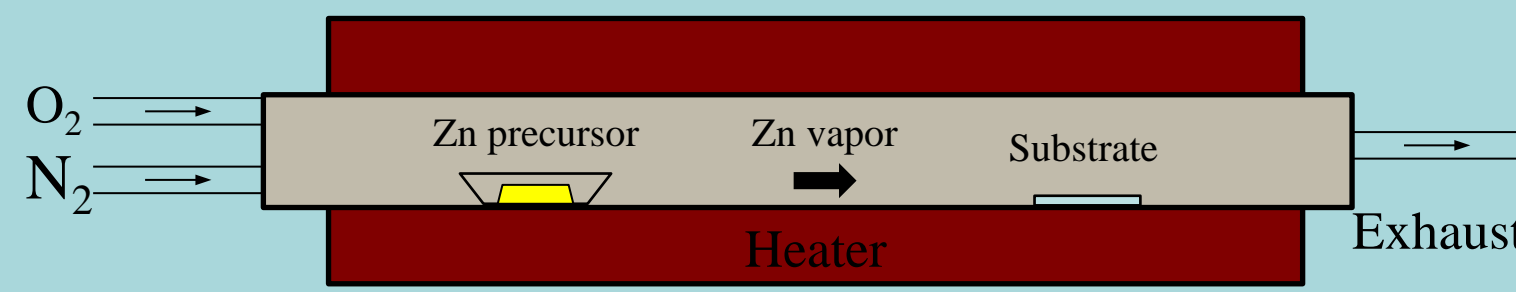


Figure 3. Schematic drawing of CVD synthesis system

Table 1. List of controllable parameters and range of CVD synthesis

Controllable parameters	Conditions range
Precursor temperature	400 - 600°C
Ramping/Reaction time	10 - 60 min / 30 - 120 min
Pressure	Atmospheric pressure ~ 10 ⁻³ torr
Carrier gas	Nitrogen (N ₂)
Oxidizer	Oxygen (O ₂)
Flow rate	N ₂ : 50 - 200 SCCM / O ₂ : 0 - 10 SCCM
Precursor-Substrate distance	0 - 2 cm
Substrate orientation	Parallel to gas flow

Experimental condition

Table 2. List of synthesis condition and results in terms of transparency and fluorescence emission

Sample #	Temperature (°C)	Position (inch)	N ₂ rate (SCCM)	O ₂ rate (SCCM)	Reaction time (min)	Transparency	Fluorescence
1	550	+0.5	90	10	30/30	NO	NO
2	550	+0.5	92	8	30/30	NO	NO
3	550	+0.5	94	6	30/30	NO	NO
4	550	+0.5	96	4	30/30	NO	YES
5	550	+0.5	98	2	30/30	NO	YES
6	550	+0.5	99	1	30/30	Slightly	YES
7	550	+0.5	100	0.5	30/30	YES	YES
8	550	+0.5	100	0.3	30/30	YES	NO

Characterization

- SEM for physical structure and morphology
- XRD for crystal structure and crystallinity
- Fluorescence emission under 365 nm UV lamp

Results and discussion

Crystallinity analysis by XRD

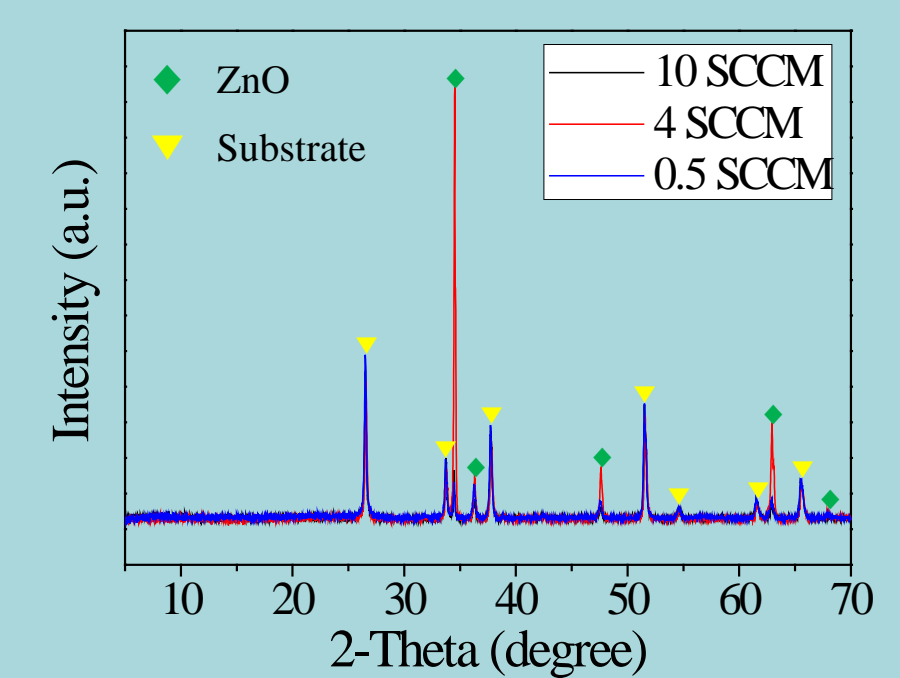


Figure 4. XRD results of synthesized ZnO on FTO substrate

The XRD results show two different sets of peaks, one from the synthesized ZnO material and one from the substrate. The peaks marked by a yellow triangle are from the substrate (F:SnO₂). The peaks marked by a green square are from the target material, ZnO. The XRD results confirms that the synthesized material is high crystalline wurtzite ZnO.

Results and discussion (cont.)

Morphology analysis by SEM

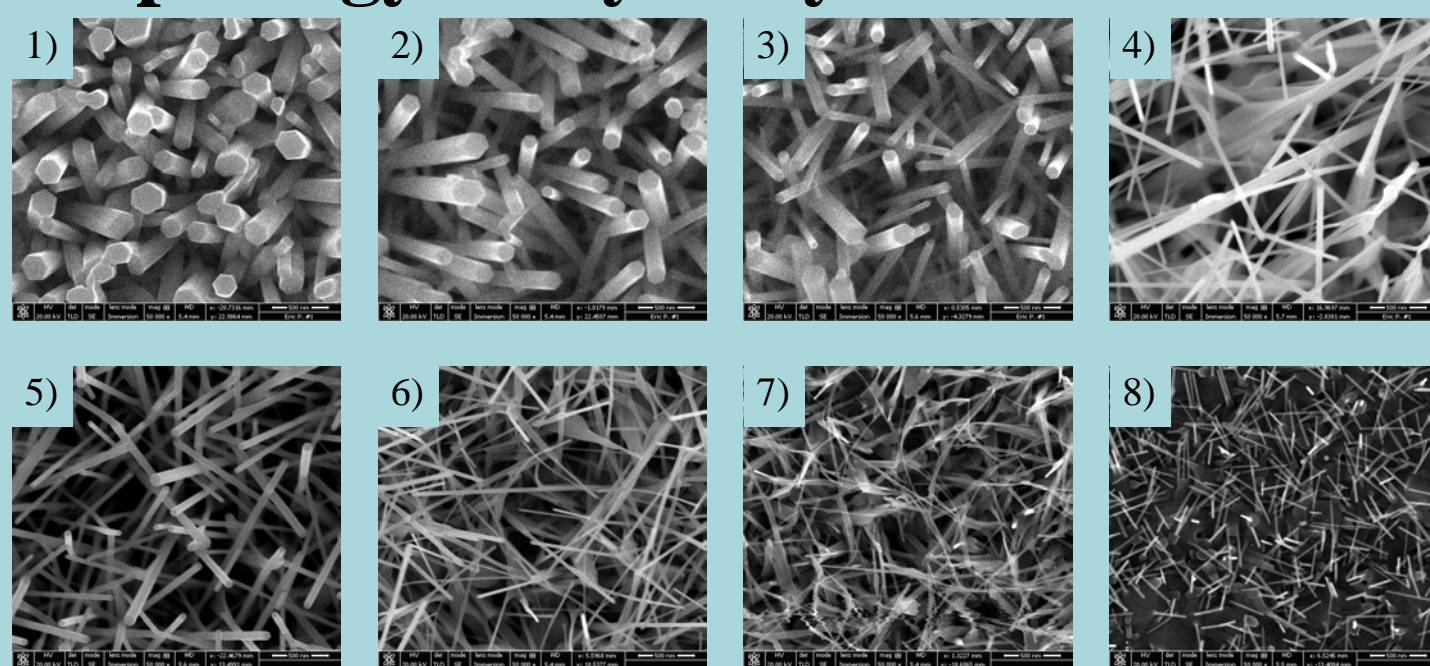


Figure 5. SEM images of ZnO synthesized under different oxygen partial pressure

The morphology and dimensions of ZnO nanostructures synthesized under different partial pressure ratios of N₂ and O₂ are observed by scanning electron microscope (SEM). As shown above, the partial pressure of oxygen plays a pivotal role in the shape and size of the synthesized product. Furthermore, when the oxygen ratio was too high, ZnO nanostructures were not grown on the surface, but could only grow near the edge of the substrate.

Transparency and fluorescence

Table 3. Transparency and fluorescence emission of synthesized ZnO under different oxygen partial pressure

Sample	1)	2)	3)	4)	5)	6)	7)	8)
Transparency								
Fluorescence								

In high oxygen conditions (samples 1-3), the ZnO layer was not evenly formed through the entire substrate surface, but rather formed on the area closest to the precursor. In medium oxygen conditions (samples 4-5), the ZnO layer was formed on the entire surface and the fluorescence emission was strong but the ZnO layers were not transparent. However, in low oxygen conditions (samples 6-7), the ZnO layer became more transparent while they still had fluorescence emission. Very thin ZnO layer was formed in extremely low oxygen condition (sample 8). ZnO synthesized at 0.5 SCCM oxygen rate (sample 7) is the most suitable for DSSC applications because it has both characteristics, transparency and fluorescence emission.

Summary

In this research, we have successfully synthesized 1-dimensional ZnO nanostructures exhibiting fluorescence emission excited from 365 nm UV light. By using the emission as a secondary light source, this ZnO nanostructures have the potential to improve photovoltaic characteristics of DSSCs. High crystallinity wurtzite crystal structure is confirmed by XRD characterization. Morphology and dimensional characteristics are observed by SEM. In conclusion, the synthesized ZnO nanostructures are suitable for a wide range of applications including photovoltaic devices, UV sensors, phototransistors, or photocatalysts.

Acknowledgements

The SEM characterization was conducted at the Central Facility for Advanced Microscopy and Microanalysis (CFAMM). This research was partially funded by the American Public Power Association (APPA) Demonstration of Energy & Efficiency Developments (DEED) Program, the UC MEXUS-CONACYT Program, the 2014 MacREU Program at UCR, and the University of California Advanced Solar Technologies Institute (UC Solar).