Portland State University

PDXScholar

Dissertations and Theses

Dissertations and Theses

Winter 2-12-2016

One-Dimensional Nanostructure and Sensing Applications: Tin Dioxide Nanowires and Carbon Nanotubes

Hoang Anh Tran
Portland State University

Follow this and additional works at: https://pdxscholar.library.pdx.edu/open_access_etds

Part of the [Chemistry Commons](#), [Materials Science and Engineering Commons](#), and the [Nanoscience and Nanotechnology Commons](#)

Let us know how access to this document benefits you.

Recommended Citation

Tran, Hoang Anh, "One-Dimensional Nanostructure and Sensing Applications: Tin Dioxide Nanowires and Carbon Nanotubes" (2016). *Dissertations and Theses*. Paper 2689.
<https://doi.org/10.15760/etd.2685>

This Dissertation is brought to you for free and open access. It has been accepted for inclusion in Dissertations and Theses by an authorized administrator of PDXScholar. Please contact us if we can make this document more accessible: pdxscholar@pdx.edu.

One-Dimensional Nanostructure and Sensing Applications:

Tin Dioxide Nanowires and Carbon Nanotubes

by

Hoang Anh Tran

A dissertation submitted in partial fulfillment of the
requirements for the degree of

Doctor of Philosophy
in
Chemistry

Dissertation Committee:
Shankar Rananavare, Chair
Carl Wamser
Albert Benight
Raj Solanki
Erik Sanchez
James Blackwell

Portland State University
2016

Abstract

The key challenge for a nanomaterial based sensor is how to synthesize in bulk quantity and fabricate an actual device with insightful understanding of operational mechanisms during performance. I report here effective, controllable methods that exploit the concepts of the “green approach” to synthesize two different one-dimensional nanostructures, including tin oxide nanowires and carbon nanotubes. The syntheses are followed by product characterization and sensing device fabrications as well as sensor performance understanding at the molecular level. Sensor-analyte response and recovery kinetics are also presented.

The first part of the thesis describes bulk-scale synthesis and characterization of tin oxide nanowires by the molten salt synthetic method and the nanowire doping with antimony (n-types) and lithium. The work builds on the success of using n-doped SnO_2 nanoparticles to selectively detect chlorine gas at room temperature. Replacing n-doped nanoparticles with n-doped nanowires reduces the number of inter-particle electron hops between sensing electrodes. The nanowire based sensors show unprecedented 5 ppb detectability of corrosive Cl_2 gas concentration in air. At the higher range, 10 ppm of Cl_2 gas leads to a 250 fold increase in the device resistance. During sensor recovery, FT-IR studies show that dichlorine monoxide (Cl_2O) and chlorine dioxide (ClO_2) are the desorbing species. Long term stability of devices is affected by lattice oxygen vacancies replaced by chlorine atoms.

Bulk-scale synthesis of multiwall carbon nanotube (MWCNTs) was achieved by a novel inexpensive synthetic method. The green chemistry method uses the non-toxic

and easy to handle solid carbon source naphthalene. The synthesis is carried out by simply heating naphthalene and organometallic precursors as catalysts in a sealed glass tube. Synthesis at 610° C leads to MWCNTs of 50 nm diameter and lengths exceeding well over microns. MWCNT doping is attempted with nitrogen (n-type) and boron (p-type) precursors. Palladium nanoparticles decorated on as-synthesized MWCNTs are employed for specific detection of explosive hydrogen gas with concentrations far below the explosive concentration limits. During performance, the sensor exhibits abnormal response behaviors at hydrogen gas concentrations higher than 1%. A model of charge carrier inversion, brought about by reduction of MWCNT by hydrogen molecules dissociated by Pd nanoparticles is proposed.

To my beloved Dad, I miss you here on Earth.

This dissertation is dedicated to you.

Acknowledgements

It gives me a pleasure to gratefully and sincerely thank my advisor, Professor S.B Rananavare. I have been extremely fortunate to have an exceptional advisor who gave me invaluable guidance, constant inspiration and continued support during my academic career at Portland State University. All that has been achieved in this thesis was made possible by him.

I am also grateful to Professors Carl Wamser, Albert Benight, Raj Solanki, and Erik Sanchez for their time and invaluable inputs in my dissertation committee. I also would like to express my gratitude to Dr. James Blackwell for not only serving in my dissertation committee but also giving me a chance to work with different collaborators and sponsoring me through Intel's funding.

I am indebted to Professor Andrea Goforth for photoluminescence analysis, Professor Andres La Rosa for the usage of the Raman system. Especially, I am thankful to Dr. Daner Abdulla for his intellectual help on Raman spectroscopy as well as uncountable advice on my thesis. Many thanks go to Dr. Joo Chan and Dr. Allen Chapadraza for the treasure of knowledge I had inherited when I was still an undergraduate student. I also would like to thank Professor Rananavare's research group Ryan Lerud, Srikar Rao, Kayode Morakinyo, Sayan just to mention a few, for their support and help during my PhD studies.

Most importantly, none of this would have been possible without the support, love and patience of my family. I would like to take this opportunity to thank my uncle Le's

family, who has given me all the supports and foundation in this country. Without them I would not be the person that I am today.

To my wife Leyna, I know you suffer from my imperfections and my shortcomings largely during the challenging moments. Your sweetness, your kindness, the beauty of your heart and your thoughtfulness never cease to amaze me. You are the motivation of my life.

To my mom, I know I owe you everything that I am. I could not ask for a better mom than you. Thank-you mom for your patience, your wisdom and most important, your love.

Table of Contents

Abstract	i
Dedication	iii
Acknowledgements	iv
List of Tables	x
List of Figures	xi
CHAPTER 1. SENSOR BASED ON ONE DIMENSIONAL NANOMATERIALS.....	1
1.1 General Introduction Nanomaterials.....	1
1.2 Electronic Properties of Bulk and Nanomaterials.....	2
1.3 One Dimensional Nanomaterials for Sensing Applications	6
1.3.1 Chlorine Sensor.....	7
1.3.2 Hydrogen Sensor.....	9
CHAPTER 2. INSTRUMENTATION, DEVICE CHARACTERIZATION AND NANOSTRUCTURE SYNTHESIS	13
2.1 Synthesis and Sensor Fabrication	13
2.1.1 Sample Preparation for SnO ₂ NWs.....	13
2.1.2 Sample Preparation for CNTs.....	15
2.2 Electron Microscopy.....	21
2.3 Physical Vapor Deposition (PVD).....	22
2.4 Infrared Spectroscopy (FTIR).....	22
2.5 UV-Visible Spectroscopy (UV-Vis).....	22
2.6 Photoluminescence Spectroscopy (PL).....	22

2.7 Raman Spectroscopy.....	23
2.8 Sensor Testing System.....	25
CHAPTER 3. TIN OXIDE NANOWIRES (SnO ₂ NWs)	28
3.1 Background and Significance	28
3.1.1 Crystal Structure	28
3.1.2 Stoichiometry, Defect Structure and Conductivity	29
3.1.3 Raman Active Vibrational Modes of SnO ₂ Rutile Structure	31
3.1.4 Growth of SnO ₂ NWs	33
3.1.5 SnO ₂ NWs as Sensing Material	35
3.2 Results and Discussion	39
3.2.1 Electron Microscopy	39
3.2.2 Photoluminescence Characterization	44
3.2.3 UV-Vis Spectroscopy	47
3.2.4 Raman Spectroscopy	51
3.2.5 Electrical Properties of Thick Film Using n-doped SnO ₂ NWs	53
3.3 Sensing Application: Chlorine Resistive Gas Sensor Based on n-doped SnO ₂ NWs	55
3.3.1 Sensor Response, Response Time and Recovery Time	55
3.3.2 Sensor Sensitivity	58
3.3.3 Sensor Stability and Understanding Sensing Mechanism	63
3.4 Conclusion	75
CHAPTER 4. CARBON NANOTUBES (CNTs)	77
4.1 Background and Significance	77

4.1.1 Electronic Properties	77
4.1.2 Growth of CNTs	79
4.1.3 Raman Spectroscopy of CNTs.....	82
4.1.4 Doping CNTs	85
4.1.5 CNTs Based Gas Sensor	87
4.2 Results and Discussion	90
4.2.1 Electron Microscopy	90
4.2.2 Raman Spectroscopy.....	98
4.2.3 Palladium Nanoparticles Attached MWCNTs for Hydrogen Sensor	103
4.3 Sensing Application: Hydrogen Gas Sensor Based on Pd NPs Attached MWCNTs	107
4.3.1 Sensing Performance	107
4.3.2 Understanding Sensing Mechanism.....	113
4.4 Conclusion	125
CHAPTER 5. FUTURE WORK	126
5.1 Future Work of SnO ₂ NWs	126
5.2 Future Work of CNTs	127
References	130
Appendices	
Appendix A. Chemical Reagents.....	150
Appendix B. Characterization of SnO ₂ NWs at Different Synthetic Conditions	151
Appendix C. Synthesis and Characterization of Heterojunctions of Silicon Nanowires (SiNWs) and Carbon Nanotubes (CNTs)	158

Appendix D. Negative Differential Resistance Observed from Pd NPs –CNTs	161
---	-----

List of Tables

Table 1: A brief summary of nanomaterial based chlorine sensors.....	8
Table 2: A brief summary of metal oxide based hydrogen sensors.....	10
Table 3: Phonon symmetry of SnO ₂ rutile structure.....	32
Table 4: The effect of synthetic conditions on SnO ₂ NWs morphology	43
Table 5: Optical bandgap determined from UV absorption edge and PL.....	50
Table 6: Raman data analysis of different SnO ₂ samples	52
Table 7: Kinetic analysis of response curves during the exposure process.	70
Table 8: Kinetic analysis of response curves during the recovery process.....	72
Table 9: IR observation of dichlorine monoxide (Cl ₂ O) and chlorine dioxide (ClO ₂).....	74
Table 10: The non-linear least squares analysis of D and G bands from Raman spectroscopy of as-synthesized MWCNTs	99
Table 11: G' bands from Raman spectroscopy of as-synthesized MWCNTs	101
Table 12: Summary kinetic constants from fitted response curves	115
Table 13: Summary of kinetic constants from fitted recovery curves	118

List of Figures

Figure 1: Illustration of band diagram for an intrinsic semiconductor.	4
Figure 2: The density of states $D(E)$ (left) for a 3D semiconductor and Fermi-Dirac distribution $f(E)$ (right) at different temperatures for intrinsic semiconductor.....	4
Figure 3: Schematic showing density of state of low dimensional nanostructures.	5
Figure 4: Photographs of 1%, 5% and 10% Sb-doped SnO_2 NWs based sensors	15
Figure 5: An optical image of duo-chamber glass reaction vessel for the synthesis of MWCNTs.....	16
Figure 6: Schematic illustration of CNTs synthesis from three different synthetic methods.....	18
Figure 7: The photograph of the custom-built CNT synthesis system housed inside a tube furnace.....	18
Figure 8: Molecular structure of doping agents for CNTs.....	19
Figure 9: A photograph of fabricated CNT based sensor using method C	21
Figure 10: Raman transitional scheme.....	24
Figure 11: Schematic showing the home-built Raman system with major components. .	25
Figure 12: A schematic view of the gas test chamber	26
Figure 13: Schematic of gas diluting system (left) and the photograph of home-built sensor testing system (right).	27
Figure 14: Unit cell of the crystal structure of SnO_2	29
Figure 15: Illustration of ionic lattice of MO_{2-x}	31
Figure 16: Raman active vibrational modes of SnO_2 rutile structure.	33

Figure 17: A. Energy levels of oxidizing and reducing gases in relation to band edges of SnO ₂ . B. Location and position Sb reduction potential.	37
Figure 18: Schematic diagrams of porous sensing layers.	38
Figure 19: Optical images of undoped (left) and Sb doped NWs (right)	39
Figure 20: SEM images of undoped (left) and Sb doped NWs (right)	41
Figure 21: TEM images of n-doped SnO ₂ NWs with low (left) and high magnifications along with electron diffraction in the inset (right).	41
Figure 22: Optical image and SEM image of Li-doped SnO ₂ NWs	42
Figure 23: SEM image of Li-doped SnO ₂ NWs synthesized at 600°C and 700°C	43
Figure 24: A) HR-TEM of SnO _{2-x} coated Sn NP precursors and B) TEM of Li-doped SnO ₂ NWs	44
Figure 25: Photoluminescence of undoped SnO ₂ NWs and NPs.	45
Figure 26: Photoluminescence of undoped and n-doped SnO ₂ NWs	46
Figure 27: Photoluminescence of undoped, 5% n-doped and 5% Li-doped.	47
Figure 28: UV-Vis spectra of different SnO ₂ NW samples: undoped, 5% n-doped and 5% Li-doped.	48
Figure 29: Determination of optical bandgap from the UV absorption edge.	49
Figure 30: Raman spectrum of SnO ₂ NWs samples.	51
Figure 31: Illustration of curve fitting of 2% n-doped SnO ₂ NWs sample.	52
Figure 32: I-V curves of different dopant concentrations of n-doped NW based sensors	54
Figure 33: The sensor resistance (A) and recovery time (B) at six different Cl ₂ concentrations from 5 ppb to 10ppm	56

Figure 34: Response and recovery time (minute) as a function of temperature (°K)	56
Figure 35: Response as a function of chlorine concentration at room temperature.....	59
Figure 36: Response of the sensor as a function of $\gamma^{1/2}$	61
Figure 37: Response of the sensor as a function of doping concentration.....	62
Figure 38: Schematic diagram of the porous sensing layer composed from nanowires...	65
Figure 39: Response of the sensor from 4 continuous cycles of exposure and recovery. 65	
Figure 40: Illustration of mechanism (IV).	66
Figure 41: Sensor resistance baseline as a function of cycle time	67
Figure 42: Sensor response curves after storing in high Cl ₂ concentration	68
Figure 43: An example of fitting sensor response resistance as a function of time.....	70
Figure 44: An example of fitting curve.....	72
Figure 45: Infrared spectrum composed from blank sample (A), mixture of chlorine and air (B), reaction products during sensor recovery process (C)	73
Figure 46: (n,m) coordinate system for a graphene sheet.	77
Figure 47: Schematic diagram of a DC-arc discharge system for CNT synthesis.....	80
Figure 48: Schematic diagram of a laser ablation system to synthesize CNTs	81
Figure 49: Schematic diagram of a CVD system to synthesize CNTs	82
Figure 50: An example of a typical Raman of SWCNTs	83
Figure 51: The RBM mode of a SWNT.....	83
Figure 52: Vibrational modes of G ⁺ (A) and G ⁻ (B) bands.....	84
Figure 53: Doping CNTs through many different pathways	86
Figure 54: Various structural defects on CNT	89

Figure 55: An example of NH_3 molecule adsorption on defective CNT	89
Figure 56: Formation of Ni nanoparticle catalysts after annealing 5nm and 20nm thick Ni thin films at 680°C	90
Figure 57: Method A for the synthesis of CNTs at 590°C , 600°C , 610°C and 620°C	91
Figure 58: HRTEM of CNT synthesized at 600°C in 2hrs (top) and EDX of catalyst (bottom) and STEM of a CNT.	94
Figure 59: Method B for the synthesis of CNTs in uncontrolled manner at 590°C , 600°C , 610°C and 620°C	95
Figure 60: Optical images of method C for the synthesis of CNTs ferrocene (top) and cobaltocene (bottom)	96
Figure 61: SEM images of CNTs by method C.	96
Figure 62: TEM images of CNTs by method C.....	97
Figure 63: Raman spectroscopy of as-synthesized MWCNTs	98
Figure 64: An example of fitting using the undoped CNT sample.....	99
Figure 65: Raman Spectroscopy of as-synthesized MWCNTs.....	101
Figure 66: The band intensity ratios of D/G and G'/G of p-doped CNTs (left) and n-doped CNTs (right).	103
Figure 67: SEM images of commercial CNTs and palladium nanoparticle decorated CNTs.	104
Figure 68: Illustrative concept for the Decoration of CNTs with Pd Nanoparticles	105
Figure 69: SEM of thin film of CNTs deposited on the wall of glass tube before and after the attachment of Pd nanoparticles.	106

Figure 70: Low and high magnification TEM images of as-synthesized CNTs decorated with Pd nanoparticles	106
Figure 71: Response of testing sensor upon the exposure of 4% H ₂ with the inset showing the initial stage of H ₂ exposure with the flow rate of 1L/minute.....	107
Figure 72: Response of Pd nanoparticles attached as-synthesized CNTs sensor to different H ₂ concentrations from 10ppm to 4% (40,000 ppm).	109
Figure 73: An example of fitted curve to 10ppm H ₂	110
Figure 74: Slope of the response curve as a function of H ₂ concentrations.	111
Figure 75: Abnormal behaviors of sensor response and recovery at high H ₂ concentrations.	112
Figure 76: Electrochemical energy level diagram of hydrogen, palladium and oxygen in relation to the Fermi level of CNTs	114
Figure 77: Illustration of the response mechanism of Pd Nps decorated CNTs upon the exposure with H ₂	114
Figure 78: Sensor response at 10ppm H ₂ concentration	115
Figure 79: H ₂ molecular dissociation constant (left) and electrons transferred to nanotube constant (right) change as a function of H ₂ concentration.....	116
Figure 80: Sensor recovery after 10ppm H ₂ exposure	117
Figure 81: a) 5 stages of charge carrier inversion process b) Time dependence of excess hole and electron concentration in CNT.	119
Figure 82: Mechanism of 5 stages of charge carrier inversion process (left) and electronic band diagram of sensing process (right)	121

Figure 83: Kinetic of charge carrier inversion and slow recovery using diluted air.....	123
Figure 84: Response of n-doped SnO ₂ /Pd (right) sensors to different hydrogen concentrations H ₂	127
Figure 85: The sketch illustrating experiment for future study of NDR-like behavior using two individuals Pd Np attached CNTs.....	128

CHAPTER 1. SENSOR BASED ON ONE DIMENSIONAL NANOMATERIALS

1.1 General Introduction Nanomaterials

Nanomaterials are cornerstone of nanoscience and nanotechnology. In the past few decades, a broad and interdisciplinary area of nanostructure research and development has been growing explosively due to their novel properties and their emerging new applications. However, nanomaterials have a long scientific history, which began after the first research report by Michael Faraday in 1857 with the introduction of colloidal gold nanoparticle (NP) synthesis [1]. One of the major industrial applications of nanomaterials emerged in the early 1940s in the United States of America and Germany with the replacement of ultrafine carbon black by precipitated and fumed silica nanoparticles for rubber reinforcements [2]. Another prime example is the investigation of nanostructural aspects of catalysts over the last 150 years [3]. Today, nanophase engineering has expanded in a rapidly growing number of structural and functional materials, both inorganic and organic, allowing for the manipulation of mechanical, catalytic, electric, magnetic, optical and electronic functions [3].

Nanomaterials are defined as the materials whose structures have at least one dimension less than 100 nm. Depending on how many dimensions are less than 100 nm, nanomaterials are classified into three different categories: zero (0D, nanoparticles), one (1D, nanowires /nanotubes), and two dimensional (2D, nanofilms) nanostructures [3]. The size and dimension of nanomaterials lead to some remarkable properties that may be significantly different from the properties of bulk materials. Some such peculiar properties have been well-known and studied, but there is surely a plethora more to be

discovered. Many physical properties of nanomaterials are related to the increased surface area and quantum confinement effects [4]. Compared to their bulk counterparts, nanomaterials have a much greater surface area to volume ratio, which can provide greater chemical reactivity and can provide better mechanical strength in composites. Quantum effects at the nanoscale determine the material's properties and characteristics, leading to novel optical (bandgap engineering e.g., CdSe, Si NPs), electrical and magnetic characteristics different from their conventional bulk forms. Currently, intense worldwide research has been focused on the synthesis, characterization, assembly and manipulation of nanostructural blocks for their device applications [5].

In this thesis, sensors based on one dimensional nanomaterials are developed that exploit their unusual electrical properties, especially conductivity, which is reviewed in the following section.

1.2 Electronic Properties of Bulk and Nanomaterials

A fundamental electronic property of materials is its conductivity which varies dramatically for metals, semiconductors and insulators. The conductivity σ_{tot} of a general semiconductor crystal can be described as the sum due to electronic (ionic σ_{ion} , electron σ_e and hole σ_p) carriers if the associated conduction processes are considered independent [6]:

$$\sigma_{tot} = \sigma_e + \sigma_p + \sum \sigma_{ion,i} \approx \sigma_e + \sigma_p = n \cdot \mu_e \cdot e + p \cdot \mu_p \cdot e \quad (\text{eq. 1.1})$$

Where n , p and e are electron concentration, hole concentration and elementary charge (1.6×10^{-19} coulombs), respectively. μ_i denotes the mobility of carrier i . In

chemical sensing carrier concentrations or mobilities of the carriers are modulated by interaction with analyte.

Bulk material:

The resistance, an experimentally measured quantity of a homogeneous bulk material, can be related to its bulk conductivity σ_b , and geometric features. Geometrical features that affect resistance are its length l and cross section area A . Thus for a bulk semiconductor:

$$R_b = \frac{l}{\sigma_b A} \text{ with } \sigma_b = \sigma_e + \sigma_p = n \cdot \mu_e \cdot e + p \cdot \mu_p \cdot e \quad (\text{eq. 1.2})$$

The charge carrier concentrations of electrons (n) and holes (p) for an intrinsic semiconductor can be calculated [6]:

$$n = \int_{E_c}^{\infty} D(E) f(E) dE \quad ; \quad p = \int_{-\infty}^{E_v} D(E) (1 - f(E)) dE \quad (\text{eq. 1.3})$$

With the density of states $D(E)$ and Fermi-Dirac distribution $f(E)$ are defined as [7]:

$$D(E) = \frac{1}{2\pi^2} \left(\frac{2m_e}{\hbar^2} \right)^{\frac{3}{2}} (E - E_c)^{\frac{1}{2}} ; f(E) = \frac{1}{1 + \exp\left(\frac{E - E_F}{kT}\right)} \quad (\text{eq. 1.4})$$

Where E_C , E_V and E_F are conduction band and valence band edges and Fermi energy level, respectively. k , m_e , h are Boltzmann constant, mass of carrier, and Planck's constant respectively. These band edges are depicted along with band gap E_g in Figure 1:

Figure 1: Illustration of band diagram for an intrinsic semiconductor.

The variation of density of state for a bulk 3D semiconductor and the Fermi functions for a bulk semiconductor are illustrated below in Figure 2:

Figure 2: The density of states $D(E)$ (left) for a 3D semiconductor and Fermi-Dirac distribution $f(E)$ (right) at different temperatures for intrinsic semiconductor [8].

Low-dimensional nanostructures: 0, 1, and 2 dimensions:

In cases of low-dimensional nanostructures, the conductivity is different from its counterpart in bulk 3D material. It can also be defined as equation (1.1), and the charge carrier concentrations electrons (n) and holes (p) can be calculated by equation (1.3). The Fermi-Dirac distribution function $f(E)$ stays the same, but the density of states $D(E)$ depends on the dimensionality.

A general method to calculate the density of states in energy and wavevector space (E versus k space) of these low-dimensional nanostructures follows equation (1.5) with the representative density of states for 0, 1 and 2 dimensional nanostructures are illustrated in Figure 3:

$$D(E) = 2 \frac{V}{(2\pi)^3} \frac{1}{\frac{dE_k}{dk}} \quad (\text{eq. 1.5})$$

Where V is the volume of a spherical shell in k space

In one-dimensional nanostructure the density of state is calculated as [8]:

$$D(E) = \frac{L}{2\pi} \left(\frac{\hbar}{2m^*} \right) \frac{1}{\sqrt{E}} \equiv \frac{L}{2\pi} \left(\frac{\hbar}{2m^*} \right) \frac{1}{\sqrt{E-E_c}} \quad (\text{eq. 1.6})$$

Where m^* and L are the effective mass of electron and the width barrier of 1D quantum well, respectively. E and E_c are the electron energy in and at the bottom of conduction band, respectively.

Figure 3: Schematic showing density of state of low dimensional nanostructures [9].

For one dimensional structures the $D(E)$ has singularities implying high conductivity at these points. The modification of conductivity of sensors based on the 1D system occurs through changes in carrier concentration brought about by electron transport between analyte and the nanostructure. In addition, geometric factors due to

creation of the depletion layer also affect sensor resistance. Therefore, 1D nanostructures are an ideal candidates for sensing material. This is the primary hypothesis behind the work developed in this thesis. Its validity is tested for SnO_x nanowires and multiwall carbon nanotubes based sensors for selective and sensitive detection of chlorine and hydrogen gases, respectively.

1.3 One Dimensional (1D) Nanomaterials for Sensing Applications

In the last fifteen years, the number of publications involving 1D nanostructures has exploded due to their unique electronic properties and also due to development of methods for their bulk-scale syntheses. With the lowest dimension for efficient electrical carrier transport, it is ideally utilized for the task of moving charges and photons, i.e., for applications involving nanoelectronics and nanophotonics. They have found wide ranging applications in devices such as transistors, diodes, circuits or as light emitters. In addition, 1D nanostructure, including tubular structures with the unique features such as low mass density, high porosity and extremely large surface area to volume ratios, can provide superior performance as catalysts, fuel cell electrodes, and chemical/biochemical sensors. Out of these, sensor devices based on 1D nanostructures have attracted much greater attention. A comprehensive review of current research activities of chemical sensors on nanotubes, nanorods, nanobelts and nanowires, focusing on experimental principles, design of sensing devices and sensing mechanisms has recently appeared [10]. For example, silicon nanowires based gas sensors have been fabricated for the detection of ammonia and water vapor at room temperature [11]. Sensors utilizing tin oxide nanobelts, operating at 400°C, can detect carbon monoxide as low as 0.5 ppm [12], . Zinc

oxide nanorods, grown by plasma-enhanced chemical vapor deposition (PECVD) on polycrystalline Al_2O_3 have been used as a chemical sensor for nitrogen dioxide gas detection [13]. Carbon nanotubes have been well-known as not only a chemical sensor for NO_2 and NH_3 [14], but also a biological sensor [15]. The fundamental sensing mechanism of these gas sensors involves changes in conductivity of nanostructures as result of gas adsorption and interaction between the underlying semiconductor and surface complexes (such as O^- , H^+ , OH^-), reactive chemical species. The adsorption and charge transport (i.e., change in the n or p the carrier concentrations) from the analyte brings the underlying semiconductor in either the depletion or accumulation region leading to changes in channel conductivity.

However, so far, the synthesis and integration for practical, inexpensive sensor devices based on 1 D nanostructures, in term of commercialization, has not received much attention. Integration requires a complete understanding of the device operational parameters, reasonable lifetime and sustained high sensitivity. Therefore, the work presented in this thesis explored ways in which different 1D nanostructures which are easily synthesized and employed for sensing applications for detection of chlorine and hydrogen as an oxidizing and a reducing gases, respectively.

1.3.1 Chlorine sensor

Recently, the usage of gas sensors has been on the rise due to their wide array of applications, ranging from safety equipment (explosion, leakage, fire, contamination of poisoning protection) to emission and air quality monitoring, quality assurance, process instrumentation, and measurement technology [16]. In principle, materials that change

their properties upon exposure to ambient gas can be used as gas sensing materials. Many metal oxides are suitable for gas resistive sensors. These sensors incorporate nanostructures in porous a thick film which provides a high surface to volume ratio enhancing the detection sensitivity. These device operate at elevated temperatures around 300°C [17] to enable rapid response time.

In order to detect oxidizing and highly corrosive gases or operate in harsh conditions, appropriate sensing materials must be selected. A prime example of one such gas is chlorine, which is useful in many industrial processes, but is deadly toxic gas at 5-10 ppm level. A few oxide based Cl₂ gas sensor have been developed and summarized in Table 1 below:

Table 1: A brief summary of nanomaterial based chlorine sensors. Adapted and modified from ref [18]

Sensor type	Detection limit (ppm)	Operation temperature (°C)
ZnO	300	400
WO ₃ /FeNbO ₄	2	230
CdIn ₂ O ₄	0.2	250
SnInO	3	300
CdSnO ₃	0.1	250
CuO-CdIn ₂ O ₄	400	250
In ₂ O ₃	0.5	250
Zn ₂ In ₂ O ₅ -MgIn ₂ O ₄	0.01	300
Mg-phthalocyanine	0.18	200
FEP/polyaniline	100	25
Cu-phthalocyanine	0.18	170
WO ₃	1	300
In ₂ O ₃ -Fe ₂ O ₃	0.2	250
NiFe ₂ O ₄	1000	250

However, the most commonly used in gas sensing materials is nanostructured SnO₂ due to its several key advantages over other metal oxides such as low cost of production and bulk-scale availability [19]. Moreover, SnO₂ is chemically stable and is

readily regenerated making it an ideal candidate for detection of such a highly corrosive gas. Additionally, we recently developed an Sb-doped SnO₂ NPs sensor, which operates even at room temperature and can detect Cl₂ gas at ppm level. As discussed earlier, in the research presented here replaces the n-doped SnO₂ NPs by n-doped SnO₂ NWs. The use of NWs, as opposed to NPs, reduces inter-particles electron hopping thereby leading to better sensor performance. This thesis focuses on the study of a highly sensitive, thick-film Cl₂ resistive sensor fabricated from Sb-doped SnO₂ NWs. A novel feature of this system is that it also works at room temperature providing good sensor response (50ppb to 10 ppm) and sensitivity. The work presented here explores the mechanisms of sensor stability and develops chemical models for the sensing mechanism.

1.3.2 Hydrogen sensor

Hydrogen is not only one of the fundamental building blocks of living things in nature, but also it is a key starting material in the chemical industry. Recently, hydrogen has emerged as an alternate energy source that can replace dwindling fossil fuel reserves. Additional advantages of its use are that it provides energy security, and it helps reduce global warming. As a colorless, odorless and tasteless flammable gas, hydrogen cannot be detected by human senses, even at concentrations at which it poses an explosion risk. Therefore, a rapid and accurate hydrogen gas sensor with long term stability and wide dynamic range of concentration is desirable.

Numerous approaches are currently being investigated to develop effective hydrogen sensors. These sensors employ thermoelectric, electrochemical, optical and acoustic methods of detection [20]. However, the most commercially available hydrogen

sensors are based on metal oxide materials such as tin oxide. Typically, these types of sensors operate at elevated temperatures to maintain proper performance [21]. Thus, these kinds of sensors still have poor energy efficiency and require complicated fabrication methods [21, 22].

Table 2: A brief summary of metal oxide based hydrogen sensors. Adapted and modified from refer [21]

Sensor type	Detection limit (ppm)	Operation temperature (°C)
SnO ₂ nanowires	10	300
SnO ₂ nanorod	500	230
SnO ₂ nanobelts	30,000	80
SnO ₂ film	10	300
Sn _{0.9} Ti _{0.1} O film	1000	450
Pt-Ag-SnO ₂ film	1000	250
Pd-ZnO nanorods	100	25
WO ₃ film	1000	110
Pt/Li-NiO film	30,000	100
Pt/NiO film	30,000	100
Pt/Ga ₂ O ₃	10,000	610

Recently, palladium has been reported as one of the most promising materials for the detection of H₂. Different Pd nanostructures have been proposed as an alternative to the metal oxide to achieve high sensitivity and selectivity toward hydrogen. A thin film of Pd nanoparticle based alloy was employed to detect relative high concentrations of hydrogen at room temperature [23]. The detection principle was based on the formation of Pd hydride, resulting from hydrogen atom incorporation into the Pd lattice structure which changes its overall resistance. These so-called chemiresistor sensors have been further modified to use Pd nanowire arrays for the specific sensing of H₂ [24].

Along with Pd, other one-dimensional nanostructure based sensors of other materials have also attracted a great attention due to their extraordinary properties such as size, unique electronic properties and thus superior sensitivity, and power consumption. One such a sensor employs titanium dioxide nanotubes [25]. Furthermore, one-

dimensional nanostructures have been incorporated in hybrid structure sensors. For example, Pd-functionalized tin oxide nanowires exhibited a dramatic improvement in sensitivity toward hydrogen [26]..

It is well-known that carbon nanotubes (CNTs) have become one of the most exciting one-dimensional nanomaterials due to their extraordinary electrical properties that have been exploited in a wide variety of applications. One of these is the sensing application mentioned above. However, very weak adsorption of hydrogen gas results in poor electronic interaction with CNT and even with defective CNTs, poor sensitivity was observed [27]. Therefore, CNTs composites were proposed as a sensing material for hydrogen detection and demonstrated through many studies. Enhanced H₂ sensitivity was observed when a composite based on Pd evaporated on single wall carbon nanotubes was used as hydrogen sensing material [27, 28]. Pd nanoparticles were attached to single walled CNTs for a better sensor performance [29]. Even though there was some preliminary explanation about the electronic interaction between adsorbates (H₂), catalysts (Pd) and sensing materials (CNTs) in this type of structure, the understanding about this sensing mechanism is still not fully established. Especially, in the case of replacing semiconducting SWNTs by multiwall CNTs, not much work has been conducted.

In this study, a better hydrogen gas sensor performance operating at room temperature and using hybrid structure of MWCNTs and Pd nanoparticles is reported. The work presented here provides a new green synthetic method developed to produce defective MWCNTs in a simple evacuated glass tube. These CNTs, decorated with Pd

NPs, show remarkably wide dynamic range for H₂ detection from a few ppm to 4%. The sensing mechanism during chemiresistive process is also established that entails a carrier inversion in the p-type nanotubes and even observation of negative differential resistance. These phenomena can be explained in terms simple reduction of CNTs by atomic/anionic hydrogen generation catalyzed by Pd NPs.

CHAPTER 2. INSTRUMENTATION, DEVICE CHARACTERIZATION AND NANOSTRUCTURE SYNTHESIS

2.1 Synthesis and Sensor Fabrication

The synthesis of nanostructures includes bulk scale preparation of SnO₂ NWs, CNTs, and their doped counterparts. Sensor fabrications based on these nanomaterials for H₂ and Cl₂ detection are also presented in this section.

2.1.1. Sample preparation for SnO₂ NWs

2.1.1.1 Synthesis of undoped and doped SnO₂ NWs:

A. Undoped SnO₂ NWs

Method I: The precursors Sn NPs were synthesized by reducing a thoroughly mixed solution of 50 ml of 0.05 M SnCl₄·5H₂O and 0.5 g of 1, 10-phenanthroline with 100ml of 0.1 M NaBH₄ aqueous solution. The reducing solution was introduced drop wise to synthesize phenanthroline capped Sn nanoparticles [30]. NPs were separated from the reaction mixture by centrifuging (10,000 rpm for 15 minutes) after two hours of reaction time. The precipitate was dried at 50°C in 2hrs on a hotplate. 0.2 g of 1,10-phenanthroline capped Sn NP powder was mixed with a mixture of 0.4 g of NaCl and 0.6 g of KCl (eutecting point at 657C°), grounded into a fine powder, and heated at 750 °C for 2 h in a furnace in air. The molten mixture was then slowly cooled to room temperature. The solidified product was washed several times with deionized (DI) water, to remove KCl and NaCl. Their removal was tested with 0.05mM AgNO₃ solution for any residual Cl⁻ anion. The wet powder was dried overnight in an oven kept at 100°C to remove water.

Method II: A modified synthetic method could be adapted to produce much higher quantity products of SnO₂ NWs. 5g of SnCl₄·5H₂O was dissolved in 150mls of water. The solution was stirred for 15 minutes before drop wise addition of 50ml of 2g NaBH₄ dissolved in water. The mixture turned black, indicating formation of the oxide coated Sn NPs. The precipitate was collected by filtering and left to dry in air overnight. 2g of the above precursor was mixed with 0.2g of 1,10-phenanthroline, 4 g of NaCl and 6 g of KCl, ground into a fine powder, and kept at 750⁰C for 2hr in an oven in air.

B. n-doped SnO₂ NWs

For n-doping, varying amount of antimony chloride (1-5% at. wt. in relation to Sn) was added to the initial solution of SnCl₄·5H₂O. The rest of the synthetic procedure was identical to the synthesis of undoped NWs as described in method II. Note that the actual amount of dopant inserted in NWs could be different than the concentration of dopant atoms used during synthesis.

C. Li-dopedSnO₂ NWs

Method III: For Li-doping, LiCl replaced antimony chloride as a dopant. The amounts of lithium chloride added to the initial reaction mixture ranged from 1 to 5 wt%. The precursors, Sn NPs coated with a thin layer of tin oxide, were synthesized without capping agents by adding 0.35g SnCl₄·5H₂O with 0.1 M NaBH₄ aqueous solution containing LiCl. The mixture turned black, indicating formation of oxide coated Sn NPs. The precipitate was collected by filtering and left to dry in air over night. 0.2g of the above precursor was mixed with 0.2g of 1,10-phenanthroline, 0.31g of LiCl and 0.6g of KCl (eutectic point at 353 C°) , ground into a fine powder, and kept at 480 C° for 2hr.

The molten mixture was then slowly cooled to room temperature at a cooling rate of $4^{\circ}\text{C}/\text{minute}$. The products were collected and purified as in the case of undoped NWs by multiple washing with DI water till Cl^{-} ions were not detected by the AgNO_3 test.

2.1.1.2 Sensor fabrication using n-doped SnO_2 NWs:

To fabricate the sensor, a simple powder compression technique was employed. For the chlorine sensor, 2g of Sb-doped SnO_2 were mixed with 0.5 ml of 2% ethyl silicate in acidified ethanol. The paste was compressed using a pressure of $5\,463\text{ kg cm}^{-2}$ to yield a 13 mm pellet with a thickness of $800\text{ }\mu\text{m}$. The pellet was heated at 450°C for 30 minutes. It was then mounted onto a glass substrate and copper contacts attached using conductive silver epoxy.

Figure 4: Photographs of 1%, 5% and 10% Sb-doped SnO_2 NWs based sensors (from left to right). Note the increase in the dopant concentration leads to increasingly blueish color to the pellets. Electrical contacts (silver epoxy and copper wire) and compressed nanoparticle pellet are mounted on a microscope glass slide.

2.1.2. Sample preparation for CNTs

2.1.2.1 Synthesis of undoped MWCNTs:

A. Dual chamber method (method A):

This method was carried out through two steps including the synthesis of Ni nanoparticles as catalyst precursors, which are used as a catalyst for the synthesis of CNTs through vapor-liquid-solid (VLS) mechanism.

Synthesis of Ni nanoparticles: A 5 nm thin film of Nickel on silicon substrate was deposited by using the physical vapor deposition (PVD) system (see below). The thin film covered silicon substrate was then placed in a glass tube, which was evacuated to 80mTorr pressure. After evacuation, the glass tube was sealed off. The nickel nanoparticle catalysts were formed by annealing at 675⁰C for 15 minutes which breaks the Ni thin film into nano-size islands by dewetting.

Synthesis of undoped CNTs: 10 mg of Naphthalene was first placed into the bottom of a borosilicate glass tube (0.8 cm O.D. X 0.6 cm I.D.), ensuring that the solid did not adhere to the sidewalls. The tube was sectioned into two chambers by thinning its diameter to a neck with a propane torch at approximately 5 cm away from the bottom of the tube. The catalytic Ni nanoparticle coated substrate was then inserted into the top half of the tube. The thinned neck prevents the substrate from falling into the precursor, naphthalene. The top of the tube was thinned to a neck with a propane torch approximately 25 cm from the bottom of the tube for ease of sealing. The whole tube was connected to glass vacuum line at pressure of 80 mTorr. After evacuation, the glass tube was sealed off resulting in a 20 cm duo-chamber glass reaction vessel as shown in Figure 5 below

Figure 5: An optical image of duo-chamber glass reaction vessel for the synthesis of MWCNTs. Catalyst is deposited on the silicon substrate in the right side chamber; and Naphthalene, the CNT feed stock, is kept in the left chamber.

The two chambers were heated to different temperatures using separate heating tapes. To ensure that the Si substrate reaches the desired temperature, the substrate chamber temperatures was adjusted first. The naphthalene containing glass chamber temperature was adjusted once the substrate temperature reached the desired range. . The naphthalene containing chamber temperature was kept at 100°C at the beginning of the reaction. Over the course of reaction (about 2 hours) the chamber temperature rose to about 200°C since the substrate side is kept at about 600°C. This is termed controlled synthesis and illustrated in Figure 6a.

B. Single Chamber method (method B):

In this case the procedure was carried out similarly to method A, but the length of glass tube was changed to 10cm after sealing. Both Ni nanoparticles coated substrate and naphthalene were located in single chamber. The whole tube was then inserted at once into the furnace kept at desired temperature. This is termed as uncontrolled synthesis. See Figure 6b.

Figure 6: Schematic illustration of CNTs synthesis from three different synthetic methods. a) Dual chamber separating catalyst from the reactant. Independent temperature control on the two chambers was obtained using 2 heating sources (heating tapes, Omega, STH051-080). b) Single chamber with one heating source. c) Single chamber reactor where catalyst and reactant are mixed.

Figure 7: The photograph of the custom-built CNT synthesis system housed inside a tube furnace.

C. Using Ferrocene and Cobaltocene as catalyst generators (method C):

The preparation method was similar to the previous methods using vacuum sealed glass tubes. However, no substrate coated with catalysts was utilized. CNTs were grown directly on the inner wall of the glass tube (Figure 6c). A mixture of 15mg of naphthalene mixed thoroughly with ferrocene or cobaltocene with mass ratio 5:1 was placed in one side of the tube and vacuumed down to 60 mTorr followed by sealing. The temperature of this side was ramped up from 100°C to 200°C with the heating rate of 1°C/minute while the other side was kept at 625°C.

2.1.2.2 Synthesis of doped MWCNTs:

In this work, the “green approach” was demonstrated by not using toxic gases as precursors or dopants. Dopants were based on the use of boron and nitrogen containing organic compounds, which are in the solid phase at room temperature. Triphenylborane and benzylamine (Figure 8) are considered as p- and n- type substitutional dopants for carbon, respectively.

Figure 8: Molecular structure of doping agents for CNTs

Boron- and Nitrogen-doped CNTs were synthesized using various levels of triphenylborane and benzylamine as p- and n-type doping sources, respectively. The

syntheses were carried out using a non-substrate pathway (method C). Different amounts of dopant were added to the mixture of cobaltocene and naphthalene to alter the doping concentrations.

2.1.2.3 Synthesis of palladium nanoparticles attached MWCNTs

The attachment of palladium nanoparticles on the walls of CNTs was adapted from Karousis's method [31]. 5 mg of commercial MWCNTs were suspended in 10 ml of 0.05 M aqueous sodium dodecyl sulfate (SDS) solution along with the 10 mg of palladium acetate. This mixture was ultrasonicated for 20 minutes and refluxed at 90°C for 6 hrs. The palladium nanoparticle decorated CNTs were collected and separated from the reaction mixtures by centrifugation and repeated washings with ethanol to remove the excess adsorbed surfactant. The precipitate was then suspended in dimethylformamide solvent (DMF).

The sensor was constructed by depositing a thin film of the above suspension. One ml of the final solution was slowly deposited drop wise and evaporated at 40°C on a 5mm x 10 mm silicon substrate. Conductive silver epoxy was used as the material for electrodes.

2.1.2.4 Sensor construction from as-synthesized MWCNTs and Palladium nanoparticles

The sensor construction for as synthesized-undoped CNTs by the sealed glass tube method was also investigated for hydrogen detection. In this case, the CNT product synthesized from method C was employed. A 3 cm glass tube which has CNTs deposited was cut and gently placed into 10 ml of 0.05 M aqueous sodium dodecyl sulfate (SDS) solution containing 10 mg of palladium acetate. Attachment of Pd nanoparticles onto

CNTs was achieved similarly through the above synthetic process, but without the sonication. This portion of glass tube after synthesis was thoroughly washed many times with water and completely dried in the oven at 100°C in 2hrs. Conductive silver epoxy was deposited on both edges of the tube and used as electrodes for sensor measurements.

Figure 9: A photograph of fabricated CNT based sensor using method C. Electrical contact to the two sides was provided by using silver epoxy and copper wire as shown in Figure 4.

2.2 Electron Microscopy

Scanning electron microscopy (SEM)

An FEI Siron XL30 model SEM was used for electron microscopy imaging. The imaging studies were performed at an accelerating voltage of 5 kV, and beam current of ~300 μ A.

Transmission electron microscopy

The TEM microscope employed in this study was a FEI Technai F-20 TEM operating at an accelerating voltage of 200 kV. A column vacuum of less than 0.10 μ Torr was routinely achieved. TEM studies included bright field, dark field, selected area diffraction, EDX analysis and high resolution lattice plane imaging.

2.3 Physical Vapor Deposition (PVD)

Physical Vapor Deposition (PVD) system was made by Lesker Thin Film Deposition System Model KJLC AXXIS. Various film thicknesses from 5 to 20 nm of Ni catalyst on Si substrate were deposited to optimize the growth conditions for CNTs.

2.4 Infrared Spectroscopy (FTIR)

Fourier transform infra-red (FTIR) spectra were acquired on a Perkin-Elmer RXI instrument. Desorbed gases from the sensor chambers were captured in a gas sampling cell having transparent salt plate windows for Infra-red spectroscopic analysis.

2.5 UV-Visible Spectroscopy

Optical absorption properties NWs were investigated by UV-Vis light using a Perkin Elmer spectrometer model UV-3600 UV-Vis-NIR. The spectra were collected in the range from 250nm to 900nm wavelength. The analysis of absorption yielded the band-gap of as-synthesized SnO₂ NWs.

2.6 Photoluminescence Spectroscopy

In this dissertation work, the optical photoluminescence studies were performed on a Shimadzu spectrofluorophotometer RF-5301PC, using a xenon light source with 3 nm bandwidth. The detector was placed on an axis at 90° angle from that of the incident beam. The sample holder was a rectangular quartz cell with all 4 polished vertical slides. This holder was adjusted at right angle of the incident beam and detector to permit the transmitted radiation to pass through the test solution without contaminating the output signal received by the detector. A sample with SnO₂ NWs dispersed in deionized water was first scanned at different excitation wavelengths, then at a fixed emission wavelength, to determine the optimal excitation wavelength.

2.7 Raman Spectroscopy

Raman spectroscopy is a spectroscopic technique based on the shifts in wavelength of the inelastically scattered radiation. In principle, when a monochromatic laser beam is incident on a material of choice, photons of the laser are absorbed. Majority of the light is Rayleigh/elastically scattered, wherein the energy of emitted photons is equal to incident photons from the laser. However, some photons interact inelastically with the material and undergo a change in energy. Frequency, ν , of the reemitted photons are shifted down (Stokes scattering) or shifted up (anti-Stokes scattering) in comparison to incident photon frequency. These shifts are called “Raman Shifts” and provide a wealth of information about phonon/vibrational energies [32]. A simplified energy diagram illustrating these concepts is shown on the Figure 10.

The primary purpose of conducting Raman spectroscopic studies was to get an indirect indication of sample doping. Both CNT and SnO₂ have well characterized Raman spectra which have been used in the past to interpret the extent of doping by considering the shift in the spectral peak positions as a function of dopant concentration. This was the only method available to characterize the extent of doping on a molecular scale in the synthesized samples.

Figure 10: Raman transitional scheme: different types of scattering are achieved depending on energy of emitted photon ($h\nu'$) relative to the energy of absorbed photon ($h\nu_0$). The bottom two vibrational energy levels as $v=0$ (ground state) and first excited vibrational state $v=1$ are shown.

Raman spectroscopic measurements were carried out using a home-built system illustrated in Figure 11. It consisted of four major components including excitation source (laser), samples illuminating system and light collection optics, wavelength selector (spectrometer) and detector charge-couple device (CCD). Light from an Argon ion laser at 532nm wavelength passed through a laser line filter making the beam highly monochromatic. The beam was then reflected through a couple of beam scattering systems (BS) and focused onto the sample through an objective lens. The Rayleigh and Raman scattered light was collected through the same objective lens and passed through a notch filter to selectively block the Rayleigh scattering. The remaining Raman scattered light passed through a spectrometer (diffraction grating) and collected by a detector CCD to produce Raman spectra on a computer after appropriate signal averaging.

Since a multichannel dispersive spectrometer was used, Raman spectra were collected as peak positions in pixel unit versus peak intensity. The data was analyzed

after converting the unit of pixel to unit of energy (cm^{-1}). Further refinement in the precise peak position was achieved through curve fitting, which is demonstrated in the following sections. In addition, the Si peak was used as a reference to calibrate the system. It has been reported that the first-order Raman spectrum of silicon yields a strong signal at $\sim 520 \text{ cm}^{-1}$ [33]. Therefore, the offset of peak positions on Raman spectra were corrected based on the position of the Si Raman peak, from the silicon substrate bearing CNT or SnO_2 samples.

Figure 11: Schematic showing the home-built Raman system with major components including excitation source (Lighthouse Photonic, model Sprout-G), optic system (Thorlabs, model VM1), wavelength selector (Spectra Pro, model 2150i) and detector (Princeton Instrument, model 7527-0001).

2.8 Sensor testing system

The sensor performance was tested in a home-built test chamber at room temperature [34]. A schematic view of the chamber made from glass is shown in Figure 12. A two wire resistance of the sensor material was monitored by a Fluke 8840A multimeter which was interfaced to a computer on a National Instrument's GPIB-PCII interface. In this method, a continuous flow of diluted test gas was employed as opposed to fixed dosage of analyte gas to the chamber. This was done to prevent the problem of

gas adsorption on the chamber wall which reduces the concentration analyte. The sensor response time was measured by analyzing resistance versus time data. To measure recovery time, a flow of fresh air was maintained in the chamber and the sensor temperature was adjusted. The sample temperature can be adjusted between 25°C and 120°C by varying the resistive heater voltage (0-5volts). A Cr–Al thermocouple was mounted on the heater to measure the sensor operating temperature. The output of the thermocouple was fed to a temperature indicator Mastech (MAS-345) multimeter. The output of the thermocouple was fed to a temperature indicator Mastech (MAS-345) multimeter.

Figure 12: A schematic view of the gas test chamber

The exhaust gas was bubbled through alkali solution to trap the toxic Cl_2 gas. For the hydrogen source, forming gas (4% H_2 in 96% N_2) was used as a primary source of hydrogen. Diluted H_2 gas was directly released in the chemical hood. The test gas diluting system, illustrated in Figure 13 (right), was built on analyte and diluting gas sources as well as flow regulators (Dwyer A25431, Aalborg PMR1011029).

Figure 13: Schematic of gas diluting system (left) and the photograph of home-built sensor testing system (right).

CHAPTER 3. TIN OXIDE NANOWIRES (SnO₂ NWs)

It is well-known that metal oxides have a wide range of conductivity ranging from wide band gap insulators to metal conductors and superconductors. Amongst metal oxide materials, tin oxide SnO₂ is commonly used in electronic applications. A key benefit of this semiconductor over the contemporary semiconductors such as silicon or gallium arsenide is that it is chemically corrosion resistant. Thus, this wide band-gap semiconductor (3.56eV) has found uses in electronics such as transparent conductors [35-38] for flat panel displays, solar cells, and in chemistry as heterogeneous catalysts [39-44] and solid state gas sensing materials (O₂, CO, Cl₂, NO₂, NH₃ etc.) [45-50]. Sensors of SnO₂ nanowire provide a useful comparison with the CNT nanotube based sensors as discussed below. In this chapter, background and significance of SnO₂ nanomaterials are reviewed. Following these, studies of synthesis, characterization and sensor performance of n-doped SnO₂ NWs for chlorine gas detection are presented.

3.1 Background and Significance

The review herein describes physical properties of SnO₂ nanomaterial including crystal structure, stoichiometry and defect structure as well as Raman characteristics of SnO₂ rutile structure.

3.1.1 Crystal Structure

SnO₂ crystallizes in a tetragonal rutile structure with space group D_{4h}¹⁴ [29, 30]. The unit cell contains six atoms, two tin and four oxygen atoms, as illustrated in Figure 14. Each tin atom is at the center of six oxygen atoms placed approximately at the corners of a regular octahedron, and every oxygen atom is surrounded by three tin atoms

approximately at the corners of an equilateral triangle. Thus, it is a structure of 6:3 coordination. The lattice parameters determined by Baur are $a = b = 4.737\text{\AA}$ and $c = 3.185\text{\AA}$ [16, 51]. The ionic radii for O^{2-} and Sn^{4+} are 1.40 and $0.71\text{\AA}$, respectively. Metal atoms (cations) are located at positions $(0,0,0)$, $(0,1,0)$, $(1,0,0)$, $(0,0,1)$, $(1,1,0)$, $(1,0,1)$, $(0,1,1)$, $(1,1,1)$ and $(\frac{1}{2},\frac{1}{2},\frac{1}{2})$ in the unit cell, and the oxygen atoms (anions) at $\pm (u,u,0)$, $(u,u,1)$, and $\pm (\frac{1}{2}+u,\frac{1}{2}-u,\frac{1}{2})$, where the internal parameter, u , takes the value 0.307 . Each cation has two anions at a distance (d_1) of $(2ua)^{\frac{1}{2}}$ ($2.053\text{\AA}$) and four anions at distance (d_2) $[2(\frac{1}{2}-u)^2 + (\frac{c}{2a})^2 a]^{\frac{1}{2}}$ ($2.597\text{\AA}$). Each anion is seen to be bonded to cations in a planar trigonal configuration.

Figure 14: Unit cell of the crystal structure of SnO_2 : large circles indicate tin atoms and the small circles indicate oxygen atoms.

3.1.2 Stoichiometry, defect structure and conductivity

It is well-known that physical properties of oxides strongly depend on the deviation from the stoichiometric composition (SnO_{2-x} as opposed to SnO_2 native disorder) and on the nature and concentrations of foreign atoms incorporated into the

crystal lattice [52]. It is generally agreed that SnO₂ in its undoped form is an n-type, wide-band gap semiconductor with a bandwidth of $E_g = 3.6\sim 4.0$ eV

Therefore, creation of oxygen vacancies ($V_{\ddot{O}}$) as a result of O₂ leads to electron donation to the conduction band [53]

Here Kröger–Vink notation [54] is used , wherein superscripted dots and primes indicate positive and negative charges respectively. The doubly ionized (passively charged) oxygen vacancies ($V_{\ddot{O}}$) serve as main the intrinsic defects and determine the electric properties of the material, and excess electrons (eq. 3.1) donated to the conduction band making the tin oxide an n type conductor. The oxygen vacancies maintain overall charge neutrality on the macroscopic scale. The structure of tin oxide can be represented as SnO_{2-x} ($V_{\ddot{O}}\rangle_x e_{2x}$) ($0 < x < 10^{-5}$). Thus, the conductivity of SnO₂ is ascribed to oxygen vacancies that result from its incomplete oxidation.

An alternative explanation of SnO₂ conducting properties can be based on the diffusion of O⁻² and electron-hopping between metal cations in the metal oxide lattice. In principle, due to non-stoichiometry and oxygen deficiency, the structure of metal oxide semiconducting materials such as TiO₂ and SnO₂ can be represented as MO_{2-x} ($0 < x < 10^{-5}$), which is a combination of MO and MO₂. The ionic lattice is illustrated in Figure 15:

Figure 15: Illustration of ionic lattice of MO_{2-x} where small, big circles and squares represent metal cations, oxygen anions and oxygen vacancies, respectively.

As can be seen from Figure 15, there are a few M^{2+} ions in a sea of M^{4+} ions and a few oxygen vacancies in the lattice. Therefore, the conduction can be achieved by the diffusion of O^{2-} ions through the vacancies or the electrons hopping from M^{2+} to M^{4+} . The resulting oxide is an n-type due to the domination of electron-hopping [55]

3.1.3 Raman Active Vibrational Modes of SnO_2 Rutile Structure

Raman spectra of SnO_2 can be used to evaluate the effect of doping that enhances the conductivity. A brief discussion of its Raman spectrum is provided in this section.

Since there are 6 atoms in its unit cell, SnO_2 lattice structure possesses a total of 18 vibrational modes in the first Brillouin zone. However, just only 4 modes are Raman active including 3 non-degenerate modes A_{1g} , B_{1g} , B_{2g} and one doubly degenerate mode E_g [56]. Their group theoretical symmetry designations with their irreducible representations are presented in Table 3 below.

Table 3: Phonon symmetry of SnO₂ rutile structure with basic functions of the irreducible representation of Raman active modes. Table is adapted and modified from Ref [57]

Phonon Symmetry	Activity	Raman peak position	Irreducible representation
A _{1g}	Raman Active	634 cm ⁻¹	x ² + y ² , z ²
B _{1g}	Raman Active	123 cm ⁻¹	x ² - y ²
B _{2g}	Raman Active	773 cm ⁻¹	xy
E _g	Raman Active	473 cm ⁻¹	zx, zy

In these Raman active modes, oxygen atoms vibrate while Sn atoms are at rest. Specifically, the non-degenerate modes vibrations take place in the plane perpendicular to the c axis while doubly degenerate mode vibrates in the direction of the c axis (Figure 16). The Raman spectrum of rutile-type SnO₂ single crystal shows four peaks corresponding to A_{1g} (634 cm⁻¹), B_{1g} (123 cm⁻¹), B_{2g} (773 cm⁻¹) and E_g (473 cm⁻¹). A_{1g} and B_{2g} are related to bond expansion and contraction. In the A_{1g} mode, two Sn-O bonds contract and the other four Sn-O bonds expand or contract in opposite ways. In B_{2g} mode, all six Sn-O bonds, which are around the Sn ion, contract and expand cooperatively, leading to a larger repulsive force of the O-O bonds than in A_{1g} mode. The B_{1g} mode consists of rotation of the oxygen atoms around the c axis, with all six oxygen atoms of the octahedral participating in the vibration. E_g mode relates to the vibration of oxygen in the direction of the c axis [58].

Figure 16: Raman active vibrational modes of SnO_2 rutile structure from the projection images in the x-y plane [57]. 6 big circles represent 6 oxygen atoms in two different planes (the first plane contains 2 shaded circles and the second contains 4 unshaded circles). Small circles indicate Sn atoms. (+) and (-) signs illustrate the in and out of plane motions, respectively.

3.1.4 Growth of SnO_2 nanowires

During the last decade, methods of synthesis of SnO_2 NWs have been well established. Different morphologies of 1D SnO_2 have been reported such as nanowires, nanorods, nanobelts and nanotubes. The preparation methods can be classified into two categories: vapor phase deposition and solution-based crystal growth. Many common methods include laser ablation [59, 60], thermal evaporation [61, 62], vapor-liquid-solid (VLS) growth [63, 64], carbothermal reduction [65, 66] and rapid oxidation [67]. The vapor phase deposition methods produce high quality nanostructures with well-defined diameter and length distributions. In the other case, solution based growth techniques use molten salt [68, 69] as a solvent to grow nanostructures at high temperature. Furthermore, solvothermal/hydrothermal methods [62, 70] provide bulk synthesis at a lower growth temperature. Even though aspect ratios and morphological features are not

as precisely controlled, as in the case of vapor phase deposition methods, the advantage of solution-based syntheses is bulk scale production to generate materials for sensing application. Therefore, this synthetic pathway was employed in this work to synthesize undoped and doped SnO₂ NWs, which are the focus for the sensing application in this chapter.

In the most general sense, the growth mechanism of one-dimensional nanostructures through solution based synthetic methods follows the Oswald ripening process assisted by anisotropic-directing growing factors. In particular, the precursors of crystal growth process are nanoscopic morphologies or seeds. They are formed under supersaturated condition in the dissolved in a medium due to critical conditions such as pressure concentration and temperature [71]. Following this nucleation stage is the crystal growth phase where the Oswald ripening mechanism prevails. During Oswald ripening the larger crystal structures grow at the expenses of smaller crystals [72]. During this process, morphologies of crystal structure formation can be controlled by varying the parameters in the reaction systems that affect the kinetic and thermodynamic properties of the system through choices of solvents, capping reagents, control of reaction temperatures, change of pH values, etc [73].

Of these, the most important factor in controlling the formation of one dimensional nanostructures is the coordination of surfactants/capping agents [73]. Selective adsorption of surfactant in a particular single crystal plane affects its surface energy. According to the Gibbs- Wulff facet theorem, a nanocrystal, which is bounded by facets, has a minimum total surface energy at equilibrium [74]:

$$\int_i^n \gamma_i dS_i = 0 \quad (\text{eq. 3.2})$$

Where:

$\gamma(i)$: the specific surface free energy of facet i

$S(i)$: surface area of facet i

In other words, the equilibrium shapes of a nanocrystal are constructed by the facets and their corresponding areas so that the total surface energy is minimized. The morphology of nanostructures is determined by the relatively specific surface energies of the facets. Thus, it is possible to manipulate the shape of a crystal by controlling the free energies of crystallographic surfaces [75]. The introduction of appropriate surfactants or capping agents can be employed for this purpose during the crystal growth processes through the facet selectively passivating process. Therefore, by appropriately choosing the surfactant, solution based method could be employed to synthesize different one dimensional nanostructure materials [73]. Recently a new mode of NW growth through development screw dislocations has been established [76]. The establishment of this mechanism requires sophisticated TEM analysis and hence was not pursued.

3.1.5 SnO₂ NWs as sensor material

As discussed before, oxygen vacancies transform SnO₂, a wide band gap semiconductor, to an n-type semiconductor. SnO₂ based sensors function through changes in conductivity brought about by electron transport to/from analyte to the semiconductor. There are two kinds of sensing responses observed. The first mechanism is that the electrons are withdrawn from the conduction band upon the adsorption of charge accepting molecules such as NO, Cl₂ and O₂ (oxidizing agents). The negatively

charged adsorbed species repel the majority carriers from the surface. This leads to the depletion region near the surface that is devoid of carriers, i.e., electrons with concomitant reduction of conductivity. On the other hand, as a second mechanism, some molecules such as CO or H₂ (reducing agents) could react with the surface adsorbed oxygen anion and release electrons to the conduction band to cause an increase in conductance [77] by driving the semiconductor in accumulation mode.

Figure 17 below illustrates the adsorption of chlorine gas on the surface of SnO₂ nanoparticles. The respective energy levels with respect to vacuum and the Nernst hydrogen electrode levels are shown. Since the redox energy level of Cl₂ is lower than oxygen, Cl⁻ ions can be generated even in the presence of adsorbed oxygen anions. As Cl⁻ ions are formed by the abstraction of electrons from the conduction band, they repel the majority carriers which are electrons in the n-type semiconductor from the gas-solid interface, leading to formation of a depletion layer as showed in Figure 18. As a result, this layer provides an inter-nanoparticle potential barrier that hinders electron transport and hence increases the sensor resistance [48]. An oxidizing species increases the sensor resistance while a reducing species decreases it resistance for an n-type semiconductor. Complementary behavior is expected for a p-type semiconductor. This generic response can be made selectively if a proper dopant with specific affinity for the analyte can be found. An example for this type was demonstrated in case of Cl₂ detection using antimony doping [34] (Figure 17 B).

Figure 17: A. Energy levels of oxidizing and reducing gases in relation to band edges of SnO₂. B. Location and position Sb reduction potential which enable it to act as a catalyst/dopant conferring selectivity for Cl₂ detection.

Primary driving force for detection of oxidizing species is the location of conduction band edge of n-type SnO₂. Any oxidizing species whose reduction potential is below the edge can extract electrons and reduce the conductivity of the SnO₂. While reducing species, whose reduction potential is above the filled states of the conduction band, can donate electrons and increase conductivity of the semiconductor. This is dictated by the minimization electron energy which requires that it seeks the lowest available energy state. Given that the reduction potential of antimony is between the conduction band edge and Cl₂ reduction potential, the latter is highly and selectively detected with respect to other gases such as NO and H₂ whose reduction potential are greater than antimony. While gases such as NO₂ and Br₂ are detected but with lower degree of selectivity as their reduction potential falls between chlorine and antimony,

Figure 18: Schematic diagrams of porous sensing layers. a) No or very small depletion layer (high conductivity) in absence of Cl_2 , (b) in chlorine, depletion layer thickness increases. (c) In nanowires, the number of electron hops amongst the charge carrying particles would be reduced leading to an expectation of lower resistance.

Additionally it can be surmised from Figure 18b and 18c, that NWs based sensor should have lower resistance than its counterpart composed from NP as fewer inter-particle electron hops are required for electron transport between electrodes. Nevertheless, the actual relative resistance between the NP-based and NW-based sensor might be different. For compressed powders, used in these studies, it is clear that the electrical conductivity of as-fabricated sensors not only depends on extrinsic factors such as the applied pressure but also on the intrinsic factors such as morphological properties [78]. The difference in morphologies, which determine the degree of packing of the material or the density, leads to the change in contact resistance between NWs and NPs in compressed powder. Since the same preparation method was employed the extrinsic factors can be neglected. Intrinsic factors play an important role in the conductivity of as-fabricate sensors. These effects are discussed in later in the discussion section. Furthermore, it should be noted that the contacting electrode material, silver, has its

Fermi level near the conduction band edge, Figure 17 A. This leads to a desirable Ohmic contact for a resistive sensor.

3.2 Results and Discussion

Figure 19: Optical images of undoped (left) and Sb doped NWs (right)

3.2.1 Electron Microscopy

A. n-doped SnO_2 NWs

The colors of undoped and doped products are illustrated in Figure 19. Different colors white and dark green correspond to undoped and n-doped nanowires. The green color becomes darker for higher dopant concentrations.

The morphology of as-prepared un-doped and Sb-doped SnO_2 NWs was characterized by SEM. The majority of the un-doped NWs showed an average NW diameter ranging from 20 to 50 nms and the length from 1 to 5 μms , (Figure 20 left). It was noticed that NWs were mixed with the whisker-like nanorods. In the n-doped NWs, the size distribution was much broader (from a few tens of nanometers corresponding to length of a nano-whisker up to 10 μms for the nanowire length), perhaps due to incorporation and morphological effects of dopant atoms. Also, note that the smaller

diameter NWs tended to fuse with one another (Figure 20 right). Such structures are rarely observed in the nucleation and growth of NWs mediated by the vapor-liquid solid (VLS) mechanism. TEM and high resolution TEM of n-doped NWs along with electron diffraction in inset (Figure 21) indicate the single crystalline nature of NWs with a prominent peak corresponding to the interplanar spacing of 0.34 nm reflecting a well-known crystallographic plane [110] of SnO_2 . The crystal structure of these 20 nm diameter NWs was a characteristic cassiterite type (details are not shown here) which is commonly observed upon calcination at 500°C in SnO_2 NPs synthesized by sol-gel method [79]. The NWs showed a very straight and smooth surface morphology suggesting a 1-D Ostwald-ripening mechanism [68] by which the formation of NWs (the larger crystal structures) could have grown at the expense of NPs (smaller crystals). Since the synthesis was carried out at elevated temperature (750°C), 1, 10-phenanthroline has a very minor effect as a surface free energy controller on the growth mechanism. However, it is a crucial factor as an interfacial capping agent in the formation of precursors. During the synthesis of NPs, 1,10-phenanthroline could cap to the surface of Sn NPs and prevent their agglomeration as well as the formation of SnO_x . The Sn NPs can grow in spherical shape and more uniform size distribution [80], resulting in better morphology of NWs. This is supported by the experiments summarized in Table 4. Specifically, the higher aspect ratio of SnO_2 NWs was achieved with the presence of 1,10 phenanthroline in the same synthetic condition of temperature and medium. Perhaps, during 1-D Ostwald-ripening process, this capping agent has reduced the formation of SnO_x on the surface of Sn NPs and facilitated the growth of SnO_2 NWs.

Figure 20: SEM images of undoped (left) and Sb doped NWs (right)

Figure 21: TEM images of n-doped SnO_2 NWs with low (left) and high magnifications along with electron diffraction in the inset (right).

B. Li-doped SnO_2 NWs:

The color of Li-doped nanowires is more yellowish than undoped nanowires as showed in Figure 22.

Figure 22: Optical image and SEM image of Li-doped SnO_2 NWs synthesized at 490°C

SEM images of Li-doped NWs synthesized at 490°C showed a broad size distribution for length and diameter. The length of wires varied from 200 nm up to 1 μm while the diameter of single wires ranged from 50 nm to hundreds of nanometers. Like the n-doped NWs, whisker-like nanorods with smaller diameters and shorter lengths were also observed along with the tendency of the NWs to fuse. Thus, preformed NPs transformed into NWs through recrystallization from alkali halide salts exhibited similar characteristics regardless of the nature (n or p type) of the dopant. The morphology of NW nanostructure was influenced by a variety of factors, such as precursors, the moisture content of grounded mixture before the recrystallization and growth temperature. Different experiments (Table 4) have been carried out to investigate the effects of synthetic conditions on the formation of SnO_2 NWs. Results indicated that with the same synthetic conditions, better morphology in terms of the aspect ratio of SnO_2 NWs product were achieved by using Sn nanoparticles compared to their oxide capped counterparts. Besides, it was observed that the pre-synthesized powder, when dried to get rid of moisture, gave a better yield of wires and more uniform dimensional distributions

(data not shown here). Also, the NWs formed at temperatures higher than 490°C, gave longer wires (>1µm in length) with a more uniform length distribution as shown in Figure 23. Replacing coated Sn NP precursors by SnO NPs [81] did not produce NWs at 490°C. However, recrystallization at 750°C did produce NWs with low yield.

Figure 23: SEM image of Li-doped SnO₂ NWs synthesized at 600°C and 700°C

Table 4: The effect of synthetic conditions on SnO₂ NWs morphology (O and NO stand for observed and non-observed, respectively)

Synthetic condition of SnO ₂ NWs	Length (nm)	Diameter (nm)
SnO _x nps + N ₂ + 490°C	NO	NO
SnO _x nps + N ₂ + Phen + 490°C	O	20±2
Dilute SnO + 490°C	925	125±6
Sn nps + N ₂ + 490°C	240±3	46±5
Sn nps + Air+ 490°C	292±3	21±3
Sn+ Air+ Phen+490°C	407±4	129±3
Sn nps + Air + 700°C	800±20	72±2

In addition, HR-TEM was also employed to characterize the precursors and resulting NWs. SnO coated Sn NPs were synthesized by methods B and C, described in chapter 2 (section 2.1.1.1). Figure 24 shows the morphology of nanoparticles with

spherical shape and mean diameter around 10 nm. They show a tendency to agglomerate perhaps due to the absence of interfacial capping agent 1,10 phenanthroline.

Figure 24: A) HR-TEM of SnO_{2-x} coated Sn NP precursors and B) TEM of Li-doped SnO_2 NWs

The agglomeration of NPs was possibly responsible for fused NWs during the growth process (Figures 20, 22 and 23). The HR-TEM of Li-doped SnO_2 NWs shown in Figure 24B indicated their single crystalline nature with lattice fringe characteristics. The lattice interplanar spacing is about 0.34 nm corresponding to the [110] plane of the cassiterite crystal structure.

3.2.2 Photoluminescence Characterization

A. n-doped SnO_2 NWs:

Room-temperature photoluminescence spectra of undoped SnO_2 NPs and NWs appear in Figure 25. The excitation wavelength used in this study was 310 nm corresponding to energy greater than the bulk bandgap energy for SnO_2 . Two different PL characteristics were observed for NPs and NWs. Both nanostructures gave a UV emission at 360 nm, corresponding to the band gap of SnO_2 (3.56 eV), partially overlapped

with Raman (stokes shift) scattering of water (this also occurs at 360 nm). The latter peak appears at a constant frequency different from the excitation frequency.

Figure 25: Photoluminescence of undoped SnO₂ NWs and NPs (excitation wavelength 310 nm). A new broad peak around 460 nm (2.7eV) was observed for NWs.

Bulk SnO₂ is an indirect band-gap semiconductor so it is not expected to have a strong PL corresponding to its bandgap energy, as it would violate the principle of momentum conservation. Observation of PL in these nano-geometries (i.e. in both NPs and NWs) implies a PL characteristic of a direct band-gap type semiconductor. Similar effects are well known in silicon NPs, porous silicon and silicon NWs [82] but are not still well understood. Interestingly, the NWs have an additional broad emission peak near 460nm (2.7eV). The precise understanding of this PL peak currently does not exist; although explanations in terms of defects including vacancies of oxygen, dopant segregation, and lattice disorders inside the lattice of SnO₂ [60, 68, 83, 84] or near the surface can be advanced.

High concentration of dopants can lead to an impurity band near the conduction (in the case of n-type dopant) or the valence (in the case of Li dopant) bands. The resulting band gap narrowing could result in a red shift in the PL. As shown below in Figure 26, under the same excitation wavelength 280nm, both un-doped and n-doped SnO_2 NWs gave two UV emission peaks, one, at 310 nms, is due to solvent Raman scattering and the other broad peak appearing at $\sim 345\text{nm}$ wavelength (3.59 eV). For the latter peak a red shift of 20nm was observed for Sb-doped NWs, with peak $\sim 365\text{ nm}$ (3.4eV), in comparison with the undoped NWs. This is consistent with a dopant induced band-gap narrowing concept [85]. These estimate give an approximate bandgap which is larger than the actual bandgap as the precise method for determining band gap is unavailable. Using the PL band edge at high wavelength can have contribution from midgap or surface states so its use would report lower band gap, hence the emission peak maximum was chosen as a crude compromise.

Figure 26: Photoluminescence of undoped and n-doped SnO_2 NWs (excitation wavelength 280 nm). The red shift of 20nm is observed. Note Solvent scattering at 310nm

B. Li-doped SnO₂ NWs:

A comparison of photoluminescence spectra, excited with 280nm, in undoped, 5% n- and 5% Li-doped SnO₂ NWs appears in Figure 27. Two emission peaks were observed for all different types (undoped, n- and Li-doped) of NWs. The first peak at 310 nm is due to solvent. The PL of Li-doped NWs also showed a red shift (≈ 23 nm) for the broad emission peak occurring at ~ 368 nm (3.37eV). This shift could be explained similarly to the case of n-doped using the induced band-gap narrowing concept.

Figure 27: Photoluminescence of undoped (peak 345nm or 3.59 eV), 5% n-doped (peak at 365 nm or 3.4 eV) and 5% Li-doped (peak at 368 nm or 3.38 eV) SnO₂ NWs. Solvent peak is at 310 nm

3.2.3 UV-Vis Spectroscopy

The optical bandgap of SnO₂ nanostructure samples including undoped, doped SnO₂ NWs and n-doped SnO₂ NPs were investigated using the UV-visible absorption spectrometer in the wavelength range of 200–800 nm as shown in Figure 28. The

wavelength on the X-axis was converted to energy unit, (eV) for the bandgap determination as following:

Figure 28: UV-Vis spectra of different SnO₂ NW samples: undoped, 5% n-doped and 5% Li-doped. 5% n-doped NP sample was also investigated for comparison.

It is suggested that the bandgap is direct in the rutiled-nanostructure of SnO₂ [86].

As the direct interband transition takes place, the optical absorption coefficient α follows as a function of incident photon energy E exceeding the band gap [87].

$$\alpha = [A(E - E_0)]^{\frac{1}{2}} \quad (\text{eq. 3.3})$$

Where:

E_0 is optical bandgap and A is a constant.

Since the optical absorption coefficient α and transmittance T are related as:

$$T = e^{-\alpha L} \text{ or } \alpha = -\frac{\ln T}{L} \quad (\text{eq. 3.4})$$

Where

L is the thickness of the sample.

Therefore the transmittance and incident photon energy are correlated as:

$$-\frac{\ln T}{L} = [A(E - E_0)]^{\frac{1}{2}} \text{ or } (\ln T)^2 = AL^2(E - E_0) \quad (\text{eq. 3.5})$$

This linear relationship is confirmed in the photon energy range of 3.5 – 4.5 eV depicted in Figure 29, and its extrapolation to the E axis gives E_0 values for different SnO_2 samples.

Figure 29: Determination of optical bandgap from the UV absorption edge. Points and dashed curve are experimental and linear fitting/extrapolating, respectively.

Table 5: Optical bandgap determined from UV absorption edge and photoluminescence

Sample	Undoped SnO ₂ NWs	5% Li-doped SnO ₂ NWs	5% n-doped SnO ₂ NWs	5% n-doped SnO ₂ NPs
Band edge from UV-VIS E ₀ (eV) (λ nm)	3.43 $\pm$ 0.07 (362)	3.32 $\pm$ 0.06 (373)	3.29 $\pm$ 0.06 (377)	3.11 $\pm$ 0.08 (398)
Band peak from PL E ₀ (eV)	3.6 $\pm$ 0.1	3.38 $\pm$ 0.06	3.4 $\pm$ 0.04	N/A

The evaluated bandgaps of different SnO₂ nanostructure samples are summarized in Table 5. The optical bandgap of undoped SnO₂ NWs was determined to be 3.43 $\pm$ 0.07 eV, which is close to the bandgap of bulk SnO₂ 3.6 eV [88]. Upon the introduction of impurities into the lattice, the bandgap of SnO₂ NWs decreases to 3.32 $\pm$ 0.06 eV and 3.29 $\pm$ 0.06 eV corresponding to 5% Li-doped and 5% n-doped, respectively. Specifically, donor and acceptor energy levels have been added by the impurities into the band gap of SnO₂ inducing the shift of the Fermi level closer to band edges (CB for n-type and VB for p-type). With an increase in the amount of doping, the density of states generated by impurities increases and forms a continuum of states below or above the conduction or valence bands leading to narrowing of the bandgap [89]. In addition, the decrease of the bandgap upon doping, and determined herein, is consistent with what has been observed from photoluminescence analysis, by which 3.59 eV, 3.40 eV, 3.38 eV are the bandgaps of undoped, 5% n-doped and 5% Li-doped SnO₂ NWs, respectively.

It was observed that with the same doping concentration of antimony, doped SnO₂ nanoparticles synthesized from sol-gel methods have a bandgap energy 3.11 $\pm$ 0.08 eV lower than doped SnO₂ NWs synthesized by the molten salt method. This could be the result of different concentration of impurities inserting into the lattice during the doping processes by two different synthetic pathways. However no further quantitative analysis was performed to verify this hypothesis.

3.2.4 Raman Spectroscopy

The Raman of SnO₂ nanostructure samples including undoped, doped SnO₂ NWs and doped SnO₂ NPs were investigated using a custom-fabricated Raman system. The data were recorded in the wavelength range 550- 850 nm as shown in Figure 30.

Figure 30: Raman spectrum of SnO₂ NWs samples. Three peaks at around 633 (cm⁻¹) corresponding A_{1g} symmetry vibration, 697 (cm⁻¹) and 775 (cm⁻¹) corresponding to B_{2g} symmetry were observed. Points and solid curve are experimental and fitting, respectively.

Each curve in the spectrum was fitted using three Lorentzians and one tilted baseline. An example of fitting for 2% n-doped NWs is illustrated in Figure 31. The complete data analysis of Raman peaks from the curve fittings is showed in Table 6

Figure 31: Illustration of curve fitting of 2% n-doped SnO₂ NWs sample. Points and solid curve (yellow) are experimental and fitting, respectively. The complete fitted curve (yellow) is composed from three Lorentzian (blue, green and red solid) curves and one tilted baseline.

Table 6: Raman data analysis of different SnO₂ samples

Sample/ vibrational mode	Undoped SnO ₂ NWs	2% n-doped SnO ₂ NWs	5% n-doped SnO ₂ NWs	5% Li-doped SnO ₂ NWs	10% Li-doped SnO ₂ NWs
A _{1g} (cm ⁻¹)	632±2	622±3	617±3	627±6	623±3
A _{2u} (cm ⁻¹)	N/A	698±6	671±24	702±7	704±7
B _{2g} (cm ⁻¹)	794±5	768±6	N/A	777±12	768±6

Raman spectrum from the analysis show three peaks consistent with the previous report for rutile-type SnO₂ single crystals [58]. Peaks at around 633 (cm⁻¹) and 775 (cm⁻¹) could be assigned to A_{1g} and B_{2g}, respectively. Due to apparatus limitation, B_{1g} and E_g modes were not resolved from the spectrum. Interestingly, the intense peak around 697 cm⁻¹ which corresponds to the Raman-forbidden mode (A_{2u}) is also observed. This mode, which is only infrared active, is now seen in the Raman spectrum of SnO₂ NWs. This unexpected peak could be associated with a breakdown of the selection rules, or new

modes called ‘volume modes’. This Raman activity is induced by disorder [56, 90, 91] as the result of sensitivity to local geometric disorientation and to the neighboring disorder (particularly atoms from other sub-lattices or electric defects associated with substitutions or vacancies) [92].

It is also observed that the insertion of impurities into the SnO₂ NW lattice leads to the Raman peak shift (A_{1g} mode) toward lower frequency. Compared to Li doped, n-type SnO₂ NWs have more noticeable red shifts upon the introduction of dopant. This phenomenon is consistent with Mo-doped SnO₂ NWs, observed by Yi’s group [93] and explained by lattice distortion or lattice symmetry lowering.

3.2.5 Electrical properties of thick film using n-doped SnO₂ NWs

In this study, silver epoxy was used as the electrical contacting electrode, and plays an important role in forming an Ohmic contact with semi-conductive Sb-doped SnO₂ nanomaterial compressed to form a thick film (see chapter 2). Figure 32 also shows I-V characteristics of NWs as function of dopant concentration. The linear dependence between current and voltage from the I-V measurement of each sample shows the Ohmic contact characteristic as expected. This good apparent ohmic contact takes place perhaps due to degenerate doping [94]. As impurity concentration is increased, the as-fabricated sensor resistance significantly decreases and achieves its minimum value at 5% dopant concentration. However, at higher dopant concentration an increase in the resistance was observed. The resistances of 3%, 5% and 10% doped NW based sensors are ~270 kΩ, 0.2 kΩ and 2.1 kΩ, respectively. This characteristic is derived from the phase separation

upon high doping concentration of Sb, which generates an insulating layer Sb_2O_3 on the surface of NWs and increases inter-nanowires potential barrier.

Figure 32: I-V curves of different dopant concentrations of n-doped NW based sensors. The resistances of 3%, 5% and 10% doped NW based sensors are $\sim 270 \text{ k}\Omega$, $0.2 \text{ k}\Omega$ and $2.1 \text{ k}\Omega$, respectively.

3.3 Sensing Application: Chlorine Resistive Gas Sensor Based on n-doped SnO₂ NWs

In this section, the as-fabricated Sb-doped SnO₂ NW based sensor is employed for chlorine gas detection. The sensor performance is investigated through its response, sensitivity and stability toward air-diluted chlorine gas. The study is based on direct current conductivity measurements.

3.3.1 Sensor response, response time and recovery time

Typically, the gas response (*S*) of a resistive sensor is defined as the ratio of the change in resistance of the sensor on exposure to the target gas to the original resistance in air:

$$S = \frac{R_{gas} - R_{air}}{R_{air}} \quad (\text{eq. 3.6})$$

Where *R_{gas}* and *R_{air}* are the resistance of the sensor in a target gas medium and in air, respectively

The response time is defined as the time needed for the sensor resistance to change by 90% of the difference from the maximum value to the minimum after the test gas injection. The recovery time is the time required for the sensor resistance to change by 90% of the difference from the minimum value to the maximum after releasing the test gas [95]. The response and recovery time at room temperature of 10% Sb-doped SnO₂ NW based sensor, at six different Cl₂ concentrations, are shown in Figure 33. The response and recovery time at five different temperatures (25°C, 35°C, 45°C, 90°C, 110°C) are summarized in Figure 34.

Figure 33: The sensor resistance (A) and recovery time (B) at six different Cl₂ concentrations from 5 ppb to 10ppm. The sensor was recovered by introducing fresh air and heating at 45°C. (Note: Data in Figure 33A does not include sensor response during recovery process)

Figure 34: Response and recovery time (minute) as a function of temperature (°K). The slope of a linear fitting from ln(time) vs 1/T for recovery gives desorption activation energy of 82±3 kJ/mol.

The response times of the device are short at room temperature but the recovery times appear to be quite long. Specifically, it takes close to 10 minutes for the sensor to respond but about 60 minutes to recover at room temperature after exposure to 1 ppm chlorine. However, it was found that the response kinetics of the device is affected by the kinetics of the gas flow through the testing chamber. As long as the chlorine concentration in the testing chamber still remains in the high ppm range, the response and recovery times cannot be determined accurately. This impact is even more dominated when switching from high to low chlorine concentration detections.

The response and recovery times are temperature dependent as expected from analyte adsorption and desorption processes, respectively. The response time exhibited a minimum around 45°C while the recovery time decreased as a function of temperature and followed Arrhenius type behavior (eq. 3.7) [96]. From this, the thermal activation energy from desorption process was determined to be 82±3 kJ/mole as shown in Figure 34 above. The value of E_a is suggestive of chemisorption of chlorine.

$$Rate = Ae^{-\frac{E_a}{RT}}$$

$$\text{Or } \ln(Rate) = -\frac{E_a}{RT} + B \quad (\text{eq. 3.7})$$

Where:

A and B are constants; R is gas constant

E_a is activation energy (KJ/mole)

T is absolute temperature (°K).

Data presented in Figure 33 A above also show high response ($S \approx 125$ for 1ppm of Cl_2 concentration) and able to detect chlorine at room temperature with relatively fast

response time compared to current known metal-oxide based sensors for chlorine detection [97-100]. Remarkably, the sensor exhibits a very high sensitivity to chlorine concentration (5 ppbs). This characteristic has been significantly improved from our recent report of a Sb doped-SnO₂ NPs based chlorine sensor [34]. Similarly, the undoped SnO₂ NW based sensor shows a relatively poor sensitivity compared to its Sb-doped counterparts reconfirming a potential catalytic effect of Sb, as shown in Figure 17, in improving sensitivity.

3.3.2 Sensor Sensitivity

A. Sensor Sensitivity as a function of chlorine concentration

Figure 35 shows the response of the sensor fabricated from 10% Sb doped SnO₂ NWs to different Cl₂ concentrations in air from 5ppb to 10ppm. In this study, the sensor was continuously renewed after every 2 cycles of exposure and recovery. This has minimized the effect of adsorption and desorption processes on the sensor stability, which is discussed in the following section. The response increases non-linearly with increasing chlorine concentrations and tends to get saturated after 1ppm. The continuous flow of analyte can eliminate the effect of adsorption of Cl₂ on the chamber wall and contributes to the ability to detect target at very low concentrations. Significantly, chlorine gas can be unprecedentedly detected at ppb level at room temperature.

Figure 35: Response as a function of chlorine concentration at room temperature. The sensor response was investigated in the range from 5ppb to 10ppm of Cl₂ concentrations.

A model used to describe this observed non-linear relationship between concentration and response can be based on a surface-trap-limit model, proposed by Dawson [101]. The equilibrium of a dissociative chemisorption of the gas with sites, S, on the oxide surface is formulated as:

These reactions are formulated as a competitive adsorption of chlorine and oxygen:

$$\frac{\theta_{Cl} P_{O_2}^{\frac{1}{2}}}{\theta_O P_{Cl_2}^{\frac{1}{2}}} = \frac{\theta_{Cl}}{\theta_O} \gamma^{-\frac{1}{2}} = K_{ads} \quad (\text{eq. 3.8})$$

Where θ is the fractional coverage over the surface, P is the partial pressure and $\gamma = \frac{P_{Cl_2}}{P_{O_2}}$

In the chemisorption process, the uncharged adsorbed species on the surface of metal oxide acts as a surface electron trap. The equilibriums can be represented as:

$$\frac{\theta_{Cl^-}}{n' \theta_{Cl}} = K_{Cl}$$

$$\frac{\theta_{O^-}}{n' \theta_O} = K_O$$

If it is assumed that the uncharged and charged species are the major and minor surface coverage species, respectively, then at equilibrium:

$$\theta_O + \theta_{Cl} \approx 1$$

$$\theta_{O^-} + \theta_{Cl^-} = q$$

Where q is a constant with $q \ll 1$. From eq. 3.8, we have:

$$\left(\frac{q}{K_O} \right) \left(\frac{1}{n'} \right) = \frac{1 + \frac{K_{Cl} K_{ads} \gamma^{1/2}}{K_O}}{1 + K_{ads} \gamma^{1/2}} \quad (\text{eq. 3.9})$$

Since the left-hand side of eq. 3.9 is proportional to the resistance R of the sensor, by considering the limit of low and high concentration of chlorine, we have:

$$\frac{R}{R_0} = \frac{1 + \frac{K_{Cl} K_{ads} \gamma^{1/2}}{K_O}}{1 + K_{ads} \gamma^{1/2}} = \frac{1 + K_{rel} K_{ads} \gamma^{1/2}}{1 + K_{ads} \gamma^{1/2}} \quad (\text{eq. 3.10})$$

Where $\frac{K_{Cl}}{K_O} = K_{rel}$

If R_∞ and R_0 denote the resistance response of the sensor in the limit of high and low concentration of chlorine, then we will have $K_{ads} \gamma^{1/2} \gg 1$:

$$\frac{R_{\infty}}{R_0} = \frac{K_{Cl}}{K_O} = K_{rel} \text{ and } K_{rel} \gg 1$$

The equation can be written as:

$$\frac{R-R_0}{R_0} = \frac{(K_{rel}-1)K_{ads}\gamma^{1/2}}{1+K_{ads}\gamma^{1/2}} \approx \frac{K_{rel}K_{ads}\gamma^{1/2}}{1+K_{ads}\gamma^{1/2}} \quad (\text{eq. 3.11})$$

The resulting expression bears similarity to the Langmuir model. The curve fitting of sensor response as a function of $\gamma^{1/2}$ based on eq. 3.11 is shown in Figure 36

Figure 36: Response of the sensor as a function of $\gamma^{1/2}$. Red points and black solid curve are experimental and fitting data, respectively. K_{rel} and K_{ads} determined from the curve fitting were 450 ± 50 and 0.2 ± 0.1 , respectively.

Since $\frac{K_{Cl}}{K_O} = K_{rel}$, $K_{rel} \gg 1$ has indicated that the energy of the state represented by Cl_{ads} is much lower than the energy of the state of O_{ads} as anticipated from the redox levels for oxygen and chlorine (Figure 17). Upon doping SnO_2 by Sb, the chemisorption activation energy of chlorine compared to oxygen on Sb-doped SnO_2 is significantly reduced. Furthermore, since K_{rel} is the ratio of the resistance of sensor in the high and low

concentration of chlorine, the value of K_{rel} also provides the estimation of the maximum sensor response, which is around 450 ± 50 .

B. Sensor Sensitivity as a function of doping concentration:

It is well-known that a large amount of Sb can be incorporated into a SnO_2 lattice to enhance its conductivity [102] leading to a change of sensor performance. The response of the sensor as a function of dopant concentration (Sb) was investigated to determine the optimal dopant concentration for highly sensitive devices. The data in Figure 37 shows that the response of the sensor, and thus sensitivity, is optimal at the doping level of the synthetic condition of about 10% (mole Sb/mole Sn). At higher doping levels, both conductivity and response decrease.

Figure 37: Response of the sensor as a function of doping concentration (Sb/Sn mole ratio). The sensor response is optimal at 10% doping. At very high doping concentration, sensor response decrease perhaps due to the formation of Sb_2O_3 on the surface of NWs.

It has been established that the incorporation of high concentration of Sb into SnO_2 , dopant atoms are concentrated near the surface of the nanostructures and induces

surface segregation of Sb to Sb^{3+} and Sb^{5+} [103-105]. These surface segregated Sb atoms may form oxygen vacancy complexes at the surface and may be important active sites for adsorption of electron-withdrawing gas molecules [106]. Therefore, the mechanism for the adsorption of chlorine of the surface of Sb-doped SnO_2 NWs at room temperature can be proposed as [34]:

Since the response of the sensor is proportional to concentration of segregated Sb atoms on the surface of NWs, the response increases along with increases of the doping concentration. However, at very high doping concentration, formation of insulating oxide Sb_2O_3 , increases the barrier for abstraction of an electron from semiconductor nanowires, and leads to the decrease of conductivity as well as response of the sensor as showed in Figure 37.

3.3.3 Sensor Stability and Understanding the Sensing Mechanism

A. More specific model of sensing mechanism

The chlorine gas sensing response of metal oxide sensor is based on the chemisorption model. In particular, chlorine is presumed to react with metal oxides to form chlorides or displace lattice oxygen at the interface, or possibly adsorb onto surface oxygen vacancies. The chemisorption chlorine involves mainly four kinds of adsorption behavior in the reactions below [101, 107, 108]:

Where subscripts, surf and o, mean the species adsorbed on the surface and the species occupying lattice oxygen sites, respectively. $V_O^{\bullet\bullet}$ is an oxygen vacancy bearing double positive charge, and x denotes charge neutrality and e represents the conduction electron. Positive charge and negative charge are marked by a point (e.g. $A^{\bullet}$) and a hyphen (e.g. $A^{\bullet-}$), respectively. In the mechanism (I) and (II), chlorine substitutes for adsorbed oxygen and lattice oxygen to form $Cl^{\bullet}$ and donates electrons to the conduction band of the semiconductor, which should result in a decrease of resistance, and thus inconsistent with the observed results in this study. On the other hand, in the mechanism (III), chlorine is adsorbed on the surface and draws electrons from the semiconductor to form $Cl^{\bullet}$, leading to an increase of resistance. Similarly for the mechanism (IV), absorbed chlorine occupies the oxygen vacancy along with drawing electrons to become chlorine ions. This mechanism also results in the increase of resistance.

Intuitively, both of these mechanisms (III) and (IV) are probable upon the exposure of Cl_2 gas, corresponding to the reaction of Cl_2 on the surface and inside the lattice of Sb-doped SnO_2 NWs. This model was already incorporated at the beginning of section 3.4.2. With the mechanism (III), creation of the chloride ion formed on the oxide surface would lead to a depletion layer by repelling majority carrier away from the surface as illustrated in Figure 38. A depletion layer as well as inter-nanowire barrier that hinders electron transport hence increases the sensor resistance.

Figure 38: Schematic diagram of the porous sensing layer composed from nanowires (each circle illustrates NW cross section). There is no depletion layer in the absence of Cl_2 . Upon exposure to Cl_2 , the depletion layer is generated as a result of Cl' formation on the surface of NWs.

To investigate the effect of mechanism (IV) on the response of the sensor, the experiments based on multiple times of repeated exposure and refresh cycles on the same sensor were carried out. Data below show the response of the sensor upon the exposure to 1ppm Cl_2 gas, the sensor recovery was achieved by a flow of air combining with heating the sensor at 45°C .

Figure 39: Response of the sensor from 4 continuous cycles of exposure and recovery. Sensor resistance (R) and cycle time (t) were recorded to investigate the effect of mechanism (IV).

Figure 40: Illustration of mechanism (IV). The square inside the SnO_2 lattice represents oxygen vacancy, which is occupied by diffused $\text{Cl}\cdot$. This process leads to electron deficiency inside SnO_2 , causing sensor resistance baseline drifting.

It can be seen that the baseline resistance drifts with the increase of initial sensor's resistance (R_{air}) after every cycle of exposure and recovery. The increase of R_{air} has led to the sensor response (S), which is defined as $(R_{\text{gas}} - R_{\text{air}}) / R_{\text{air}}$, to decrease over time. Explanation for this characteristic can be based on mechanism (IV) and illustrated in Figure 40. Particularly, chlorine gas molecules are absorbed on the surface of doped SnO_2 NWs and dissociate into two chlorine atoms, which subsequently diffuse into to the bulk SnO_2 NW lattice. These chlorine atoms replace oxygen vacancies along with drawing electrons from the conduction band of Sb-doped SnO_2 to become chloride ions occupying oxygen vacancies. This mechanism contributes to the drifting of the sensor resistance baseline after every cycle of exposure and recovery.

The drift of baseline resistance as a function of chlorine exposure time can be modeled as first order kinetic process described by follows equation:

$$R = R_0 (1 - e^{-kt}) \quad (\text{eq. 3.12})$$

Where R_0 is the maximum resistance, resulting from complete saturation of oxygen vacancies by chloride ions

The plot in Figure 41 shows resistance drifting as a function of time. From this, the maximum drifting resistance value R_0 ($14000 \pm 500 \Omega$) was achieved by curve fitting based on eq. 3.12.

Figure 41: Sensor resistance baseline as a function of cycle time. 5 black data points in the graph correspond to t_0 , t_1 , t_2 , t_3 , t_4 . Solid red curve shows the fitting following eq. 3.12.

In order to further confirm the effect of chloride ions occupying oxygen vacancies, which led to the shift of the sensor resistance baseline, we have carried out an experiment by storing the sensor in very high concentration of chlorine gas (99%). After 8 hours of storage, maximum sensor resistance was achieved to be $\sim 59.5 \text{ k}\Omega$. Following this, the sensor was refreshed with air over night at room temperature. At this point, sensor resistance was measured to be around $8 \text{ k}\Omega$ s. The sensor performance was then investigated (data presented in Figure 42) by exposing to 1 ppm chlorine concentration

and recovered by a flow of dried air along with heating at 100°C in 8 hours. The stabilized sensor resistance after cooling down to room temperature was measured to be ~ 5.1 kΩs.

Figure 42: Sensor response curves after storing in high Cl_2 concentration, the sensor performance was investigated by exposing to 1 ppm Cl_2 gas. There was not much shift of sensor resistance baseline in this case.

As can be observed from Figure 42, when the sensor was stored under high chlorine concentration, mechanism(III) and (IV) has taken place, and oxygen vacancies were completely replaced by chloride ions, leading to maximum sensor resistance (59.5 kΩ). After refreshing with air for next the performance investigation, the resistance baseline drifting was significantly reduced. The resistance after recovery stabilized at almost the same level as before the exposure (~8 kΩs). At this point, the exposure and recovery were simply based on the adsorption and desorption processes (mechanism III), which took place only on the surface of SnO_2 NWs, leading to a better sensor performance characteristic. When the sensor was refreshed with air under high

temperature 100°C in 8 hours, the reverse mechanism (IV) took place for oxygen vacancy recovery, leading to the decrease of sensor resistance.

B. Sensor response kinetics

For every cycle of the exposure and recovery process, the sensor response was investigated in terms of a simple successive reaction kinetic model. The kinetic response during the exposure process can be assumed to follow two steps. Chlorine gas molecules firstly approach the surface of SnO₂ and dissociate into chlorine atoms, which subsequently extract electrons from SnO₂ conduction band. This can be presented in equation below:

Where k_1 and k_2 are dissociation and electron extraction kinetic constants of chlorine atoms on the surface of SnO₂, respectively.

Therefore, the change of sensor resistance as a function of time can be written as:

$$R = R_0 \left(1 - \frac{k_1}{k_1 - k_2} (e^{-k_1 t} - e^{-k_2 t}) - e^{-k_1 t} \right) + C \quad (\text{eq. 3.13})$$

Where R_0 and C are the maximum sensor resistance and the average value of baseline resistance during a cycle, respectively

The fitting using eq. 3.12 for the first exposure cycle is illustrated in Figure 43.

¹The breakdown of Cl-Cl bond may proceed through 2 steps:

However, the detection of $Cl \cdot$ was not pursued.

Figure 43: An example of fitting sensor response resistance as a function of time following (eq. 3.13)

Table 7: Kinetic analysis of response curves during the exposure process.

	1 st cycle	2 nd cycle	3 rd cycle	4 th cycle
$k_1 (s^{-1}) (x10^3)$	6 ± 2	5.3 ± 0.5	4.0 ± 0.3	4.0 ± 0.5
$k_2 (s^{-1}) (x10^3)$	6 ± 2	5.3 ± 0.5	4.0 ± 0.3	2.5 ± 0.3
$R_0 (ohm) (x10^{-3})$	7.0 ± 2	7.0 ± 1	6.0 ± 1	6.0 ± 1
$C (ohm) (x10^2)$	2.0 ± 0.5	9.0 ± 0.9	2.7 ± 0.1	3.3 ± 0.1

The results of kinetic constants of chlorine dissociation and electron abstraction on the surface of Sb-doped SnO₂ are summarized in Table 7. As can be seen, there is not much change in the dissociation rate constant (k_1) with successive cycles of adsorption and desorption. The difference in rate constant (k_2) is more obvious between the first cycles corresponding to “fresh surface” and 4th cycle corresponding to “unclean surface”. The almost two times decrease in the rate constant suggests that the oxygen vacancies replaced by chlorine slow down the electron transfer rate from the conduction band of SnO₂ to the adsorbed chlorine atom.

When the sensor is recovered, reverse reactions including surface desorption and oxygen vacancy recovery are expected. Trapped electrons during the chlorine absorption process could be released and donated back to the conduction band of Sb-doped SnO₂ by

reaction between chlorides and oxygen in air, as represented by eq. 3.14 and eq.3.15. This would cause the increase of electron concentration, and thus conductivity. However, complete refresh is just only achieved at elevated temperatures upon the reversed reaction of mechanism (IV) or the recovery of oxygen vacancies. Because the diffusion of chlorine out of defect is absolutely negligible at the temperature used in this experiment [109], chloride species still remain on the near surface and inside the SnO₂ lattice, leading to the depletion of charge carrier concentration, causing drifting of sensor resistance baseline as analyzed above.

Understanding of kinetic mechanism during this recovery process was based on the continuous cycles of exposure and recovery data. Reactions, causing the decrease of sensor resistance, between chloride ions absorbed on the surface of Sb doped SnO₂ NWs as well as with oxygen gas in air can be represented as:

$$\frac{d[Cl^-]}{dt} = -k_1[Cl^-] - k_2[Cl^-]^2$$

$$-dt = \frac{d[Cl^-]}{[Cl^-](k_1 + k_2[Cl^-])}$$

$$[Cl^-] = \frac{k_1 e^{-k_1 t + c}}{1 - k_2 e^{-k_1 t + c}}$$

$$R = R_0 \frac{k_1 e^{-k_1 t + c}}{1 - k_2 e^{-k_1 t + c}} \quad (\text{eq. 3.16})$$

Where C is a constant. k_1 and k_2 are the reaction kinetic constants.

These kinetic constants (eq. 3.14 and eq. 3.15) can be evaluated from fitting of the sensor resistance change as a function of time followed eq. 3.16. An example of this fitting is illustrated in Figure 44 below:

Figure 44: An example of fitting curve follows eq. 3.16: sensor resistance decays as a function of time during the first sensor recovery process

Table 8: Kinetic analysis of response curves during the recovery process.

	1 st curve	2 nd curve	3 rd curve	4 th curve
$k_1 (x10^{-5})$	10 ± 1	9 ± 1	8 ± 1	7 ± 1
k_2	0.30 ± 0.01	0.65 ± 0.01	0.67 ± 0.08	0.75 ± 0.1

C. Reaction products of recovery process

Detection of ClO_2 and Cl_2O gases as reaction products of the sensor recovery process was also attempted. During the sensor refreshing process, a flow of air through the exposed sensor was captured and collected in an IR gas cell and analyzed by FT-IR spectroscopy.

Figure 45: Infrared spectrum composed from blank sample (A), mixture of chlorine and air (B), reaction products during sensor recovery process (C)

Three additional absorption peaks were observed in IR spectrum of the recovery process sample (C) in the region from 700 cm^{-1} to 1500 cm^{-1} . These peaks can be ascribed to dichlorine monoxide (Cl_2O) and chlorine dioxide (ClO_2). Specifically, the absorption near 980 cm^{-1} peak shows a hybrid band ($\nu_2 + \nu_3$), and the peak at 1250 cm^{-1} corresponds to overtone ($2\nu_1$) of the vibrational modes of dichlorine monoxide where ν_1 ($\sim 630\text{ cm}^{-1}$) and ν_2 (270 cm^{-1}) are stretching modes while ν_3 (cm^{-1}) is a bending mode [110]. Due to instrument limitations, absorptions below 650 cm^{-1} were not detected. The origin of these bands could be rationalized from the following reaction:

Additionally, the $\text{Cl}=\text{O}$ stretching mode of ClO_2 , which is around 1040 cm^{-1} is also observed from the result. This is consistent with the observations from many studies on the vibration of the ClO_2 molecule [111-113] noted in Table 9 below:

Table 9: IR observation of dichlorine monoxide (Cl_2O) and chlorine dioxide (ClO_2)

Molecule	Vibrational Mode	Experimental observation	Literature values
Cl_2O	Overtone of stretching mode	$\sim 980 \text{ cm}^{-1}$	977 cm^{-1} [110]
	Hybrid of stretching and bending modes	$\sim 1250 \text{ cm}^{-1}$	1260 cm^{-1} [110]
ClO_2	Stretching mode	$\sim 1040 \text{ cm}^{-1}$	1040 cm^{-1} [112], 1039 cm^{-1} [111], 1040.2 cm^{-1} [113]

3.4 Conclusion

Single crystalline n-doped and Li-doped SnO₂ NWs structures exhibit two strong emission bands in their fluorescence spectra. The PL spectra of doped materials also show a red shift of the emission wavelength compared to their undoped counterpart. Raman peaks in the regions from 550 cm⁻¹ to 850 cm⁻¹ were observed and attributed to the A_{1g}, B_{2g} and A_{2u} Raman shifts. The band-gap measurement using optical spectroscopy has shown the dopant induced bandgap narrowing characteristic upon doping, implying higher carrier concentrations of both n- and Li NWs. Availability of this synthetic method for n- and Li NWs opens up new opportunities for crafting transparent nanoelectronic devices, and sensors.

A thick film chlorine sensor of Sb-doped SnO₂ SW material has been fabricated and demonstrated. Replacing n-doped nanoparticles with n-doped NWs has shown a better sensor performance, in term of sensitivity. The detection limit of chlorine gas has been extended to the ppb level at ambient condition. The chlorine detection sensitivity has been measured and explained in terms of a surface-trap-limit model. It also reconfirmed the role of Sb dopant used to achieve high sensitivity of SnO₂ nanostructures toward Cl₂ gas.

Additionally, the mechanisms of sensor response and recovery have been studied and well established. The transient responses of the sensor, i.e. response and recovery of the device as a function of time were kinetically investigated. The sensing mechanism was interpreted in terms of a chemisorption model accompanied by sensor baseline drifting explanation. The occupancy of chlorine atoms in oxygen vacancies of an SnO₂

lattice, affecting sensor stability, was demonstrated. Dichlorine monoxide (Cl_2O) and chlorine dioxide (ClO_2) are the main products of sensor recovery process

CHAPTER 4. CARBON NANOTUBES (CNTs)

In this chapter, some background and significance of carbon nanotubes (CNTs) are reviewed. Following these, studies of synthesis, characterization and sensor performance of as-synthesized MWCNTs for hydrogen gas detection are presented.

4. 1 Background and significance

4.1.1 Electronic properties of CNTs

Carbon nanotubes have a cylindrical structure. The cylinder is formed from graphene – a honey-comb lattice of covalently bonded carbon atoms. There are many possible nanotube (NT) geometries, depending on how graphene is rolled into a cylinder. Subtle geometric variables, such as the alignment between the cylinder axis and the graphene crystal axes, strongly influence the electrical properties of a NT. The structure of CNTs may be completely characterized by the unit chiral vector coordinates $\bar{a}_1$ and $\bar{a}_2$ multiplied by their respective periodicity indices n and m and mapped on a graphite lattice (eq. 4.1).

Figure 46: (n,m) coordinate system for a graphene sheet. For values of $n=m$ (a) the corresponding SWNTs are metallic “armchair.” SWNTs with $m=0$ (b) are called “zigzag” and are also metallic. All other combinations (c) are chiral and may be metallic or semiconducting [5].

Thus carbon nanotubes are named for their specific (n, m) components of $\vec{a}_1$ and $\vec{a}_2$. Three different types of carbon nanotubes are: armchair, zigzag and chiral. Using the parameters n and m, one can directly calculate the single wall CNT diameter using equation 4.2 and the so called chiral angle for chiral CNTs by using eq. 4.3

$$\text{Tube diameter} \quad d_t = 0.246 \left[\frac{\sqrt{n^2 + nm + m^2}}{\pi} \right] \quad (\text{eq. 4.2})$$

$$\text{Chiral angle} \quad \phi = \sin^{-1} \frac{\sqrt{3}m}{2\sqrt{(n^2 + nm + m^2)}} \quad (\text{eq. 4.3})$$

One of the remarkable properties of carbon nanotubes is that they may behave as a conductor or semiconductor as determined by the (n, m) parameters in the tight binding approximation [114]. When $n-m=3j$ (where j is an integer,) the valence and conduction bands meet at 6 points in 2D k-space. These six points are energetically located at the Fermi level and if the wave vector for the first Brillouin zone passes directly through them making a zero band gap or metallic tube. This is also true for the condition where $n = m$. When $n - m \neq 3j$ the wave vector for the first Brillouin zone no longer pass through the conduction and valence band contact points. In this case there is a band gap, and the tubes are semiconducting.

It is interesting to note that the metallic nature of a SWNT is not solely dependent on the relative magnitudes of n and m, but also the tube geometry plays an important role. An experiment by Yildirin and Gulseren [115] at the NIST Center for Neutron Research have shown that by compressing a semiconducting SWNT in to an elliptical shape the

tube will undergo a transition where the band gap vanishes. At this point the otherwise semiconducting tube shows metallic behavior. Generally the bandgap of CNTs depends inversely on the diameter of the tube, and for CNTs with diameters greater than 14 nm thermal energy exceeds the bandgap. A consequence of this is that most MWCNTs as well as larger CNTs exhibit metal-like conductivity.

4.1.2 Growth of CNTs

Historically, the electric arc discharge is the oldest synthetic method for carbon nanotube production including MWNT and SWNT. Two other synthetic methods are laser evaporation/ablation and chemical vapor deposition (CVD).

The arc discharge technique uses two high-purity graphite electrodes. The anode is graphite containing catalyst metals. The catalysts usually are Fe, Ni, Co, Mo, Y either alone or in mixtures. Better results are obtained using bimetallic catalysts. The synthesis is carried out in vacuum chamber containing inert gas at low pressures around 600 mbar. A strong bias is applied across electrodes, which are momentarily brought into contact with each other to strike an arc. The temperature in the inter-electrode zone is so high that carbon sublimates from the positive electrode (anode) and is consumed. For this method, various kinds of products are formed in different parts of the reactor: large quantities of rubbery soot on the reactor walls; web-like structures between the cathode and the chamber walls; grey hard deposits at the end of cathode; and spongy collaret around the cathodic deposits. Amorphous carbon, encapsulated metal nanoparticles, polyhedral carbon are also present in the products [116, 117].

Figure 47: Schematic diagram of a DC-arc discharge system for CNT synthesis [118]

The laser ablation method is technically similar to the arc discharge counterpart. The difference between them is the quality and purity of obtained products. In 1995, Smalley's group [119] at Rice University utilized the pulsed laser-ablation process for the production of single-wall carbon nanotubes. In this method, a laser beam was focused and scanned across a metal-graphite composite target which is placed in a high temperature furnace around 1200°C. The products were swept by a flow of Argon gas (inert gas) and collected by a water-cooled copper collector positioned downstream. A configuration test has indicated that a composite ablation target consisting of 1% each of Ni and Co metal uniformly mixed with graphite gave the best purity of product and highest yield of SWNTs. Impurities in the product were amorphous carbon, graphite particles, fullerenes and metal catalysts.

Figure 48: Schematic diagram of a laser ablation system to synthesize CNTs [118]

Along with the capability of producing large quantities of unpurified nanotubes by arc discharge, chemical vapor deposition is another advanced synthetic method of CNT not only in terms of bulk synthesis but also providing control over morphology and location of the synthesized product [120]. Today, the catalytic chemical vapor deposition (CCVD) method is considered as the only economically viable process for large-scale CNT production and the integration of CNTs into various devices [121]. In principle, chemical vapor deposition is the catalytic decomposition of hydrocarbon or carbon monoxide feedstock with the aid of supported transition metal catalysts. Generally, the experiment is carried out in a flow-tube furnace at atmospheric pressure. The reaction mixture containing a source of hydrocarbon and an inert gas (carrier gas) is passed over the catalyst bed at temperatures ranging from 500 °C to 1100 °C. Ultrafine metal catalyst particles are either introduced into the reactor directly or formed *in situ* using precursors such as metallic thin films and then annealed to form particles. The general nanotube growth mechanism in CVD process involves the decomposition of hydrocarbon molecules into carbon atoms, which subsequently diffuse into catalysts. The saturation of

this system leads to the formation of tubular carbon solids with sp^2 hybrid structure. The characteristics of the carbon nanotubes produced by the CVD method depend on the working conditions such as the temperature, pressure, concentration and type of hydrocarbon. In addition to the size and nature of metallic catalyst, substrate and reaction time [122] play important roles.

Figure 49: Schematic diagram of a CVD system to synthesize CNTs. Figure was adapted and modified from ref [123]

4.1.3 Raman Spectroscopy of CNTs

Raman is one of the most important techniques to characterize CNTs, providing further understanding about properties of this nanomaterial such as structural, optical and electronic properties. Herein, Raman spectroscopy characteristics of CNTs are presented:

A. Single Wall CNT (SWCNT):

A typical Raman spectrum of a SWCNT is illustrated in Figure 50 below. The most important Raman features of these nanostructures are Raman-active breathing modes (RBMs), the high frequency disordered-induced band (D), graphite band (G band), and second-order Raman scattering from the D-band occurring at twice the frequency of the D band (2D).

Figure 50: An example of a typical Raman of SWCNTs [32] with 4 bands RBM, D, G and 2D observed in the spectrum.

RBM mode: This mode is a confirmation for the presence of CNTs in the sample and located between 75 and 400 cm^{-1} from the excitation line. It corresponds to an in-phase stretch of carbon along the circumference of a nanotube as illustrated in Figure 51. Therefore, its magnitude changes as a function of the nanotube diameter [124] and can be used to determine the diameter distribution of the nanotubes in a bundle [125]:

$$\omega_{RBM} = \frac{A}{d_t} + B \quad (\text{eq. 4.4})$$

Where A and B are constants and vary between individual tubes

Figure 51: The RBM mode of a SWNT [126]. Small solid circles are carbon atoms in SWNT structure (left) and the arrows indicate the direction of the C-atom displacements (right).

G-band: The G-band is an intrinsic feature of carbon nanotubes that is closely related to vibrations in all sp^2 carbon materials. In simple graphite, a single mode is observed at around 1580 cm^{-1} [127]. In SWCNT, this mode occurs between $1500 - 1605\text{ cm}^{-1}$ and transforms into two different modes called G^+ and G^- . This is a result of the confinement of wave-vectors along the circumference [128]. The lower frequency component [129] is associated with vibrations along the circumferential direction, and thus depends on the curvature or diameter of nanotube. The higher frequency component (G^+) corresponds to vibrations along the direction of the nanotube axis and does not depend on the diameter of nanotube. These split modes also play an important role in determining whether the nanotube is semiconducting or metallic. Specifically, metallic tubes are easily recognized by the broad and asymmetric line shape of the G^- band compared to its counterpart in semiconducting tubes. The G^+ frequency remains essentially a sharp peak and constant with diameter of either metallic or semiconductor tubes [128].

Figure 52: Vibrational modes of G^+ (A) and G^- (B) bands [126]. Small solid circles and arrows indicate carbon atoms and their vibrational displacement directions, respectively.

The D band and G' (2D) bands: These bands are both observed in the Raman spectra of semiconducting and metallic SWNT. The D band is known as a disordered or

defect mode because it involves scattering from a defect, which breaks the basic symmetry of a graphene sheet [128]. On the other hand, the second-order G'-band does not require an elastic defect-related scattering process, and it is an intrinsic property of the nanotube and graphite, and present even in defect-free nanotubes for which the D band is completely absent. In Raman spectra, the disordered D-band is located 1250-1450 cm^{-1} , and its second-order overtone G'-band is 2500 – 2900 cm^{-1} .

B. Multi-walled Carbon Nanotubes:

Compared to SWCNTs, which are seamless cylinders comprised of only a single layer of graphene, MWCNTs have higher structural complexity. Because MWCNTs consist of multi rolled layers of graphene, most of the characteristic differences that distinguish the Raman spectra in SWCNT from the spectra for graphite are not so evident in MWCNT. For example, the RBM mode of MWCNTs, which associates with the diameter in the case of SWCNT, is usually too weak to be observed and has broad Raman signals. These characteristics result from large diameter outermost tubes and the ensemble average diameters of tubes in MWCNT [130]. Similarly, the G-band in SWCNT is a diameter dependence band mode and splits into G^+ and G^- modes for small diameter tubes. The corresponding splitting modes in MWCNTs are small in intensity and are smeared out due to the effect of the diameter distribution within individual MWCNT. Therefore, the G-band of MWCNTs in Raman spectra is observed as a single broad peak around 1582 cm^{-1} , just like in graphite [130].

4.1.4 Doping of CNTs

It is well known that CNTs have a very wide range of application in almost every field of nanotechnology. One of the most crucial applications of this nanomaterial is in nanoelectronics, requiring n and p type doping, which can be attained by doping with electron donors and electron acceptors. Many routes to achieve the dopant containing CNTs are listed in Figure 53.

Figure 53: Doping CNTs through many different pathways. The figure is adapted and slightly modified from ref [131]

- a) Substitutional doped single-wall nanotubes through post synthesis modification of CNT with dopant containing organic molecule or incorporation during the synthetic process
- b) Nanotube bundles intercalated with atom or ions
- c) CNTs are functionalized with dopant containing molecules that change the properties of CNTs
- d) CNTs are filled with fullerenes and inorganic compounds

However, conventional pathways to manipulate doping in semiconducting CNTs are based on the intercalation of alkaline metals such as K, Rb [132] [133] or Li [134, 135] for n-type doping. Besides, accumulation in the form of carbon-halogen complex using fluorine [136] or bromine [137] is also reported. More recently and commonly, studies on boron and nitrogen substitutionally doped CNTs have showed that their electronic properties strongly depend on the composition, where the dopants including nitrogen and boron tend to produce electronic properties of metallic CNTs [138-141].

The most common pathway to dope CNTs is through classical CVD and aerosol assisted CVD methods. The doping is routinely carried out *in situ*, in a CVD reactor or enclosed chamber, by flowing volatile gases containing dopants such as acetonitrile [142], NH_3 [143], melamine [144], tri-isopropyl borate [145] etc., via mass flow regulators that control the precise flow rate. The overall cost of system complexity and handling toxic gases are large obstacles for the synthesis of doped-CNTs, and these methods are considered to be “non-green approach”.

4.1.5 Carbon nanotube based gas sensors

Kong and coworkers [14] demonstrated that upon exposure to gaseous molecules such as nitrogen dioxide (NO_2), ammonia (NH_3) or oxygen (O_2) conductivity of semiconducting SWCNTs are significantly altered with high sensitivity and fast response time. Modi's group [146] has developed better sensors that have superior selectivity and are insensitive to effects of environmental conditions such as moisture, temperature and gas flow rate. They used thin devices utilized thin film of MWCNTs and employed electrical break down voltage for sensing different gases on the tips of nanotubes. In

another study, Jang et al [129] developed a chemical sensor employing MWCNTs as the active sensing element for detection of NH_3 in air at different flow rates. Further studies for other gas detections were also achieved such as the work done by Penza's group [147]. In this work, vapors of volatile organic compounds were detected at ppm level concentrations by a CNT-coated surface acoustic wave sensor.

In general, the CNT sensing mechanism of many gases is based on the charge transfer between defect sites on CNTs and the adsorbed analyte. The transfer of either electrons or holes from adsorbed molecules to CNT or vice versa will change the conductance of CNTs. Previous theoretical studies have indicated that due to the weak CNT-analyte adsorption interaction in defect free CNTs leads to a large activation barrier for charge transport for NH_3 , CO or NO_2 gases even at high temperatures [14, 148, 149]. However, the adsorption of gas molecules is much easier on defective CNTs [150, 151] which leads to lower adsorption barrier and promotes the charge transfer rate. Oxygen is known to adsorb and dissociate on the defective sites of CNTs. Furthermore, dissociated oxygen extracts an electron from CNT doping it to a p-type semiconductor [152]. This is because the reduction potential of oxygen is lower than the commonly accepted Fermi level in CNTs as will be discussed later. The concentration of the ionized oxygen can be influenced by reducing as well as oxidizing gaseous species thereby affecting conductivity of CNTs. Multiple oxygen molecules could potentially dissociate on the same CNT defect, providing more dissociation sites for gaseous molecules and leading to much higher net charge transfer to CNTs [153].

Figure 54: Various structural defects on CNT including defect-free tube (a), vacancy (b), interstitial (c), straight Stone-Wales (d) O_2 molecule pre-dissociated defect (e), and slanted stone-Wales defect (f)

Figure 55: An example of NH_3 molecule adsorption on defective CNT for different types of defect illustrated on figures above.

Work presented here shows a new method to synthesize CNTs and also a mean of derivatizing them for use as sensors. The primary focus here is understanding the detailed molecular mechanism of analyte and CNT interactions.

4.2 Results and Discussions

4.2.1 Electron microscopy

A. Ni nanoparticle precursor synthesis:

In CNT synthesis methods (A and B; see chapter 2) we generate Ni nanoparticles on substrate by annealing a thin film of Ni deposited through physical vapor deposition. The effect of catalyst layer thickness to nanoparticle formation was explored. As the thickness of Ni film changes, the morphology of nanoparticles also changes. Figure 56 below shows SEM images of Ni nanoparticle catalysts, annealed from two different Ni film thicknesses. As analyzed from the images, when the annealing film thickness increases from 5nm to 20nm, the nanoparticle size increases from 50 ± 20 nm to 340 ± 150 nm. In addition, formation of completely separated individual particle cannot be achieved if the Ni film is too thick. This has been observed in the 20nm thick Ni film case where the film still remains with generated catalysts trapped inside the film after the annealing process.

Figure 56: Formation of Ni nanoparticle catalysts after annealing 5nm and 20nm thick Ni thin films at 680°C .

B. CNT synthesis

Method A: The synthesis of CNTs was carried out using a 5 nm thick Ni thin film to form nanoparticle precursors. From dual chamber synthesis (method A described in chapter 2, section 2.2.1), different results were observed at different synthesis temperatures. Formations of carbon nanotube/nanowhisker like structures using Ni nanoparticle catalysts took place at temperatures between 600°C and 610°C with best results at 600°C. Lower or higher synthetic temperatures resulted in non-CNT products. It was also observed that the diameter of CNTs increases as the reaction temperature increases. At 590°C nucleation of CNT appears to collapse. On the other hand, at much higher temperature, 620°C carbon nanoparticles were generated.

Figure 57: Method A for the synthesis of CNTs at 590°C, 600°C, 610°C and 620°C

These observations could be rationalized using a model developed for silicon nanowire/nanoparticle synthesis by Chan [154] and it is illustrated below.

The model assumes that decomposition of naphthalene results in the formation of an unstable intermediate of carbon (C). The concentration of carbon could be consumed in two competing pathways, either CNTs or C particles/fibers. Therefore, there are three dominant reaction rates affecting the formation of products, including the rate of naphthalene decomposition (R_1), the rate of CNT (R_2) and the rate of carbon nanoparticle growth (R_3).

At the elevated temperature, k_1 is large and not a rate determining step. The rate constants k_2 and k_3 are inherently temperature dependent and change with temperature. Thus, the dominating product depends on the ratio of k_2 to k_3 .

The rate of formation of CNTs can be written for equation as:

$$R_2 = \frac{d[\text{CNT}]}{dt} = k_2[\text{Ni}^0] \cdot [\text{C}]^n \quad (\text{eq. 4.5})$$

The rate of formation of C particle can be written for equation as:

$$R_3 = \frac{d[\text{C (particle)}]}{dt} = k_3 \cdot [\text{C}]^m \quad (\text{eq. 4.6})$$

The ratio of the rate of reactions is:

$$\frac{R_2}{R_3} = \frac{k_2[Ni^0].[C]^n}{k_3.[C]^m} = K.[C]^{n-m} \quad (\text{eq. 4.7})$$

If $m \gg n$;

$$\frac{R_2}{R_3} = K \cdot \frac{1}{[C]^m} \quad (\text{eq. 4.8})$$

Therefore, high carbon concentration in the vapor phase at high temperature leads to $R_2 \ll R_3$ meaning that the rate of C particle formation would dominate the rate of CNT formation. In other words, the heterogeneous nucleation rate would exceed the rate of nanotube growth mediated through Ni nanoparticles. On the other hand, at lower temperatures, relative low carbon concentration in the vapor leads to $R_2 \gg R_3$, resulting in a higher density of CNTs and lower quantity of C particles.

Since the atomic rearrangement is the main the characteristic of CNT growth mechanism, which involves a thermally activated elementary process, the growth is diffusion controlled, and thus its rate is temperature dependent, according to the Arrhenius equation (eq. 4.9). The rupture carbon structures at 590°C could be associated with the slowdown of growth rate at such lower temperatures.

$$D = D_0 e^{-E_A/kT} \quad (\text{eq. 4.9})$$

Where

D and D_0 are the diffusion coefficient at temperature T and at infinite temperature

E_A is the activation energy for diffusion

k is the Boltzmann constant

In order to confirm the formation of CNTs which have the hollow structure, high resolution TEM studies were undertaken. The data below show the morphology of as-synthesized products at 600°C. From Figure 58, the diameter of CNTs is about 20nm with

length of a few microns. It also shows that there are about 20 to 30 concentric shells within each MWCNT. The MWCNT growth mechanism was the tip based as confirmed from EDX elemental composition analysis showing Ni nanoparticles at the tip of the CNT.

Figure 58: HRTEM of CNT synthesized at 600°C in 2hrs (top) and EDX of catalyst (bottom) and STEM of a CNT.

Method B: In this method, the catalysts and carbon source (naphthalene) were heated up to the desired temperature simultaneously in single glass chamber. Similar to the previous process using dual chamber synthesis, four different products were observed at four different temperatures. The wire-like products were formed at 600°C and 610°C.

At a lower temperature 590°C and higher temperature 620°C, carbon sheathed nanoparticles were observed

Figure 59: Method B for the synthesis of CNTs in uncontrolled manner at 590°C, 600°C, 610°C and 620°C

Method C: Optical images of glass tubes shown below are from results of synthesis, using ferrocene and cobaltocene as catalysts with naphthalene as a carbon source. Black powder like residue coated the glass tube walls. The temperature range, monitored by a temperature probe along the glass tube, indicated that CNTs just grow in the temperature window from 590°C to 620°C. This range was comparable with the observation in the case of Ni nanoparticle catalysts, annealed from thin film (Figure 56) using methods A and B. Further characterizations were carried out by SEM with the data below showing the observation of non-optimized products of CNT synthesis. SEM

images also revealed the presence of catalyst nanoparticles (Fe and Co) at the tip of CNTs. With identical growth conditions including temperatures and pressures (same amounts of precursors), cobaltocene as the catalyst precursor provided smaller diameter CNTs with higher density. Besides CNTs, carbon fiber-like nanostructures of larger diameters were also present in the products.

Figure 60: Optical images of method C for the synthesis of CNTs ferrocene (top) and cobaltocene (bottom). Synthesis using cobaltocene gives higher density of CNT products.

Figure 61: SEM images of CNTs by method C: ferrocene (left) and cobaltocene (right). Smaller diameter CNTs were observed using cobaltocene.

Further confirmation of CNT formation was carried out using HR-TEM. Similarly, tip growth mechanism using cobaltocene as catalyst precursor was observed in Figure 62. The image shows 20 to 30 concentric CNTs in each MWCNT. More

importantly, CNTs, grown by this method, seem to be highly defective. It has been confirmed that the substrate material, its surface morphology and textural properties greatly affect the yield and quality of the resulting CNTs. The chemical state and structure of the substrate are more important than that of the metal catalyst [155]. The weak metal (Ni catalysts)-support (silica substrate) interaction allows high metal dispersion, resulting in metal catalyst aggregation and a high density of catalytic sites [156]. This could facilitate the growth of more defective CNTs and carbon nano-fiber [157].

However, defective characteristic of the as-synthesized CNT product plays an extremely crucial role in the sensing application, which is based on the electrical response in the presence of chemical vapor. The defect sites can dominate the electrical response by forming low energy adsorption sites that also serve as sites for analyte condensation [151]. Therefore, defective CNTs synthesized from this method are employed for the sensing application.

Figure 62: TEM images of CNTs by method C using cobaltocene as catalyst precursor. Tip-growth mechanism and about 20-30 concentric CNTs in each MWCNT are observed.

4.2.2 Raman Characterization of undoped and doped CNTs

Raman spectroscopy was employed to investigate the structure of as-synthesized undoped and doped CNTs [158]. Different n-type and p-type doping concentration CNT samples were collected and characterized. The data was analyzed to confirm the insertion of impurities (dopants) into the lattice structure of CNTs upon doping. In this experiment, five different samples undoped, 5% p-doped, 10% p-doped, 5% n-doped and 10% n-doped CNTs were investigated

Figure 63: Raman spectroscopy of as-synthesized MWCNTs in the region 1200-1800 cm⁻¹. Two peaks corresponding to D band and G band are observed. The data was fitted following eq. 4.10. The curves are shifted vertically to better display the results.

The curve fitting was based on two Lorentzians and one linear baseline:

$$Y = \frac{I_1}{1+B_1(X-X_{01})^2} + \frac{I_2}{1+B_2(X-X_{02})^2} + CX + D \quad (\text{eq. 4.10})$$

Where

X and Y: wavenumber and intensity, respectively.

I: peak intensity

B: Reciprocal of peak FWHM (full width at half maximum)

X_{0i} : peak positions

C, D: fitting constants for modeling background

Figure 64: An example of fitting using equation (eq. 4.10) using the undoped CNT sample data. Blue, red, grey and black curves correspond to D band, G band, linearly tilted base line and entire Raman spectrum of sample, respectively.

Table 10: The non-linear least squares analysis of D and G bands from Raman spectroscopy of as-synthesized MWCNTs

	Undoped	5%P	10%P	5%N	10%N
$I_D \pm 3\sigma$	1970 ± 30	2180 ± 50	4250 ± 70	2140 ± 20	2560 ± 20
$I_G \pm 3\sigma$	3600 ± 30	3560 ± 40	6210 ± 50	3220 ± 20	3210 ± 20
$B_D \pm 3\sigma$ ($\times 10^4$)	2.4 ± 0.1	2.2 ± 0.1	1.0 ± 0.1	1.8 ± 0.1	1 ± 0.1
$X_{0D} \pm 3\sigma$ (cm^{-1})	1368 ± 1	1371 ± 1	1394 ± 1	1368 ± 1	1392 ± 1
$X_{0G} \pm 3\sigma$ (cm^{-1})	1567 ± 1	1558 ± 1	1594 ± 1	1569 ± 1	1598 ± 1
$I_D/I_G \pm 3\sigma$	0.54 ± 0.04	0.61 ± 0.06	0.68 ± 0.05	0.66 ± 0.03	0.80 ± 0.03

Note: σ is standard deviation. Subscripts D and G denote D and G band, respectively.

Raman spectra in the 1200-1800 cm^{-1} region of multiple doped and undoped MWCNTs samples are showed in Figure 63 with peak positions and analysis summarized in Table 10. The first peak is located around 1370 cm^{-1} , denoting disorder-induced in graphitic material D band. The intensity of this peak is strongly related to the impurities and defects present in CNT structure. The feature with the frequency around 1580 cm^{-1} indicates the graphitic G band, which relates to a lattice tangential vibration of CNTs. To determine D and G band frequency, the raw data of each Raman curve is fitted using two Lorentzians and linear baseline. An example illustrated in Figure 64 showing their relative contributions.

Another Raman peak was found in the region 2500-2900 cm^{-1} depicted in Figure 65 with peak positions and analysis summarized in Table 11. This is the second-order Raman peak named G', which locates at $\sim 2700 \text{ cm}^{-1}$. G' band is an overtone mode of the D band due to the frequency close to twice that of the D-band frequency.

Figure 65: Raman Spectroscopy of as-synthesized MWCNTs in the region 2500-2900 cm^{-1}

Table 11: G' bands from Raman spectroscopy of as-synthesized MWCNTs

	Undoped	5%P	10%P	5%N	10%N
$A_{G'} \pm 3 \sigma$ (a.u)	1740 ± 40	1560 ± 60	1740 ± 30	960 ± 90	800 ± 50
$B_{G'} \pm 3 \sigma$ ($\times 10^4$)	4.8 ± 0.3	3.3 ± 0.4	3.4 ± 0.3	3.7 ± 0.2	3.1 ± 0.3
$X_{0G'} (\text{cm}^{-1})$	2710 ± 1	2711 ± 1	2727 ± 1	2710 ± 4	2732 ± 1

It has been well-known that the ratio of intensity of the D over the G band is an important characteristic to examine the defect concentration of CNTs [159-161]. A higher ratio corresponds to lower crystallinity of the tube due to increases of the D band intensity. Data from Table 10 show that this characteristic increases as a function of doping level. Significant increases were observed at doping concentrations 10% for both p-doped and n-doped samples (Figure 63). Compared to undoped CNTs, which has a ratio 0.54 ± 0.04 , 10% n-doped and 10% p-doped counterparts provided relatively higher values of 0.68 ± 0.05 and 0.80 ± 0.03 , respectively. Also, as can be seen, the full half width

at maximum ($\text{FWHM} = 1/B$) of the D band increases with increasing dopant concentration for both n- and p-doping. This value trend was expected since FWHM relates to a measure of the concentration of defects and crystallinity of the tubes [162]. Therefore, these characteristics are a confirmation of impurity insertion of as-synthesized doped CNT samples. Upon the introduction of triphenylborane and benzylamine as p- and n-type doping sources, boron and nitrogen atoms are expected to be impurities, inserting into the lattice system of CNTs. However, actual quantitative doping concentration as well as type of lattice defects needs to be further addressed. Characterization techniques for such information should include X-ray photo electron (XPS) and scanning tunneling electron microscopy (STM).

Interestingly, also from Table 10, upon the introduction of defects into the lattice of CNTs by doping, the band frequencies of doped samples show noticeable shifts in both the D and G bands. Compared to undoped, the Raman frequency up-shift of the D band is more obvious in the case of 10% doping for both p-doped ($\sim 26\text{cm}^{-1}$) and n-doped ($\sim 24\text{cm}^{-1}$). Similarly, for the G band, the up-shift is $\sim 27\text{cm}^{-1}$ and $\sim 31\text{cm}^{-1}$ for 10% p-doped and 10% n-doped, respectively. These observations are consistent with previous Raman studies of p-doped [163, 164] and n-doped [165, 166] CNTs. The up-shift frequency of the D band is a characteristic that indicates the appearance of new types of disorder in doped-CNTs relative to pure CNTs. Meanwhile, the up-shift frequency of G band could be attributed to a modification in the electronic structure of CNTs upon doping [165, 167, 168].

Another most pronounced difference between Raman spectra of undoped- and doped- CNTs was observed through the G' band (Figure 65). Data from our experiments show that the intensity, shape and position of this band are very sensitive to doping level. From Table 11, broader peak and red shift in frequency of G' bands are observed for both n-type and p-type doping. Additionally, there is a strong decrease in intensity ratio between the G and G' bands (Figure 66). This observation is consistent with previous observations in graphene [169] and CNT [165].

Figure 66: The band intensity ratios of D/G and G'/G of p-doped CNTs (left) and n-doped CNTs (right).

4.2.3. Pd nanoparticle attached MWCNTs for hydrogen sensor

A. Attachment of Pd NPs on commercial CNTs

Commercial CNTs used in this experiment had average diameters from 20 -40 nm and length varying from 5-10 μm (Figure 67A). The *in-situ* Pd nanoparticle decorated CNTs were synthesized according to section 2.1.2.3. An SEM image of CNT with attached Pd nanoparticles on commercial CNTs is showed in Figure 67B. It shows 5-10 nm size Pd NPs are attached to CNTs with different densities.

Figure 67: SEM images of commercial CNTs and palladium nanoparticle decorated CNTs.

The mechanism of Pd nanoparticle decoration is presented in Figure 68. In general, the attachment involves two steps including wetting the surface of CNTs by SDS during the sonication and the reduction of palladium acetate to Pd nanoparticles during the reflux process. SDS plays a role as a surface wetting agent, and 1-dodecanol generated from thermal conversion of SDS acts as a reducing agent. Pd-CNTs are well dispersed in aqueous medium by stabilizer components, constituted mainly of 1-dodecanoic acid, 1-dodecanol and unconverted SDS.

Figure 68: Illustrative concept for the Decoration of CNTs with Pd Nanoparticles. This figure is adapted and modified from Ref [31].

B. Attachment of Pd nanoparticles on as-synthesized CNTs

CNTs synthesized by method C using cobaltocene (Figures 69A and 69B) were employed for the attachment of Pd nanoparticles. After the synthesis, hybrid structure of Pd NPs and CNTs is shown in Figures 69C and 69D. As observed, much higher density of Pd nanoparticle attachment compared to its counterpart using commercial CNTs. The walls of CNT are decorated with different sizes of Pd nanoparticles, shown in TEM images (Figure 70). This could be the result of an as-synthesized CNT characteristic, which have more defective sites for the attachment reaction. Higher density of Pd-Nanoparticles turns out to be a better hybrid material for H₂ detection as shown later.

Figure 69: SEM of thin film of CNTs deposited on the wall of glass tube before (A and B) and after (C and D) the attachment of Pd nanoparticles.

Figure 70: Low and high magnification TEM images of as-synthesized CNTs decorated with Pd nanoparticles

4.4 Sensing Application: Hydrogen Gas Sensor Based on Pd NPs Attached MWCNTs

In this section, the sensing performance and mechanism of sensors, using a hybrid structure of Pd nanoparticle attached CNTs, toward hydrogen gas is presented. Similar to the Cl_2 gas sensor, the study is based on measurements of direct current conductivity to establish the sensing model.

4.4.1 Sensing Performance

Initial testing and understanding about the response of Pd-CNTs hybrid structure toward hydrogen was carried out using commercial MWCNTs. The response of sensor fabricated from Pd nanoparticles decorating commercial CNTs is showed in Figure 71. It was observed that the exposure of the as-constructed sensor to continuous flow of 4% of H_2 diluted in nitrogen resulted in a slow increase in sensor resistance. It took about 15 minutes for the sensor to achieve the maximum resistance and about 1.5 hours to recover upon refreshing with air at room temperature.

Figure 71: Response of testing sensor upon the exposure of 4% H_2 with the inset showing the initial stage of H_2 exposure with the flow rate of 1L/minute

Previous studies have showed that attachment of Pd nanoparticles impart remarkable sensitivity on the electrical properties of nanotubes toward molecular hydrogen. With its ability to split hydrogen molecules into hydrogen atoms, Pd nanoparticles play a role as a catalyst in the interaction between H₂ molecules and nanotubes [29, 170]. Specifically, hydrogen molecules split into atomic hydrogen spill-over from Pd nanoparticles to CNTs. This is followed by electron transfer to CNT surface and changes in carrier density leading to changes in the resistance ($R \propto 1/n$, where n is the carrier density) of the CNT. Therefore, as can be seen in Figure 70, the slow response in this case could be a result of a slow electrons transfer rate to nanotubes, caused by low density of Pd nanoparticles attached on MWCNTs. Such response also makes the as-fabricated sensor impractical, and thus a higher density of attached Pd nanoparticles is required to improve sensor performance.

Replacing commercial CNTs by as-synthesized CNTs, a similar response characteristic was obtained. The sensor resistance increased upon the exposure to H₂ and quickly reverses when hydrogen gas is turned off, followed by an introduction of dry air into the testing chamber. However, a much faster response time as well as a higher detection limit was achieved in this case. Response and recovery of the sensor were orders of magnitude faster than what was observed from commercial CNTs. The result of H₂ detection by CNTs, synthesized by the sealed glass tube growth method C following attachment of Pd nanoparticles, is showed in Figure 72. The investigation was carried by exposing the sensor with different H₂ concentrations diluted in N₂ from 10ppm to 4%.

Figure 72: Response of Pd nanoparticles attached as-synthesized CNTs sensor to different H₂ concentrations from 10ppm to 4% (40,000 ppm).

Sensor resistance changed about 200Ω during the response to such a range of H₂ concentrations. During exposures to 10 ppm H₂, the sensor response time was ~ 10 minutes. This time became shorter with higher H₂ concentration. Full sensor recovery time was much quicker than response time, in the range of 30-50 seconds. This result clearly shows that the sensor composed of Pd NPs decorated as-synthesized CNTs provides much better sensor performance compared to its counterpart made using commercial MWCNTs.

To understand the sensor response kinetics, the sensor response curves were fitted to a third order polynomial. Figure 73 shows an example of fitting at 10ppm H₂ concentration.

$$R = a_3t^3 + a_2t^2 + a_1t + R_0 \quad (\text{eq. 4.11})$$

Where:

R is the sensor resistance response during the exposure (Ω)

t is the exposure time

a_1, a_2, a_3 are fitting constants

R_0 is the sensor resistance baseline

Figure 73: An example of fitted curve to 10ppm H₂ concentration following (eq. 4.11)

Instead of using conventional response (the ratio of the change in sensor resistance on exposure to target gas versus original sensor resistance in air), the slopes at inflection point of fitting curves were employed to investigate the relationship between sensor response and exposed hydrogen concentration. These slopes were plotted as a function of H₂ concentration with the data shown in Figure 74.

Figure 74: Slope of the response curve as a function of H_2 concentrations with slope = $26.2 \log [H_2] + 53$. Small black circles represent raw data, and the red line corresponds to fitted data as a linear line.

Figure 74 shows that the slope of the sensor response curve increases approximately linearly with the log of the hydrogen concentration (ppm), slope = 26.2. Since the slope is related to the sensor response time, the increase of slope value is the result of faster sensor response. This has opened a new way to correlate sensor response and target gas concentration, in the case of the magnitude of conventional sensor response (i.e change in resistance) does not vary significantly upon exposure. However, due to logarithmic concentration, dependence of slope makes differentiation of two close exposure concentrations could be difficult. Comparison with commercial CNTs and CNTs synthesized in this study indicate that Pd nanoparticles density is proportional to sensor response time, and thus the slope of response curve. This also could be a factor leading to better sensor performance as observed.

An interesting phenomenon was observed at higher concentrations of H_2 in Figure 75. Specifically, the sensor shows two different response characteristics corresponding to different levels of H_2 concentration exposure (above and below 1% hydrogen concentration). At 1% of H_2 concentration, normal sensing behavior was similar to low H_2 concentrations. Continuous exposures to higher than 1% H_2 in N_2 showed that the sensor resistance still varies in the range of 200Ω , but its initial sharp rise is followed by an abrupt decrease and followed by a stabilization of resistance during the exposure process. During the recovery process by refreshing with air, unusual behavior was also observed. The sensor resistance increased quickly to a certain level of resistance and then went back down to almost the original resistance. Further analysis to understand this behavior will be devoted in following sections.

Figure 75: Abnormal behaviors of sensor response and recovery at high H_2 concentrations.

4.4.2 Understanding Sensing Mechanism:

A. General sensing mechanism

Since the size of the nanotubes is greater than 7 nm, synthesized nanotubes are metallic. The CNT Fermi level as well as the reduction potential of H_2 and O_2 are presented in Figure 76. The underlying mechanism of sensing is attributed to the interaction between target gas (H_2), catalysts (Pd nanoparticles), and nanotubes with a model illustrated in Figure 77. It is well-established that Pd nanoparticles catalytically activate the dissociation of molecular hydrogen [170, 171], and CNTs are p-type semiconductor material in ambient condition due their oxidation (see Figure 76) by atmospheric oxygen [172-174]. Upon adsorption on the surface of Pd nanoparticles, H_2 molecules undergo dissociative chemisorption to split into atomic hydrogen atoms. The resulting atomic hydrogen atoms could either dissolve into Pd with high solubility or diffuse away from receptor sites of Pd and move onto the surface of the CNTs. Since the dissolution of atomized hydrogen into Pd lowers the work function of Pd, [28, 175], spilled-over hydrogen atoms donate electrons to nanotubes [171] through receptor sites or defects of CNTs thereby annihilating holes. This leads to the decrease in charge carrier concentration.

Figure 76: Electrochemical energy level diagram of hydrogen, palladium and oxygen in relation to the Fermi level of CNTs (dashed blue line). Dissolution of hydrogen reduces the Pd work function shifting it upwards as indicated. Donation of electron from the spilled hydrogen atom changes the Fermi level in the same direction as indicated by arrows, reducing the hole concentration in the p-type CNT. Green horizontal lines indicate electrochemical energy states of the analyte where electron/hole transfer could take place

Donation of an electron the from hydrogen atom results in formation of protons on the CNT surface which repels the majority carrier, holes. Consequently, a depletion layer at the CNT interface is established. Reduction in carrier concentration and formation of depletion layer lead to sensor resistance increase.

Figure 77: Illustration of the response mechanism of Pd Nps decorated CNTs upon the exposure with H₂.

In summary, the kinetics of this overall process can be illustrated as:

Therefore, the fitting of each sensor response curve to show sensor resistance as a function of exposure time is based on following equation (eq. 4.12) which reflects decreased hole concentration in CNT as resulting from successive two kinetic steps:

$$R = R_0 \left(1 - \frac{k_1}{k_1 - k_2} (e^{-k_1 t} - e^{-k_2 t}) - e^{-k_1 t} \right) + C \quad (\text{eq. 4.12})$$

Where:

k_1 and k_2 are kinetic constants corresponding to H_2 molecular dissociation by Pd, and electrons transferred to nanotube, respectively.

R_0 is the fitting constant

C is the fitting resistance baseline

Figure 78: Sensor response at 10ppm H_2 concentration: sensor resistance changes as a function of exposure time. Red line and blue line correspond to raw and fitted data, respectively.

Table 12: Summary kinetic constants from fitted response curves

	10ppm	40ppm	400ppm	1000ppm	10000ppm
$k_1 \times 10^2$	1.1 ± 0.1	0.9 ± 0.1	5.4 ± 1	61 ± 5	16000 ± 100
$k_2 \times 10^2$	1.1 ± 0.1	0.9 ± 0.1	4.2 ± 1	9 ± 2	16.2 ± 3
R_0	124 ± 10	180 ± 41	283 ± 30	281 ± 30	285 ± 50

The investigation and curve fitting were based on 5 different concentrations with an example shown in Figure 78, corresponding to 10ppm H₂ exposure. As can be seen, both kinetic constants increase nonlinearly with the increase of H₂ concentration indicating that the other kinetic steps need to be included to properly fit the data. At very low H₂ concentrations such as 10 ppm and 40ppm, these constants seem to be stable at a value around $\sim 10^{-2}\text{s}^{-1}$. They rapidly increase at high H₂ concentrations, especially at 1%, at which the sensor starts showing abnormal response and recovery behaviors. It was observed that the values of k_2 and $\log$ of k_1 increased exponentially as a function of $\log$ of exposed H₂ concentration. These relationships are shown in Figure 79.

Figure 79: H₂ molecular dissociation constant (left) and electrons transferred to nanotube constant (right) change as a function of H₂ concentration: $\log k_1 = 0.03(\log C)^{3.5}$ and $k_2 = 0.4(\log C)^{2.7}$ where C is H₂ concentration (ppm).

Reversibility of sensor resistance during the recovery process with air would involve the effect of O₂. Ultimately O₂⁻ species on the CNT surface can be regenerated by removing hydrogen from the surface. Hydrogen atoms, including dissolved in and spilled over Pd nanoparticles, could combine with oxygen in air along with a withdrawing electron from the conduction band of CNTs, departing the hybrid system in the form of volatile hydrogen di-oxide vapor. Also a hydrogen free surface can

host O_2 and further depletes electrons from the VB as showed in Figure 76. As a result, the p-type nanotube interface is recovered, and the sensor resistance goes back to the original resistance level.

This process can be kinetically represented as:

Thus, each recovery curve can be fitted according to equation below:

$$R = R_0 e^{-k_3 t} + C \quad (\text{eq. 4.13})$$

Where:

R is the sensor resistance

k_3 is the desorption kinetic constant.

$k_{3'}$ is rate of electron abstraction from hydrogen depleted CNTs

R_0 is the fitting constant

C is the fitting resistance baseline

Figure 80: Sensor recovery after 10ppm H_2 exposure: sensor resistance change as a function of recovery time. Red line and blue line correspond to raw and fitted data, respectively.

Table 13: Summary of kinetic constants from fitted recovery curves

	10ppm	40ppm	400ppm	1000ppm	10000ppm
k_3	0.23 ± 0.01	0.11 ± 0.01	0.17 ± 0.01	0.08 ± 0.01	0.08 ± 0.01
R_0	136 ± 4	127 ± 3	248 ± 4	208 ± 5	210 ± 5

From the data, it has indicated that the desorption kinetic constant during the recovery process has a consistent value $\sim 0.1s^{-1}$ and does not depend on hydrogen exposure concentration.

Additionally, as can be seen on Figure 75, the sensor resistance baseline tends to drift to lower resistance after many cycles of exposure and recovery processes. It is believed that this behavior relates to the restoration of $-CH_2$ in CNT structure. As showed in the morphological characterizations, as-synthesized CNTs using this synthetic method are highly defective and heavily oxygenated. Defects could be derived from mostly oxygen functional groups including OH- and $-COOH$ decorating CNT surface and edges [176]. During the exposure, H_2 molecules are dissociated by Pd particles to atomic hydrogen. Via spillover and surface diffusion [177], these species could remove the defective groups in the form of water vapor and CO_2 [178] as well as subsequently restore $-CH_2$ composition. Through many reduction (exposure) and oxidation (recovery) cycles, higher degree of crystallinity CNTs is recovered and leads to defect-free CNTs, decreasing the overall sensor resistance.

B. Sensing mechanism of abnormal response behavior at high concentration exposure:

In order to understand better the abnormal behavior of sensor resistance change, experiment involving a single cycle of exposure to high concentration H_2 and recovery in dry air was carried out as shown in Figure 81.

Figure 81: a) 5 stages of charge carrier inversion process b) Time dependence of excess hole and electron concentration in CNT. Zero on the vertical axis indicates intrinsic CNT. Red and Purple line colors indicate n-type and p-type CNT, respectively.

The experimental data was collected by exposing the sensor to two different hydrogen concentrations and finally refreshing the sensor with a flow of 1% of air in nitrogen. The entire exposure and regeneration cycle of the sensor is illustrated through 5 stages. Initially, low concentration of H₂ 400ppm was used to bring and stabilize the sensor resistivity to maximum response resistance (stage 1). It was then exposed to a high

concentration of 2% H_2 . Such a concentration initially increased sensor resistance and was followed by an abnormal resistance decrease response behavior as mentioned above. The sensor resistance, after reaching the maximum value (stage 2), decreased thereafter to lower level of resistivity and eventually stabilized (stage 3). During the recovery process, 1% of air in N_2 was used instead of air to reduce the rate of recovery reaction. With such refreshing, behavior of this process can be more readily observable. The sensor resistance went back up to almost the same level of maximum resistance (stage 4) followed by a resistance decrease as in normal recovery behavior (stage 5).

The observed abnormal behavior could be rationalized in terms of charge carrier inversion. It assumes that CNTs are converted from p-type to n-type and then back to original the p-type carriers during a cycle of exposure to H_2 and recovery with O_2 in air. The scheme to explain 5 stages of this process is illustrated in Figure 82.

Figure 82: Mechanism of 5 stages of charge carrier inversion process (left) and electronic band diagram of sensing process (right)

Initially, in an ambient environment exposed to air, MWCNTs are p-type semiconductors with respect to the electronic property of the outermost layer. During the first stage of exposure of low H_2 concentration (stage 1), the reduction of CNT by hydrogen brings the hole concentration down (Figure 82 left). Surface bound protons repel the majority carriers away generating a depletion layer. In this process the Fermi level of CNT begins to shift from valance band towards the conduction band due to accumulation of electrons in the VB. This layer is built up during high H_2 concentration exposure in stage 2 due to electron transfer from split hydrogen atoms to CNT the interface to neutralize p-type charge carriers. A maximum resistance is expected when

the excess hole concentration is quenched, corresponding to the shift of CNT Fermi level between VB and CB. Ultimately in stage 3, the value of electron concentration increases along with the increase of high H_2 concentration exposure time. As a result, the Fermi level is shifted close to CB, making CNT n-type. Also, in this stage, the sensor resistance stabilizes at the lowest level resistivity corresponding to saturated electron concentration on surface of nanotube. Ultimately, the Fermi level should lines up with hydrogen redox potential showed in Figure 82 (right).

The reversal of charge carrier inversion occurs in stage 4 during the recovery process. During this stage, O_2 in air react with H^+ on the nanotube surface along with withdrawing electrons from CNT to form volatile dihydrogenoxide molecules. This process reduces and eliminates the excess electron concentration built up during stage 3. The maximum resistance value in this stage is consistent with its counterpart in the stage 2 derived from maximum depletion layer and minimum charge carrier concentration. At this point the Fermi level should be at the center of band gap. When air continued flowing into the sensor in stage 5, O_2 physisorbs on the nanotube wall as well as diffuses into the edge, inducing p-type doping as discussed before. The sensor resistance in this stage decreases due to increase in the hole concentration.

The understanding kinetic of charge carrier inversion and slow recovery process using diluted air is shown in Figure 83 below:

Figure 83: Kinetic of charge carrier inversion and slow recovery using diluted air. The inset shows kinetic fitting curves of stage 3 and 5.

During the charge carrier inversion (stage 3), after reaching maximum depletion thickness corresponding to highest sensor resistance, split atomized hydrogen continues transferring electrons onto nanotube interface. Thus, the interaction and reaction kinetics on the nanotube surface can be represented as:

$$R = R_0 e^{-k_4 t} \quad (\text{eq. 4.14})$$

The rate of electron transfer on the nanotube surface has been reported to be very fast in nanotube based microelectrodes, especially when the electrode has a porous structure and is well wetted by the electrolyte medium. However, it also has been observed that when using individual nanotubes as electrodes, nanotubes do not exhibit good wettability leading to a significant decrease of electron transfer rate [179]. In this

study, the curve fitting followed equation provides the kinetic constant of electron transfer during this stage $\sim 0.0095 \text{ s}^{-1}$. This is considered to be relatively small. In addition, compared to its counterpart (k_2) during the exposure process, this value is much smaller. It could be a result of multiple electron transfer stages as well as electronic structure of nanotube layers [180]. In this case, the electrons spilled over and diffusion on nanotube surface is perhaps the rate determining step accounting for a much slower electron transfer rate. Besides, a higher energy barrier attributed to the formation of depletion layer could also be a reason for significant reduction of this rate constant.

Similarly, reaction between protons and O_2 molecules by refreshing sensor with air results in resistance decrease in stage 5.

$$R = R_0 e^{k_5 t} + C$$

The rate constant of this reaction from curve fitting is $\sim 0.0092 \text{ s}^{-1}$. This value is about 10 fold less than its counterpart (k_3) from previous recovery process using 100% air. Since this constant change as a function of O_2 concentration, they can be represented as:

$$k_3 = K([O_2]_{100\%})^m$$

$$k_5 = K([O_2]_{1\%})^m$$

The calculation using values of k_3 and k_5 provides $m = \sim 1/2$. In other words, the general desorption rate constant of recovery process changes as a function of oxygen concentration and follows: $k_{gen} = K([O_2])^{1/2}$

4.5 Conclusion

The current research project provides a simple glass tube method for synthesis of CNTs. The synthetic method involves a clean chemistry whereby no toxic or combustible hydrocarbon gases are used. The method also provides another way to achieve CNTs by using an easy handling solid carbon feed source and offers a promising way to synthesize doped CNTs. In general, naphthalene derivatives which contain the dopant heteroatoms (e.g. B, N etc) atoms were used as targets for the doping. Raman spectroscopy has showed the insertion of impurities into the CNT lattice leads to shifts of D and G bands as well as D/G ratio change upon doping.

The sensor, constructed from the incorporation of Pd nanoparticles with as-synthesized CNTs, has shown the ability to detect H₂ at ppm levels with a wide dynamic range of detection. The sensing mechanism was based on electron transfer from the interaction between atomized hydrogen and surface of CNTs.

Abnormal sensing behaviors of as-fabricated sensors took place when high H₂ concentrations (>1%) were used. The characteristic of response curves were associated with charge carrier inversion during the exposure process, by which holes on CNT surface were neutralized by electrons donated from split atomized hydrogen atoms, and then further converted p-type to n-type CNTs.

CHAPTER 5. FUTURE WORK

5.1 Future work of SnO₂ NWs

In this study, the sensor response was based on the change of resistance, and fixed frequency (DC conductance) was used throughout all the measurements. In order to gain deeper understanding about how electrons hop between nanowires, frequency dependent electrical impedance studies should be conducted to probe electron movement in nanoparticles and in nanowires. In addition, there is also a need to look at the possibility to detect other oxidizing gases (such as NO₂ and SO₂) using n-doped SnO₂ nanomaterials.

The hybrid structure of Pd nanoparticles and doped SnO₂ nanoparticle/nanowires has been investigated as another material for hydrogen sensors. Preliminary data in Figure 84 shows the detection of this explosive gas at different concentrations at room temperature. The sensors do not display any memory effects observed in sensors, as evidenced by the return of resistance to the baseline after every fresh air purge cycle. The resistance of the sensor quickly decreases upon exposure to hydrogen, followed by a slow increase and slow stabilization and then returns to the initial resistance when exposed to air. Further investigation to understand these interesting response behaviors is needed.

Figure 84: Response of n-doped SnO_2/Pd (right) sensors to different hydrogen concentrations H_2 .

5.2 Future work of CNTs

Even though Raman spectroscopy shows characteristics of impurities inside a doped CNT lattice, further characterization techniques are still needed to confirm the insertion and determine the percentage of the dopants. Such techniques include electron energy loss spectroscopy (EELS) and X-ray photoelectron spectroscopy. In addition, the dopant concentration in doped CNTs can be detected by Time-of-Flight Secondary Ion Mass Spectroscopy (TOF SIMS). Finally, conductivity measurement using a probe station should be performed to compare the undoped and doped CNTs.

Successful detection of H_2 at ppm levels was achieved by the as-constructed sensor. A model to understand the interaction mechanism between exposure gas and the surface of CNTs was established. However, there is still a need to detect and confirm the products of chemical reaction during the recovery process. Besides, in this experiment, the carrier gas was N_2 , although for a practical sensor H_2 diluted in air needs to be

detected. Such studies should be carried out to investigate the sensor response behaviors with oxygen present at high concentration during the exposure.

Interesting charge carrier inversion was observed from the hybrid structure of Pd NPs and CNTs. The study was carried out based on the sensor system, which is composed from multiple CNTs packed together. Further investigation through studies of individual hybrid structures of Pd NPs-CNT are needed to fully isolate effects of electron flow in single vs collective transport in MWCNTs. Specifically, two as-synthesized Pd NPs decorated MWCNTs can be separated from the bulk material and brought together horizontally on a glass substrate partially sputtered with indium tin oxide (ITO). The local electrical measurement could be performed through atomic force microscopy (AFM) associated with the conducting probe tip, which is brought in contact to one of the tubes. With the setup as showed in Figure 85, a stable DC bias voltage can be applied to the device and measure the current flowing through the tip as H_2 with desired concentrations flows into the system. The resulting I-V should be collected and analyzed.

Figure 85: The sketch illustrating experiment for future study of NDR-like behavior using two individuals Pd Np attached CNTs.

The studies as illustrated from Figure 85 would be a good method to confirm charge carrier inversion observation. It also leads to a better understanding how p and n layers are formed within single Pd NPs attached MWCNT during the exposure process, testing the as-proposed mechanism of NDR-like behavior (appendix section D).

References

1. Faraday, M., *The Bakerian Lecture: Experimental Relations of Gold (and Other Metals) to Light*. Phil. Trans. R. Soc. Lond, 1857. **147**: p. 145-181.
2. Alagarasi, A., *Nanomaterials*. VISWANATHAN B ed. 2009.: Narosa Publishing House.
3. Ashby, M.F., P.J. Ferreira, and D.L. Schodek, *Chapter 1 - Nanomaterials and Nanotechnologies: An Overview*, in *Nanomaterials, Nanotechnologies and Design*, M.F.A.J.F.L. Schodek, Editor. 2009, Butterworth-Heinemann: Boston. p. 1-16.
4. Sutariya, V.B. and Y. Pathak, *Biointeractions of Nanomaterials*. 1st ed. 2014, Boca Raton: CRC Press.
5. Harris, P., *Carbon Nanotubes and Related Structures*. 2001, Cambridge University Press: Cambridge.
6. Sirdeshmukh, D. and L. Sirdeshmukh, *Electrical, Electronic and Magnetic Properties of Solids* 1st ed. 2014, New York: Springer.
7. Rumyantseva, M.N., et al., Russian Chemical Bulletin, 2003. **V52**: p. 1217-1238.
8. Zeghbroeck, B.V., *Principles of Semiconductor Devices*. 2011.
9. Amin, G., *ZnO and CuO Nanostructures: Low Temperature Growth, Characterization, their Optoelectronic and Sensing Applications* in *Department of Science and Technology* 2012, Linköping University.
10. Huang, X.-J. and Y.-K. Choi, *Chemical sensors based on nanostructured materials*. Sensors and Actuators B: Chemical, 2007. **122**(2): p. 659-671.

11. Zhou, X.T., et al., *Silicon nanowires as chemical sensors*. Chemical Physics Letters, 2003. **369**(1–2): p. 220-224.
12. Comini, E., et al., *Stable and highly sensitive gas sensors based on semiconducting oxide nanobelts*. Applied Physics Letters, 2002. **81**(10): p. 1869-1871.
13. Barreca, D., et al., *Urchin-like ZnO nanorod arrays for gas sensing applications*. CrystEngComm, 2010. **12**(11): p. 3419-3421.
14. Kong, J., et al., *Nanotube Molecular Wires as Chemical Sensors*. Science, 2000. **287**(5453): p. 622-625.
15. Cullinan, M.A. and M.L. Culpepper, *Carbon nanotubes as piezoresistive microelectromechanical sensors: Theory and experiment*. Physical Review B, 2010. **82**(11): p. 115428.
16. Jarzebski, Z.M. and J.P. Marton, *Physical Properties of SnO Materials*. Journal of The Electrochemical Society, 1976. **123**: p. C199-C205.
17. Batzill, M. and U. Diebold, *The surface and materials science of tin oxide*. Progress in Surface Science, 2005. **79**(2–4): p. 47-154.
18. Chapadraz, A., *Electron Transport and Sensor Characteristics of Tin Dioxide Based Nanocrystalline Materials in Chemistry*. 2009, Portland State University: Portland.
19. Gopel, W., *Chemisorption and charge transfer at ionic semiconductor surfaces: Implications in designing gas sensors*. Progress in Surface Science, 1985. **20**(1): p. 9-103.

20. Hübert, T., et al., *Hydrogen sensors – A review*. Sensors and Actuators B: Chemical, 2011. **157**(2): p. 329-352.
21. Wang, B., et al., *Fabrication of a SnO₂ Nanowire Gas Sensor and Sensor Performance for Hydrogen*. The Journal of Physical Chemistry C, 2008. **112**(17): p. 6643-6647.
22. Katsuki, A. and K. Fukui, *H₂ selective gas sensor based on SnO₂*. Sensors and Actuators B: Chemical, 1998. **52**(1–2): p. 30-37.
23. Ramanathan, M., et al., *Crossover behavior in the hydrogen sensing mechanism for palladium ultrathin films*. Nanotechnology, 2010. **21**(12): p. 125501.
24. Fournier, C., et al., *Discontinuous Palladium Nanostructures for H₂ Sensing*. ECS Meeting 2008. **50**: p. 3147-3147.
25. Maggie, P., et al., *Unprecedented ultra-high hydrogen gas sensitivity in undoped titania nanotubes*. Nanotechnology, 2006. **17**(2): p. 398.
26. Kolmakov, A., et al., *Enhanced Gas Sensing by Individual SnO₂ Nanowires and Nanobelts Functionalized with Pd Catalyst Particles*. Nano Letters, 2005. **5**(4): p. 667-673.
27. Khalap, V.R., et al., *Hydrogen Sensing and Sensitivity of Palladium-Decorated Single-Walled Carbon Nanotubes with Defects*. Nano Letters, 2010. **10**(3): p. 896-901.
28. Kong, J., M.G. Chapline, and H. Dai, *Functionalized Carbon Nanotubes for Molecular Hydrogen Sensors*. Advanced Materials, 2001. **13**(18): p. 1384-1386.

29. Mubeen, S., et al., *Palladium Nanoparticles Decorated Single-Walled Carbon Nanotube Hydrogen Sensor*. The Journal of Physical Chemistry C, 2007. **111**(17): p. 6321-6327.
30. Wang, Y., J.Y. Lee, and T.C. Deivaraj, *Controlled Synthesis of V-shaped SnO₂ Nanorods*. The Journal of Physical Chemistry B, 2004. **108**(36): p. 13589-13593.
31. Karousis, N., et al., *Carbon Nanotubes Decorated with Palladium Nanoparticles: Synthesis, Characterization, and Catalytic Activity*. The Journal of Physical Chemistry C, 2008. **112**(35): p. 13463-13469.
32. Abdula, D., *Functional Surface Chemistry of Carbon-Based Nanostructures*, in *Materials Science and Engineering*. 2011, University of Illinois at Urbana-Champaign: Urbana.
33. Parker, J.H., D.W. Feldman, and M. Ashkin, *Raman Scattering by Silicon and Germanium*. Physical Review, 1967. **155**(3): p. 712-714.
34. Allen, C. and B.R. Shankar, *Room temperature Cl₂ sensing using thick nanoporous films of Sb-doped SnO₂*. Nanotechnology, 2008. **19**(24): p. 245501.
35. Hamberg, I. and C.G. Granqvist, *Evaporated Sn-Doped In₂O₃ Films - Basic Optical-Properties and Applications to Energy-Efficient Windows*. Journal of Applied Physics, 1986. **60**(11): p. R123-R159.
36. Hoffman, R.L., B.J. Norris, and J.F. Wager, *ZnO-based transparent thin-film transistors*. Applied Physics Letters, 2003. **82**(5): p. 733-735.
37. Presley, R.E. and et al., *Tin oxide transparent thin-film transistors*. Journal of Physics D: Applied Physics, 2004. **37**(20): p. 2810.

38. Wager, J.F., *Transparent Electronics*. Science, 2003. **300**(5623): p. 1245-1246.
39. Batley, G.E., A. Ekstrom, and D.A. Johnson, *Studies of topochemical heterogeneous catalysis : 3. Catalysis of the reduction of metal oxides by hydrogen*. Journal of Catalysis, 1974. **34**(3): p. 368-375.
40. Bond, G.C., L.R. Molloy, and M.J. Fuller, *Oxidation of carbon monoxide over palladium-tin(IV) oxide catalysts: an example of spillover catalysis*. Journal of the Chemical Society, Chemical Communications, 1975(19): p. 796-797.
41. Croft, G. and M.J. Fuller, *Water-promoted oxidation of carbon monoxide over tin(IV) oxide-supported palladium*. Nature, 1977. **269**(5629): p. 585-586.
42. Harrison, P.G., C. Bailey, and W. Azelee, *Modified Tin(IV) Oxide (M/SnO₂M=Cr, La, Pr, Nd, Sm, Gd) Catalysts for the Oxidation of Carbon Monoxide and Propane*. Journal of Catalysis, 1999. **186**(1): p. 147-159.
43. Matsui, T., et al., *Effect of reduction-oxidation treatment on the catalytic activity over tin oxide supported platinum catalysts*. Science and Technology of Advanced Materials, 2006. **7**(6): p. 524-530.
44. Solymosi, F. and J. Kiss, *Adsorption and reduction of NO on tin(IV) oxide doped with chromium(III) oxide*. Journal of Catalysis, 1978. **54**(1): p. 42-51.
45. Allegretti, F., et al., *A tin oxide semiconductor sensor for oxygen determination in the sub-ppm range*. Sensors and Actuators B: Chemical, 1993. **10**(3): p. 191-195.
46. Buttà, N., et al., *A family of tin oxide-based sensors with improved selectivity to methane*. Sensors and Actuators B: Chemical, 1992. **6**(1-3): p. 253-256.

47. Cané, C., et al., *Detection of gases with arrays of micromachined tin oxide gas sensors*. Sensors and Actuators B: Chemical, 2000. **65**(1-3): p. 244-246.
48. Chaparadza, A. and S.B. Rananavare, *Room temperature Cl₂ sensing using thick nanoporous films of Sb-doped SnO₂*. Nanotechnology, 2008. **19**(24): p. 245501.
49. Tamaki, J., et al., *CuO-SnO₂ element for highly sensitive and selective detection of H₂S*. Sensors and Actuators B: Chemical, 1992. **9**(3): p. 197-203.
50. Wu, J.M., *A room temperature ethanol sensor made from p-type Sb-doped SnO₂ nanowires*. Nanotechnology, 2010. **21**(23): p. 235501.
51. Baur, W.H., 1956. **9**: p. 515-529.
52. Jarzebski, Z.M., *Oxide Semiconductors*,. 1973, Oxford-Newyork: Pergamon Press.
53. Amma, D.S.D., V.K. Vaidyan, and P.K. Manoj, *Structural, electrical and optical studies on chemically deposited tin oxide films from inorganic precursors*. Materials Chemistry and Physics, 2005. **93**(1): p. 194-201.
54. Kröger, F.A. and H.J. Vink, *Relations between the Concentrations of Imperfections in Crystalline Solids*, in *Solid State Physics*, S. Frederick and T. David, Editors. 1956, Academic Press. p. 307-435.
55. Braithwaite, N. and G. Weaver, *Electronic Materials*. 1990, Oxford: Butterworth-Heinemann.
56. Dieguez, A., et al., *The complete Raman spectrum of nanometric SnO₂ particles*. Journal of Applied Physics, 2001. **90**(3): p. 1550-1557.

57. Merle, P., et al., *Uniaxial-stress dependence of the first-order Raman spectrum of rutile. I. Experiments*. Physical Review B, 1980. **21**(4): p. 1617-1626.
58. Sangeetha, P., V. Sasirekha, and V. Ramakrishnan, *Micro-Raman investigation of tin dioxide nanostructured material based on annealing effect*. Journal of Raman Spectroscopy, 2011. **42**(8): p. 1634-1639.
59. Hu, J.Q., et al., *Laser-Ablation Growth and Optical Properties of Wide and Long Single-Crystal SnO₂ Ribbons*. Advanced Functional Materials, 2003. **13**(6): p. 493-496.
60. Liu, C.M., et al., *Fabrication and characterization of wire-like SnO₂*. Journal of Physics D: Applied Physics, 2006. **39**(12): p. 2494.
61. Kim, H., J. Lee, and C. Lee, *Growth of tin oxide rod-like and sheet-like structures*. Journal of Materials Science: Materials in Electronics, 2009. **20**(2): p. 99-104.
62. Pan, Z.W., Z.R. Dai, and Z.L. Wang, *Nanobelts of Semiconducting Oxides*. Science, 2001. **291**(5510): p. 1947-1949.
63. Mathur, S. and S. Barth, *Molecule-Based Chemical Vapor Growth of Aligned SnO₂ Nanowires and Branched SnO₂/V₂O₅ Heterostructures*. Small, 2007. **3**(12): p. 2070-2075.
64. Chen, Y., et al., *Bulk-quantity synthesis and self-catalytic VLS growth of SnO₂ nanowires by lower-temperature evaporation*. Chemical Physics Letters, 2003. **369**(1-2): p. 16-20.

65. Nguyen, P., et al., *Epitaxial Directional Growth of Indium-Doped Tin Oxide Nanowire Arrays*. Nano Letters, 2003. **3**(7): p. 925-928.
66. Wang, J.X., et al., *Growth of SnO₂ nanowires with uniform branched structures*. Solid State Communications, 2004. **130**(1-2): p. 89-94.
67. Ma, X.L., Y. Li, and Y.L. Zhu, *Growth mode of the SnO₂ nanobelts synthesized by rapid oxidation*. Chemical Physics Letters, 2003. **376**(5-6): p. 794-798.
68. Wang, W., J. Niu, and L. Ao, *Large-scale synthesis of single-crystal rutile SnO₂ nanowires by oxidizing SnO nanoparticles in flux*. Journal of Crystal Growth, 2008. **310**(2): p. 351-355.
69. Wang, Y. and J.Y. Lee, *Molten Salt Synthesis of Tin Oxide Nanorods: Morphological and Electrochemical Features*. The Journal of Physical Chemistry B, 2004. **108**(46): p. 17832-17837.
70. Yu, C., et al., *Growth of single-crystalline SnO₂ nanocubes via a hydrothermal route*. CrystEngComm, 2010. **12**(2): p. 341-343.
71. Yu, S.-H., et al., *A Novel Solventothermal Synthetic Route to Nanocrystalline CdE (E = S, Se, Te) and Morphological Control*. Chemistry of Materials, 1998. **10**(9): p. 2309-2312.
72. Adamson, A.W., *Physical Chemistry of Surfaces*. 1997. 808.
73. Guifu, Z., et al., *Solvothermal/hydrothermal route to semiconductor nanowires*. Nanotechnology, 2006. **17**(11): p. S313.
74. Herring, C., *Some Theorems on the Free Energies of Crystal Surfaces*. Physical Review, 1951. **82**(1): p. 87-93.

75. Venables, J.A., *Introduction to Surface and Thin Film Processes*. 2000, Cambridge: Cambridge University Press. 4.
76. Morin, S.A., et al., *Mechanism and Kinetics of Spontaneous Nanotube Growth Driven by Screw Dislocations*. *Science*, 2010. **328**(5977): p. 476-480.
77. Zemel, J.N., *Theoretical description of gas-film interaction on SnO_x*. *Thin Solid Films*, 1988. **163**: p. 189-202.
78. Sánchez-González, J., et al., *Electrical conductivity of carbon blacks under compression*. *Carbon*, 2005. **43**(4): p. 741-747.
79. Chaparadza, A., S.B. Rananavare, and V. Shutthanandan, *Synthesis and characterization of lithium-doped tin dioxide nanocrystalline powders*. *Materials Chemistry and Physics*, 2007. **102**(2-3): p. 176-180.
80. Jung-Sub, L., S. Vaidyanathan, and D.Y. Jeon. *Formation of Sn-multiwalled carbon nanotube composite layer for the application of thermal interface materials*. in *Electronic Materials and Packaging, 2007. EMAP 2007. International Conference on*. 2007.
81. Majumdar, S., et al., *Room temperature synthesis of nanocrystalline SnO through sonochemical route*. *Materials Letters*, 2008. **62**(8-9): p. 1249-1251.
82. Chan, J.C., et al., *Facile pyrolytic synthesis of silicon nanowires*. *Solid-State Electronics*, 2010. **54**(10): p. 1185-1191.
83. Kim, T.W., D.U. Lee, and Y.S. Yoon, *Microstructural, electrical, and optical properties of SnO₂ nanocrystalline thin films grown on InP (100) substrates*

- for applications as gas sensor devices*. Journal of Applied Physics, 2000. **88**(6): p. 3759-3761.
84. Wang, B., et al., *Field emission and photoluminescence of SnO₂ nanograss*. Journal of Applied Physics, 2005. **98**(12): p. 124303-4.
 85. Tran, H.A. and S.B. Rananavare, *Synthesis and characterization of N- and P-doped tin oxide nanowires*, in *2011 11th IEEE International Conference on Nanotechnology (2011)*. 2011, IEEE: Portland, OR. p. 144-149.
 86. Chopra, K.L., S. Major, and D.K. Pandya, *Transparent conductors—A status review*. Thin Solid Films, 1983. **102**(1): p. 1-46.
 87. Wang, Y., I. Ramos, and J.J. Santiago-Aviles, *Optical bandgap and photoconductance of electrospun tin oxide nanofibers*. Journal of Applied Physics, 2007. **102**(9): p. 093517-5.
 88. Ji, Z., et al., *A novel transparent pn⁺ junction based on indium tin oxides*. Thin Solid Films, 2004. **460**(1-2): p. 324-326.
 89. Ahmed, A.S., et al., *Band gap narrowing and fluorescence properties of nickel doped SnO₂ nanoparticles*. Journal of Luminescence, 2011. **131**(1): p. 1-6.
 90. Johnson, P.L. and D. Teeters, *Formation and characterization of SnO₂ nanobaskets*. Solid State Ionics, 2006. **177**(26-32): p. 2821-2825.
 91. Abello, L., et al., *Structural Characterization of Nanocrystalline SnO₂ by X-Ray and Raman Spectroscopy*. Journal of Solid State Chemistry, 1998. **135**(1): p. 78-85.

92. Gouadec, G.e. and P. Colomban, *Raman Spectroscopy of Nanomaterials: How Spectra Relate to Disorder, Particle Size and Mechanical Properties*. Crystal Growth and Characterization of Materials, 2007. **53**: p. 1-56.
93. Luo, Z.-H., et al., *Controlling the electronic structure of SnO₂ nanowires by Mo-doping*. Chinese Physics B, 2010. **19**(2): p. 026102.
94. Chapadraz, A., *Electron Transport and Sensor Characteristics of Tin Oxide Based Nanocrystalline Material*, in *Chemistry*. 2009, Poatland State University: Portland.
95. Jain, G.H., L.A. Patil, and V.B. Gaikwad, *Studies on gas sensing performance of (Ba_{0.8}Sr_{0.2})(Sn_{0.8}Ti_{0.2})O₃ thick film resistors*. Sensors and Actuators B: Chemical, 2007. **122**(2): p. 605-612.
96. Somorjai, G.A. and Y. Li, *Introduction to Surface Chemistry and Catalysis*. 2010: John Wiley and Sons.
97. Bender, F., et al., *Characterization of a WO₃ thin film chlorine sensor*. Sensors and Actuators B: Chemical, 2001. **77**(1–2): p. 281-286.
98. Zhao, W., et al., *Sensitive properties of In-based compound semiconductor oxide to Cl₂ gas*. Journal of Semiconductors, 2009. **30**(3): p. 034010.
99. Joshi, A., et al., *ZnO-nanowires modified polypyrrole films as highly selective and sensitive chlorine sensors*. Applied Physics Letters, 2009. **94**(10): p. -.
100. Patil, D.R. and L.A. Patil, *Room temperature chlorine gas sensing using surface modified ZnO thick film resistors*. Sensors and Actuators B: Chemical, 2007. **123**(1): p. 546-553.

101. Dawson, D.H. and D.E. Williams, *Gas-sensitive resistors: surface interaction of chlorine with semiconducting oxides*. Journal of Materials Chemistry, 1996. **6**(3): p. 409-414.
102. Chan, J.C., et al., *Mechanisms of Aging of Antimony Doped Tin Oxide Based Electrochromic Devices*. Japanese Journal of Applied Physics, 2006 **45**(49): p. L1300-L1303.
103. Szczuko, D., et al., *XPS investigations of surface segregation of doping elements in SnO₂*. Applied Surface Science, 2001. **179**(1–4): p. 301-306.
104. Egdell, R.G., W.R. Flavell, and P. Tavener, *Antimony-doped tin(IV) oxide: Surface composition and electronic structure*. Journal of Solid State Chemistry, 1984. **51**(3): p. 345-354.
105. McGinley, C., et al., *Pure and Sb-doped SnO₂ nanoparticles studied by photoelectron spectroscopy*. The European Physical Journal D - Atomic, Molecular, Optical and Plasma Physics, 2001. **16**(1): p. 225-228.
106. Dusastre, V. and D.E. Williams, *Sb(III) as a Surface Site for Water Adsorption on Sn(Sb)O₂, and Its Effect on Catalytic Activity and Sensor Behavior*. The Journal of Physical Chemistry B, 1998. **102**(35): p. 6732-6737.
107. Kamble, R.B. and V.L. Mathe, *Nanocrystalline nickel ferrite thick film as an efficient gas sensor at room temperature*. Sensors and Actuators B: Chemical, 2008. **131**(1): p. 205-209.

108. Zhao, X., et al., *Room-temperature chlorine gas sensor based on CdSnO₃ synthesized by hydrothermal process*. Journal of Advanced Ceramics, 2013. **2**(1): p. 31-36.
109. Inderwildi, O.R. and M. Kraft, *Adsorption, Diffusion and Desorption of Chlorine on and from Rutile TiO₂{110}: A Theoretical Investigation*. ChemPhysChem, 2007. **8**(3): p. 444-451.
110. Rochkind, M.M. and G.C. Pimentel, *Infrared Spectrum and Vibrational Assignment for Chlorine Monoxide, Cl₂O*. The Journal of Chemical Physics, 1965. **42**(4): p. 1361-1368.
111. Schack, C.J. and D. Pilipovich, *Chlorine perchlorate*. Inorganic Chemistry, 1970. **9**(6): p. 1387-1390.
112. Christe, K.O., C.J. Schack, and E.C. Curtis, *Halogen perchlorates. Vibrational spectra*. Inorganic Chemistry, 1971. **10**(8): p. 1589-1593.
113. Mueller, H.S.P. and H. Willner, *Vibrational and electronic spectra of chlorine dioxide, OClO, and chlorine superoxide ClOO, isolated in cryogenic matrixes*. The Journal of Physical Chemistry, 1993. **97**(41): p. 10589-10598.
114. Avouris, P., *Carbon nanotube electronics*. Chemical Physics, 2002. **281**(2-3): p. 429-445.
115. Dag, S., et al., - *Electronic structure of the contact between carbon nanotube and metal electrodes*. 2003.
116. Journet, C., et al., *Large-scale production of single-walled carbon nanotubes by the electric-arc technique*. Nature, 1997. **388**(6644): p. 756-758.

117. Fonseca, A. and J.B. Nagy, *Carbon Nanotubes Formation in the Arc Discharge Process*, in *Carbon Filaments and Nanotubes: Common Origins, Differing Applications?*, L.P. Biró, et al., Editors. 2001, Springer Netherlands. p. 75-84.
118. Suzuki, S., *Syntheses and Applications of Carbon Nanotubes and Their Composites*. 2013: InTech.
119. Guo, T., et al., *Self-Assembly of Tubular Fullerenes*. The Journal of Physical Chemistry, 1995. **99**(27): p. 10694-10697.
120. Sinnott, S.B. and R. Andrews, *Carbon Nanotubes: Synthesis, Properties, and Applications*. Critical Reviews in Solid State and Material Sciences, 2001. **26**(3): p. 145-249.
121. Seo, J.W. and et al., *Catalytically grown carbon nanotubes: from synthesis to toxicity*. Journal of Physics D: Applied Physics, 2007. **40**(6): p. R109.
122. Paradise, M. and T. Goswami, *Carbon nanotubes - Production and industrial applications*. Materials & Design, 2007. **28**(5): p. 1477-1489.
123. Mathur, R.B., S. Chatterjee, and B.P. Singh, *Growth of carbon nanotubes on carbon fibre substrates to produce hybrid/phenolic composites with improved mechanical properties*. Composites Science and Technology, 2008. **68**: p. 1608-1615.
124. Schmidt, V., et al., *Silicon Nanowires: A Review on Aspects of their Growth and their Electrical Properties*. Advanced Materials, 2009. **21**(25-26): p. 2681-2702.

125. Doorn, S.K., et al., *Resonant Raman excitation profiles of individually dispersed single walled carbon nanotubes in solution*. Applied Physics A, 2004. **78**(8): p. 1147-1155.
126. Rao, A.M., et al., *Diameter-Selective Raman Scattering from Vibrational Modes in Carbon Nanotubes*. Science, 1997. **275**(5297): p. 187-191.
127. Tuinstra, F. and J.L. Koenig, *Raman Spectrum of Graphite*. The Journal of Chemical Physics, 1970. **53**(3): p. 1126-1130.
128. Dresselhaus, M.S., et al., *Raman spectroscopy on isolated single wall carbon nanotubes*. Carbon, 2002. **40**(12): p. 2043-2061.
129. Jang, Y.-T., et al., *A simple approach in fabricating chemical sensor using laterally grown multi-walled carbon nanotubes*. Sensors and Actuators B: Chemical, 2004. **99**(1): p. 118-122.
130. Dresselhaus, M.S., et al., *Raman spectroscopy of carbon nanotubes*. Physics Reports, 2005. **409**(2): p. 47-99.
131. Ewels, C. (2007) *Doping of Carbon Nanotubes*. **Volume**, DOI: <http://www.ewels.info/science/publications/papers/2008.DopingChapter.pdf>
132. Zhou, O., et al., *Defects in Carbon Nanostructures*. Science, 1994. **263**(5154): p. 1744-1747.
133. Mordkovich, V.Z., et al., *Intercalation into carbon nanotubes*. Carbon, 1996. **34**(10): p. 1301-1303.
134. Nalimova, V.A., et al., *Lithium interaction with carbon nanotubes*. Synthetic Metals, 1997. **88**(2): p. 89-93.

135. Maurin, G., et al., *Electrochemical intercalation of lithium into multiwall carbon nanotubes*. Chemical Physics Letters, 1999. **312**(1): p. 14-18.
136. Hamwi, A., et al., *Fluorination of carbon nanotubes*. Carbon, 1997. **35**(6): p. 723-728.
137. Jin, Z.-x., G.Q. Xu, and S.H. Goh, *A preferentially ordered accumulation of bromine on multi-wall carbon nanotubes*. Carbon, 2000. **38**(8): p. 1135-1139.
138. Stephan, O., et al., *Doping Graphitic and Carbon Nanotube Structures with Boron and Nitrogen*. Science, 1994. **266**(5191): p. 1683-1685.
139. Kim, S.Y., et al., *X-ray Photoelectron Spectroscopy and First Principles Calculation of BCN Nanotubes*. Journal of the American Chemical Society, 2007. **129**(6): p. 1705-1716.
140. Hsu, W.K., et al., *Metallic behaviour of boron-containing carbon nanotubes*. Chemical Physics Letters, 2000. **323**(5-6): p. 572-579.
141. Terrones, M., et al., *N-doping and coalescence of carbon nanotubes: synthesis and electronic properties*. Applied Physics A: Materials Science & Processing, 2002. **74**(3): p. 355-361.
142. Kudashov, A.G., et al., *Influence of Ni–Co Catalyst Composition on Nitrogen Content in Carbon Nanotubes*. The Journal of Physical Chemistry B, 2004. **108**(26): p. 9048-9053.
143. Maldonado, S., S. Morin, and K.J. Stevenson, *Structure, composition, and chemical reactivity of carbon nanotubes by selective nitrogen doping*. Carbon, 2006. **44**(8): p. 1429-1437.

144. Liu, J., S. Webster, and D.L. Carroll, *Highly aligned coiled nitrogen-doped carbon nanotubes synthesized by injection-assisted chemical vapor deposition*. Applied Physics Letters, 2006. **88**(21): p. -.
145. Ruiz-Soria, G., et al., *On the purification of CVD grown boron doped single-walled carbon nanotubes*. physica status solidi (b), 2011. **248**(11): p. 2504-2507.
146. Modi, A., et al., *Miniaturized gas ionization sensors using carbon nanotubes*. Nature, 2003. **424**(6945): p. 171-174.
147. Penza, M., F. Antolini, and M.V. Antisari, *Carbon nanotubes as SAW chemical sensors materials*. Sensors and Actuators B: Chemical, 2004. **100**(1–2): p. 47-59.
148. Peng, N., et al., *Sensing Mechanisms for Carbon Nanotube Based NH₃ Gas Detection*. Nano Letters, 2009. **9**(4): p. 1626-1630.
149. Shu, P. and C. Kyeongjae, *Chemical control of nanotube electronics*. Nanotechnology, 2000. **11**(2): p. 57.
150. Feng, X., et al., *Sensitivity of Ammonia Interaction with Single-Walled Carbon Nanotube Bundles to the Presence of Defect Sites and Functionalities*. Journal of the American Chemical Society, 2005. **127**(30): p. 10533-10538.
151. Robinson, J.A., et al., *Role of Defects in Single-Walled Carbon Nanotube Chemical Sensors*. Nano Letters, 2006. **6**(8): p. 1747-1751.
152. Avouris, P., M. Freitag, and V. Perebeinos, *Carbon-nanotube photonics and optoelectronics*. Nat Photon, 2008. **2**(6): p. 341-350.
153. Andzelm, J., N. Govind, and A. Maiti, *Nanotube-based gas sensors – Role of structural defects*. Chemical Physics Letters, 2006. **421**(1–3): p. 58-62.

154. Chan, J.C., *A novel approach to the synthesis of silicon nanowires*. 2008, Portland State University: Portland.
155. Kumar, M. and Y. Ando, *Chemical vapor deposition of carbon nanotubes: a review on growth mechanism and mass production*. J Nanosci Nanotechnol., 2010. **10**(6): p. 3739-58.
156. Nagaraju, N., et al., *Alumina and silica supported metal catalysts for the production of carbon nanotubes*. Journal of Molecular Catalysis A: Chemical, 2002. **181**(1–2): p. 57-62.
157. Ago, H., et al., *Roles of Metal–Support Interaction in Growth of Single- and Double-Walled Carbon Nanotubes Studied with Diameter-Controlled Iron Particles Supported on MgO*. The Journal of Physical Chemistry B, 2004. **108**(49): p. 18908-18915.
158. Dresselhaus, M.S., et al., *Perspectives on Carbon Nanotubes and Graphene Raman Spectroscopy*. Nano Letters, 2010. **10**(3): p. 751-758.
159. Ceragioli, H.J., et al., *Synthesis and characterization of boron-doped carbon nanotubes*. Journal of Physics: Conference Series, 2008. **100**(5): p. 052029.
160. Maciel, I.O., et al., *Boron, nitrogen and phosphorous substitutionally doped single-wall carbon nanotubes studied by resonance Raman spectroscopy*. physica status solidi (b), 2009. **246**(11-12): p. 2432-2435.
161. Sharifi, T., et al., *Nitrogen doped multi walled carbon nanotubes produced by CVD-correlating XPS and Raman spectroscopy for the study of nitrogen inclusion*. Carbon, 2012. **50**(10): p. 3535-3541.

162. Yadav, R., et al., *Effect of Growth Temperature on Bamboo-shaped Carbon-Nitrogen (C-N) Nanotubes Synthesized Using Ferrocene Acetonitrile Precursor*. Nanoscale Research Letters, 2008. **4**(3): p. 197 - 203.
163. McGuire, K., et al., *Synthesis and Raman characterization of boron-doped single-walled carbon nanotubes*. Carbon, 2005. **43**(2): p. 219-227.
164. Shim, M., et al., *Spectral Diversity in Raman G-band Modes of Metallic Carbon Nanotubes within a Single Chirality*. The Journal of Physical Chemistry C, 2008. **112**(33): p. 13017-13023.
165. Bulusheva, L.G., et al., *Effect of nitrogen doping on Raman spectra of multi-walled carbon nanotubes*. physica status solidi (b), 2008. **245**(10): p. 1971-1974.
166. Fagan, S.B., et al., *Electronic properties of Ag- and CrO₃-filled single-wall carbon nanotubes*. Chemical Physics Letters, 2005. **406**(1-3): p. 54-59.
167. Pisana, S., et al., *Breakdown of the adiabatic Born-Oppenheimer approximation in graphene*. Nat Mater, 2007. **6**(3).
168. Yan, J., et al., *Electric Field Effect Tuning of Electron-Phonon Coupling in Graphene*. Physical Review Letters, 2007. **98**(16): p. 166802.
169. Das, A., B. Chakraborty, and A.K. Sood, *Raman spectroscopy of graphene on different substrates and influence of defects*. Bull. Mater. Sci, 2007. **31**(3).
170. Tierney, H.L., et al., *Hydrogen Dissociation and Spillover on Individual Isolated Palladium Atoms*. Physical Review Letters, 2009. **103**(24): p. 246102.

171. Parambath, V.B., et al., *Investigation of Spillover Mechanism in Palladium Decorated Hydrogen Exfoliated Functionalized Graphene*. The Journal of Physical Chemistry C, 2011. **115**(31): p. 15679-15685.
172. Donghun, K., et al., *Oxygen-induced p-type doping of a long individual single-walled carbon nanotube*. Nanotechnology, 2005. **16**(8): p. 1048.
173. Derycke, V., et al., *Controlling doping and carrier injection in carbon nanotube transistors*. Applied Physics Letters, 2002. **80**(15): p. 2773-2775.
174. Collins, P.G., et al., *Extreme Oxygen Sensitivity of Electronic Properties of Carbon Nanotubes*. Science, 2000. **287**(5459): p. 1801-1804.
175. Mandelis, A. and C. Christofides, *Physics, Chemistry and Technology of Solid State Gas Sensor Devices*. 1993, New York: Wiley.
176. Kaniyoor, A., T.T. Baby, and S. Ramaprabhu, *Graphene synthesis via hydrogen induced low temperature exfoliation of graphite oxide*. Journal of Materials Chemistry, 2010. **20**(39): p. 8467-8469.
177. Nikitin, A., et al., *Hydrogen Storage in Carbon Nanotubes through the Formation of Stable C-H Bonds*. Nano Letters, 2007. **8**(1): p. 162-167.
178. Gao, W., et al., *New insights into the structure and reduction of graphite oxide*. Nat Chem, 2009. **1**(5): p. 403-408.
179. Nugent, J.M., et al., *Fast Electron Transfer Kinetics on Multiwalled Carbon Nanotube Microbundle Electrodes*. Nano Letters, 2001. **1**(2): p. 87-91.
180. Ajayan, P.M., *Nanotubes from Carbon*. Chemical Reviews, 1999. **99**(7): p. 1787-1800.

Appendix A. Chemical reagents

Tin chloride pentahydrate	Sigma-Aldrich, CAS No. 10026-06-9, >99.8%
Sodium borohydride (NaBH ₄)	Sigma-Aldrich, CAS No. 16940-66-2, powder, >98.5%
Sodium chloride	Sigma-Aldrich, CAS No. 7647-14-5, powder, >99%
Potassium chloride	Sigma-Aldrich, CAS No. 7447-40-7, powder, >99%
Lithium chloride	Sigma-Aldrich, CAS No. 7447-41-8, powder, >99%
1,10 Phenanthroline monohydrate	Sigma-Aldrich, CAS No. 5144-89-8, powder, 99%
Naphthalene	Sigma-Aldrich, CAS No. 91-20-3, crystal, 99%
Ferrocene	Sigma-Aldrich, CAS No. 102-54-5, powder, 99%
Cobaltocene	Acros Organic, CAS No. 1277-43-6, 98%
Dimethylformamide (DMF)	Sigma-Aldrich, CAS No. 68-12-2, >99.8%
Antimony trichloride (SbCl ₃)	Acros Organic, CAS No. 10025-91-9, 99.5%
Triphenylborane (TPB)	Sigma-Aldrich, CAS no. 960-71-4, <2% H ₂ O
Benzylamine	Sigma-Aldrich, CAS no. 100-46-9, 99%
Sodium dodecyl sulfate	Sigma-Aldrich, CAS No. 151-21-3, powder, 98%
Palladium acetate	Sigma-Aldrich, CAS No. 3375-31-3, powder, 98%

Appendix B. Characterization of SnO₂ NWs at different synthetic conditions

The morphology of SnO₂ NWs was investigated at different synthetic conditions including temperature, environment and the presence of surfactant, 1,10 phenanthroline. In this study, the syntheses were carried out using method C (section 2.7.1.1). SEM was employed for the characterization, examining diameter and length of SnO₂ NW formation.

Size histograms of diameter and length of nanowires were analyzed based on standard Gaussian distribution function. The results are showed in the following table B.1

$$G(x) = a \cdot e^{-b(x-c)^2} + d$$

Where

c is the fitting parameter (length or diameter)

a, b, d are fitting constants

Table B.1: The effect of synthetic conditions on SnO₂ NWs morphology (O and NO stand for observed and non-observed, respectively)

Synthetic condition of SnO ₂ NWs	Length (nm)	Gaussian Distribution (b x 10 ⁶)	Diameter (nm)	Gaussian Distribution (b x 10 ⁶)
SnO _x nps + N ₂ + 490C	NO	NO	NO	NO
SnO _x nps + N ₂ + Phen + 490°C	O	0	20±2	47±1
Dilute SnO + 490°C	925	80±5	125±6	300±50
Sn nps + N ₂ + 490°C	240±3	81±9	45.8±5	210±20
Sn nps + Air+ 490°C	292±3	400±100	21.3±3	2100±300
Sn+ Air+ Phen+490°C	407±4	81±9	129±3	210±20
Sn nps + Air + 700°C	800±20	12±4	72±2	120±20

In the Gibbs nucleation model, the term b is related to the surface energy of nucleating cluster. Therefore, the surface free energies of the clusters corresponding to two dimensions (length and diameter) can be estimated.

$$b \sim \frac{4\pi\gamma}{3k_B T}$$

Table B.2: Calculation of the surface free energies of cluster nucleation

Synthetic condition of SnO ₂ NWs	Gaussian Distribution of length (b x 10 ⁶)	γ_L (Nm ⁻¹) ($\gamma \times 10^{26}$)	Gaussian Distribution of diameter (b x 10 ⁶)	γ_D (Nm ⁻¹) ($\gamma \times 10^{26}$)
SnO _x nps + N ₂ + Phen + 490°C	0	0	47±1	11±1
Dilute SnO + 490°C	80±5	20±1	300±50	75±1
Sn nps + N ₂ + 490°C	81±9	20±2	210±20	52±5
Sn nps + Air+ 490°C	400±100	100±3	2100±300	528±75
Sn+ Air+ Phen+490°C	81±9	20±2	210±20	210±5
Sn nps + Air + 700°C	12±4	4±1	120±20	120±5

Figure B1: SnO₂ NWs synthesized from SnO_x NPs at 490°C in an N₂ environment with the presence of 1,10 phenanthroline. Nanowires with small length were observed from this synthetic condition, but not fitted by Gaussian distribution.

Figure B2: SnO_2 NWs synthesized from SnO_x NPs at 490°C in air environment without 1, 10 phenanthroline along with Gaussian distributions of length and diameter. In this synthetic condition, the ratio between precursors SnO_x NPs and mixture of LiCl/KCl was decreased 10 times (0.02g of SnO_x NPs was mixed with 0.31g of LiCl and 0.6g of KCl, see method C, section 2.7.1.1)

Figure B3: SnO₂ NWs synthesized from Sn NPs at 490°C in N₂ environment without 1,10 phenanthroline along with Gaussian distributions of length and diameter

Figure B4: SnO_2 NWs synthesized from Sn NPs at 490°C in air environment without 1,10 phenanthroline along with Gaussian distributions of length and diameter

Figure B5: SnO_2 NWs synthesized from Sn NPs at 490°C in air environment with the presence of 1,10 phenanthroline along with Gaussian distributions of length and diameter

Figure B6: SnO_2 NWs synthesized from Sn NPs at 700°C in air environment without the presence of 1,10 phenanthroline along with Gaussian distributions of length and diameter

Appendix C. Synthesis and characterization of heterojunctions of silicon nanowires (SiNWs) and carbon nanotubes (CNTs)

The aim of this study was to grow heterojunctions of SiNWs and CNTs using a low cost and versatile synthetic method. This includes the attachment of Au nanoparticles on functionalized MWCNTs using the ion exchange method. The hybrid structure of Au NPs/CNTs was used as a template to synthesize SiNWs, employing a novel synthetic method proposed by Chan etc (Chan, 2010 #70). Heterostructure formation of SiNWs/CNT was characterized by electron microscopy (SEM and TEM).

Figure C.1: Schematic showing synthesis steps of heterostructures of SiNWs and CNTs.

C.1. High density attachment of Au nanoparticles on amine functionalized MWCNTs.

0.01g of amine functionalized MWCNTs (nano-lab, catalog NO. PD30L5-20-NH₂) was dispersed in 20ml DI H₂O by sonication and added by 0.5ml of Au colloid solution (Tedpella, catalog NO. 15702-20). The mixture was stirred for 24 hours at room temperature to obtain MWCNT-Au NPs hybrid structures. The black solid was collected and fully washed with DI water and ethanol for several times by centrifugation at 10,000 rpm. The final black precipitate was dried at 60 °C in a vacuum oven overnight. To

prepare the stock solution for further syntheses, 0.001 g of this precipitate was dispersed in 20 ml of water and followed by sonication.

Results

Figure C.2: SEM (left) and TEM (right) images of hybrid structure Au NPs /CNTs. The inset of TEM image show high density of Au NPs with the average diameter 5 nm attached on CNTs.

C.2. SiNW/CNT heterojunction growth

Au NPs attached MWCNTs were used as a template to grow heterostructures of CNT and SiNW. The synthesis was carried out using the dual chamber synthetic method (Chan, 2008 #142). One drop of the above MWCNT-Au NPs stock solution was pipetted on a silicon substrate, followed by heating at 100 °C to completely evaporate the solvent. The substrate was inserted into the dual chamber glass reaction tube, which contains 7 μ L of diphenyl silane at the bottom. The tube was then sealed with a propane torch to form an approximate 13 cm reaction tube. Similarly, the reaction was carried out at 500 °C, using a custom-built glass furnace made from heating tape.

Results

Figure C.3 shows a SEM image of the as grown heterojunctions of SiNWs/CNTs. It also can be seen that a high density of SiNW growing from the sides of CNTs with Au NPs still attached on nanowire tips. Further characterization of heterostructure was obtained by HRTEM shown in Figure C.3 (right). From this, lattice fringes of SiNWs can be observed, confirming the formation of single crystalline SiNWs. More importantly, from the inset of Figure C.3 (right), it reveals the nucleation of heterojunctions of the hybrid structure, where SiNW with Au NP on the tip nucleated from the wall of CNT.

Figure C.3: SEM (left) and HRTEM (right) reveal the heterojunctions of SiNWs and CNTs

Appendix D. Negative differential resistance observed from Pd NPs attached CNTs.

Further characterization of abnormal response behavior of the sensor toward H₂ gas was carried out using the I-V characteristic. The sensor current was monitored as a function of applied voltage during H₂ gas exposure process. In the experiment, the bias was applied and current was collected after introducing H₂ gas into sensor testing system.

Results

Figure D.1: Raw data of I-V curves showing abnormal behavior at different H₂ concentration exposures.

The current from raw data in Figure D.1 was normalized as:

$$I' = \frac{V}{\frac{V}{I} - \alpha R_0}$$

Where

I' is the normalized current

α is the normalization coefficient

R_0 is sensor resistance in air

Figure D.2: Normalization of I' -V curves from Figure D.1 where $\alpha = 0.95$.

Figure D.2 shows the sensor current reading at different H_2 gas concentration exposures when the bias was applied from -0.1V to 0.5V at the rate of 0.01V/s. The sensor current experienced three different responses throughout such an applied voltage range. From -0.1V to ~ 0.17 V, Ohmic resistance of the sensor was achieved. However, when the bias reached 0.17V, negative differential resistance-like behavior was observed. At this point, the voltage across and the current through the sensor changed in opposite direction. An increase in voltage from 0.17V to 0.22V caused a decrease in current from ~ 0.6 mA to ~ 0.48 mA. After 0.22V, normal Ohmic characteristic of sensor resistance was recovered. It also can be seen from Figure D.2, such behavior is more noticeable at high H_2 concentration exposures, especially above 1% where charge carrier inversion was observed (Chapter 4, section 4.4.2).

It was believed that this phenomenon is similar to negative differential resistance (NDR), which is a characteristic of a tunnel diode. Therefore, the data analysis was based on the S.M Sze model, by which the NDR region was focused and analyzed similarly to Figure D.3 below:

Figure D.3: Current-voltage characteristic of NDR behavior. I_P and V_P are the peak current and peak voltage, respectively. I_V and V_V are the valley current and valley voltage, respectively.

Figure D.3: Experimental I-V curves retrieved from Figure D.2. The curve data from 2% and 3% H₂ exposures were achieved from NDR-like behavior region and normalized by shifting to X0Y coordinate.

According to the S.M Size model, the total current in negative differential resistance behavior is the sum of tunneling, excess and diffusion currents. These currents are represented by following equations:

$$I_{tunnel} = I_p \left(\frac{V}{V_p} \right) \exp \left(1 - \frac{V}{V_p} \right)$$

$$I_{excess} = I_0 \exp \left(\frac{q}{KT} V \right)$$

$$I_{diffusion} = I_v \exp(B(V - V_v))$$

Where I_p , I are the peak and valley current peak. V_p and V_v are the peak and valley voltage. I_0 and B are the excess current constant and diffusion current coefficient, respectively.

Therefore:

$$I_{Total} = I_P \left(\frac{V}{V_P} \right) \exp \left(1 - \frac{V}{V_P} \right) + I_0 \exp \left(\frac{q}{KT} V \right) + I_V \exp(B_2(V - V_V)) \quad (\text{eq. D.1})$$

Table D.1: Fitting parameters following equation D.1

Fitting parameter	Value
I_P	0.07 (mA)
V_P	25mV
I_V	0.005 (mA)
V_V	75 (mV)
B	$31.2 \left(\frac{1}{V} \right)$
I_0	78 aA
KT	25.8 eV

Figure D.4: I' -V curve from fitted data following eq. D.1. NDR-like behavior was observed with 2% and 3% H_2 concentration exposures.

It was believed that such observed NDR-like behavior relates to the formation of p-n junctions, derived from charge carrier inversion. As discussed in chapter 4, under high H_2 concentration exposure and the H_2 molecule splitting process by Pd NPs, MWCNTs are converted from p-type to n-type semiconducting material, with respect to their outermost layers. In addition to the formation of a depletion layer, p-type

semiconducting layers still remain inside the tube core. As a result, a single MWCNT is separated into three portions, which play a role as a p-n junction of a tunnel diode.

Figure D.5: Model illustrating formation of a p-n junction in a single MWCNT along with the tube outermost layer band diagrams. The tube converted from p-type to n-type semiconducting material with respect to the outermost layers, but still remains the p-type portion inside the core. The generation of a depletion layer leads to the formation of a p-n junction.

As can be seen from the model above, both p and n sides are considered to be degenerate. When a bias is applied the electrons can tunnel from the n-side to the other due to available unoccupied states in p-side. This causes the increase of current and decrease of unoccupied state concentration. When reaching the peak value at I_p , if the voltage is further increased, there are fewer available unoccupied states on the p-side leading to the current decrease, and eventually this tunnel current stops flowing. As still increasing the bias, the normal thermal current will flow along with diffusion current.

Figure D.6. Band diagram of a tunnel diode with E_C , E_V and E_F are the conduction band, valence band and Fermi level, respectively.

- a) At zero bias, there is no current flow
- b) With small bias applied, electrons in the conduction band of the n-side tunnel to available states of the valence band in the p-side
- c) Increasing the voltage leads to maximum tunnel current, corresponding to a majority of electrons in the n-side equal to that of available states in the p-side
- d) Further increasing bias, the current decreases due to the number of electrons in the n-side decreasing
- e) As higher voltage is applied, the tunnel current drops to zero. Regular diode current (thermal and diffusion currents) takes place.