

# **UNIVERSITI PUTRA MALAYSIA**

# FEASIBILITY STUDIES OF POLYANILINE NANOSTRUCTURES COATED ON TAPERED OPTICAL FIBER FOR AMMONIA SENSING

SITI AZLIDA BINTI IBRAHIM @ GHAZALI

FK 2017 21



# FEASIBILITY STUDIES OF POLYANILINE NANOSTRUCTURES COATED ON TAPERED OPTICAL FIBER FOR AMMONIA SENSING

By

SITI AZLIDA BINTI IBRAHIM @ GHAZALI

Thesis Submitted to the School of Graduate Studies, Universiti Putra Malaysia, in Fulfilment of the Requirements for the Degree of Doctor of Philosophy

January 2017

# COPYRIGHT

All material contained within the thesis, including without limitation text, logos, icons, photographs and all other artwork, is copyright material of Universiti Putra Malaysia unless otherwise stated. Use may be made of any material contained within the thesis for non-commercial purposes from the copyright holder. Commercial use of material may only be made with the express prior, written permission of Universiti Putra Malaysia.

Copyright © Universiti Putra Malaysia



Abstract of the thesis presented to the Senate of Universiti Putra Malaysia in fulfilment of the requirement for the degree of Doctor of Philosophy

# FEASIBILITY STUDIES OF POLYANILINE NANOSTRUCTURES COATED ON TAPERED OPTICAL FIBER FOR AMMONIA SENSING

By

## SITI AZLIDA BINTI IBRAHIM @ GHAZALI

January 2017

Chairman Faculty :

:

Mohd. Adzir Mahdi, PhD Engineering

Polyaniline (PANI) has been used for ammonia (NH<sub>3</sub>) sensing for several decades, mostly in electrical based sensors due to its variation in conductivity during the interaction. Optical sensors are known to have advantages in certain aspects as compared to electrical sensors, but this area of research has not been fully explored. Therefore, this research project aims to explore optical based NH<sub>3</sub> sensor with the incorporation of PANI nanostructure. Tapered multimode fiber (MMF) was chosen as the transducing platform for the sensor because of ease in fabrication, high sensitivity and its suitability for remote sensing applications. The sensitivity of tapered fiber based sensors can be improved by reducing the waist diameter. The influence of PANI nanostructure morphology, thickness and dopants on the sensing performance was also studied in this thesis.

In this project, PANI nanostructures were synthesized and deposited on tapered MMF using two methods. The first method was in-situ deposition method, where poly(methyl vinyl ether-alt-maleic acid)(PMVEA)-doped PANI (PANI-PMVEA) nanogranules and nanofibers were grown on tapered MMF surface during polymerization process. For the second method, camphorsulfonic acid (CSA)-doped PANI (PANI-CSA) nanofibers were spray coated on the tapered MMF. Three processes were involved in the preparation of PANI-CSA, namely the synthesis of hydrochloric acid-doped PANI (PANI-HCl), the dedoping of PANI-HCl to obtain emeraldine base PANI (PANI-EB) powder and the redoping of PANI-EB with CSA to obtain PANI-CSA nanostructures. The thickness and morphology of PANI nanostructures were characterized using scanning electron microscopy (SEM) and atomic force microscopy (AFM). The formation of PANI nanostructures in emeraldine salt form was confirmed from molecular vibrational analysis using Raman Spectroscopy and Fourier Transform Infrared Spectroscopy (FTIR). The response of the sensors towards NH<sub>3</sub> at different concentration was measured using cumulative absorbance change within the wavelength range of 500 - 800 nm. The sensing performance was evaluated in terms of response time, recovery time, sensitivity and repeatability.

PANI-PMVEA coating thickness and morphology was varied by varying the deposition duration and PMVEA/aniline ratio. The highest sensitivity was achieved by the most uniformly-distributed nanogranules PANI-PMVEA coating with thickness of approximately 913 nm. The achieved sensitivity, response and recovery time was 2.19/vol%, 2.82 minutes and 11.52 minutes, respectively. PANI-CSA nanofibers were coated on tapered MMF with different waist diameters. The highest sensitivity of 2.44/vol% was attained by the smallest diameter (20  $\mu$ m) sensor with PANI-CSA coating of approximately 700 – 850 nm. The response and recovery time are 1.73 minutes and 12 minutes, respectively. The sensor using PANI-CSA nanofibers have higher sensitivity and faster response than the sensor using PANI-PMVEA nanostructures at 1% NH<sub>3</sub>.



Abstrak tesis yang dikemukakan kepada Senat Universiti Putra Malaysia sebagai memenuhi keperluan untuk ijazah Doktor Falsafah

# KAJIAN KELAYAKAN POLYANILINE BERSTRUKTUR NANO DI SALUT PADA FIBER OPTIK TIRUS SEBAGAI PENDERIA AMMONIA

Oleh

# SITI AZLIDA BINTI IBRAHIM @ GHAZALI

Januari 2017

Pengerusi Fakulti :

:

Mohd. Adzir Mahdi, PhD Kejuruteraan

Polianilina (PANI) telah digunakan untuk penderiaan ammonia (NH<sub>3</sub>) selama beberapa dekad, kebanyakannya dalam penderia berasaskan elektrik disebabkan oleh konduktivitinya yang berubah-ubah ketika bertindak balas dengan gas NH<sub>3</sub>. Penderia optik diketahui mempunyai kelebihan dalam aspek tertentu berbanding dengan penderia elektrik tetapi masih belum diterokai secara meluas. Oleh itu, projek penyelidikan ini bertujuan untuk meneroka penderia NH<sub>3</sub> berasaskan optik dengan menggabungkan nano struktur PANI. Gentian berbilang mod (MMF) tirus dipilih sebagai platform transduksi untuk penderia kerana mudah direka, kepekaan yang tinggi dan sesuai untuk penggunaan penderiaan jauh. Kepekaan penderia berasaskan gentian tirus boleh ditingkatkan dengan mengecilkan diameter tengah. Pengaruh morfologi, ketebalan dan bahan dop nano struktur PANI terhadap prestasi penderiaan juga dikaji dalam tesis ini.

Dalam projek ini, nano struktur disintesiskan dan terendap pada MMF tirus menggunakan dua kaedah. Kaedah pertama ialah kaedah pengendapan in-situ iaitu poli(metil vinil eter-alt-asid maleik) (PMVEA) terdop granul nano dan gentian nano PANI (PANI-PMVEA) terhasil pada permukaan tirus MMF semasa proses pempolimeran. Bagi kaedah kedua, asid kamforsulfonik (CSA) terdop gentian nano PANI (PANI-CSA) disembur menyaluti MMF tirus. Tiga proses yang terlibat dalam penyediaan PANI-CSA, iaitu sintesis asid hidroklorik terdop PANI (PANI-HCl), pengedopan PANI-HCl untuk mendapatkan serbuk PANI-EB dan pengedopan semula PANI-EB dengan CSA untuk mendapatkan struktur nano PANI dalam bentuk garam yang berwarna hijau zamrud disahkan melalui analisis getaran molekul menggunakan Spektroskopi Raman dan Spektroskopi Inframerah Transformasi Fourier. Tindak balas penderia terhadap NH<sub>3</sub> pada kepekatan berbeza diukur menggunakan perubahan

keserapan kumulatif dalam julat jarak gelombang 500-800 nm. Prestasi penderiaan dinilai dari segi masa tindak balas, masa pemulihan, kepekaan dan keterulangan.

PANI-PMVEA Ketebalan dan morfologi salutan dipelbagaikan dengan mempelbagaikan masa pengendapan dan nisbah PMVEA\anilina. Kepekaan tertinggi dicapai dengan taburan granul nano salutan PANI-PMVEA paling sekata dengan ketebalan kira-kira 913 nm. Kepekaan, tindak balas dan masa pemulihan yang dicapai adalah masing-masing 2.19/vol%, 2.82 minit dan 11.52 minit. Gentian nano PANI-CSA disalutkan pada MMF tirus dengan diameter tengah yang berbeza. Kepekaan tertinggi ialah 2.44/vol% dicapai dengan penderia diameter terkecil (20 µm) dengan salutan PANI-CSA kira-kira 700 – 850 nm. Masa tindak balas dan pemulihan adalah masing-masing 1.73 minit dan 12 minit. Penderia yang menggunakan gentian nano PANI-CSA mempunyai kepekaan lebih tinggi dan tindak balas lebih cepat berbanding dengan penderia yang menggunakan nano struktur PANI-MVEA pada 1% NH<sub>3</sub>.



### ACKNOWLEDGEMENTS

First and foremost, all praise to Allah, the Almighty God for His mercy in giving me the good health, strength, patience and guidance to complete this PhD journey. Secondly, a special thanks to my main supervisor, Prof. Dr. Mohd. Adzir Mahdi, for his invaluable support, guidance, encouragement and intellectual input in conducting this research. His trust and confidence on me motivated me to keep going. A special gratitude to all my co-supervisors Dr. Mohd. Hanif Yaacob, Dr. Norizah Abdul Rahman, and Dr. Muhammad Hafiz Abu Bakar for their guidance, advice and contributions towards the success of this research.

I also would like to take this opportunity to express my sincerest love to my husband, Armizy Abdullah Shukri and my two lovely sons, Khairul Anwar Najmi bin Armizy and Khairul Akmal Badri bin Armizy for their unconditional love, support and patience. I thank my beloved parents, Ibrahim Ismail and Siti Asiah Abdullah and for their encouragements and prayers. A special thanks also goes to my parents in law, Abdullah Shukri Haji Yusoff and Norizan Mohd Noor for their supports and love.

I wish to thank all my colleagues and friends at the Photonic Lab and Chemistry Lab for their help in providing a friendly and inspiring environment to conduct research.

I would also like to thanks Ministry of Higher Education Malaysia for awarding me the PhD scholarship. My thanks extended to my colleagues in Faculty of Engineering, Multimedia University for their friendship and encouragements. This thesis was submitted to the Senate of Universiti Putra Malaysia and has been accepted as fulfilment of the requirement for the degree of Doctor of Philosophy. The members of the Supervisory Committee were as follows:

## Mohd Adzir Mahdi, PhD Professor Faculty of Engineering Universiti Putra Malaysia (Chairman)

# Mohd Hanif Yaacob, PhD Lecturer Faculty of Engineering Universiti Putra Malaysia (Member)

### Norizah Abdul Rahman, PhD

Senior Lecturer Faculty of Science Universiti Putra Malaysia (Member)

## Muhammad Hafiz Abu Bakar, PhD

Senior Lecturer Faculty of Engineering Universiti Putra Malaysia (Member)

### **ROBIAH BINTI YUNUS, PhD**

Professor and Dean School of Graduate Studies Universiti Putra Malaysia

Date:

# **Declaration by Members of Supervisory Committee**

This is to confirm that:

 $\bigcirc$ 

- the research conducted and the writing of this thesis was under our supervision;
- supervision responsibilities as stated in the Universiti Putra Malaysia (Graduate Studies) Rules 2003 (Revision 2012-2013) are adhered to.

Signature: Name of Chairman of Supervisory Committee:	Prof. Dr. Mohd Adzir Mahdi
Signature: Name of Member of Supervisory Committee:	Dr. Mohd. Hanif Yaacob
Signature: Name of Member of Supervisory Committee:	Dr. Norizah Abdul Rahman
Signature: Name of Member of Supervisory Committee:	Dr. Muhammad Hafiz Abu Bakar

# TABLE OF CONTENTS

			Page
AB AC AP DE LIS LIS	STRACT STRAK KNOWLED PROVAL CLARATIO ST OF TABL ST OF FIGU ST OF ABBF	N JES	i iii v vi viii xiii xiii xvii
СН	IAPTER		
1	INTRODU	ICTION	1
1		rerview	1
		blem Statement	3
		jectives	4
		ope and Limitation	5
		esis Organisation	6
2	LITERAT	URE REVIEW	7
-		roduction	7
		tical-based Techniques in Gas Sensing	, 7
		pered Optical Fiber-based Gas Sensors	12
		sic Properties of Conducting Polymers	14
		nducting Polymer Gas Sensors	17
		lyaniline Nanostructure Properties and Synthesis	19
		ethods	
	2.7 Re	view of Ammonia Sensors based on Polyaniline	23
	2.7	1.1 Electrical-based Ammonia Sensors using	23
		Polyaniline	
	2.7	2. Optical-based Ammonia Sensors using Polyaniline	25
	2.8 Int	roduction to Characterization Techniques used in this	27
	Th	esis	
	2.8		27
	2.8	1 17	28
		8.3 Fourier Transform Infrared Spectroscopy (FTIR)	28
		3.4 Atomic Force Microscopy	29
	2.9 Su	mmary	30
3	AMMONI	A SENSOR DEVELOPED BY IN-SITU	31
	DEPOSIT	ION OF PMVEA-DOPED POLYANILINE	
		RUCTURES ON TAPERED MMF	
		roduction	31
		nsor Fabrication	31
	3.2	1	31
	3.2	1	33
		Tapered MMF	

3.3	Sensor Response Measurement	36
3.4	PANI-PMVEA Characterization Results	38
	3.4.1 PANI-PMVEA Layer Thickness	38
	3.4.2 PANI-PMVEA Morphology	42
	3.4.3 Raman Spectrum of PANI-PMVEA	46
	3.4.4 FTIR Spectrum of PANI-PMVEA	47
3.5	Sensing Results	48
	3.5.1 Ammonia Sensing Response of the Sensors	49
	Prepared from Solution with PMVEA/Aniline	
	Ratio of 33%	
	3.5.2 Ammonia Sensing Response of the Sensors	55
	Prepared from Solution with PMVEA/Aniline	
	Ratio of 27%	
	3.5.3 Sensing Mechanism	58
3.6	Summary	58
4 AMN	IONIA SENSOR DEVELOPED BY SPRAY COATING	59
OF	CSA-DOPED POLYANILINE NANOFIBERS ON	
	ERED MMF	
4.1	Introduction	59
4.2	Sensor Fabrication	60
	4.2.1 Synthesis of PANI-EB Powder	60
	4.2.2 Preparation of PANI-CSA Solution	63
	4.2.3 Deposition of PANI-CSA Nanofibers on Tapered	64
	MMF	01
4.3	PANI-CSA Characterization Results	65
1.5	4.3.1 PANI-CSA Morphology	65
	4.3.2 PANI-CSA Thickness and Surface Roughness	67
	4.3.3 Raman Spectrum of PANI-CSA	68
	4.3.4 FTIR Spectrum of PANI-CSA	69
4.4	Sensing Results	70
4.5	Summary	76
4.5	Summary	70
5 CON	CLUSION AND FUTURE WORKS	77
5 0010	Conclusion	77
5.2	Future Works	78
5.3	Outcome and Author's Achievements	78 79
5.5	Outcome and Author's Achievements	13
REFEREN	ICES	80
APPENDI		80 87
	A OF STUDENT	87 88
LIST OF P	UBLICATIONS	89

xi

# LIST OF TABLES

Table	2	Page
3.1	Summary results on the influence of deposition duration on PANI-	41
	PMVEA coating thickness	
3.2	Band assignment of the Raman spectrum of PANI [92][60]	47
3.3	Summary results of sensor response towards NH <sub>3</sub> at 1% for sensor	53
	A6, A2 and A1.	
4.1	Response, recovery time and sensitivity for sensors with different	74
	diameters as the sensor was exposed to 1% ammonia.	
	-	



# LIST OF FIGURES

Figure	;	Page
2.1	Absorption spectra for five gases in the Mid-IR region [35].	8
2.2	Experimental setup for hydrogen sensor using WO <sub>3</sub> coated transparent substrate [37].	9
2.3	Evanescent field at the core cladding interface of an optical fiber	10
2.4	[48]. Modified optical fibers. (a) Side-polished fiber, (b) tapered fiber,	11
	and (c) D-shaped fibers [28].	
2.5	Illustration of tapered optical fiber.	12
2.6	Basic chemical structure of (a) PPy, (b) PTh and (c) PANI.	14
2.7	Energy level diagram of conducting polymer, showing the band gap and energy levels of its different oxidation states [60].	
2.8	Oxidative doping of PPy [16]	16
2.9	Protonic acid doping of PANI [16]	17
2.10	General structure of PANI. (a) Benzenoid unit, (b) Quinonoid unit, (c) PANI repeat unit	19
2.11	Chemical structure of PANI at different oxidation states [60].	20
2.12	Absorbance spectrum of PANI-EB (dedoped) and PANI-ES (doped) films [76].	21
2.13	Reaction between PANI-ES with NH <sub>3</sub> [66].	22
2.14	Schematic diagram of SEM [86].	27
2.15	AFM operation principle [88].	29
3.1	Flow chart of experimental work.	31
3.2	Vytran GPX-3400 Optical Glass Fiber Processor.	32
3.3	GUI of Vytran GPX-3400.	32
3.4	SEM image of (a) transition region and (b) waist region of the tapered MMF.	33
3.5	Two solutions prepared (a) aniline and PMVEA dissolved in deionized water, (b) APS dissolved in deionized water.	34
3.6	Colour changes observed after (a) 2 minutes, and (b) 4 minutes after	34
	the oxidant solution was added dropwise into the aniline/PMVEA solution, indicating the occurrence of polymerization process.	
3.7	Tapered fibers were hanged on a steel stand and the tapered part immersed in the polymerizing solution. Colour change can be	35
	observed after (a) 5 minutes and (b) 15 minutes.	26
3.8	Ammonia gas sensing experimental setup.	36
3.9	Customized gas chamber.	37
3.10	SEM images of PANI-coated bare optical fiber (cross-sectional view), prepared by in-situ deposition with deposition time of 16 hours (Sample A16).	38
3.11	SEM images of PANI-coated bare optical fiber (cross-sectional	39
5.11	view), prepared by in-situ deposition with deposition duration of 6 hours. (Sample A6).	J
3.12	SEM image of PANI-coated bare optical fiber (cross-sectional	40
5.12	view), prepared by in-situ deposition with deposition duration of 3 hours. (Sample A3).	10

3.13	SEM image of PANI-coated bare optical fiber (cross-sectional view), prepared by in-situ deposition with deposition duration of 2 hours. (Sample A2).	40
3.14	SEM image of PANI-coated bare optical fiber (cross-sectional view), prepared by in-situ deposition with deposition duration of 1 hour. (Sample A1).	41
3.15	SEM images of tapered MMF coated with PANI-PMVEA, for sample (a) A6, (b) A2, and (c) A1.	43
3.16	FESEM images of PANI-PMVEA on glass substrate at magnification of (a) 30,000 times and (b) 80,000 times.	44
3.17	SEM images of tapered MMF coated with PANI-PMVEA, for sample (a) B20, (b) B40, and (c) B1.	45
3.18	Raman spectrum of PANI-PMVEA.	46
3.19	FTIR Spectrum of PANI-PMVEA.	47
3.20	Illustration of (a) response time and (b) recovery time.	49
3.20		50
5.21	Absorbance vs wavelength of sensor (a) A6, (b) A2 and (c) A1, exposed to ammonia gas from 0.125% to 1% for 5 minutes at each concentrations.	30
3.22	Dynamic responses of sensor (a) A6, (b) A2 and (c) A1, exposed to different concentrations of NH <sub>3</sub> gas.	52
3.23	The error bar plot of cumulative absorbance changes as a function of $NH_3$ concentration for Sensor (a) A6 (b) A2 and (c) A1. The error bar was calculated from 3 cycles of the sensor testing results.	54
3.24	Absorbance vs wavelength of sensor (a) B20, (b) B40 and (c) B1, exposed to ammonia gas from 0.125% to 1% for 5 minutes at each concentrations.	55
3.25	Dynamic response of sample of sensor (a) B20, (b) B40 and (c) B1, exposed to different concentrations of NH <sub>3</sub> gas.	56
3.26	The changes in cumulative absorbance vs ammonia concentration for sensors A1, A2, A6, B20 and B40.	57
4.1	Experimental work flow in developing ammonia sensor using spray coating of PANI-CSA on tapered MMF.	59
4.2	The PANI-HCl synthesis process. The HCl containing APS was added dropwise into HCl containing aniline while stirring using magnetic stirrer (a) at the beginning, (b) after a few drops, and (c) after continuous droppings. (d) The solution was stirred for 24 hours at room temperature.	61
4.3	PANI-HCl dedoping process. (a) Precipitated PANI-HCl collected on filter paper. (b) The collected PANI-HCl transferred into glass beaker to be mixed with NH <sub>4</sub> OH. (c) Mixture of PANI-HCl and NH <sub>4</sub> OH stirred for 4 hours.	62
4.4	PANI-EB (a) on filter paper after dried in oven for 48 hours at 40°C and (b) after grinded.	63
4.5	PANI-CSA during (a) stirring using magnetic stirrer and (b) sonication using ultrasonic bath and (c) ready to be sprayed.	63
4.6	(a) Spray coating setup. (b) Air pump and spray gun used for spray coating.	64
4.7	SEM images of (a) Bare glass slide, PANI-CSA spray coated on glass slide at magnification factor of (b) 15000 (c) 10000 and (c) 6000.	66

- 4.8 SEM image of (a) bare tapered fiber waist, (b),(c) tapered fiber 67 coated with PANI-CSA at magnification of 2000 and (d) tapered fiber coated with PANI-CSA at magnification of 4000.
- 4.9 (a) 3D AFM image of boundary area between bare glass and PANI CSA coated glass. (b) Optical image captured during AFM analysis, showing the scan area.

69

70

71

72

- 4.10 Raman spectrum of PANI-CSA.
- 4.11 FTIR Spectrum of PANI-CSA.
- 4.12 Absorbance responses for Sensor (a) D20, (b) D30, (c) D40 and (d) D50.
- 4.13 Dynamic responses for Sensor (a) D20, (b) D30, (c) D40, and (d) D50 as the sensors were alternately exposed to ammonia of different concentrations and purified air.
- 4.14 Cumulative absorbance changes as a function of NH<sub>3</sub> concentration 73 for the Sensors (a) D20, (b) D30, (c) D40, and (d) D50.
- 4.15 Error bar plot of cumulative absorbance change as a function of NH<sub>3</sub>
  75 concentration for Sensor (a) D20, (b) D30, (c) D40, and (d) D50. The error bar were calculated from the 3 cycles of the sensor testing results.

# LIST OF ABBREVIATIONS

AFM	Atomic Force Microscopy
APS	Ammonium persulfate
Au	Gold
BTB	Bromothymol blue
BTC	benzene-1,3,5-tricarboxylate
BTEX	Benzene, toluene, ethylbenzene, and xylene
CSA	Camphorsulfonic acid
CH <sub>4</sub>	Methane
СО	Carbon monoxide
CPR	chlorophenol red
DBSA	Dodecylbenzenesulfonate
DiOHP	bis(2-ethylhexyl) hydrogen phosphate
EB	Emeraldine base
ES	Emeraldine salt
FBG	Fiber Bragg Grating
FESEM	Field Emission Scanning Electron Microscopy
FTIR	Fourier Transform Infrared
FTO	Fluorine-doped tin oxide
GO	Graphene oxide
He-Ne	Helium-neon
HCl	Hydrochloric acid
HClO <sub>4</sub>	Perchloric acid
HF	Hydrofluoric acid
Hydrazine	$N_2H_4$
$H_2S$	Hydrogen sulphide
$H_2SO_4$	Sulphuric acid
IR	Infrared
ITO	Indium tin oxide
I <sub>2</sub>	Iodine
LEB	Leucoemeraldine base
Mid-IR	Mid-Infrared
MMF	Multimode Fiber
NH <sub>3</sub>	Ammonia

NH4Cl	Ammonium chloride
NIR	Near Infra-red
NO <sub>x</sub>	Nitric oxides
O <sub>3</sub>	Ozone
PANI	Polyaniline
PANI-CSA	Camphorsulfonic acid-doped polyaniline
PANI-EB	Emeraldine base polyaniline
PANI-ES	Emeraldine salt polyaniline
PANI-HC1	Hydrochloric acid-doped polyaniline
PANI-PMVEA	Poly(methyl vinyl ether-alt-maleic acid)-doped polyaniline
PEB	Pernigraniline base
PMMA	polymethyl methacrylate
PMVEA	Poly(methyl vinyl ether-alt-maleic acid)
PA6	polyamide 6
PPy	Polypyrrole
PTh	Polythiophene
P4PV	Poly(4-vinylpyridine)
SMF	Single mode fiber
$SnO_2$	Tin oxide
$SO_2$	Sulfur dioxide
SPR	Surface plasmon resonance
TiO <sub>2</sub>	Titanium dioxide
UV-Vis	Ultraviolet-visible
VOC	Volatile organic compound
WO <sub>3</sub>	Tungsten trioxide

### **CHAPTER 1**

#### **INTRODUCTION**

#### 1.1 Overview

Silica optical fiber is well-known as a medium for data transmission in the telecommunication industry, which enables high speed data communications. In the past three decades, optical fiber has been studied; not only for communication but also for many other applications. One of the most interesting areas is for sensing applications. Even though electrical-based sensor is a well-established technology, there are some limitations that these electrical sensors face, which can be overcomed by using optical fiber sensors. Electrical sensors are not suitable to be used in harsh environments such as in high voltage machinery or in chemical interaction medium. Optical fiber can stand those harsh conditions because it is a dielectric medium that does not conduct electrical signal. It is immunes to electromagnetic interference or any electrical signal disturbance and it can stands high temperature up to 1200°C before it start to soften [1]. In addition, the fiber is inexpensive, small size and not susceptible to corrosion.

Fiber optic sensors can be designed either to replace existing sensors or to be applied in new applications. In order for optical fiber sensors to be able to replace conventional sensors, they have to show a significant improvement in terms of performance, accuracy, reliability, safety and cost. Optical fiber gyroscopes is one of the important example of a sensor that has been successfully replaced the conventional sensor. This sensor are used in automobile navigation system, inertial measurement system for aircraft, and backup guidance system for Boeing 777 [2]. Applications of fiber optic sensors are very broad. It ranges from supporting process control in manufacturing system, medical applications, building and structures monitoring system, remote monitoring in harsh environment, natural hazard emergency response system, marine system monitoring, oil exploration and many more that are impossible to be covered here [3]. The number of applications keep increasing because of the rapid research and development in this area, and decreasing in the cost factor. The broad range of applications is also due to the various configurations and parameters that can be measured including pressure, acoustic waves, vibration, position, strain, liquid level, flow, velocity, acceleration, rotation rate, temperature, electrical current, magnetic field, electromagnetic field, chemical process, and the surrounding refractive index [4].

The sensors can be classified based on the working principle such as interferometric sensor. This class of sensor obtained the information from the interference between two optical signals. Well-known sensors under this category are Mach-Zehnder and Michelsen interferometer [2]. Besides that, fiber sensor can be designed based on microbending [5], grating [6], or evanescent wave [7][8]. Evanescent wave sensor can

be fabricated by removing some part of the cladding or by tapering the conventional optical fiber. Recently, research on sensors based on the tapered optical fiber has attracted great attention due to various advantages in terms of compactness, robustness, strong evanescent field, and simple fabrication method. Tapered optical fiber has been studied to measure physical parameters such as temperature [9], humidity [10], strain [11], refractive index [12], to detect chemicals species [13] and also for biosensors [14]. Great potentials of tapered fiber sensors are now recognized by the research community.

Detection of chemical species using fiber sensors has been studied since 1970s. There are two main approaches to sense chemical using fiber sensor [15]. One of the approaches is by measuring the intrinsic optical properties of the analyte such as its refractive index, emission or absorption. The other one is done by monitoring the change in optical properties of the immobilized indicator, or sensing layer that is deposited on the fiber. The second approach usually incorporates some materials as sensing layer that can react with the target analyte. Materials for sensing layer are typically a material that has the switching capability such as conducting polymers (organic), semiconductor metal oxides (inorganic), and composite materials.

Conducting polymer is an organic material that has semiconductor or metal-like conductivity. Its synthesis method and processing is easier than metals, making it very attractive as sensing layer. It has served as sensing layer in electronic gas sensors since 1980s. The sensors were reported to have high sensitivities, short response time, and the most attractive feature is good response at room temperature [16]. Conducting polymers that have been investigated as sensing layer are polyaniline (PANI), polypyrrole (PPy), and polythiophine (PTh). Among these polymers, PANI has been widely studied because of its simple synthesis, low cost, high conductivity and high environmental stability [17]. It has been used to sense nitrogen dioxide (NO<sub>2</sub>), hydrogen sulphide (H<sub>2</sub>S), sulfur dioxide (SO<sub>2</sub>), and most widely for ammonia (NH<sub>3</sub>). Recent development on the nanostructure PANI has increase the interest to incorporate PANI as sensing layer, since nanostructure materials have shown much higher sensitivity compared to bulk material.

Applying optical fiber sensors for gas sensing applications has open up new possibilities of in-situ monitoring on various types of gases at remote or hard-to-reach areas. Real-time and continuous monitoring of certain gas species is in huge demand in process control, automotive, medical, agricultural and many more. Combining the advantages of optical fiber sensors with sensitive nanomaterials for gas sensing application is an interesting research direction to be explored.

### **1.2 Problem Statement**

NH3 is a natural gas that exists in our atmosphere. The main sources include decomposition of manure in agricultural and wildlife, chemical plants (fertilizer & refrigerator manufacturing industries), natural bacterial nitrification activities, and motor vehicles. In industries that use pure NH<sub>3</sub> such as fertilizer and refrigerator industries, leakage in the system may lead to life threatening situations. Exposure to high NH<sub>3</sub> concentrations is hazardous to human health; at concentration of 500 ppm, the person exposed will experience breathing difficulties, and irritation to the nose, eyes and throats. At concentration higher than 1000 ppm can cause pulmonary oedema, long term lung disorder and can be fatal. Exposure to extremely high concentration (5000 - 10,000 ppm) is deadly within 5 - 10 minutes [18]. More worrying, the gas is flammable at 50°C at very high concentration (150,000 ppm) [19]. In history, one of the worst disasters related to NH<sub>3</sub> happened in April 1947, when a ship that carried solid fertilizers exploded while it was docking in Texas City [20] The incident had damaged more than 1,500 buildings and killed almost 600 people. Therefore, it is extremely important to develop an NH<sub>3</sub> gas monitoring system with good sensors to avoid such safety hazards.

Research on electrical-based NH<sub>3</sub> sensors employing PANI is well established. The sensing mechanism is based on measuring changes in conductance (conductometry) or resistance (chemresistors) as PANI chemically reacts towards NH<sub>3</sub>. However, as mentioned earlier, electrical-based sensors are not suitable to be deployed in certain environment especially when there is risk of explosion. PANI has also been used in optical-based sensors by measuring changes in light absorbance, reflectance or transmittance. The sensors were realized by depositing thin PANI film on waveguides [21], glass substrate [22][23][24], and optical fibers [25][26][27]. The use of waveguide as the transducing platform in [21] requires the use of optical bench setup, which is quite bulky and sensitive to misalignment problem. The technique which uses glass substrate in [22] and [24] is more suitable for remote monitoring application because it is less bulky than the waveguide bench setup. However, misalignment of the two fibers that carry light to and from the sensing glass surface is a big challenge that needs to be addressed. The use of optical fibers as the transducing platform overcome the misalignment problem, and enables the sensor to be used for remote monitoring applications.

Evanescent-based fiber sensor is very attractive due to its simple and cheap fabrication methods. It can be realized using side-polished, D-shaped, removed-clad or tapered fibers [28]. Tapered fiber was shown to be more sensitive than side-polished and D-shape fiber [29]. This is due to the presence of evanescent wave surrounding the taper waist region, while D-shape or side polished fiber only has evanescent at one side of the fiber. Usually, to fabricate removed-clad fiber, the cladding of silica optical fibers were removed using chemical etching method [30]. The major drawback with chemical etching method is that it is difficult to control; hence cause low reproducibility. With the advanced in fiber tapering technology, a more reproducible and controllable process to produce evanescent fiber sensors is realized. Tapering does

not involve the removal of the fiber cladding, but it reduces the fiber diameter by heating while pulling the fiber. Since the transition between the original fiber and the tapered fiber waist is smooth, it is less fragile than remove-clad fiber. Due to various advantages of tapered fiber compared to other types of evanescent fiber, it was chosen to be used in this thesis. To the best of author's knowledge, there is no work reported on NH<sub>3</sub> sensor developed by using PANI on tapered optical fiber. Therefore, in this thesis, a new NH<sub>3</sub> sensor deploying PANI as sensitive layer, coated on tapered MMF was developed.

Currently, research on gas sensors using nanostructure materials is a hot topic due to the impressive sensitivity improvement. High surface to volume ratio increases the adsorption and fusion rate of gas molecules into the nanostructures, which leads to faster response and higher sensitivity. Deployment of PANI nanostructures (nanogranules, nanofibers, nanotubes, etc) in electrical-based sensors has been widely studied [31] [32], [33]. On the other hand, study on optical sensor characteristics that employ PANI nanostructure is still at the infancy stage. Only few optical sensing research works are found in literature that deposited PANI nanostructure on glass substrate and observed the absorbance response towards NH<sub>3</sub> [34][23]. Since the sensor in [34] used the most similar measurement setup and unit, it can be used as the benchmark for this work. Their sensor has achieved the sensitivity of 0.1018/vol%. From the author's thorough literature survey, there is no work reported so far regarding the study on the NH<sub>3</sub> optical fiber sensor by employing PANI nanostructures as sensitive sensing layer.

Factors that influence the PANI sensing performance includes its morphology and type of dopants. For tapered fiber sensor, the tapered fiber dimension such as its waist diameter determines the penetration depth and the strength of evanescent field surrounding it; hence influence the sensor's sensitivity. In addition, PANI deposition methods will determine its morphology and thickness, which also affects the sensor's performance. In this thesis, detail investigation on the influence of these parameters on the sensor's absorbance response is carried out systematically.

### 1.3 Objectives

The aim of this thesis is to develop optical fiber  $NH_3$  sensors by integrating PANI nanostructures with tapered optical fibers. The specific objectives are listed as follows:

- To synthesis and deposit PANI nanostructure on tapered optical fiber.
- To characterize the PANI nanostructure material properties and analyse its relation with the sensing performance in terms of sensitivity, response time, recovery time and repeatability.

- To analyse the influence of tapered fiber geometrical properties on the sensing performance.
- To understand and explain the optical sensing mechanism to sense the interaction between PANI nanostructures and NH<sub>3</sub> gas molecules.

In order to achieve these objectives, the following research questions are outlined:

- What are the suitable synthesis and deposition methods to coat PANI nanostructures on tapered optical fiber?
- How do the synthesis and deposition methods affect the sensing layer's morphology and thickness?
- How different are the sensing performances of PANI nanostructures with different morphologies and thickness?
- How different are the sensing performances of tapered fiber sensors with different waist diameters?

#### **1.4** Scope and limitation

This thesis is focusing on evanescence-based optical fiber sensor using tapered multimode fiber. Two types of PANI nanostructures were used as the sensing layer, which are PANI-CSA nanofibers and PANI-PMVEA nanogranules. The sensing mechanism is based on evanescent wave absorption. The sensors were tested at room temperature (26°C) to sense NH<sub>3</sub> at four different concentrations (0.125%, 0.25%, 0.5% and 1%). The sensors were not tested for different temperature because it is purposely designed for room temperature applications. PANI, a type of conducting polymer, was chosen as the sensing layer because it is known to have good response at room temperature. Besides, the polymer is not suitable to be used at high temperature because it will decompose at temperature of more than  $100^{\circ}$ C.

The tapered fiber used in this thesis was fabricated from conventional graded index multimode fiber with core/cladding ratio of 62.5  $\mu$ m/125  $\mu$ m. The tapered fiber has fixed down taper and up taper length of 2mm, and fixed waist length of 10 mm. In the method that uses PANI-CSA as sensing layer, the waist diameter was varied between 20  $\mu$ m to 50  $\mu$ m. The diameter was not reduced below 20  $\mu$ m because the fiber becomes very fragile and difficult to handle. A preliminary work using tapered fiber with waist diameter of 15 $\mu$ m was done, but it was not continued because the fiber was broken. So, it was decided not to use the fiber with diameter of less than 20  $\mu$ m. Meanwhile, the sensor with 50  $\mu$ m showed a very low sensitivity due to very small evanescent penetration depth. Therefore, it is useless to further increase the diameter. For sensor with PANI-PMVEA coating, the diameter was not varied because the focus of that chapter is to study the effect of sensing layer morphology and thickness.

 $\bigcirc$ 

This thesis also does not focus on the doping level of the CSA and PMVEA into PANI. In Chapter 3, PMVEA was only varied by 5%, which does not really affect the doping level, because the amount of polymer coated on the fiber is very small relative to the total amount of polymer produced in 100 ml of solution. Meanwhile, in Chapter 4, the amount of CSA was not varied.

## 1.5 Thesis Organisation

This research work is reported in a thesis that consists of 5 chapters. Chapter 1 provides a brief overview of the tapered fiber sensor fields followed by the problem statements and objectives. Theoretical background and review on the previous reported research findings related to this work are presented in Chapter 2. In Chapter 3, the sensor fabrication process that involves in-situ deposition of PANI-PMVEA nanogranules on tapered fiber is elaborated in details. The analyses on the PANI coating properties together with the sensing results are included in the same chapter. Chapter 4 discusses the sensor fabrication method that involves synthesis and spray coating of PANI-CSA nanofibers together with the PANI coating properties and sensing performance. Finally, research findings and outlines of future recommendations for this research are concluded in Chapter 5.

#### REFERENCES

- [1] K. L. Loewenstein, *The Manufacturing Technology of Continuous Glass Fibres*, 3rd ed. Elsevier, 1993.
- [2] E. Udd, *Fiber Optic Sensors*, Second. New York: CRC Press, 2008.
- [3] G. D. Pitt, P. Extance, R. C. Neat, D. N. Batchelder, R. E. Jones, J. A. Barnett, and R. H. Pratt, "Optical-fibre sensors," *Optoelectron. IEE Proc. J*, vol. 132, pp. 214–248, 1985.
- [4] Y. Shi-Kay and C. Asawa, "Fiber Optical Intensity Sensors," *Sel. Areas Commun. IEEE J.*, vol. 1, pp. 562–575, 1983.
- J. W. Berthold, "Optical Fiber Sensor Technology: Applications and Systems,"
   K. T. V Grattan and B. T. Meggitt, Eds. Boston, MA: Springer US, 1999, pp. 225–240.
- [6] Y. J. Rao, "Recent progress in applications of in-fibre Bragg grating sensors," *Opt. Lasers Eng.*, vol. 31, pp. 297–324, 1999.
- [7] P. Wang, G. Brambilla, M. Ding, Y. Semenova, Q. Wu, and G. Farrell, "Highsensitivity, evanescent field refractometric sensor based on a tapered, multimode fiber interference.," *Opt. Lett.*, vol. 36, no. 12, pp. 2233–5, Jun. 2011.
- [8] A. G. Mignani, R. Falciai, and L. Ciaccheri, "Evanescent Wave Absorption Spectroscopy by Means of Bi-tapered Multimode Optical Fibers SPECTROSCOPY USING A MULTIMODE," *Appl. Spectrosc.*, vol. 52, no. 4, pp. 546–551, 1998.
- [9] M. Barbu, K. Jovanovich, R. Trahan, and P. Chirlian, "The Use of Tapered Fibers in an Intensity Based Single Mode Temperature Sensor," pp. 47–50, 2000.
- [10] L. Zhang, F. Gu, J. Lou, X. Yin, and L. Tong, "Fast detection of humidity with a subwavelength-diameter fiber taper coated with gelatin film.," *Opt. Express*, vol. 16, no. 17, pp. 13349–53, Aug. 2008.
- [11] S. Mas, J. Mart, and J. Palac, "Biconical tapered fibers manipulation for refractive index and strain sensing applications," *IEEE Sens. J.*, vol. 15, no. 3, pp. 1331–1335, 2015.
- [12] W. Bin Ji, H. H. Liu, S. C. Tjin, K. K. Chow, and A. Lim, "Ultrahigh Sensitivity Refractive Index Sensor Based on Optical Microfiber," *IEEE Photonics Technol. Lett.*, vol. 24, no. 20, pp. 1872–1874, Oct. 2012.
- [13] J. Villatoro, D. Luna-Moreno, and D. Monzón-Hernández, "Optical fiber hydrogen sensor for concentrations below the lower explosive limit," *Sensors Actuators B Chem.*, vol. 110, no. 1, pp. 23–27, Sep. 2005.
- [14] A. Leung, K. Rijal, P. M. Shankar, and R. Mutharasan, "Effects of geometry on transmission and sensing potential of tapered fiber sensors.," *Biosens. Bioelectron.*, vol. 21, no. 12, pp. 2202–9, Jun. 2006.
- [15] O. S. Wolfbeis, "Fibre-optic chemical sensors and biosensors," *Anal. Chem.*, vol. 78, no. 12, pp. 3859–74, 2006.
- [16] H. Bai and G. Shi, "Gas Sensors Based on Conducting Polymers," *Sensors*, vol. 7, no. 3, pp. 267–307, Mar. 2007.
- [17] G. Ciric-Marjanovic, "Recent advances in polyaniline research: Polymerization mechanisms, structural aspects, properties and applications," *Synth. Met.*, vol. 177, no. 3, pp. 1–47, 2013.

- [18] B. Timmer, W. Olthuis, and A. Van Den Berg, "Ammonia sensors and their applications—a review," *Sensors Actuators B Chem.*, vol. 107, no. 2, pp. 666– 677, Jun. 2005.
- [19] Praxair, "Ammonia , anhydrous Ammonia , anhydrous Safety Data Sheet P-4562," pp. 1–9, 2015.
- [20] "1947 Texas City Disaster," *Moore Memorial Public Library*. [Online]. Available: http://www.texascity-library.org/disaster/fire.php.
- [21] A. Airoudj, D. Debarnot, B. Bêche, and F. Poncin-Epaillard, "A new evanescent wave ammonia sensor based on polyaniline composite.," *Talanta*, vol. 76, no. 2, pp. 314–9, Jul. 2008.
- [22] S. Christie, E. Scorsone, K. Persaud, and F. Kvasnik, "Remote detection of gaseous ammonia using the near infrared transmission properties of polyaniline," *Sensors Actuators B Chem.*, vol. 90, no. 1–3, pp. 163–169, 2003.
- [23] P. Stamenov, R. Madathil, and J. M. D. Coey, "Dynamic response of ammonia sensors constructed from polyaniline nanofibre films with varying morphology," *Sensors Actuators B Chem.*, vol. 161, no. 1, pp. 989–999, Jan. 2012.
- [24] J. Castrellon-Uribe, M. E. Nicho, and G. Reyes-Merino, "Remote optical detection of low concentrations of aqueous ammonia employing conductive polymers of polyaniline," *Sensors Actuators B Chem.*, vol. 141, no. 1, pp. 40– 44, Aug. 2009.
- [25] M. El-Sherif, "Fiber Optic Sensors For Detection of Toxic and Biological Threats," vol. 7, no. 12, pp. 3100–3118, Dec. 2007.
- [26] L. Ai, J. C. Mau, W. F. Liu, T. C. Chen, and W. K. Su, "A volatile-solvent gas fiber sensor based on polyaniline film coated on superstructure fiber Bragg gratings," *Meas. Sci. Technol.*, vol. 19, no. 1, p. 17002, Jan. 2008.
- [27] S. K. Mishra, D. Kumari, and B. D. Gupta, "Surface plasmon resonance based fiber optic ammonia gas sensor using ITO and polyaniline," *Sensors Actuators B Chem.*, vol. 171–172, pp. 976–983, Aug. 2012.
- [28] W. Jin, H. L. Ho, Y. C. Cao, J. Ju, and L. F. Qi, "Gas detection with microand nano-engineered optical fibers," *Opt. Fiber Technol.*, vol. 19, no. 6 PART B, pp. 741–759, 2013.
- [29] W. Henry, "Evanescent field devices: a comparison between tapered optical fibres and polished or D-fibres," *Opt. Quantum Electron.*, vol. 26, no. 3, pp. 261–272, 1994.
- [30] M. El-Sherif, "Fiber-optic chemical sensor using polyaniline as modified cladding material," *IEEE Sens. J.*, vol. 3, no. 1, pp. 5–12, Feb. 2003.
- [31] I. Fratoddi, I. Venditti, C. Cametti, and M. V. Russo, "Chemiresistive polyaniline-based gas sensors: A mini review," *Sensors Actuators, B Chem.*, vol. 220, pp. 534–548, 2015.
- [32] A. L. Sharma, K. Kumar, and A. Deep, "Nanostructured polyaniline films on silicon for sensitive sensing of ammonia," *Sensors Actuators A Phys.*, vol. 198, pp. 107–112, Aug. 2013.
- [33] V. Talwar, O. Singh, and R. C. Singh, "ZnO assisted polyaniline nanofibers and its application as ammonia gas sensor," *Sensors Actuators B Chem.*, vol. 191, pp. 276–282, Feb. 2014.
- [34] H. Kebiche, D. Debarnot, A. Merzouki, F. Poncin-Epaillard, and N. Haddaoui, "Relationship between ammonia sensing properties of polyaniline nanostructures and their deposition and synthesis methods," *Anal. Chim. Acta*, vol. 737, pp. 64–71, 2012.

- [35] J. Hodgkinson and R. P. Tatam, "Optical gas sensing: a review," *Meas. Sci. Technol.*, vol. 24, no. 1, p. 12004, Jan. 2013.
- [36] O. S. Wolfbeis, "Fiber-optic chemical sensors and biosensors," *Analytical Chemistry*, vol. 80, no. 12. pp. 4269–4283, 2008.
- [37] M. H. Yaacob, M. Breedon, K. Kalantar-zadeh, and W. Wlodarski, "Absorption spectral response of nanotextured WO3 thin films with Pt catalyst towards H2," *Sensors Actuators B Chem.*, vol. 137, no. 1, pp. 115–120, Mar. 2009.
- [38] A. Airoudj, D. Debarnot, B. Bêche, and F. Poncin-Epaillard, "Development of an optical ammonia sensor based on polyaniline/epoxy resin (SU-8) composite.," *Talanta*, vol. 77, no. 5, pp. 1590–6, Mar. 2009.
- [39] Y. Zhao, X. Li, X. Zhou, and Y. Zhang, "Review on the graphene based optical fiber chemical and biological sensors," *Sensors Actuators B Chem.*, vol. 231, pp. 324–340, 2016.
- [40] M. H. Yaacob, M. Z. Ahmad, A. Z. Sadek, J. Z. Ou, J. Campbell, K. Kalantar-Zadeh, and W. Wlodarski, "Optical response of WO3 nanostructured thin films sputtered on different transparent substrates towards hydrogen of low concentration," *Sensors Actuators, B Chem.*, vol. 177, pp. 981–988, 2013.
- [41] A. Y. Mironenko, A. A. Sergeev, A. E. Nazirov, E. B. Modin, S. S. Voznesenskiy, and S. Y. Bratskaya, "H2S optical waveguide gas sensors based on chitosan/Au and chitosan/Ag nanocomposites," *Sensors Actuators, B Chem.*, vol. 225, pp. 348–353, 2016.
- [42] A. Yimit, K. Itoh, and M. Murabayashi, "Detection of ammonia in the ppt range based on a composite optical waveguide pH sensor," *Sensors Actuators, B Chem.*, vol. 88, no. 3, pp. 239–245, 2003.
- [43] M.-K. Bae, J. A. Lim, S. Kim, and Y.-W. Song, "Ultra-highly sensitive optical gas sensors based on chemomechanical polymer-incorporated fiber interferometer.," *Opt. Express*, vol. 21, no. 2, pp. 2018–23, Jan. 2013.
- [44] J. Homola, "Present and future of surface plasmon resonance biosensors.," *Anal. Bioanal. Chem.*, vol. 377, no. 3, pp. 528–39, Oct. 2003.
- [45] A. K. Sharma, R. Jha, and B. D. Gupta, "Fiber-Optic Sensors Based on Surface Plasmon Resonance: A Comprehensive Review," *IEEE Sens. J.*, vol. 7, no. 8, pp. 1118–1129, Aug. 2007.
- [46] R. Kashyap and G. Nemova, "Surface Plasmon Resonance-Based Fiber and Planar Waveguide Sensors," *J. Sensors*, vol. 2009, pp. 1–9, 2009.
- [47] B. Lee, S. Roh, and J. Park, "Current status of micro- and nano-structured optical fiber sensors," *Opt. Fiber Technol.*, vol. 15, no. 3, pp. 209–221, 2009.
- [48] C. J. Gouveia, J. M. Baptista, and P. S. Jorge, "Refractometric Optical Fiber Platforms for Label Free Sensing," *Curr. Dev. Opt. Fiber Technol.*, no. May 2016, pp. 345–373, 2013.
- [49] Z. L. Poole, P. Ohodnicki, R. Chen, Y. Lin, and K. P. Chen, "Engineering metal oxide nanostructures for the fiber optic sensor platform," vol. 22, no. 3, pp. 2009–2013, 2014.
- [50] X. Chong, K. J. Kim, P. R. Ohodnicki, E. Li, C. H. Chang, and A. X. Wang, "Ultrashort Near-Infrared Fiber-Optic Sensors for Carbon Dioxide Detection," *IEEE Sens. J.*, vol. 15, no. 9, pp. 5327–5332, 2015.
- [51] G. Brambilla, V. Finazzi, and D. J. Richardson, "Ultra-low-loss optical fiber nanotapers," *Opt. Express*, vol. 12, pp. 2258–2263, 2004.
- [52] L. S. Grattan and B. T. Meggit, *Optical Fiber Sensor Technology*. Springer US, 1999.

- [53] H. Tai, H. Tanaka, and T. Yoshino, "Fiber-optic evanescent-wave methanegas sensor using optical absorption for the 3.392-Mm line of a He-Ne laser," vol. 12, no. 6, pp. 437–439, 1987.
- [54] J. Villatoro and D. Monzón-Hernández, "Fast detection of hydrogen with nano fiber tapers coated with ultra thin palladium layers.," *Opt. Express*, vol. 13, no. 13, pp. 5087–92, Jun. 2005.
- [55] R. Jarzebinska, S. Korposh, S. James, W. Batty, R. Tatam, and S.-W. Lee, "Optical Gas Sensor Fabrication Based on Porphyrin-Anchored Electrostatic Self-Assembly onto Tapered Optical Fibers," *Anal. Lett.*, vol. 45, no. 10, pp. 1297–1309, Jul. 2012.
- [56] S. O. Korposh, N. Takahara, J. J. Ramsden, S.-W. Lee, and T. Kunitake, "Nano-assembled thin film gas sensors. I. Ammonia detection by a porphyrinbased multilayer film," *J. Biol. Phys. Chem.*, vol. 6, no. August, pp. 125–132, 2006.
- [57] S. Korposh, S. Kodaira, W. Batty, S. W. James, and S. W. Lee, "Nanoassembled Thin-Film Gas Sensor II. An Intrinsic Highly Sensitive Fibre Optic Sensor for Ammonia Detection," *Sensors Mater.*, vol. 21, no. 4, pp. 179– 189, 2009.
- [58] L. I. Espada, M. Shadaram, J. Robillard, and K. H. Pannell, "Ferrocenylenesilylene Polymers as Coatings for Tapered Optical-Fiber Gas Sensors," J. Inorg. Organomet. Polym., vol. 10, no. 4, pp. 169–176, 2000.
- [59] G. B. Bredas, J. L. and Street, "Polarons, Bipolarons, Solitons in Conducting Polymers," *Acc. Chem. Res.*, vol. 18, pp. 309–315, 1985.
- [60] N. Abdul Rahman, "Electrospun Conducting Polymer Nanofibers for Biomedical Applications," The University of Auckland, New Zealand, 2012.
- [61] P. R. T. Gordon G. Wallace, Geoffrey M. Spinks, Leon A. P. Kane-Maguire, Conductive Electroactive Polymers: Intelligent Polymer Systems, 3rd ed. CRC Press, 2009.
- [62] J. Stubb, H., Punkka, E., Paloheimo, "Electronic and optical properties of conducting polymer thin films," *Mater. Sci. Eng.*, vol. 10, pp. 85–140, 1993.
- [63] A. Partridge, M. Jansen, and W. Arnold, "Conducting polymer-based sensors," *Mater. Sci. Eng. C*, vol. 12, no. 1–2, pp. 37–42, Aug. 2000.
- [64] U. Lange, N. V Roznyatovskaya, and V. M. Mirsky, "Conducting polymers in chemical sensors and arrays.," *Anal. Chim. Acta*, vol. 614, no. 1, pp. 1–26, Apr. 2008.
- [65] S. L. Patil, M. a. Chougule, S. Sen, and V. B. Patil, "Measurements on room temperature gas sensing properties of CSA doped polyaniline–ZnO nanocomposites," *Measurement*, vol. 45, no. 3, pp. 243–249, Apr. 2012.
- [66] D. Nicolas-Debarnot and F. Poncin-Epaillard, "Polyaniline as a new sensitive layer for gas sensors," *Anal. Chim. Acta*, vol. 475, no. 1–2, pp. 1–15, Jan. 2003.
- [67] W. Prissanaroon, L. Ruangchuay, A. Sirivat, and J. Schwank, "Electrical conductivity response of dodecylbenzene sulfonic acid-doped polypyrrole films to SO2–N2 mixtures," *Synth. Met.*, vol. 114, no. 1, pp. 65–72, Jul. 2000.
- [68] L.-X. Wang, X.-G. Li, and Y.-L. Yang, "Preparation, properties and applications of polypyrroles," *React. Funct. Polym.*, vol. 47, no. 2, pp. 125–139, Mar. 2001.
- [69] L. Al-Mashat, H. D. Tran, W. Wlodarski, R. B. Kaner, and K. Kalantar-Zadeh, "Conductometric Hydrogen Gas Sensor Based on Polypyrrole Nanofibers," *IEEE Sens. J.*, vol. 8, no. 4, pp. 365–370, Apr. 2008.
- [70] D. Verma and V. Dutta, "Role of novel microstructure of polyaniline-CSA thin

film in ammonia sensing at room temperature," *Sensors Actuators B Chem.*, vol. 134, no. 2, pp. 373–376, Sep. 2008.

- [71] V. C. Gonçalves and D. T. Balogh, "Optical chemical sensors using polythiophene derivatives as active layer for detection of volatile organic compounds," *Sensors Actuators B Chem.*, vol. 162, no. 1, pp. 307–312, Feb. 2012.
- [72] B. Li, S. Santhanam, L. Schultz, M. Jeffries-EL, M. C. Iovu, G. Sauvé, J. Cooper, R. Zhang, J. C. Revelli, A. G. Kusne, J. L. Snyder, T. Kowalewski, L. E. Weiss, R. D. McCullough, G. K. Fedder, and D. N. Lambeth, "Inkjet printed chemical sensor array based on polythiophene conductive polymers," *Sensors Actuators B Chem.*, vol. 123, no. 2, pp. 651–660, May 2007.
- [73] J. Cerón Solís, E. De la Rosa, and E. Peña Cabrera, "Absorption and refractive index changes of poly (3-octylthiophene) under NO2 gas exposure," *Opt. Mater. (Amst).*, vol. 29, no. 2–3, pp. 167–172, Nov. 2006.
- [74] X. Guo, Y. Kang, T. Yang, and S. Wang, "Low-temperature NO2 sensors based on polythiophene/WO3 organic-inorganic hybrids," *Trans. Nonferrous Met. Soc. China*, vol. 22, no. 2, pp. 380–385, Feb. 2012.
- [75] S. Virji, J. Huang, R. B. Kaner, and B. H. Weiller, "Polyaniline Nanofiber Gas Sensors: Examination of Response Mechanisms," *Nano Lett.*, vol. 4, no. 3, pp. 491–496, Mar. 2004.
- [76] A. Airoudj, D. Debarnot, B. Bêche, and F. Poncin-Epaillard, "Design and sensing properties of an integrated optical gas sensor based on a multilayer structure.," *Anal. Chem.*, vol. 80, no. 23, pp. 9188–94, Dec. 2008.
- [77] S. Sinha, S. Bhadra, and D. Khastgir, "Effect of Dopant Type on the Properties of Polyaniline," *Appl. Polym. Sci.*, vol. 112, pp. 3135–3140, 2009.
- [78] Y.-Z. Long, M.-M. Li, C. Gu, M. Wan, J.-L. Duvail, Z. Liu, and Z. Fan, "Recent advances in synthesis, physical properties and applications of conducting polymer nanotubes and nanofibers," *Prog. Polym. Sci.*, vol. 36, no. 10, pp. 1415–1442, 2011.
- [79] A. H. Navarchian, Z. Hasanzadeh, and M. Joulazadeh, "Effect of Polymerization Conditions on Reaction Yield, Conductivity, and Ammonia Sensing of Polyaniline," *Adv. Polym. Technol.*, vol. 32, no. 3, p. n/a-n/a, 2013.
- [80] V. V. Chabukswar, S. Pethkar, and A. A. Athawale, "Acrylic acid doped polyaniline as an ammonia sensor," *Sensors Actuators, B Chem.*, vol. 77, no. 3, pp. 657–663, 2001.
- [81] M. Matsuguchi and T. Asahi, "Properties and stability of polyaniline nanofiber ammonia sensors fabricated by novel on-substrate method," *Sensors Actuators, B Chem.*, vol. 160, no. 1, pp. 999–1004, 2011.
- [82] Z. Pang, J. Fu, P. Lv, F. Huang, and Q. Wei, "Effect of CSA Concentration on the Ammonia Sensing Properties of CSA-Doped PA6/PANI Composite Nanofibers," *Sensors*, vol. 14, no. 11, pp. 21453–21465, 2014.
- [83] T. Mérian, N. Redon, Z. Zujovic, D. Stanisavljev, J. L. Wojkiewicz, and M. Gizdavic-Nikolaidis, "Ultra sensitive ammonia sensors based on microwave synthesized nanofibrillar polyanilines," *Sensors Actuators B Chem.*, vol. 203, pp. 626–634, Nov. 2014.
- [84] K. Crowley, a Morrin, a Hernandez, E. Omalley, P. Whitten, G. Wallace, M. Smyth, and a Killard, "Fabrication of an ammonia gas sensor using inkjet-printed polyaniline nanoparticles," *Talanta*, vol. 77, no. 2, pp. 710–717, Dec. 2008.
- [85] Y. Huang, L. Wieck, and S. Tao, "Development and evaluation of optical fiber

NH3 sensors for application in air quality monitoring," *Atmos. Environ.*, vol. 66, pp. 1–7, 2013.

- [86] Y. Leng, "Scanning Electron Microscopy," in *Materials Characterization*, John Wiley & Sons (Asia) Pte Ltd, 2008, pp. 121–144.
- [87] Y. Leng, "Vibrational Spectroscopy for Molecular Analysis," in *Materials Characterization*, John Wiley & Sons (Asia) Pte Ltd, 2008, pp. 253–300.
- [88] Y. Leng, "Scanning Probe Microscopy," in *Materials Characterization*, John Wiley & Sons (Asia) Pte Ltd, 2008, pp. 145–170.
- [89] L. Zhang, H. Peng, C. F. Hsu, P. a Kilmartin, and J. Travas-Sejdic, "Selfassembled polyaniline nanotubes grown from a polymeric acid solution," *Nanotechnology*, vol. 18, no. 11, p. 115607, Mar. 2007.
- [90] C. Koo, B. Jeon, and I. Chung, "The Effect of Poly(methyl vinyl ether-altmaleic acid) Stabilizer on the Stability of Polyaniline-Poly(methyl vinyl etheralt-maleic acid) Dispersions.," J. Colloid Interface Sci., vol. 227, no. 2, pp. 316–321, 2000.
- [91] M. Ahmad and L. L. Hench, "Effect of taper geometries and launch angle on evanescent wave penetration depth in optical fibers.," *Biosens. Bioelectron.*, vol. 20, no. 7, pp. 1312–9, Jan. 2005.
- [92] C. Liu, J. Zhang, G. Shi, and F. Chen, "Doping level change of polyaniline film during its electrochemical growth process," *J. Appl. Polym. Sci.*, vol. 92, no. 1, pp. 171–177, 2004.
- [93] H. Tai, Y. Jiang, G. Xie, J. Yu, X. Chen, and Z. Ying, "Influence of polymerization temperature on NH3 response of PANI/TiO2 thin film gas sensor," *Sensors Actuators, B Chem.*, vol. 129, no. 1, pp. 319–326, 2008.
- [94] H. Tai, Y. Jiang, G. Xie, and J. Yu, "Preparation, Characterization and Comparative NH3-sensing Characteristic Studies of PANI/inorganic Oxides Nanocomposite Thin Films," J. Mater. Sci. Technol., vol. 26, no. 7, pp. 605– 613, 2010.
- [95] N. Abdul Rahman, M. Gizdavic-Nikolaidis, S. Ray, A. J. Easteal, and J. Travas-Sejdic, "Functional electrospun nanofibres of poly(lactic acid) blends with polyaniline or poly(aniline-co-benzoic acid)," *Synth. Met.*, vol. 160, no. 17–18, pp. 2015–2022, Sep. 2010.
- [96] Z. Wu, X. Chen, S. Zhu, Z. Zhou, Y. Yao, W. Quan, and B. Liu, "Enhanced sensitivity of ammonia sensor using graphene/polyaniline nanocomposite," *Sensors Actuators B Chem.*, vol. 178, pp. 485–493, Mar. 2013.
- [97] V. J. Babu, S. Vempati, and S. Ramakrishna, "Conducting Polyaniline-Electrical Charge Transportation," *Mater. Sci. Appl.*, vol. 4, no. 1, pp. 1–10, 2013.
- [98] B. Min, "SnO2 thin film gas sensor fabricated by ion beam deposition," *Sensors Actuators B Chem.*, vol. 98, no. 2–3, pp. 239–246, Mar. 2004.
- [99] J. Chen, B. Winther-Jensen, Y. Pornputtkul, K. West, L. Kane-Maquire, and G. G. Wallace, "Synthesis of Chiral Polyaniline Films via Chemical Vapor Phase Polymerization," *Electrochem. Solid-State Lett.*, vol. 9, no. 1, p. C9, 2006.
- [100] M. C. Bernard and A. Hugot-Le Goff, "Quantitative characterization of polyaniline films using Raman spectroscopy," *Electrochim. Acta*, vol. 52, no. 2, pp. 595–603, Oct. 2006.
- [101] Sudha, D. Kumar, and M. Iwamoto, "Investigation of the chiroptical behavior of optically active polyaniline synthesized from naturally occurring amino

acids," Polymer (Guildf)., vol. 45, no. 2, pp. 160–165, 2013.

- [102] A. B. Socorro, I. Del Villar, J. M. Corres, I. R. Matias, and F. J. Arregui, "Lossy mode resonances dependence on the geometry of a tapered monomode optical fiber," *Sensors Actuators A Phys.*, vol. 180, pp. 25–31, Jun. 2012.
- [103] N. G. Deshpande, Y. G. Gudage, R. Sharma, J. C. Vyas, J. B. Kim, and Y. P. Lee, "Studies on tin oxide-intercalated polyaniline nanocomposite for ammonia gas sensing applications," *Sensors Actuators, B Chem.*, vol. 138, no. 1, pp. 76–84, 2009.
- [104] P. Kilmartin, "Polymeric Acid Doped Polyaniline Nanotubes for Oligonucleotide Sensors," *Electroanalysis*, 2007.

