

# **UNIVERSITI PUTRA MALAYSIA**

# PREPARATION AND CHARACTERIZATION OF POLYPYRROLE/MONTMORILLONITE CLAY CONDUCTING POLYMER

FARIZ BIN ADZMI

FS 2007 45



# PREPARATION AND CHARACTERIZATION OF POLYPYRROLE/MONTMORILLONITE CLAY CONDUCTING POLYMER

FARIZ BIN ADZMI

# MASTER OF SCIENCE

2007



# PREPARATION AND CHARACTERIZATION OF POLYPYRROLE/MONTMORILLONITE CLAY CONDUCTING POLYMER

By

# FARIZ BIN ADZMI

Thesis Submitted to the School of Graduate Studies, Universiti Putra Malaysia, in Fulfilment of the Requirement for the Degree of Master of Science

January 2007



Abstract of thesis to the Senate of Universiti Putra Malaysia in fulfilment of the requirement for the degree of Master of Science

## PREPARATION AND CHARACTERIZATION OF POLYPYRROLE/MONTMORILLONITE CLAY CONDUCTING POLYMER

By

#### FARIZ BIN ADZMI

#### January 2007

#### Chairman: Professor Anuar Kassim, PhD

Faculty: Science

In this study, Ppy/MMT nanocomposites have been prepared by polymerization of pyrrole in the presence of FeCl<sub>3</sub> as oxidation agent and MMT clay in aqueous medium through the chemical polymerization.

The electrical conductivity of the Ppy/MMT nanocomposites was measured by using four-point probe technique. Conductivity of the Ppy/MMT nanocomposites increased from 0.45 to 2.25 S/cm. The increase of conductivity depended on the percentages of Ppy loading in the nanocomposites. Ppy/MMT nanocomposites exhibited similar infrared (IR) absorption peaks as Ppy and MMT clay. Thereby, confirming the presence of the Ppy and MMT in the products. The X-Ray Diffraction (XRD) patterns revealed, MMT strong peak at  $2\theta = 6.68^{\circ}$  (*d*-spacing 13.22 Å) and for Ppy/MMT nanocomposites is at  $2\theta = 4.92^{\circ}$  (*d*-spacing 17.94 Å). The shifting of the angle  $2\theta$  is indicated that the insertion of Ppy has occurred. As determined from XRD analyses the crystalline size of MMT is 17.31 Å and is 25.42 Å for Ppy/MMT nanocomposites.



Thermal gravimetric analysis (TGA) revealed the improvement of degradation temperature of Ppy is at 157.36 to 177.21 °C for Ppy/MMT nanocomposites. Scanning electron micrographs (SEM) revealed some interesting morphological differences between the pure MMT clay and Ppy/MMT nanocomposites. The Ppy/MMT nanocomposite exhibits a denser and more compact morphology. Particle formation of irregular and smaller size was clearly revealed by SEM.

The effects of several parameters in producing Ppy/MMT nanocomposites were also studied. When percentage of MMT was increased from 1 to 7% the conductivity was decreased from 2.25 to 0.38 S/cm. Result shows higher conductivity for sample prepared at low temperature (~5°C) compared to room temperature (25°C). The surface morphology of the Ppy/MMT nanocomposites at low temperature is more compact and denser compared to Ppy/MMT nanocomposites prepared at room temperature. FeCl<sub>3</sub> is the suitable oxidation agent to produce Ppy/MMT nanocomposites in aqueous medium. While 6 hours is the suitable preparation time for completion of polymerization of pyrrole.



Abstrak yang dikemukakan kepada Senat Universiti Putra Malaysia sebagai memenuhi keperluan untuk ijazah Master Sains

## PENYEDIAAN DAN PENCIRIAN TERHADAP POLIPIROL/MONTMORILLONITE KLAY POLIMER PENGALIR

Oleh

#### FARIZ BIN ADZMI

#### Januari 2007

#### Pengerusi: Professor Anuar Kassim, PhD

Fakulti: Sains

Dalam kajian ini, Ppy/MMT nanokomposit telah dihasilkan melalui proses pempolimeran pirol dengan menggunakan FeCl<sub>3</sub> sebagai agen pengoksidaan dan MMT klay dalam larutan akueus melalui proses pempolimeran kimia.

Keberaliran elektrik bahan Ppy/MMT nanokomposit ini diukur menggunakan teknik prob empat titik. Didapati keberaliran bahan nanokomposit ini bertambah daripada 0.45 kepada 2.25 S/cm. Pertambahan keberaliran ini bergantung kepada peratusan polipirol masuk ke dalam bahan Ppy/MMT nanokomposit. Ppy/MMT nanokomposit menunjukkan penyerapan sinar infra merah (IR) sama dengan penyerapan di dalam polipirol dan MMT klay. Ini membuktikan kewujudan polipirol dan MMT klay di dalam Ppy/MMT nanokomposit. Paten pembelauan sinar-X (XRD) menunjukkan puncak penyerapan tajam MMT klay pada  $2\theta = 6.68^{\circ}$  (jarak-*d* 13.22 Å). Manakala bagi Ppy/MMT pada pada  $2\theta = 4.92^{\circ}$  (jarak-*d* 17.94 Å). Perubahan pada sudut 20



menunjukkan kemasukkan polimer polipirol ke dalam dalam MMT klay. Saiz hablur untuk MMT adalah 19.31 Å, manakala untuk Ppy/MMT nanokomposit adalah 25.42Å. Analisis terma gravimetri (TGA) telah menunjukkan peningkatan suhu degradasi daripada 157.36°C untuk polipirol kepada 177.21°C untuk Ppy/MMT nanokomposit. Analisis mikroskop pengimbas electron (SEM) menunjukkan keadaan permukaan yang berbeza diantara MMT and Ppy/MMT nanokomposit. Ia telah menujukkan permukaan Ppy/MMT nanokomposit adalah padat dan rapat. Formasi partikel adalah tidak sekata dan bersaiz kecil dapat dilihat dengan jelas.

Kajian ini juga mengkaji kesan-kesan dan faktor-faktor lain yang mempengaruhi penghasilan Ppy/MMT nanokomposit. Apabila peratusan MMT ditingkatkan daripada 1 kepada 7% didapati keberaliran Ppy/MMT nanokomposit semakin menurun daripada 2.25 kepada 0.38 S/cm. Kajian menunjukkan keberaliran pasa suhu rendah (~5°C) adalah lebih tinggi darapada suhu bilik (25°C). Perbezaan juga terdapat dalam morfologi permukaan Ppy/MMT nanokomposit di mana pada suhu rendah permukaan Ppy/MMT nanokomposit menjadi semakin padat dan rapat. Seterusnya FeCl<sub>3</sub> adalah agen pengoksidaan yang paling sesuai digunakan untuk penghasilan Ppy/MMT dalam medium akueus. Sementara 6 jam adalah masa yang sesuai untuk melengkapkan proses pempolimeran pirol.



#### ACKNOWLEDGEMENTS

#### Alhamdulillah to Almighty God.

First and foremost, I would like to express my deepest appreciation to my supervisor, Prof. Dr. Anuar Kassim and members of the committee, Prof. Dr. Zulkarnain Zainal and Assoc. Prof. Dr. Mohamad Zaki Ab Rahman for their guidance, advice and suggestion throughout the duration of this study.

I would like to extend my tremendous thanks and greatest gratitude to the lecturers and supported staff at Chemistry Department especially to Mr. Abas, Mr. Khidir, Mr Nordin and others who have directly or indirectly contributed their part. My sincere thanks also to my laboratory mate Mr. Ikramul, Mrs. Rozita, Ayu and Shidat. I'm also very thankful to the financial support from MARA.

My special thanks to my parent Adzmi b. Othman and Maznah bt. Yaacob for their support and encourage.

Last but not least, I'm also indebted to my beloved friends Norhafeza, Khairul, Pian, Itha, Ezamy and others, whose friendship I'll forever treasure.



This thesis submitted to the Senate of Universiti Putra Malaysia and has been accepted as fulfillment of the requirement for the degree of Master of Science. The members of the supervisory committee are follows:

# Anuar Kassim, Ph. D.

Professor Chemistry Department Faculty of Science Universiti Putra Malaysia (Chairman)

## Zulkarnain Zainal, Ph. D.

Professor Chemistry Department Faculty of Science Universiti Putra Malaysia (Member)

## Mohamad Zaki Ab Rahman, Ph. D.

Associate Professor Chemistry Department Faculty of Science Universiti Putra Malaysia (Member)



## DECLARATION

I hereby declare that the thesis is based on my original work except for quotations and citations which have been duly acknowledged. I also declare that is has not been previously or concurrently submitted for any degree at UPM or other institutions.

FARIZ BIN ADZMI Date: 23 April 2007



# TABLE OF CONTENTS

APPROVAL DECLARATI LIST OF TAR LIST OF FIG	LEDGEMENTS L FION ABLES	
Ι	INTRODUCTION Declaration	17
	Background Conducting Delemen	17
	Conducting Polymer Problems Statement	19 22
		22
	Objectives of the Study	23
II	LITERATURE REVIEW	
	Polypyrrole	24
	Properties of Polypyrrole	26
	Montmorillonite Clay	
	The Structure	35
	Physical Properties	36
	Polymer Nanocomposites	38
	Polypyrrole/montmorillonite Nanocomposites	39
	Polypyrrole/Al <sub>2</sub> O <sub>3</sub> Nanocomposites	40
	Polypyrrole/zeolite Nanocomposites	41
	Polypyrrole/Fe <sub>3</sub> O <sub>4</sub> Nanocomposites	41
	Epoxy-clay Nanocomposites	41
	Polyurethane-clay Nanocomposites	42
	Polypropylene-clay Hybrids	42
	Polystyrene-clay Nanocomposites	43
	Rubber-clay Hybrids	44
	Applications	45
III	MATERIALS AND METHODS	
	Chemicals	46
	Experimental	
	Effect of the Concentration Oxidant and Pyrrole	46
	Effect of Percentage MMT	47
	Effect of Temperature	48
	Effect of Oxidant	48



	Effect of Preparation Time	48	
	Characterization Methods	40	
	Electrical Conductivity Measurement X-ray Diffraction Analysis (XRD)	48 49	
	Analysis of Surface Area and Porousimetry (ASAP)	49 50	
	Fourier Transfer Infrared Spectroscopy (FTIR)	50	
	Scanning Electron Microscopy (SEM)	50	
	Thermal Gravimetric Analysis (TGA)	52	
IV	RESULTS AND DISCUSSION		
	Conductivity Measurement		
	Effect of Concentration FeCl <sub>3</sub> and Pyrrole	54	
	Effect of Percentage MMT	56	
	Effect of Preparation Temperature	58	
	Effect of Preparation Time	58	
	Effect of Temperature	58	
	X-Ray Diffraction Analysis (XRD)	61 65	
	Effect of Percentage MMT	68	
	Effect of Preparation Temperature	70	
	Effect of Oxidant	71	
	Scanning Electron Microscopy (SEM)	73	
		75	
	Effect of Percentage MMT	76	
	Effect of Preparation Temperature	81	
	Fourier Transfer Infrared Spectroscopy (FTIR)	84	
	Thermal Gravimetric analysis (TGA)	87 89	
	Effect of Percentage MMT	07	
	Effect of Preparation Temperature		
	Surface Area and Porosity Analysis (ASAP)		
V	CONCLUSION AND RECOMMENDATION		
	Conclusion	92	
	Recommendations	93	
REFERENCES		94	
APPENDICES		99	
<b>BIODATA OF T</b>	HE AUTHOR	101	
	a	102	
PUBLICATION		103	
CONFERENCE AND SEMINAR			



# LIST OF TABLES

Table		Page
1	Typical conducting polymer structure.	20
2	Properties changes typically observed upon electrical stimulation to switch conducting polymer between oxidized and reduce state.	21
3	The mole ratio of FeCl <sub>3</sub> and pyrrole monomer.	47
4	Preparation of samples for characterization method.	53
5	The conductivity of the Ppy/MMT nanocomposites by increasing concentration of $FeCl_3$ with fix concentration of pyrrole (0.1M) monomer and percentage of MMT (1%).	55
6	The conductivity of the Ppy/MMT nanocomposites by increasing concentration of pyrrole monomer with fix concentration of FeCl <sub>3</sub> $(0.1M)$ and percentage of MMT (1%).	56
7	The conductivity of Ppy/MMT nanocomposites by increasing percentage of MMT with fix concentration of pyrrole monomer (0.1 M) and FeCl <sub>3</sub> (0.4 M).	57
8	The conductivity of Ppy/MMT nanocomposites prepared in different preparation temperature contains 1% of MMT, 0.1 M pyrrole and 0.4 M FeCl <sub>3</sub> .	59
9	The conductivity of Ppy/MMT nanocomposites against temperature.	60
10	XRD analysis for Ppy/MMT by increasing the percentage amount of MMT.	67
11	XRD observation for Ppy/MMT nanocomposites prepared at different temperature.	68
12	Assignment of FTIR absorption bands of Ppy, MMT and Ppy/MMT nanocomposites.	79
13	TGA analysis for Ppy, MMT, Ppy/MMT nanocomposites.	83
14	TGA analysis for pure Ppy, pure MMT and Ppy/MMT (contain 1% - 7% of MMT).	85



15 alysis for Ppy/MMT nanocomposites with different preparation temperature.



# LIST OF FIGURES

Figure		Page
1	Number of scientific papers published and patents issued in the area of conducting polymers for the past 25 years.	18
2	Scientific papers published on conducting polymers in the last 10 years categorized into various topics.	18
3	Polypyrrole formation.	26
4	Variation of tensile properties of Ppy film with phthalate concentration.	34
5	Variation of tensile properties of Ppy/PTS with applied potential.	34
6	Crystalline structure of montmorillonite showing intermeller water layers.	35
7	The conductivity of Ppy/MMT nanocomposites at different temperature.	61
8	XRD pattern for Ppy, MMT and Ppy/MMT nanocomposites.	63
9	Insertion Ppy in the MMT and expanding of <i>d</i> -spacing.	64
10	XRD pattern for Ppy/MMT nanocomposites by increasing percentage of MMT.	66
11	XRD pattern for Ppy/MMT nanocomposites prepared at different temperature.	69
12	XRD pattern for PpyMMT nanocomposites with different oxidant.	70
13a	SEM micrograph of MMT.	72
13b	SEM micrograph of Ppy.	72
13c	SEM micrograph of Ppy/MMT nanocomposites.	73
14a	SEM micrograph of Ppy/MMT nanocomposites with 3% of MMT.	74
14b	SEM micrograph of Ppy/MMT nanocomposites with 5% of MMT.	74
14c	SEM micrograph of Ppy/MMT nanocomposites with 7% of MMT.	75



15	SEM micrograph of Ppy/MMT nanocomposites low temperature of	
		76
	preparation (~5°C).	

16	The FTIR spectra for pure MMT, pure Ppy and Ppy/MMT nanocomposites.	77
17	FTIR spectrum for Ppy/MMT nanocomposites with different percentage of MMT.	80
18	TGA and DTA analysis for (a) Ppy (b) MMT and (c) Ppy/MMT nanocompoosites.	82
19	TGA and DTA analysis for Ppy/MMT nanocomposites with increasing amount of MMT (from 1% to 7%).	86
20	TGA and DTA of Ppy/MMT nanocomposites with different of preparation temperature between room temperature and low temperature (°C).	88
21a	Hysteresis loops of pure MMT.	90
21b	Hysteresis loops of Ppy/MMT nanocomposites.	90



# LIST OF ABBREVIATIONS

ASAP	Analysis of surface area and porosity
ATBN	Acrylonitrile copolymer
BNSA	5-buthylnaphthalene
CSA	Camphor sulfonic acid
D/M	Dopant/monomer
DMAC	Dimethylacemide
FTIR	Fourier transfer infrared spectroscopy
MBSA	ρ-methylbenzenes sulfonic acid
MMT	Montmorillonite
NaANS	Sodium alkylnaphthalene sulfonate
NaDES	Sodium alkylsulfonate
NSA	β-naphthalene
O/M	Oxidant/monomer
Рру	Polypyrrole
Ppy/MMT	Polypyrrole/montmorillonite nanocomposites
PTS	Polytoluene sulfonic
RIA	Relative humidity
SEM	Scanning electron microscopy
T <sub>g</sub>	Glass transition temperature
TGA	Thermal gravimetric analyses
XRD	X-ray diffraction spectroscopy
ESR	Electron spin resonance



Variable range hopping

VRH



#### CHAPTER I

#### **INTRODUCTION**

#### Background

Since their discovery in the mid-1970's, conducting polymers have been a hot research area for many academic institutions. This research has supported the industrial development of conducting polymer products and provided the fundamental understanding of the chemistry, physics and materials science of these materials. The impact of the field on science in general was recognized in 2000 by the awarding of Nobel Prize for Chemistry to the three discoverer of conducting polymer: Alan MacDiarmid, Alan Heeger and Hiderki Shirakawa.

A research over the past 25 years shows the levels of publications increasing rapidly after 1980 (Figure 1). The data suggest a peak in patenting in the late 1980's, while the rate of scientific publications increases unabated through the end of the 1990's. At the present time, around 40 new journal articles related to the conducting polymer are published every week.

Figure 2 shows the main areas of interest of the paper published on conducting polymer. With almost half of the publications related to the synthesis of conducting polymer. The next largest area of research has been into the physic conduction mechanism, while application for conducting polymer account for fewer than 20% of publication.



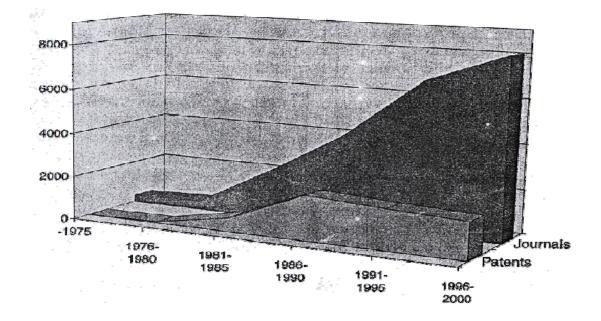


Figure 1: Number of scientific papers published and patents issued in the area of conducting polymers for the past 25 years (Gordon *et al.*, 2003).

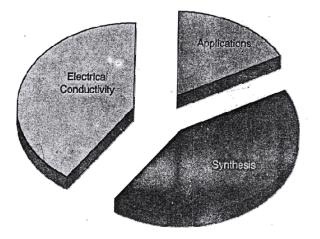


Figure 2: Scientific papers published on conducting polymers in the last 10 tears categorized into various topics (Gordon *et al.*, 2003).



## **Conducting Polymer**

It is possible to create conducting polymers with a diverse range of properties. For example, chemical properties can be manipulated to produce materials capable of trapping simple anions or to render them bioactive. Electrical properties can also be manipulated to produce materials with different conductivities and different redox properties. After synthesis, the properties of these fascinating structures can be manipulated further through redox processes. The application of electrical stimuli can result in drastic changes in the chemical, electrical and mechanical properties in conducting polymers. These complex properties can be controlled only if we understand, first the nature of processes that regulate them during the synthesis of the conducting polymers, and second, the extent to which these properties are changed by the application of an electrical stimulus.

Conducting polymers have emerged as one of most popular in materials science research. It has all the desirable properties:

- They are readily engineered at the molecular level to recognize specific stimuli.
- Because they are conductive, they are facilitating transport of electrical information.

A wide range of conducting polymers is shown in Table 1.



Conducting polymers are of class of materials that are, no doubt, destined to play a major role in intelligent materials science. As outlined in the reminder of this text, the properties of these materials are versatile.

Name	Structure
Polypyrrole (Ppy)	* [ N ]n *
Polythiophene (PTh)	* []n *
Polyaniline (PAn)	*-[-\N-]_n*
Poly(para-phenylene) (PPP)	*-[-\]n*
Poly(phenylene-vinylene) (PPV)	*-[-\
Poly(phenylenesulfide) (PPP)	*-[-{\
Poly(phenylene ethylene) (PPE)	*-[-{

# Table 1: Typical conducting polymer structures.



Table 2 shows the properties of conducting polymer and their behavior that can be manipulated *in situ* using appropriate electronic stimuli.

Property	Typical Change	Potential Application
Conductivity	From $10^{-7}$ to $10^{3}$ S/cm	Electronic components,
		Sensors
Volume	3%	Electromechanical actuators
Color	300-nm shift in absorbance	Displays, smart windows
	band	
Mechanical	Ductile-brittle transition	
Ion permeability	From zero to 10 <sup>-8</sup> molcm <sup>-2</sup> s <sup>-</sup>	<sup>1</sup> Membranes
	in solution	

# Table 2: Property changes typically observed upon electrical stimulation to switch conducting polymer between oxidized and reduced states.



#### **Problems Statement**

Poor processibility of polypyrrole (Ppy) due to its insolubility and infusibility has retarded further investigation on the structure and structure-physical properties. To improve the processibility, many researches have been engaged in the development of soluble or swollen Ppy and dispersible fine-powdered Ppy. At the same time, the electric properties and/or stability of chemically prepared Ppy have also been investigated at many laboratories because the stability of conducting polymers seems to be the main limiting factor in their practical applications.

In recent years, composites of a special category, terms "*nanocomposites*" have been studied with growing interest. For example Ppy base zeolite and Ppy base alumina nanocomposites have extensively study due to improve the processibility of the Ppy and enhance the mechanical properties of Ppy (Zhang *et al.*, 1997).

In this presence of study the Ppy/MMT nanocomposites has been prepared through the chemical polymerization by using FeCl<sub>3</sub> as the oxidation agent for polymerization of pyrrole in aqueous medium. The nanocomposites prepared are characterized in chemical and physical properties.



# **Objectives of the Study**

This thesis presents and discusses the result of series experimental and analyses study of physical and chemical characteristic of polypyrrole/montmorillonite (Ppy/MMT) nanocomposites.

To achieve this aim, the following objectives are proposed:

- 1. To prepare and characterize the Ppy/MMT nanocomposites.
- To study the effect of parameter such as concentration of pyrrole and FeCl<sub>3</sub>, increasing percentage of MMT, oxidant used, preparation temperature and time of preparation on Ppy/MMT nanocomposites.

