



**UNIVERSITI PUTRA MALAYSIA**

***ISOLATION AND STRUCTURAL CHARACTERIZATION OF  
MANNANOLIGOSACCHARIDES FROM AQUEOUS PALM KERNEL  
CAKE EXTRACT***

**NAVEENA REDDY KALIDAS**

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MANNANOLIGOSACCHARIDES FROM AQUEOUS PALM KERNEL CAKE  
EXTRACT**

**By**

**NAVEENA REDDY KALIDAS**

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

**June 2016**

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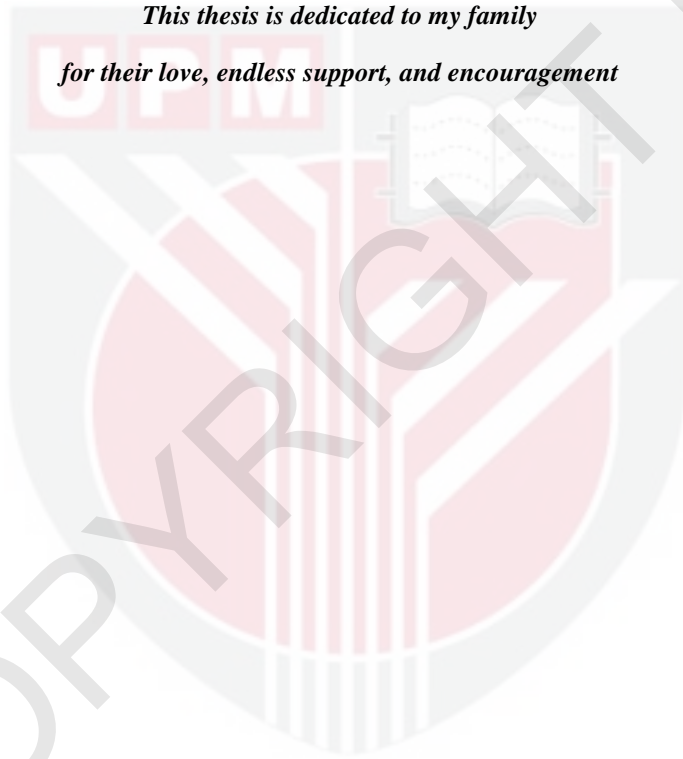
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## DEDICATION

*This thesis is dedicated to my family  
for their love, endless support, and encouragement*



Abstract of thesis presented to the Senate of Universiti Putra Malaysia in fulfilment  
of the requirements for the Degree of Master of Science

## **ISOLATION AND STRUCTURAL CHARACTERIZATION OF MANNANOLIGOSACCHARIDES FROM AQUEOUS PALM KERNEL CAKE EXTRACT**

By

**NAVEENA REDDY KALIDAS**

**June 2016**

**Chairman : Professor Khozirah Shaari, PhD**  
**Institute : Bioscience**

Palm kernel cake (PKC) is the most valuable by-product obtained from the palm kernel oil extraction process. It has been used widely as an animal feed owing to its high protein content. PKC also contains about 81% of non-starch polysaccharides, mainly in the form of mannan-based polymers. Hydrolysis of mannan into low molecular weight mannanoligosaccharides (MOS) has been claimed to have prebiotic properties. The biological mechanism and activities of the MOS are associated with its structure and molecular weight. However, very little information was available on the structural characteristics of MOS of different molecular weights from PKC. Therefore, the objectives of the present study were to isolate MOS of different degree of polymerization (DP) from the crude PKC extract and structurally characterize the MOS using chemical derivatization and spectroscopic methods. The crude PKC extract was obtained by hot water extraction method, followed by delipidation and deproteinization step. The deproteinized PKC extract containing mixtures of MOS were then separated into individual compounds based on their molecular weights using refractive index high performance liquid chromatography (RI-HPLC). The molecular weights of the MOS were determined using electrospray ionization mass spectrometry (ESI-MS/MS). The structures of the isolated MOS were investigated using a combination of chemical analyses such as sugar composition analysis and methylation analysis followed by gas chromatography mass spectrometry (GC-MS), and other spectroscopic methods such as nuclear magnetic resonance spectroscopy (NMR). The MOS mixtures were separated into four individual major compounds with different DP, designated as **MOS-III**, **MOS-IV**, **MOS-V** and **MOS-VI**. The molecular weights of the isolated MOS as determined by ESI-MS/MS were 689, 851, 1013 and 1151 Dalton (Da) corresponding to tetra-, penta-, hexa- and heptasaccharide of the **MOS-III**, **MOS-IV**, **MOS-V** and **MOS-VI**, respectively. Sugar analysis of the isolated MOS indicated the presence of mannose in each of the oligomers. Methylation and 1D/2D NMR analysis showed that the MOS have a linear structure consisting of (1→4)- $\beta$ -D-mannopyranosyl residues with DP ranging from 4 to 7. They were identified as: a)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp, b)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-

Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp, c) β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp, d) β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp-(1→4)-β-D-Manp. In conclusion, the present study revealed a successful application of chromatography, chemical analysis, ESI-MS/MS and NMR to the isolation and characterization of MOS fraction from PKC.

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

## **PENGASINGAN DAN PENCIRIAN STRUKTUR MANNANOLIGOSAKARIDA DARIPADA ISIRONG KELAPA SAWIT**

Oleh

**NAVEENA REDDY KALIDAS**

**Jun 2016**

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**Institut : Biosains**

Isirong kelapa sawit (PKC), yang diperolehi daripada proses pengekstrakan minyak isirong kelapa sawit merupakan hasil sampingan yang paling berguna. PKC digunakan secara meluas sebagai makanan haiwan kerana mempunyai kandungan protein yang tinggi. Selain itu, PKC juga mengandungi kira-kira 81% polisakarida bukan kanji dalam bentuk polimer mannan. Hidrolisis polimer mannan kepada karbohidrat berat molekul rendah seperti mannanoligosakarida (MOS) mempunyai ciri-ciri prebiotik. Mekanisme biologi dan aktiviti MOS bergantung kepada struktur dan berat molekul. Walau bagaimanapun, maklumat tentang struktur pecahan MOS yang terdiri daripada berat molekul yang berbeza daripada PKC adalah sangat sedikit. Oleh itu, objektif kajian ini adalah untuk mengasingkan MOS daripada ekstrak PKC mengikut darjah pempolimeran (DP) dan mencirikan struktur MOS dengan menggunakan kaedah kimia dan spektroskopi. Ekstrak PKC diperolehi melalui pengekstrakan air panas, dan kemudiannya diasingkan daripada lipid dan protein. Ekstrak PKC yang mengandungi campuran MOS kemudian dipisahkan kepada pecahan-pecahan MOS berdasarkan kepada berat molekul dengan menggunakan kromatografi cecair berprestasi tinggi (HPLC). Berat molekul MOS ditentukan dengan menggunakan teknik pengionan elektrosemburan spektroskopi jisim (ESI-MS/MS). Struktur MOS dikaji melalui gabungan analisis kimia seperti analisis komposisi gula dan analisis metilasi diikuti dengan kromatografi gas spektroskopi jisim (GC-MS), dan kaedah spektroskopi lain seperti spektroskopi resonans magnetik nuklear (NMR). Campuran MOS telah diasingkan kepada empat pecahan utama yang mempunyai DP berbeza, dinamakan sebagai **MOS-III**, **MOS-IV**, **MOS-V** dan **MOS-VI**. Berat molekul **MOS-III**, **MOS-IV**, **MOS-V** dan **MOS-VI** seperti yang ditentukan oleh ESI-MS/MS adalah 689, 851, 1013 dan 1151 Dalton (Da) yang dikenal pasti sebagai tetra-, penta-, hexa- dan heptasakarida, masing-masing. Analisis gula daripada MOS menunjukkan kehadiran mannosa dalam setiap satu oligomer. Metilasi dan analisis 1D/2D NMR menunjukkan bahawa MOS terdiri daripada satu struktur linear yang terdiri daripada (1→4)- $\beta$ -D-mannopiranosil yang mempunyai DP 4–7. Struktur MOS telah dikenal pasti sebagai: a)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp, b)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp, c)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-

(1→4)- $\beta$ -D-Manp, d)  $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp-(1→4)- $\beta$ -D-Manp. Kesimpulannya, hasil kajian mendapati bahawa kromatografi, analisis kimia, ESI-MS/MS dan NMR berjaya mengasingkan dan mencirikan campuran MOS daripada PKC.





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This thesis was submitted to the Senate of Universiti Putra Malaysia and has been accepted as fulfilment of the requirement for the degree of Master of Science. The members of the Supervisory Committee were as follows:

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## Declaration by Members of Supervisory Committee

This is to confirm that:

- 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.

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## TABLE OF CONTENTS

	Page
<b>ABSTRACT</b>	i
<b>ABSTRAK</b>	iii
<b>ACKNOWLEDGEMENTS</b>	v
<b>APPROVAL</b>	vi
<b>DECLARATION</b>	viii
<b>LIST OF TABLES</b>	xiii
<b>LIST OF FIGURES</b>	xiv
<b>LIST OF ABBREVIATIONS</b>	xix
 <b>CHAPTER</b>	
<b>1 INTRODUCTION</b>	<b>1</b>
<b>2 LITERATURE REVIEW</b>	
2.1 The oil palm	3
2.2 Palm kernel cake	4
2.2.1 Chemical composition of palm kernel cake	4
2.2.2 Carbohydrate content in palm kernel cake	5
2.2.2.1 Cellulose	5
2.2.2.2 Mannan	6
2.3 Mannanoligosaccharides	7
2.3.1 Beneficial effects of mannanoligosaccharides	8
2.3.1.1 Alternative to antibiotics growth promotants	8
2.3.1.2 Gastrointestinal health promotants	8
2.3.1.3 Immunomodulatory activities	9
2.3.2 Extraction of mannanoligosaccharides	9
2.3.3 Purification of mannanoligosaccharides	9
2.3.4 Fractionation of mannanoligosaccharides	10
2.3.4.1 High performance liquid chromatography	10
2.3.4.2 Gel filtration chromatography	11
2.3.4.3 Membrane Separation	12
2.4 Structural characterization of mannanoligosaccharides	12
2.4.1 Nuclear magnetic resonance spectroscopy	12
2.4.2 Mass spectroscopy	14
2.4.2.1 Matrix assisted laser desorption ionization time of flight spectroscopy	14
2.4.2.2 Electron impact ionization mass spectroscopy	15
2.4.2.3 Fast atom bombardment mass spectroscopy	15
2.4.2.4 Chemical ionization mass spectrometry	15
2.4.3 Hyphenated mass spectrometry technique	16
2.4.3.1 Gas chromatography mass	16

	spectrometry	
2.4.3.2	Liquid chromatography mass spectrometry	16
2.4.4	Chemical methods	17
2.4.4.1	Acid hydrolysis	17
2.4.4.2	Methylation	20
2.4.5	Infrared spectroscopy	22
<b>3</b>	<b>MATERIALS AND METHODOLOGY</b>	
3.1	General instrumentation	23
3.2	Sample material	23
3.3	Extraction and fractionation of PKC crude extract	23
3.4	De-proteinization of MOS fraction	25
3.5	RP-HPLC separation of the deproteinized MOS fraction	25
3.6	Analysis of monosaccharide composition	25
3.7	Analysis of glycosidic linkage	26
3.8	Physical and spectral data of the isolated oligosaccharides	26
3.8.1	$\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp ( <b>MOS-III</b> )	26
3.8.2	$\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp ( <b>MOS-IV</b> )	27
3.8.3	$\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp ( <b>MOS-V</b> )	27
3.8.4	$\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp ( <b>MOS-VI</b> )	28
<b>4</b>	<b>RESULTS AND DISSCUSSION</b>	
4.1	HPLC chromatographic profile of PKC oligosaccharides	29
4.2	Isolation and structural elucidation of MOS from PKC	32
4.2.1	Characterization of <b>MOS-III</b> as $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp	32
4.2.2	Characterization of <b>MOS-IV</b> as $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp	59
4.2.3	Characterization of <b>MOS-V</b> as $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp	69
4.2.4	Characterization of <b>MOS-VI</b> as $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp-(1 $\rightarrow$ 4)- $\beta$ -D-Manp	79
<b>5</b>	<b>CONCLUSION AND RECOMMENDATIONS FOR FUTURE RESEARCH</b>	89

<b>REFERENCES</b>	90
<b>APPENDICES</b>	100
<b>BIODATA OF STUDENT</b>	115
<b>LIST OF PUBLICATIONS</b>	116



## LIST OF TABLES

Table		Page
2.1	Chemical Compositions of PKC	5
2.2	Applications of different gel matrices	12
2.3	Different hydrolysis, reduction and acetylation conditions for a variety of polysaccharides	19
4.1	The $^1\text{H}$ NMR <sup>a</sup> and $^{13}\text{C}$ NMR <sup>b</sup> chemical shifts of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C	46
4.2	TOCSY correlation of <b>MOS-III</b>	46
4.3	HMBC correlations of <b>MOS-III</b>	47
4.4	NOESY data for <b>MOS-III</b>	57
4.5	The $^1\text{H}$ NMR <sup>a</sup> and $^{13}\text{C}$ NMR <sup>b</sup> chemical shifts of <b>MOS-IV</b> in $\text{D}_2\text{O}$ at 37 °C	68
4.6	The $^1\text{H}$ NMR <sup>a</sup> and $^{13}\text{C}$ NMR <sup>b</sup> chemical shifts of <b>MOS-V</b> in $\text{D}_2\text{O}$ at 37 °C	78
4.7	The $^1\text{H}$ NMR <sup>a</sup> and $^{13}\text{C}$ NMR <sup>b</sup> chemical shifts of <b>MOS-VI</b> in $\text{D}_2\text{O}$ at 37 °C	88

## LIST OF FIGURES

Figure		Page
2.1	A cross section of the oil palm fruit	4
2.2	Mannan polymer consisting of mannose residues linked by $\beta$ -(1 $\rightarrow$ 4) linkages	6
2.3	Galactomannan structure consisting of $\beta$ -(1 $\rightarrow$ 4) linked mannose unit with $\alpha$ -1,6 linked galactose residues in the side chains.	6
2.4	Galactoglucomannan structure consisting of $\beta$ -(1 $\rightarrow$ 4) linked mannose and glucose unit with $\alpha$ -(1 $\rightarrow$ 6) linked galactose residues in the side chains.	7
2.5	Glucomannan structure consisting of $\beta$ -(1 $\rightarrow$ 4) linked mannose and glucose residues	7
2.6	Formation of alditol acetates for monosaccharide analysis	18
2.7	Fragmentation patterns of alditol acetates	18
2.8	Formation of PMAAs for linkage analysis	21
2.9	Fragmentation pattern of PMAAs	21
3.1	General flow of extraction, purification and structural analysis of MOS from PKC	24
4.1	RI-HPLC profile of MOS fraction	29
4.2	HPLC profile of <b>MOS-III</b>	30
4.3	HPLC profile of <b>MOS-IV</b>	30
4.4	HPLC profile of <b>MOS-V</b>	31
4.5	HPLC profile of <b>MOS-VI</b>	31
4.6	GC chromatogram of alditol acetates of <b>MOS-III</b>	33
4.7	MS spectrum of alditol acetates of <b>MOS-III</b>	33
4.8	MS fragmentation of glucose/galactose/mannose sugar alditol acetate monomer.	34
4.9	Comparison of <b>MOS-III</b> with sugar alditol acetate standards (mannose, galactose and glucose)	35
4.10	GC chromatogram of PMAAs of <b>MOS-III</b>	35



4.11	MS spectrum of PMAAs of <b>MOS-III</b> (Peak 1)	36
4.12	MS spectrum of PMAAs of <b>MOS-III</b> (Peak 2)	36
4.13	$^1\text{H}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 25 °C [Expansion]	38
4.14	$^{13}\text{C}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 25 °C [Expansion]	39
4.15	HSQC spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 25 °C [Expansion]	40
4.16	$^1\text{H}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 1]	41
4.17	$^1\text{H}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 2]	42
4.18	$^{13}\text{C}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 1]	43
4.19	$^{13}\text{C}$ NMR spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 2]	44
4.20	HSQC spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 1]	48
4.21	HSQC spectrum (Anomeric region) of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 2]	49
4.22	HSQC spectrum (Non-anomeric region) of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 3]	50
4.23	HMBC spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion 1]	51
4.24	HMBC spectrum of <b>MOS-III</b> (Anomeric region) in $\text{D}_2\text{O}$ at 37 °C (Expansion 2)	52
4.25	HMBC spectrum of <b>MOS-III</b> (Non-anomeric region) in $\text{D}_2\text{O}$ at 37 °C (Expansion 3)	53
4.26	TOCSY spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion]	54
4.27	Selected HMBC ( $\rightarrow$ ) correlations of <b>MOS-III</b>	55
4.28	NOESY spectrum of <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C [Expansion]	56
4.29	ESI-MS/MS spectrum of <b>MOS-III</b>	57
4.30	Systematic nomenclature for mass fragmentation of oligosaccharides	58
4.31	Schematic fragmentation pathways of the <b>MOS-III</b>	58
4.32	ESI-MS/MS spectrum of <b>MOS-IV</b>	60
4.33	GC chromatogram of alditol acetates of <b>MOS-IV</b>	60
4.34	MS spectrum of alditol acetates of <b>MOS-IV</b>	61

4.35	Comparison of <b>MOS-IV</b> with sugar alditol acetate standards	61
4.36	GC chromatogram of PMAAs of <b>MOS-IV</b>	62
4.37	MS spectrum of PMAAs of <b>MOS-IV</b> (Peak 1)	62
4.38	MS spectrum of PMAAs of <b>MOS-IV</b> (Peak 2)	63
4.39	$^1\text{H}$ NMR spectrum of <b>MOS-IV</b> in $\text{D}_2\text{O}$ at 37 °C	64
4.40	Comparison of $^1\text{H}$ NMR spectrum of <b>MOS-IV</b> with <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C	65
4.41	$^{13}\text{C}$ NMR spectrum of <b>MOS-IV</b> in $\text{D}_2\text{O}$ at 37 °C	66
4.42	Comparison of $^{13}\text{C}$ NMR spectrum of <b>MOS-IV</b> with <b>MOS-III</b> in $\text{D}_2\text{O}$ at 37 °C	67
4.43	Schematic fragmentation pathways of <b>MOS-IV</b>	68
4.44	ESI-MS/MS spectrum of <b>MOS-V</b>	70
4.45	GC chromatogram of alditol acetates of <b>MOS-V</b>	70
4.46	MS spectrum of alditol acetates of <b>MOS-V</b>	71
4.47	Comparison of <b>MOS-V</b> with sugar alditol acetate standards	71
4.48	GC chromatogram of PMAAs of <b>MOS-V</b>	72
4.49	MS spectrum of PMAAs of <b>MOS-V</b> (Peak 1)	72
4.50	MS spectrum of PMAAs of <b>MOS-V</b> (Peak 2)	73
4.51	$^1\text{H}$ NMR spectrum of <b>MOS-V</b> in $\text{D}_2\text{O}$ at 37 °C	74
4.52	$^1\text{H}$ NMR comparison of <b>MOS-V</b> with <b>MOS-IV</b> in $\text{D}_2\text{O}$ at 37 °C	75
4.53	$^{13}\text{C}$ NMR spectrum of <b>MOS-V</b> in $\text{D}_2\text{O}$ at 37 °C	76
4.54	Comparison of $^{13}\text{C}$ NMR spectrum of <b>MOS-V</b> with <b>MOS-IV</b> in $\text{D}_2\text{O}$ at 37 °C	77
4.55	Schematic fragmentation pathways of <b>MOS-V</b>	78
4.56	ESI-MS/MS spectrum of <b>MOS-VI</b>	80
4.57	GC chromatogram of alditol acetates of <b>MOS-VI</b>	80
4.58	MS spectrum of alditol acetates of <b>MOS-VI</b>	81
4.59	Comparison of <b>MOS-VI</b> with alditol acetates standards	81

4.60	GC chromatogram of PMAAs of <b>MOS-VI</b>	82
4.61	MS spectrum of PMAAs of <b>MOS-VI</b> (Peak 1)	83
4.62	MS spectrum of PMAAs of <b>MOS-VI</b> (Peak 2)	83
4.63	$^1\text{H}$ NMR spectrum of <b>MOS-VI</b> in $\text{D}_2\text{O}$ at $37^\circ\text{C}$	84
4.64	Comparison of $^1\text{H}$ NMR spectrum of <b>MOS-VI</b> with <b>MOS-V</b> in $\text{D}_2\text{O}$ at $37^\circ\text{C}$	85
4.65	$^{13}\text{C}$ NMR spectrum of <b>MOS-VI</b> in $\text{D}_2\text{O}$ at $37^\circ\text{C}$	86
4.66	Comparison of $^{13}\text{C}$ NMR spectrum of <b>MOS-VI</b> with <b>MOS-V</b> in $\text{D}_2\text{O}$ at $37^\circ\text{C}$	87
4.67	Schematic fragmentation pathways of <b>MOS-VI</b>	88

## LIST OF APPENDICES

Appendix		Page
1	HPLC profile of <b>MOS-I</b>	100
2	HPLC profile of <b>MOS-II</b>	100
3	HPLC profile of <b>MOS-VII</b>	101
4	HPLC profile of <b>MOS-VIII</b>	101
5	<sup>1</sup> H NMR spectrum of <b>MOS-I</b> (Sucrose)	102
6	HSQC spectrum of <b>MOS-IV</b> in D <sub>2</sub> O at 37 °C	103
7	HMBC spectrum of <b>MOS-IV</b> in D <sub>2</sub> O at 37 °C	104
8	ROESY spectrum of <b>MOS-IV</b> in D <sub>2</sub> O at 37 °C	105
9	TOCSY spectrum of <b>MOS-IV</b> in D <sub>2</sub> O at 37 °C	106
10	HSQC spectrum of <b>MOS-V</b> in D <sub>2</sub> O at 37 °C	107
11	HMBC spectrum of <b>MOS-V</b> in D <sub>2</sub> O at 37 °C	108
12	ROESY spectrum of <b>MOS-V</b> in D <sub>2</sub> O at 37 °C	109
13	TOCSY spectrum of <b>MOS-V</b> in D <sub>2</sub> O at 37 °C	110
14	HSQC spectrum of <b>MOS-VI</b> in D <sub>2</sub> O at 37 °C	111
15	HMBC spectrum of <b>MOS-VI</b> in D <sub>2</sub> O at 37 °C	112
16	ROESY spectrum of <b>MOS-VI</b> in D <sub>2</sub> O at 37 °C	113
17	TOCSY spectrum of <b>MOS-VI</b> in D <sub>2</sub> O at 37 °C	114

## LIST OF ABBREVIATIONS

°C	- Degree Celcius
<sup>13</sup> C	- Carbon-13
<sup>1</sup> H	- Proton
ACN	- Acetonitrile
API	- Atmospheric pressure ionization
C <sub>5</sub> D <sub>5</sub> N	- Deuterated Pyridine
CDCl <sub>3</sub>	- Deuterated Chloroform
CI	- Chemical Ionization
D <sub>2</sub> O	- Deuterium Oxide
DMSO	- Dimethylsulfoxide
DP	- Degree of Polymerization
DSS	- 3-(trimethylsilyl) propane-1-sulfonic acid
EI	- Electron Impact Ionization
ESIMS	- Electrospray Ionization Mass Spectroscopy
eV	- Electron Volt
FAB	- Fast Atom Bombardment
FTIR	- Fourier Transform Infra-Red
GC-MS	- Gas Chromatography Mass Spectroscopy
HCl	- Acid Hydrochloric
HMBC	- Heteronuclear Multiple Bond Correlation
HPAEC	- High Performance Anion Exchange Chromatography
HPLC	- High Performance Liquid Chromatography
HSQC	- Heteronuclear Single Quantum Coherence
Hz	- Hertz
IgA	- Immunoglobulin A
IgG	- Immunoglobulin G
<i>m/z</i>	- Mass per charge
MALDI	- Matrix-Assisted Laser Desorption Ionization
MHz	- Megahertz
MOS	- Mannan oligosaccharides
MW	- Molecular Weight
NaBH <sub>4</sub>	- Sodium borohydride
NaOH	- Sodium hydroxide
NCP	- Non Cellulose Polysaccharide
NMR	- Nuclear Magnetic Resonance Spectroscopy
NSP	- Non Starch Polysaccharides
PAD	- Pulse Amperometric Detector
PKC	- Palm Kernel Cake
PMAA	- Partially Methylated Alditol Acetates

TFA	- Trifluoroacetic acid
TMS	- Tetramethylsilane
TOCSY	- Total Correlation Spectroscopy
TSP	- Trisodium Phosphate



## CHAPTER 1

### INTRODUCTION

Lignocellulosic biomass is the most abundant and renewable biopolymer in nature (Zhou *et al.*, 2011). It is composed predominantly of polysaccharides such as cellulose, hemicellulose, and lignin. It has been estimated that  $10\text{--}50 \times 10^9$  tons of lignocellulosic biomass are produced annually from agro-based industries (i.e., forestry, agricultural activities, timber industries, and pulp and paper industries) (De Vries and Visser, 2001). Much of these lignocellulosic wastes are usually disposed through burning activity, which imposes serious environmental pollution issues (Levine, 1996). However, the vast amount of plant biomass considered as waste could potentially be converted into various value added bio-products including animal feed, organic chemicals, biofuel, energy sources for fermentation and human nutrients (Howard *et al.*, 2003).

Malaysia, being the second largest palm oil producer produces huge quantities of oil palm by-products. Among the many secondary products from oil palm, palm kernel cake (PKC) has been regarded as the most promising. It has been reported that PKC contains about 81% of non-starch polysaccharides (NSPs), mainly in the form of mannan-based polymers (Knudsen, 1997). Mannans are presumably characterised as anti-nutritional polysaccharides, which are highly crystalline, extremely hard and insoluble in water (Sundu and Dingle, 2002). Moreover, other anti-nutritional substances such as galactomannan, xylan, and arabinoxylan are also most likely present in PKC. Several studies have reported that guar and wheat also contain these anti-nutritional compounds which could increase the viscosity of the animal feeds due to its high water absorbing tendency; thus causing poor nutrient absorption in the gastrointestinal tract (Dingle, 1995; Kumar *et al.*, 1997). Owing to its high fibre content, coupled with the presence of anti-nutritional compounds, application of PKC as feed supplements in non-ruminant diets is limited. However, recent studies have found that hydrolysis of mannan polymers from PKC into lower level carbohydrates such as oligosaccharides have been proven to improve the nutritive value of PKC.

Oligosaccharides are sugar polymers consisting of 3 to 10 monosaccharide residues linked by O- or N-linked glycosidic bonds (Tomomatsu, 1994). However, the degree of polymerization (DP) can go up to 60 for some oligosaccharides, for example chicory inulin, or down to 2, for example lactulose (Crittenden and Plain, 1996). Most oligosaccharides are low molecular weight carbohydrates. Oligosaccharides could be attached to amino acids or lipids, forming glycoprotein and glycolipids, respectively. Oligosaccharides have greater structural diversity due to the number of monomers, differences in the anomeric configuration, glycosidic linkages, ring size, substitution points, and branching points. As a result of the structural diversity, the flexibility, compactness, physical, biological and biochemical properties among the oligosaccharides are varied.



Oligosaccharides have been reported to exhibit significant biological properties such as immunostimulatory, antiviral, and antitumor activities (Kamasuka *et al.*, 1968; Tizard *et al.*, 1989; Ezekowitz *et al.*, 1989). However, the biological activities and mechanisms of action of oligosaccharides are not fully known but believed to be very closely related to its structure. For example, mannan from *Saccharomyces cerevisiae*, which is composed of high molecular weight partially phosphorylated mannose residues linked via  $\alpha$ -(1→6),  $\alpha$ -(1→3), and  $\alpha$ -(1→2) linkages was not active in stimulating the interferon release. Whereas, mannan from *Candida albicans*, which is composed of low molecular weight and highly branched molecule with short chains of  $\alpha$ -(1→2), linked mannose monomers, joined by  $\alpha$ -(1→6) linkages and attached to peptides, are very active in the interferon-inducing assay (DeClercq *et al.*, 1970). Suzuki *et al.* (1968) postulated that the differences in the interferon activity could be due to small structural or size variations among the polymers.

In recent years, there has been a growing interest in the use of oligosaccharides as prebiotics. Oligosaccharides are classified as prebiotics because it can withstand the digestion enzymes in the stomach and pass into the large bowel where it is selectively metabolized by beneficial bacteria. Prebiotics are defined as “selectively fermented ingredients that allow specific changes, both in the composition and/or activity in the gastrointestinal microbiota that confers benefits upon host well-being and health” (Roberfroid, 2007). Fermentation of oligosaccharides by the microbiota are strongly influenced by the chemical structures and the identity of the monomeric sugar units, DP, type of glycosidic linkages between the monomers, complexity of the fraction (branched or linear) and possible linkage to non-carbohydrates. Several types of oligosaccharide substances, which have been reported to possess prebiotic effects, such as fructooligosaccharides (FOS), galactooligosaccharides (GOS), glucooligosaccharides, mannanoligosaccharides (MOS), isomaltooligosaccharides (ISO), xylooligosaccharides (XOS), gentiooligosaccharides (GTO) and inulin (Crittenden and Playne, 1996; Rycroft *et al.*, 2001; Desai *et al.*, 2004; Pennacchia *et al.*, 2006). Recently, some studies have shown that enzyme-treated PKC releases mannose and mannose-based oligosaccharides, which could be a potential source of prebiotic (Saenphoom *et al.*, 2011; 2013). Chen *et al.* (2015) demonstrated that all three *Lactobacillus* strains showed better growth rate in the enzyme-treated crude PKC, owing to low DP oligosaccharides produced during the hydrolysis of NSPs. However, the nature and the exact type of oligosaccharides which are responsible for the growth of *Lactobacillus* could not be confirmed. Although, there have been some structural studies conducted on the mannans of PKC, the available information is only on the overall carbohydrate polymer with little information on the structural characteristics of MOS fractions of different molecular weights. Therefore, this study was conducted to provide this lack of structural information. To achieve this, the study was initiated with the following objectives:

1. To isolate and purify the oligosaccharides from the aqueous PKC extract.
2. To characterize the oligosaccharides using chemical and spectroscopic methods.



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## LIST OF PUBLICATIONS

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Naveena Reddy Kalidas, Saminathan Mookiah, Intan Safinar Ismail, Faridah Abas, Prasenjit Maity, Syed Sirajul Islam, Nurhuda Manshoor, Khozirah Shaari. (2017). Structural characterization and evaluation of prebiotic activity of oil palm kernel cake mannanoligosaccharides. *Food Chemistry*. (Under Review)

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Naveena Reddy Kalidas, Khozirah Shaari, Intan Safinar Ismail, and Faridah Abas (2015). Isolation, Fractionation, and characterization of oligosaccharides extracted from Palm Kernel Cake (PKC) using Infrared Spectroscopy (IR). International National Conference of Natural Products, pp 54.



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