



**UNIVERSITI PUTRA MALAYSIA**

**CHEMICAL CONSTITUENTS AND BIOLOGICAL ACTIVITY OF ASAM  
AUR AUR (*GARCINIA PARVIFOLIA*) AND JINGGAU (*PLOIARIUM  
ALTERNIFOLIUM*)**

**NG SOOK HAN**

**FS 2007 22**

**CHEMICAL CONSTITUENTS AND BIOLOGICAL ACTIVITY OF  
ASAM AUR AUR (*GARCINIA PARVIFOLIA*) AND JINGGAU  
(*PLOIARIUM ALTERNIFOLIUM*)**

**By**

**NG SOOK HAN**

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

**June 2007**



**CHEMICAL CONSTITUENTS AND  
BIOLOGICAL ACTIVITY OF ASAM AUR AUR  
(*GARCINIA PARVIFOLIA*) AND JINGGAU  
(*PLOIARIUM ALTERNIFOLIUM*)**

**NG SOOK HAN**

**MASTER OF SCIENCE  
UNIVERSITI PUTRA MALAYSIA**

**2007**



## DEDICATION

To my beloved parents Ng Kum Seong and Lik Chee Keon

For their endless love and concern.....

To my beloved Loke Chee Keong

For his romantic love, support, understanding and care.....

To my supervisor Assoc. Prof. Dr. Gwendoline Ee Cheng Lian

For her guidance, advice, understanding and endless support.....

To my co-supervisors Assoc. Prof. Dr. Mohd Aspollah B Hj Md Sukari

Dr. Emily Goh Joo Kheng

For their kindly advice and indispensable support.....

To my senior Dr. Lim Chan Kiang

For his wonderful encouragement and support.....

To my friends

For their wonderful love and generous moral support.....



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

**CHEMICAL CONSTITUENTS AND BIOLOGICAL ACTIVITY OF  
ASAM AUR AUR (*GARCINIA PARVIFOLIA*) AND JINGGAU  
(*PLOIARIUM ALTERNIFOLIUM*)**

By

**NG SOOK HAN**

**June 2007**

**Chairman : Associate Professor Gwendoline Ee Cheng Lian, PhD**

**Faculty : Science**

Chemical and biological studies were carried out on two plants, *Garcinia parvifolia* (Guttiferae) and *Ploiarium alternifolium* (Theaceae). The chemical investigations covered xanthenes, benzophenones, anthraquinones, and triterpenes.

The stem bark of *Garcinia parvifolia* and *Ploiarium alternifolium* were investigated and this resulted in the isolation of ten known compounds. These compounds were isolated using common chromatographic techniques and were identified by using spectroscopic experiments such as NMR, MS, IR and UV.

Detailed chemical studies on *Garcinia parvifolia* have yielded two triterpenoids, stigmasterol and  $\beta$ -sitosterol, three xanthenes, 6-deoxyjacareubin, daphnifolin and



rubraxanthone, one benzophenone, isoxanthochymol and one alkaloid, caffeine. Meanwhile, investigations on *Ploiarium alternifolium* have afforded three anthraquinones, ploiariquinone A, emodin and 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone.

The larvicidal test was carried out towards the larvae of *Aedes aegypti*. The crude chloroform, ethyl acetate and methanol extracts of *Garcinia parvifolia* were weakly active against the larvae of *Aedes aegypti* with LC<sub>50</sub> values of 204.26, 194.96 and 236.44 µg/ml respectively while the crude chloroform extract of *Ploiarium alternifolium* was weakly active against the larvae of *Aedes aegypti* with an LC<sub>50</sub> value of 159.12 µg/ml.

The antimicrobial assay was carried out towards four pathogenic bacteria, Methicillin Resistant *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Staphylococcus choleraesuis* and *Bacillus subtilis*. For the microbes MRSA, *P. aeruginosa* and *S. choleraesuis* there were no inhibition by all the crude extracts of *Garcinia parvifolia*. The crude chloroform and methanol extracts of *Ploiarium alternifolium* showed no activity towards the microbe *S. choleraesuis*.

Cytotoxic tests were performed using HL-60 Cell Line. The crude hexane, chloroform and ethyl acetate extracts of *Garcinia parvifolia* were considered to be active against the HL-60 cell line with IC<sub>50</sub> values of less than 30 µg/ml. The



crude chloroform extract of *Ploiarium alternifolium* was also considered to be active against the HL-60 cell line with an IC<sub>50</sub> value of 23.3 µg/ml. The crude methanol extract of *Ploiarium alternifolium* showed the most significant activity with an IC<sub>50</sub> value of 5.2 µg/ml.

The antifungal activity testing of the plant extracts were carried out against the fungi *Candida albican*, *Aspergillus ochraceaus*, *Sacchoromyces cerevisiae* and *Candida lypolytica*. No activity was observed for all the crude extracts.



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

**KANDUNGAN KIMIA DAN ACTIVITI BIOLOGI DARIPADA ASAM  
AUR AUR (*GARCINIA PARVIFOLIA*) DAN JINGGAU (*PLOIARIUM  
ALTERNIFOLIUM*)**

Oleh

**NG SOOK HAN**

**Jun 2007**

**Pengerusi : Profesor Madya Gwendoline Ee Cheng Lian, PhD**

**Fakulti : Sains**

Kajian kimia dan aktiviti biologi telah dijalankan ke atas dua jenis tumbuhan iaitu *Garcinia parvifolia* (Guttiferae) dan *Ploiarium alternifolium* (Theaceae). Kajian kimia terperinci merangkumi jenis sebatian seperti xanton, benzofenon, antrakuinon dan triterpena.

Kulit pokok *Garcinia parvifolia* dan *Ploiarium alternifolium* telah dikaji dan berjaya menghasilkan sepuluh sebatian yang telah dikenalpasti. Struktur sebatian-sebatian ini ditentukan dengan menggunakan eksperimen spektroskopi seperti NMR, MS, IR dan UV.



Kajian kimia terperinci ke atas *Garcinia parvifolia* telah menghasilkan dua triterpenoid, stigmasterol dan  $\beta$ -sitosterol, tiga xanton, 6-dioksijakarubin, dafnifolin dan rubraxanton, satu benzofenon, isoxantoximol dan satu alkaloid, kaffeine. Sementara itu, kajian ke atas *Ploiarium alternifolium* telah menghasilkan tiga antrakuinon, ploiarikuinon A, emodin dan 1,8-dihidroksi-3-mektosi-6-metil-antrakuinon.

Ujian larva telah dijalankan dengan menggunakan larva *Aedes aegypti*. Ekstrak mentah kloroform, etil asetat dan metanol *Garcinia parvifolia* mempunyai aktiviti yang lemah terhadap larva *Aedes aegypti* dengan nilai  $LC_{50}$  masing-masing 204.26, 194.96 and 236.44  $\mu\text{g/ml}$ . Sementara itu, ekstrak mentah kloroform *Ploiarium alternifolium* juga menunjukkan aktiviti yang lemah dengan nilai  $LC_{50}$  159.12  $\mu\text{g/ml}$ .

Ujian anti-mikrob telah dijalankan dengan menggunakan empat jenis bakteria iaitu Methicillin Resistant *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Staphylococcus choleraesuis* and *Bacillus subtilis*. Untuk bakteria MRSA, *P. aeruginosa* and *S. choleraesuis* didapati tidak aktif terhadap semua ekstrak mentah *Garcinia parvifolia*. Ekstrak mentah kloroform dan metanol *Ploiarium alternifolium* juga tidak menunjukkan sebarang aktiviti terhadap bakteria *S. choleraesuis*.

Ujian sitotoksik telah dijalankan dengan menggunakan sel HL-60. Ekstrak mentah heksana, kloroform dan etil asetat *Garcinia parvifolia* dianggap sebagai aktif ke atas sel HL-60 dengan nilai  $IC_{50}$  kurang daripada 30  $\mu\text{g/ml}$ . Ekstrak mentah kloroform *Ploiarium alternifolium* juga dianggap aktif ke atas sel HL-60 dengan nilai  $IC_{50}$  23.3  $\mu\text{g/ml}$ . Ekstrak mentah metanol *Ploiarium alternifolium* menunjukkan aktiviti yang kuat dengan nilai  $IC_{50}$  5.2  $\mu\text{g/ml}$ .

Aktiviti anti-kulat ekstrak tumbuhan telah dijalankan ke atas *Candida albican*, *Aspergillus ochraceus*, *Sacchoromyces cerevisiae* and *Candida lypolytica*. Tiada aktiviti diperhatikan ke atas semua ekstrak mentah.

## ACKNOWLEDGEMENTS

The completion of this project is not a one-man work. It is a project, which could only get completed on time with the help of many parties. Therefore I would like to take this opportunity to express my gratitude to all of them to show my appreciation for their support.

First and foremost, I wish to thank Associate Professor Dr. Gwendoline Ee Cheng Lian for being such a good supervisor and personal tutor. Her guidance leads me to some solutions, which I would never come across by myself. She is responsible for involving me in the natural product research in the first place and is always there to show me different ways to approach a research problem and the need to be persistent to accomplish any goal.

I would also like to express my sincere thanks to my co-supervisor Associate Professor Dr. Mohd Aspollah B Hj Md Sukari for his guidance, support, and encouragement throughout my study period. Special word of appreciation to Dr Emily Goh Joo Kheng, for her financial support from Monash University, her guidance, suggestion, innovative ideas and also her invaluable advice and patience throughout the duration of this project.



I am also indebted to the staffs of the Department of Chemistry, Universiti Putra Malaysia, for their help and cooperation.

A deep acknowledgment is also extended to Mr. Johadi Iskandar, Mr. Zainal Abidin Kassim and Mrs. Rusnani Amirudin for their kindness in helping me with the NMR, MS and IR spectral data. I am also grateful to Dr. Lim Chan Kiang for helping me at any time and solving any unsolvable problems during my studies. Special thanks are extended to my laboratory members Mr. Shaari Daud, Ms. Vivien Jong, Ms. Lim Chyi Mei and Ms. Wen Yen Ping who have helped me in every possible way and providing a congenial and enthusiastic atmosphere in the laboratory.

Last but not least, special greetings and thank you to my beloved family for their wonderful love and generous moral support. To a very special friend of mine, Mr Danny Loke Chee Keong, for listening to my complaints and frustrations and for support and encouragement towards the success of this research, a special thank you.



I certify that an Examination Committee has met on 1<sup>st</sup> June 2007 to conduct the final examination of Ng Sook Han on her Master of Science thesis entitled “Chemical Constituents and Biological Activity of Asam Aur Aur (*Garcinia parvifolia*) and Jingga (*Ploiarium alternifolium*)” in accordance with Universiti Pertanian Malaysia (Higher Degree) Act 1980 and Universiti Pertanian Malaysia (Higher Degree) Regulations 1981. The Committee recommends that the candidate be awarded the relevant degree. Members of the Examination Committee are as follows:

**Abdul Halim Abdullah, PhD**

Associate Professor  
Faculty of Science  
Universiti Putra Malaysia  
(Chairman)

**Mawardi Rahmani, PhD**

Professor  
Faculty of Science  
Universiti Putra Malaysia  
(Internal Examiner)

**Taufiq Yap Yun Hin, PhD**

Associate Professor  
Faculty of Science  
Universiti Putra Malaysia  
(Internal Examiner)

**Hasnah M. Sirat, PhD**

Professor  
Faculty of Science  
Universiti Teknologi Malaysia  
(External Examiner)

---

**HASANAH MOHD. GHAZALI, PhD**

Professor / Deputy Dean  
School of Graduate Studies  
Universiti Putra Malaysia

Date:



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

**Gwendoline Ee Cheng Lian, PhD**

Associate Professor  
Faculty of Science  
Universiti Putra Malaysia  
(Chairman)

**Mohd Aspollah B Hj Md Sukari, PhD**

Associate Professor  
Faculty of Science  
Universiti Putra Malaysia  
(Member)

---

**AINI IDERIS, PhD**  
Professor / Dean  
School of Graduate Studies  
Universiti Putra Malaysia

Date: 17<sup>th</sup> JULY 2007



## DECLARATION

I do 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 it has not been previously or concurrently submitted for any other degree at UPM or other institutions.

---

**NG SOOK HAN**

Date : 13<sup>th</sup> JUNE 2007



## TABLE OF CONTENTS

	<b>Page</b>
<b>DEDICATION</b>	ii
<b>ABSTRACT</b>	iii
<b>ABSTRAK</b>	vi
<b>ACKNOWLEDGEMENTS</b>	ix
<b>APPROVAL</b>	xi
<b>DECLARATION</b>	xiii
<b>LIST OF TABLES</b>	xvii
<b>LIST OF FIGURES</b>	xix
<b>LIST OF ABBREVIATIONS</b>	xxiv

### CHAPTERS

<b>1</b>	<b>INTRODUCTION</b>	1
	1.1 General Introduction	1
	1.2 Botany of Plants Studied	3
	1.2.1 The Guttiferae Family	3
	1.2.2 The Genus <i>Garcinia</i>	4
	1.2.3 The Species <i>Garcinia parvifolia</i>	5
	1.2.4 The Genus <i>Ploiarium</i>	6
	1.2.5 The Species <i>Ploiarium alternifolium</i>	6
	1.3 Biological Activity of Natural Products	8
	1.4 Objectives of Study	8
<b>2</b>	<b>LITERATURE REVIEW</b>	10
	2.1 Chemistry of <i>Garcinia</i> Species	10
	2.1.1 Previous Work on Xanthonenes in <i>Garcinia</i> species	10
	2.1.2 Previous Work on Benzophenones in <i>Garcinia</i> Species	21
	2.1.3 Previous Work on Flavonoids in <i>Garcinia</i> Species	22
	2.1.4 Previous Work on Bioactivities of <i>Garcinia</i> Species	24
	2.2 Chemistry of <i>Ploiarium</i> Species	27
	2.2.1 Previous Work on Geranyl Anthraquinones in <i>Ploiarium</i> Species	27
	2.2.2 Previous Work on Anthraquinonyl Xanthonenes in <i>Ploiarium</i> Species	28
	2.2.3 Previous Work on Bixanthone in <i>Ploiarium</i> Species	29

2.2.4	Previous Work on Triterpenoid Benzoates in <i>Ploiarium</i> Species	30
2.2.5	Previous Work on Bioactivities of <i>Ploiarium</i> Species	30
<b>3</b>	<b>EXPERIMENTAL</b>	<b>31</b>
3.1	Plant Material	31
3.2	Instruments	31
3.2.1	Infrared Spectroscopy (IR)	31
3.2.2	Ultra Violet (UV)	31
3.2.3	Mass Spectra (MS)	32
3.2.4	Melting Point	32
3.2.5	Nuclear Magnetic Resonance (NMR)	32
3.3	Chromatographic Method	32
3.3.1	Column Chromatography	32
3.3.2	Thin Layer Chromatography	33
3.3.3	High Performance Liquid Chromatography	34
3.4	Dyeing Reagents for TLC	34
3.4.1	Iron (III) Chloride Solution	34
3.4.2	Vanillin-Sulfuric Acid Solution	34
3.4.3	Copper (II) Sulfate-Sodium Citrate	35
3.5	Larvicidal Assay	35
3.6	Antimicrobial Assay	36
3.6.1	Standardization of Inocula and Media	36
3.6.2	Antibiotic Disc Assay Impregnation	36
3.6.3	Plate Inoculation, Disc Introduction, Incubation and Observation	37
3.7	Cytotoxic Assay	37
3.8	Extraction and Isolation of Compounds from <i>Garcinia</i> <i>parvifolia</i> and <i>Ploiarium alternifolium</i>	39
3.8.1	<i>Garcinia parvifolia</i>	39
3.8.1.1	Isolation of Stigmasterol ( <b>66</b> )	40
3.8.1.2	Isolation of $\beta$ -Sitosterol ( <b>67</b> )	41
3.8.1.3	Isolation of 6-Deoxyjacareubin ( <b>68</b> )	43
3.8.1.4	Isolation of Daphnifolin ( <b>69</b> )	44
3.8.1.5	Isolation of Caffeine ( <b>70</b> )	45
3.8.1.6	Isolation of Rubraxanthone ( <b>55</b> )	47
3.8.1.7	Isolation of Isoxanthochymol ( <b>49</b> )	48
3.8.2	<i>Ploiarium alternifolium</i>	50
3.8.2.1	Isolation of Ploiariquinone A ( <b>58</b> )	51
3.8.2.2	Isolation of Emodin ( <b>60</b> )	52
3.8.2.3	Isolation of 1,8-dihydroxy-3-methoxy- 6-methyl-anthraquinone ( <b>71</b> )	54

<b>4</b>	<b>RESULTS AND DISCUSSION</b>	56
4.1	Isolation of Chemical Constituents from <i>Garcinia parvifolia</i>	56
4.1.1	Characterization of Stigmasterol ( <b>66</b> )	58
4.1.2	Characterization of Sitosterol ( <b>67</b> )	65
4.1.3	Characterization of 6-Deoxyjacareubin ( <b>68</b> )	71
4.1.4	Characterization of Daphnifolin ( <b>69</b> )	85
4.1.5	Characterization of Caffeine ( <b>70</b> )	98
4.1.6	Characterization of Rubraxanthone ( <b>55</b> )	109
4.1.7	Characterization of Isoxanthochymol ( <b>49</b> )	123
4.2	Isolation of Chemical Constituents from <i>Ploiarium alternifolium</i>	139
4.2.1	Characterization of Ploiariquinone A ( <b>58</b> )	141
4.2.2	Characterization of Emodin ( <b>60</b> )	155
4.2.3	Characterization of 1,8-dihydroxy-3-methoxy- 6-methyl-anthraquinone ( <b>71</b> )	168
4.3	Bioassay Results	179
4.3.1	Larvicidal Activity	179
4.3.2	Antimicrobial Activity	180
4.3.3	Cytotoxic Activity	181
4.3.4	Antifungal Activity	182
<b>5</b>	<b>CONCLUSIONS</b>	183
	<b>REFERENCES</b>	186
	<b>APPENDICES</b>	192
	<b>BIODATA OF THE AUTHOR</b>	206



## LIST OF TABLES

Table		Page
4.1	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CHDl}_3$ ) assignments of stigmasterol ( <b>66</b> )	60
4.2	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of sitosterol ( <b>67</b> )	66
4.3	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) of 6-deoxyjacareubin ( <b>68</b> )	73
4.4	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of 6-deoxyjacareubin ( <b>68</b> )	74
4.5	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) of daphnifolin ( <b>69</b> )	87
4.6	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of daphnifolin ( <b>69</b> )	88
4.7	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of caffeine ( <b>70</b> )	100
4.8	$^1\text{H}$ NMR (400 MHz, $\text{CD}_3\text{OD}$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CD}_3\text{OD}$ ) of rubraxanthone ( <b>55</b> )	112
4.9	$^1\text{H}$ NMR (400 MHz, $\text{CD}_3\text{OD}$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CD}_3\text{OD}$ ) assignments of rubraxanthone ( <b>55</b> )	113
4.10	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of isoxanthochymol ( <b>49</b> )	126
4.11	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) of ploiariquinone A ( <b>58</b> )	143
4.12	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of ploiariquinone A ( <b>58</b> )	144
4.13	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) of emodin ( <b>60</b> )	157



4.14	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of emodin ( <b>60</b> )	158
4.15	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	170
4.16	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) and $^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) assignments of 1,8-dihydroxy-3-methoxy-6-methyl- anthraquinone ( <b>71</b> )	171
4.17	Larvicidal activity of crude extracts against the larvae of <i>Aedes aegypti</i>	179
4.18	Antimicrobial activity of crude extracts of <i>Garcinia parvifolia</i> and <i>Ploiarium alternifolium</i>	180
4.19	Cytotoxic activity of plant extracts against HL-60 Cell Line (Promyelocytic Leukemia)	181



## LIST OF FIGURES

Figure		Page
1.1	Tree of <i>Garcinia parvifolia</i>	5
1.2	Flower of <i>Garcinia parvifolia</i>	5
1.3	Fruits of <i>Garcinia parvifolia</i>	5
1.4	Stem bark of <i>Ploiarium alternifolium</i>	7
1.5	Flower of <i>Ploiarium alternifolium</i>	7
1.6	Leaves of <i>Ploiarium alternifolium</i>	7
4.1	Isolation of chemical constituents from the stem bark of <i>Garcinia parvifolia</i>	57
4.2	EIMS spectrum of stigmasterol ( <b>66</b> )	61
4.3	IR spectrum of stigmasterol ( <b>66</b> )	62
4.4	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of stigmasterol ( <b>66</b> )	63
4.5	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of stigmasterol ( <b>66</b> )	64
4.6	EIMS spectrum of sitosterol ( <b>67</b> )	67
4.7	IR spectrum of sitosterol ( <b>67</b> )	68
4.8	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of sitosterol ( <b>67</b> )	69
4.9	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of sitosterol ( <b>67</b> )	70
4.10	EIMS spectrum of 6-deoxyjacareubin ( <b>68</b> )	75
4.11	IR spectrum of 6-deoxyjacareubin ( <b>68</b> )	76
4.12	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of 6-deoxyjacareubin ( <b>68</b> )	77

4.13	$^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) spectrum of 6-deoxyjacareubin ( <b>68</b> )	78
4.14	HMBC spectrum of 6-deoxyjacareubin ( <b>68</b> )	79
4.15	HMBC spectrum of 6-deoxyjacareubin ( <b>68</b> ) (expanded)	80
4.16	DEPT spectrum of 6-deoxyjacareubin ( <b>68</b> )	81
4.17	HSQC spectrum of 6-deoxyjacareubin ( <b>68</b> )	82
4.18	HSQC spectrum of 6-deoxyjacareubin ( <b>68</b> ) (expanded)	83
4.19	$^1\text{H}$ - $^1\text{H}$ COSY spectrum of 6-deoxyjacareubin ( <b>68</b> )	84
4.20	EIMS spectrum of daphnifolin ( <b>69</b> )	89
4.21	IR spectrum of daphnifolin ( <b>69</b> )	90
4.22	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) spectrum of daphnifolin ( <b>69</b> )	91
4.23	$^1\text{H}$ - $^1\text{H}$ COSY spectrum of daphnifolin ( <b>69</b> )	92
4.24	$^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) spectrum of daphnifolin ( <b>69</b> )	93
4.25	DEPT spectrum of daphnifolin ( <b>69</b> )	94
4.26	HSQC spectrum of daphnifolin ( <b>69</b> )	95
4.27	HMBC spectrum of daphnifolin ( <b>69</b> )	96
4.28	HMBC spectrum of daphnifolin ( <b>69</b> ) (expanded)	97
4.29	EIMS spectrum of caffeine ( <b>70</b> )	101
4.30	IR spectrum of caffeine ( <b>70</b> )	102
4.31	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) spectrum of caffeine ( <b>70</b> )	103
4.32	$^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) spectrum of caffeine ( <b>70</b> )	104
4.33	HSQC spectrum of caffeine ( <b>70</b> )	105
4.34	HMBC spectrum of caffeine ( <b>70</b> )	106



4.35	$^1\text{H}$ - $^1\text{H}$ COSY spectrum of caffeine ( <b>70</b> )	107
4.36	DEPT spectrum of caffeine ( <b>70</b> )	108
4.37	EIMS spectrum of rubraxanthone ( <b>55</b> )	114
4.38	$^1\text{H}$ NMR (400 MHz, $\text{CD}_3\text{OD}$ ) spectrum of rubraxanthone ( <b>55</b> )	115
4.39	$^{13}\text{C}$ NMR (100 MHz, $\text{CD}_3\text{OD}$ ) spectrum of rubraxanthone ( <b>55</b> )	116
4.40	DEPT spectrum of rubraxanthone ( <b>55</b> )	117
4.41	$^1\text{H}$ - $^1\text{H}$ COSY spectrum of rubraxanthone ( <b>55</b> )	118
4.42	HSQC spectrum of rubraxanthone ( <b>55</b> )	119
4.43	HMBC spectrum of rubraxanthone ( <b>55</b> )	120
4.44	HMBC spectrum of rubraxanthone ( <b>55</b> ) (expanded)	121
4.45	HMBC spectrum of rubraxanthone ( <b>55</b> ) (expanded)	122
4.46	EIMS spectrum of isoxanthochymol ( <b>49</b> )	128
4.47	IR spectrum of isoxanthochymol ( <b>49</b> )	129
4.48	$^1\text{H}$ NMR (400 MHz, $\text{CDCl}_3$ ) spectrum of isoxanthochymol ( <b>49</b> )	130
4.49	$^{13}\text{C}$ NMR (100 MHz, $\text{CDCl}_3$ ) spectrum of isoxanthochymol ( <b>49</b> )	131
4.50	$^1\text{H}$ - $^1\text{H}$ COSY spectrum of isoxanthochymol ( <b>49</b> )	132
4.51	HSQC spectrum of isoxanthochymol ( <b>49</b> )	133
4.52	HSQC spectrum of isoxanthochymol ( <b>49</b> ) (expanded)	134
4.53	HMBC spectrum of isoxanthochymol ( <b>49</b> )	135
4.54	HMBC spectrum of isoxanthochymol ( <b>49</b> ) (expanded)	136



4.55	HMBC spectrum of isoxanthochymol ( <b>49</b> ) (expanded)	137
4.56	DEPT spectrum of isoxanthochymol ( <b>49</b> )	138
4.57	Isolation of chemical constituents from the stem bark of <i>Ploiarium alternifolium</i>	140
4.58	EIMS spectrum of ploiariquinone A ( <b>58</b> )	145
4.59	IR spectrum of ploiariquinone A ( <b>58</b> )	146
4.60	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of ploiariquinone A ( <b>58</b> )	147
4.61	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of ploiariquinone A ( <b>58</b> ) (expanded)	148
4.62	<sup>1</sup> H- <sup>1</sup> H COSY spectrum of ploiariquinone A ( <b>58</b> )	149
4.63	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of ploiariquinone A ( <b>58</b> )	150
4.64	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of ploiariquinone A ( <b>58</b> ) (expanded)	151
4.65	HSQC spectrum of ploiariquinone A ( <b>58</b> )	152
4.66	HMBC spectrum of ploiariquinone A ( <b>58</b> )	153
4.67	DEPT spectrum of ploiariquinone A ( <b>58</b> )	154
4.68	EIMS spectrum of emodin ( <b>60</b> )	159
4.69	IR spectrum of emodin ( <b>60</b> )	160
4.70	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of emodin ( <b>60</b> )	161
4.71	<sup>1</sup> H- <sup>1</sup> H COSY spectrum of emodin ( <b>60</b> )	162
4.72	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of emodin ( <b>60</b> )	163
4.73	DEPT spectrum of emodin ( <b>60</b> )	164
4.74	HSQC spectrum of emodin ( <b>60</b> )	165



4.75	HMBC spectrum of emodin ( <b>60</b> )	166
4.76	HMBC spectrum of emodin ( <b>60</b> ) (expanded)	167
4.77	EIMS spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	172
4.78	IR spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	173
4.79	<sup>1</sup> H NMR (400 MHz, CDCl <sub>3</sub> ) spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	174
4.80	<sup>13</sup> C NMR (100 MHz, CDCl <sub>3</sub> ) spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	175
4.81	HSQC spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	176
4.82	HMBC spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	177
4.83	DEPT spectrum of 1,8-dihydroxy-3-methoxy-6-methyl-anthraquinone ( <b>71</b> )	178

