UNIVERSITI TEKNOLOGI MARA

CYTOTOXIC ACTIVITY OF MYRICETIN AND MAHANIMBINE AGAINST A BRAIN CANCER CELL LINE (U251) AND A NORMAL CELL LINE (WRL68)

WAN IZZATI MARIAH BINTI WAN HASSAN

Dissertation submitted in partial fulfilment of the requirements for the Bachelor of Pharmacy

Faculty of Pharmacy

2012

ACKNOWLEDGEMENTS

Firstly, I would like to praise to the merciful Allah SWT, for the guidance that helped me throughout this study and finish this study on time. I also would like to thank my supervisor, Dr. Vasudevan Mani and my co-supervisor of this study, Assoc. Prof. Dr. Kalavathy A/P Ramasamy and Ms. Nurul Aqmar Mohd Nor Hazalin for their valuable guidance and advices. They had inspired me a lot and lead me to overcome problems during this study period.

I also would like to be grateful for having very supportive postgraduate students which Ms.Nur Syafiqah Bt Rahim, Ms. Siti Norshazwani Bt. Wahab, Ms. Nor Nadia Ban, Ms. Siti Munirah Bt. Jaafar and Ms. Nurul Huda Bt. Musa for their continuous guidance. I would like to thank the authority of University Teknology Mara (UiTM) for providing such a good environment and facilities to complete this study. I would also like to take this opportunity to thank to Faculty of Pharmacy for offering this subject, Research 556 since it does gives a very valuable opportunity for students to participate and learn about the operations in doing research and open up the door for students to experience in the field of research and development.

Lastly, deep appreciation to my research colleagues which are, Norhamiza Bt. Abd Hamid, Izzati Bt. Hasnul, Hazlinda Bt. Mohamad and Hosni B. Azhar for their cooperation in helping me with this study. An honourable mention goes to my families and friends for their understanding and support in completing this study. Without all the help from the particulars mentions before, I definitely would face many difficulties in completing this study. Thank you.

TABLE OF CONTENTS

TITLE	PAGE
APROVAL	
ACKNOWLEDGEMENTS	i
TABLE OF CONTENTS	ii
LIST OF TABLES	iv
LIST OF FIGURES	V
LIST OF ABBREVIATIONS	vi
ABSTRACT	vii
CHAPTER ONE (INTRODUCTION)	1
CHAPTER TWO (LITERATURE REVIEW)	
2.1 Cancer 2.1.1 Brain cancer	4 4 6
2.2 Chemotherapy	8
2.3 Anticancer from natural products	9
2.3.1 Marine	10
2.3.2 Microorganism	12
2.3.3 Plant	13
2.4 Myricetin	14
2.5 Mahanimbine	16
CHAPTER THREE (MATERIAL AND METHOD)	
3.1 Cytotoxic activity	17
3.1.1 Materials and reagents	
3.1.2 Instruments	17 17 17
3.1.3 Method	
3.2 Statistical analysis	21

CHAPTER FOUR (RESULT)

4.1 Cytotoxic activity of myricetin and mahanimbine compound	
4.1.1 Cytotoxic activity of myricetin and mahanimbine compound	22
against U251 human tumor glioma cell line	22
4.1.2 Comparison of cytotoxicity activity between myricetin and	25
mahanimbine against U251 human tumor glioma cell line	
4.2 Selectivity of myricetin and mahanimbine against U251 human tumor	27
glioma cell line when compared to WRL68 normal cell line	
4.2.1 Cytotoxic activity of myricetin and mahanimbine against WRL68	29
normal cell line	
4.2.2 Comparison of selectivity of myricetin and mahanimbine against	30
U251 human tumor glioma cell line and WRL68 normal cell line	
CHAPTER FIVE (DISCUSSION)	32
CHAPTER SIX (CONCLUSION)	36
BIBLIOGRAPHY	37

ABSTRACT

Chemotherapy has become the common treatment choice in treating cancer even it can caused undesired side effect towards the patient as well as the health care handler. Thus, a lot of studies has been conducted to discover new and more selective anticancer. Plants are one of the most potential sources in finding anticancer. Numerous drugs are marketed as anticancer drugs, hence plant have been known as a familiar sources for development of new anticancer. The present study was carried out to determine the cytotoxicity and selectivity of plant-derived compounds myricetin and mahanimbine against human glioma cell lines (U251) and human normal cell line (WRL68). Both compounds were tested for cytotoxicity against U251 and WRL68 cell lines by using the MTT assay. The IC₅₀ of both compounds were derived from the dose-response curve. In this study, both of the compounds exhibited strong cytotoxic activity against U251 with IC₅₀ of 2.52 μ g/ml and 3.98 μ g/ml respectively. Thus, compound, myricetin and mahanimbine were found to be potent against WRL68 as well, with IC50 of 2.91µg/ml and 4.03µg/ml respectively. Thus, advance studies to be needed in modifying the chemical structures of both compounds in order to reduce its toxicity towards normal cell but at the same time maintaining the effectiveness as good cytotoxic agents.