UNIVERSITI TEKNOLOGI MARA

DETERMINATION CYTOTOXICITY OF MYRICETIN AND MAHANIMBINE ON HUMAN TUMOR GLIOMA CELL LINES, SNB-19 AND SNB-75

HOSNI BIN AZHAR

BACHELOR OF PHARMACY

Faculty of Pharmacy

2012

ACKNOWLEDGEMENTS

Alhamdulillah, thanks to Allah S.W.T for all the blessing and giving me the strength and all the effort to manage and complete this project and also complete my thesis on time in accordance to the guidelines given.

Here, I want to convey my thousand appreciations and special thanks to my thesis supervisor, Assoc. Prof. Dr. Kalavathy A/P Ramasamy for all her guidance, continuous support and limitless patience in guiding me to complete this thesis. Special thank to to Mrs Nurul Akma binti Mohd Hazalin and Dr. Vasudevan Mani who continuously helped and support me in the progress of this thesis.

I also would like to express my special gratitude to Ms Nur Syafiqah binti Rahim and Ms Siti Nursyazwani binti Wahab, who continuously helped me during the practical work. Greatly thanks to all the other post-graduate students in the lab for all their kindness and willingness to lend a hand when needed.

I also like to thank my research freinds, Ms Wan Mariah binti Wan Hassan, Mrs Nurhamiza binti Abd Hamid, Ms Izzaty binti Hasnul and Nurhaslinda binti Mohamad for helping me and exchanging information in the entire progress to complete this research.

I also want to thank my family, my wife, my lecturers and also my freinds for their unlimited support throughout this research period. I acknowledge the financial support from Universiti Teknologi MARA (UiTM) Puncak Alam, recieved in form of laboratory materials and instruments for conducting my thesis.

Finally, thanks to all who were involved in giving me the information during practical work. I hope this report will fullfill the entire requirement needed to complete this Research Instrument (PHR 555) course. May Allah bless us all, InsyaAllah.

Thank You.

TABLE OF CONTENT

TIT	LE PAGE	Page
APP	ROVAL FORM	
ACKNOWLEDGEMENT		
TABLE OF CONTENTS		
LIST OF TABLES LIST OF FIGURES		
ABSTRACT		
CHA	APTER 1: INTRODUCTION	
1.1	Introduction	1
1.2	Objective	3
CHA	APTER 2: LITERATURE REVIEW	
2.1	Cancer	4
2.2	Statistic and figure	5
2.3	Risk factor	
2.4	The biology of cancer	
2.5	Current treatment of cancer	10
	2.5.1 Chemotherapy	10
	2.5.2 Radiotherapy	13
	2.5.3 Targeted Therapy	14

2.6	Others therapy		
	2.6.1	Cyberknife15	
	2.6.2	Others alternatives therapy in cancer	16
2.7	Ration	nal research on new treatment	17
2.8	Natura	al product	19
2.9	Myricetin		
2.10	Mahai	nimbine	21
CHA	PTER 3	3: MATERIALS AND METHODS	
3.1	Cytoto	oxicity activity	24
	3.1.1	Materials and reagents	24
	3.1.2	Instruments	25
	3.1.3	Methods	25
3.2	Statist	ical analysis	28
CHA	PTER 4	4: RESULTS	
4.1	Cytoto	oxic activity of myricetin and mahanimbine compound	29
	4.1.1	Cytotoxic activity of myricetin and mahanimbine compounds	29
		against human tumor glioma cell line SNB-19	
	4.1.2	Cytotoxic activity of myricetin and mahanimbine compounds against human tumor glioma cell line SNB-75	29
	4.1.3	Comparison of cytotoxic activity between myricetin and	33
		mahanimbine against human tumor glioma cell line SNB-19	
	4.1.4	Comparison of cytotoxic activity between myricetin and mahanimbine against human tumor glioma cell line SNB-75	35

ABSTRACT

Myricetin and mahanimbine widely used as anti-oxidant. Both compounds have been found potential as an anti-cancer properties against cancer cell such as Human Colon Carcinoma and Human Leukemic cell lines. However the effect of compouds againts human glioma cells (SNB-19 and SNB-75) have not been reported. In this study, the cytotoxic activity of myricetin and mahanimbine was investigate. MTT assay was used to measure the cytotoxic activity. Human tumor glioma cell line SNB-19 and SNB-75 was cultured in the optimum environment which is using Minimal Essential Medium (MEM) with Earle's Salts media and the conditions of culture were 37°C and 5% CO₂. Cell were plated into 96-flat bottom well plate with a density of 5000 cells/well and incubated for 24 hours. After incubation, the plated cell were treated with with 20uL of of test compounds at various concentrations then incubated for 72 hours. MTT solution was added and read using the ELISA® micro plate reader. IC50 (value of cytotoxic activity) were derived from dose-response curve of which the concentration of extracts required to kill 50% of cell population. Result showed that mahanimbine had IC₅₀ values of 16.56ug/ml on SNB-19 and 7.07ug/ml on SNB-75 while myricetin exhibited cytotoxic activity with IC50 values of 22.11 ug/ml on SNB-19 and 28.44ug/ml on SNB-75. Both compounds had an inhibitory effect on cancer cells and normal cells indicating that test compounds did not show selectivity. Further investigation to improve the selectivity should established for effective and safe therapy.