MICROPROPAGATION OF Lycopersicon esculentum

NUR AEZWANI BINTI MOHD HORI

Final Year Project Report Submitted in Partial Fulfillment of the Requirements for the Degree of Bachelor of Sciences (Hons.) Biology in the Faculty of Applied Sciences Universiti Teknologi MARA

JANUARY 2016

ACKNOWLEDGEMENTS

In the name of Allah, the Most Gracious and the Most Merciful Alhamdulillah, all praises to Allah for the strengths and His blessing in completing this thesis.

Special appreciation goes to my supervisor, Hjh Sarina binti Hashim, for her supervision and constant support. Her invaluable help of constructive comments and suggestions throughout the experimental and thesis works have contributed to the success of this research. She has been always there to listen and give advice. I am deeply grateful to her for the long discussions that helped me sort out the technical details of my work. I am also thankful to her for encouraging the use of correct grammar and consistent notation in my writings and for carefully reading and commenting on countless revisions of this manuscript. My lab assistance Pn Zairus for has been always there to listen and give advice. I am also thankful to the system staff who maintained all the machines in lab so efficiently.

Sincere thanks to all my friends for their kindness and moral support during my study. Thanks for the friendship and memories. I am highly indebted to the authorities of MARA University of Technology (UiTM) for providing me various infrastructure like library, Internet access and permission include copyright which have enabled me to complete my research work successfully.

Last but not least, my deepest gratitude goes to my beloved parents; Mr. Mohd Hori b. Ab Rahman and Mrs. Noriah Bt Mat Rashid and also to my sisters for their endless love, prayers and encouragement. To those who indirectly contributed in this research, your kindness means a lot to me. Thank you very much.

(Nur Aezwani Bt Mohd Hori)

TABLE OF CONTENTS

			PAGE
10	VNOU	/LEDCEMENTS	:::
ACKNOWLEDGEMENTS TABLE OF CONTENTS			iii iv
LIST OF TABLES			vi
		FIGURES	vii
		ABBREVIATIONS	viii
	STRAC		ix
	STRAK		X
CH	APTE	R 1: INTRODUCTION	1
1.1	_	ground Study	1
		em Statement	3
	_	icant of the Study	3
1.4	Object	tive of the Study	3
СН	APTEF	R 2: LITERATURE REVIEW	5
2.1	Lyco	persicon esculentum	5
	2.1.1	Types of growth	8
	2.1.2	Factors that affected grow of tomato	9
	2.1.3	The importance of tomato	11
2.2	Micro	propagation	12
	2.2.1	Sterilization Technique	13
	2.2.2	Media preparation	14
	2.2.3	Plant hormone	15
СП	A DTEI	R 3: METHODOLOGY	
3.1	Mate		
5.1		Raw material	17
	3.1.2	Chemicals	17
	3.1.3		17
		Equipment	18
3.2			18
3.2	Methods 3.2.1 Apparatus sterilization		18
	3.2.1	• •	18
	3.2.2	- 1	20
		Inoculation	21
	J.4. T	III C GIGHTON	21

	3.2.5 Incubation	22	
	3.2.6 Observations	23	
	3.2.7 Morphological identification	23	
	3.2.8 Anatomical Study	24	
3.3	Data Analysis	24	
CH	APTER 4: RESULTS AND DISCUSSION		
4.1	Sterilization of Lycopersicon esculentum	26	
	4.1.1 Direct method	26	
	4.1.2 Indirect method	28	
	4.1.3 Problem in micropropagation	29	
4.2	Propagation of Lycopersicon esculentum	31	
	4.2.1 Hormone free medium	32	
	4.2.2 Hormone treatment	33	
	4.2.3 Determination of plant growth	35	
4.3	Performance In Vitro Growth of Lycopersicon esculentum	36	
CH	APTER5: CONCLUSION AND RECOMMENDATION	37	
CITED REFERENCES			
APPENDICES			
CURRICULUM VITAE			

ABSTRACT

Micropropagation of Lycopersicon esculentum

Lycopersicon esculentum or tomato is commercially cultivated vegetable crop grown all over the world. It is very versatile and is grown either for fresh market or processing. Tomato is gaining attention because of medical benefits and as sources of food. This study was done to obtain the seed from Lycopersicon esculentum to culture under sterile condition. The objective of this study is to determine the sterilization, to propagate the seed plant and to identify the performance of in vitro growth of Lycopersicon esculentum. micropropagation technique is used with different sterilization method, followed by inoculation of explant into media with hormone treatment and without hormone treatment. The explant was cultured in incubation room for 6 week and 16:8 photoperiods. Observation was done to determine the growth performance of this plantlet. The number of leaf, the height of the plantlets and length of roots was measured at week 6. The contaminated seed from direct and indirect sterilization methods are 0% and 5%, respectively. Anyhow the germinated seed from direct and indirect sterilization method are 10% and 90% respectively. This study show that sterilization of explant indirectly (tomato pod) was better than directly sterile the seed. Direct sterilization can cause damage to explant cell. The germination rates of explant in media with hormone and without hormone are 89% and 78% respectively. Therefore, growing plantlet in media with hormone is better than media without hormone.