

UNIVERSITI TEKNOLOGI MARA

***IN VITRO* MICROPROPAGATION
OF *Musa acuminata* cv. BERANGAN
(AAA) USING NEW FORMULATION
OF BIOORGANIC MEDIA**

**NUR ATIQAH BINTI KHIRUL
ANUAR**

Msc

November 2019

ABSTRACT

Plant tissue culture provides effective production for banana's high demand supply as one of subsistence crop and commercialized product. Commonly used plant tissue culture media such as Murashige and Skoog (MS), Gamborg's (Bs) and Schenk and Hilderbrandt (SH) medium are known for their high mineral content and competent in promoting plant tissue culture. However, these media are costly, containing synthetic component and non-locally available. Therefore, this study has been conducted to develop an alternative plant tissue culture medium with high nutrient component. Fruits peel that have been formulated to develop media by using a fermentation process consisting of formula A (calamansi, key and kaffir lime peel), formula B (banana, papaya and pineapple peel), formula C (kaffir lime, apple mango and guava peel) and formula D (Banana, dragonfruit and honey mango peel) were prepared in different concentrations to study the effects of *in vitro* micropropagation of *M.acuminata*. The plantlets were cultured onto bioorganic media and incubated in culture room for six weeks. Number of shoots and height of the shoot were recorded. The highest number of shoot and root obtained in control media with 2.50 ± 0.56 and 3.17 ± 0.60 respectively while the highest plant height and number of leaves recorded in formula D with 2.78 ± 0.32 cm and 2.83 ± 0.17 respectively. Contamination rate data showed the lowest in formula D with no contamination recorded. From this study, the application of bioorganic media is not significantly different ($p \geq 0.05$) in promoting *M.acuminata* growth but significantly difference ($p < 0.05$) in reducing contamination rate in *M.acuminata*. All bioorganic media formulas showed positive response in shoots regeneration. Therefore, this bioorganic media should be considered as the alternative culture media of *M.acuminata* as it showed positive response in promoting explants growth and it has low production cost as it is made up of organic waste and locally available.

ACKNOWLEDGEMENT

First and foremost, praise to Allah S.W.T for His blessings and guidance which has given me inspiration to embark on this project and stay strong to complete this thesis. First, I would first like to thank my supervisor, Dr. Nor Azma binti Yusuf and my co-supervisor, Assoc. Profesor Dr. Asmah binti Awal, lecturers from the Faculty of Plantation and Agrotechnology, Universiti Teknologi MARA, Shah Alam. The door to Dr. Nor Azma and Assoc. Prof. Dr. Asmah Awal office were always open whenever I ran into a trouble spot or had any questions about my research and writing and who are always patiently teaches during my struggling times. They were consistently allowed this research to be my own work, but steered me in the direction where I should be.

I am also want to thanks the experts, seniors, and friends who were involved in succeeding this research projects, Ummu Hani, Farah Suhaimi, Azurin, Nazatul Ashikin, Khairunnisa, Zulaiha, and all the laboratory's friends. Without their helps and participation, this research could not have been successfully conducted. I would also like to acknowledge Ummu Hani binti Mohd Shaain of Faculty of Plantation and Agrotechnology as the second reader of this thesis, and I am gratefully indebted to her very valuable comments on this thesis.

Finally, I am must and surely expressing whole of my gratitude to both of my parents Khirul Anuar bin Abdullah and . They are always providing me with unfailing supports and continuous encouragement throughout my years of study and through the process of researching and writing this thesis. They encouraged and hold me when I were about to give up numerous times and accompanying me to most of the places where I need to be. For all family, for giving continuous supports and motivations, I am sincerely want to say thank you. This accomplishment would not have been possible without them.

TABLE OF CONTENTS

	Page
CONFIRMATION BY PANEL OF EXAMINERS	ii
AUTHOR'S DECLARATION	iii
ABSTRACT	iv
ACKNOWLEDGEMENT	v
TABLE OF CONTENTS	vi
LIST OF FIGURES	ix
LIST OF TABLES	x
CHAPTER ONE: INTRODUCTION	1
1.1 Background of Study	1
1.2 Problem Statement	3
1.3 Objectives of Research	4
1.4 Scope of Research	4
1.5 Significance of Study	4
CHAPTER TWO: LITERATURE REVIEW	5
2.1 <i>Musa</i> sp. (Banana)	5
2.1.1 Introduction	5
2.1.2 Background Study of <i>Musa</i> sp.	6
2.1.3 Biological and Morphological Characteristics of <i>M. acuminata</i>	6
2.1.4 Conventional Propagation Method	7
2.1.5 <i>In vitro</i> Tissue Culture	8
2.1.6 Micropropagation of Banana	10
2.1.7 <i>M. acuminata</i> in Plant Tissue Culture	11
2.1.8 Tissue Culture Medium	12
2.1.9 Surface Sterilization of Explants	13
2.2 Agricultural Waste	14
2.2.1 Novelty Value and Potential Content of Fruits Peel	15

CHAPTER ONE

INTRODUCTION

1.1 Background of Study

Banana, a member of Musaceae family is one of the most important crop. Banana has been reported to rank as second most important crop with 22% of world's fresh fruit production (Pua, 2007; Jafari et al., 2011). For millions of communities in humid and subhumid tropics area, banana regarded as major staple food crop (Chikezie, 2012). Therefore, it is also counted as socio-politically important in developing world (Pua, 2007). It had been placed fourth as important diets crops for people especially in South East Asian. Malaysia is one of the origin country of banana and plaintain with about 50 types expected in exportation business. However, in exportation aims, the quality needs to be further improved (Darvari et al., 2010).

As a crop that is important in commercialization aspect, banana needs continuous and constant production in achieving stable supplies (Jafari et al., 2011). However, traditional agriculture that uses conventional method is incapable to solve this high demand issue. The conventional propagation of banana were done by using the suckers. The process has various limitations either in its method and output. Conventional method consumes a long time for effective production of banana to take places (Darvari et al., 2010). While the limitations regarding the output include low propagation rates and highly disaggregating. Besides, the spreading of pest and pathogen had also further complicating this method (Vuylsteke and Ortiz, 1996; Makara et al., 2010). To conclude, time, instability and disease threat had been proven as this method disadvantages which made this method was not the best possible solution in the current issue. Threatening current production of banana which had been regarded as a highly important food source may then affecting the exports market trade balance (Becker et al., 2000).

To overcome those problems, application of advance biotechnology has been seen as the ideal solution instead of using conventional method (Altman and Hasegawa, 2012). The advanced biotechnology involved is clonal planting material which used tissue culture propagation techniques. It is necessary for continuous of good quality banana production (Jafari et al., 2011). This plant tissue culture technique had been