

DEVELOPMENT OF AN ANther CULTURE PROTOCOL FOR  
BRINJAL VARIETIES (*Solanum melongena* L.) BY VARYING  
THE HORMONE COMPOSITION OF THE  
INITIATION MEDIUM



By  
M. V. I. Lakshani



FTC279

Main Library, Eastern University, Sri Lanka

Department of Biosystems Technology  
Faculty of Technology  
Eastern University, Sri Lanka  
Chenkalady

2026

## ABSTRACT

Brinjal (*Solanum melongena*) is an important solanaceous crop cultivated extensively in tropical and subtropical regions for its nutritional and economic value. The development of pure homozygous lines through conventional breeding is laborious and time-consuming; therefore, anther culture provides an efficient alternative for the rapid production of doubled haploid plants. This study aimed to develop an efficient anther culture protocol for selected brinjal varieties by optimizing the hormonal composition of the initiation medium. Sterilized anthers from three hybrid brinjal varieties (SA-07, Amanda, and HORDI Hybrid 04) were cultured on Murashige and Skoog (MS) medium supplemented with varying concentrations of BAP, NAA, and kinetin, forming four treatments (T1-T4). The interaction between treatments and varieties was not statistically significant at the 5% level ( $p > 0.05$ ) for any measured parameter, indicating that variation was primarily due to the individual effects of genotype and medium composition. Among the treatments, T3 (3 mg L<sup>-1</sup> BAP + 1 mg L<sup>-1</sup> NAA + 3 mg L<sup>-1</sup> kinetin) resulted in the highest anther size enlargement ( $2.33 \pm 0.33\%$ ) in SA-07, while no enlargement was observed in T2 and T4 of HORDI Hybrid 04. The highest callus formation ( $16.70 \pm 10.50\%$ ) occurred in T1 (2 mg L<sup>-1</sup> BAP + 1 mg L<sup>-1</sup> NAA + 2 mg L<sup>-1</sup> kinetin) for SA-07, confirming its superior androgenic response. Although treatment effects were not statistically significant for callus formation ( $p = 0.133$ ), varietal effects were significant ( $p = 0.041$ ), highlighting the influence of genotype on in vitro response. Contamination levels did not differ significantly among treatments or varieties ( $p > 0.05$ ). Overall, the results demonstrate that hormonal balance and genotype are critical determinants of success in brinjal anther culture. The MS medium supplemented with 2 mg L<sup>-1</sup> BAP, 1 mg L<sup>-1</sup> NAA, and 2 mg L<sup>-1</sup> kinetin proved most suitable for callus induction in SA-07, suggesting its potential use in developing a standardized protocol for rapid doubled haploid production in brinjal breeding programs.

**Keywords:** Anther Culture, Androgenic, Callus Formation, Doubled Haploid, Genotype, Homozygous Lines, *Solanum melongena*, Size Enlargement.

## TABLE OF CONTENTS

DECLARATION.....	iii
DEDICATION .....	iv
ACKNOWLEDGMENT .....	v
ABSTRACT .....	vi
TABLE OF CONTENTS .....	vii
LIST OF FIGURES .....	x
LIST OF TABLES .....	xi
ABBREVIATIONS AND SYMBOLS .....	xii
CHAPTER 01 .....	1
INTRODUCTION .....	1
1.1 General Description.....	1
1.2 Problem Identification .....	4
1.3 Problem Justification .....	4
1.4 Objectives .....	5
1.4.1 Overall Objective.....	5
1.4.2 Specific Objectives .....	5
CHAPTER 02.....	6
LITERATURE REVIEW .....	6
2.1 Introduction to Brinjal ( <i>Solanum melongena</i> L.).....	6
2.1.1 Origin and Global Distribution.....	6
2.1.2 Botanical and Taxonomical Classification of Brinjal.....	6
2.1.3 Morphological Characteristics.....	7
2.1.4 Nutrient Composition and Health Benefits of Egg Plants .....	10
2.1.5 Reproduction of Brinjal .....	12
2.2 Plant Tissue Culture.....	12
2.2.1 Types of Plant Tissue Culture.....	13
2.2.2 Culture Media in Plant Tissue Culture .....	14
2.2.3 Plant Growth Regulators in Tissue Culture .....	15
2.2.4 Plant Tissue Culture Used in Brinjal .....	16
2.3 Haploid Production in Plant Breeding.....	17
2.3.1 Methods of Haploid Production .....	18
2.3.2 Applications of Haploid Production .....	19

2.3.3 Importance of Haploid Plant Breeding .....	20
2.3.4 Challenges in Haploid Production .....	20
2.4 Overview of Anther Culture .....	21
2.4.1 Mechanisms of Haploid Induction in Anther Culture .....	21
2.4.2 Anther Culture for <i>Solanum melongena</i> .....	22
2.4.3 Factors Affecting Anther Culture .....	23
2.4.4 Anther Culture Studies in Sri Lanka.....	23
2.4.5 Similar Studies Related to Brinjal Haploid Production.....	25
CHAPTER 03 .....	26
METHODOLOGY .....	26
3.1 Location of the Study.....	26
3.2 Experimental Design and Treatment .....	26
3.3 Materials .....	26
3.3.1 Plant Materials .....	26
3.3.2 Culture Media .....	27
3.4 Methodology.....	28
3.4.1Preparation of Stock Solutions .....	28
3.4.2 Preparation of Culture Media .....	30
3.4.3 Sterilization of Culture Media .....	31
3.4.4 Collection of Flower Buds.....	31
3.4.5 Pre-surface Sterilization of Flower Buds.....	32
3.4.6 Surface Sterilization of Flower Buds.....	33
3.4.7 Anthers Extraction and Sterilization.....	33
3.4.8 In-vitro Culturing of Anthers.....	34
3.5 Data Collection and Parameters.....	34
3.6 Data Analysis.....	34
CHAPTER 04 .....	35
RESULTS AND DISCUSSION.....	35
4.1 Size Enlargement of Brinjal Anthers.....	35
4.2 Callus Formation from Brinjal Anthers.....	39
4.3 Contamination of Brinjal Anthers .....	42
CHAPTER 05 .....	45
CONCLUSION .....	45

CHAPTER 06.....	46
REFERENCES .....	46
APPENDICES .....	60