

**ISOLATION OF CELLULOSE DEGRADING BACTERIA
(CDB) FROM MANGROVE SOIL.**

ADAWIAH BT ABDUL RAHMAN

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

JULY 2015

ACKNOWLEDGEMENTS

Bismillahirrahmanirrahim,

In the name of Allah SWT, the most gracious and the most merciful. Alhamdulillah, all the praises to Allah SWT for the strengths given and his blessing in completing this thesis. In this opportunity, I would like to express my highest gratitude to the supervisor and co-supervisor, Miss Siti Suhaila Bt Harith and Madam Marlina Bt Mohd Mydin, for all their endless guidance and encouragement throughout completion of this thesis.

I am also highly indebted with the lab assistants especially Mr Suhairi Bin Shuib, for the guidance and valuable information shared by them in their respective fields through conducting this thesis. Besides that, I am also indebted with Universiti Teknologi Mara (UiTM) Jengka Pahang for providing the facilities in terms of labs, machines, wireless connection and also the chemicals needed for this thesis.

Lastly, I would like to thank my family for their endless support especially in moral and financial support. I also want to thank to all my friends that had given their hands either directly or indirectly in completion of this thesis.

(AdawiahBt Abdul Rahman)

TABLES OF CONTENTS

	PAGE
ACKNOWLEDGEMENTS	i
TABLES OF CONTENTS	ii
LIST OF TABLES	iv
LIST OF FIGURES	v
LIST OF ABBREVIATIONS	vi
ABSTRACT	viii
ABSTRAK	ix
CHAPTER 1: INTRODUCTION	
1.1 Background Of Study	1
1.2 Problem Statement	2
1.3 Significant Of The Study	3
1.4 Objective Of The Study	4
CHAPTER 2: LITERATURE REVIEW	
2.1 Cellulose	5
2.2 Microbial Cellulase	7
2.2.1 Reaction Of Cellulase	8
2.2.2 Industrial Importance Of Cellulase	9
2.3 Cellulose Degrading Bacteria (CDB)	10
2.4 Mangrove Soil	12
2.4.1 Mangrove Soil Characteristics	13
CHAPTER 3: METHODOLOGY	
3.1 Materials	16
3.1.1 Raw Materials	16
3.1.2 Chemicals	16
3.1.3 Apparatus And Instruments	17
3.2 Methods	17
3.2.1 Preparation Of Agar	18
3.2.1.1 Preparation Of Nutrient Agar	18
3.2.1.2 Preparation Of CarboxyMethy Cellulose Agar (CMC)	18
3.2.2 Preparation Of Broth	19
3.2.2.1 Preparation Of Mineral Salt Medium (MSM) Broth	19
3.2.2.2 Preparation Of Nutrient Broth	19
3.2.3 Preparation Of Congo Red Stain	19

3.2.4	Preparation of trace element	20
3.2.5	Preparation of Dinitrosalicylic acid (DNS)	20
3.2.6	Sample Collection	20
3.2.7	Isolation Of Cellulose Degrading Bacteria By Enrichment Method	21
3.2.8	Screening Of Cellulose Degrading Bacteria	21
3.2.9	Cellulase Enzyme Production	22
3.2.10	Cellulase Enzyme Assays	22
3.2.11	Consequences Of The pH, incubation period And Temperature On The Development of Cellulose Degrading Bacteria	23
CHAPTER 4: RESULTS AND DISCUSSION		25
CHAPTER 5: CONCLUSION AND RECOMMENDATIONS		38
CITATED REFERENCES		39
APPENDICES		43
CURRICULUM VITAE		46

ABSTRACT

ISOLATION OF CELLULOSE DEGRADING BACTERIA FROM MANGROVE SOIL

Cellulose is the most abundant component of plant biomass and potentially used for biofuel production. This bioconversion involve cellulase enzyme. Enzymatic hydrolysis by microorganisms is the most cost effective. This study is intended to isolate the potential cellulose degrading bacteria, to assess cellulolytic potential of cellulase and to optimize the growth condition of cellulose degrading bacteria. A total of five bacteria were isolated from mangrove soil using mineral salt medium broth containing carboxymethyl cellulose(CMC). All the five bacteria do not display any hydrolysis zone in Congo Red test. Whereas, the hydrolysis zone appeared when tested with Iodine test. The isolated bacteria namely Bacteria A, Bacteria B, Bacteria C, Bacteria D and Bacteria E were undergo pre-screen using dinitrosalicylic (DNS) method. Bacteria E show high constituents of cellulolytic potential of 0.187 $\mu\text{mol/ml}$. Optimization of cultural conditions for cellulase production by Bacteria E was carried out. Optimize cultural condition at the temperature of 37°C, pH 7 and 5 days of incubation period yielded 0.368 $\mu\text{mol/ml}$.