

CULTIVATION TRIALS OF *Gracilaria* sp. (Rhodophyta)
IN POND IN PUSAT PENYELIDIKAN TERNAKAN AIR
RAYAU (PFTAP), GELANG PATAH, JOHORE

GLENDEN FLANT LANGGANGON

FAKULTI SAINS DAN TEKNOLOGI
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI
MALAYSIA
2005

LP
11
FST
2
2005

1100034622

LP 11 FST 2 2005



1100034622

Cultivation trials of *gracilaria* sp. (Rhodophyta) in pond in Pusa Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johore / Clenden Flant Langgongan.



PERPUSTAKAAN

**KOLEJ UNIVERSITI SAINS & TEKNOLOGI MALAYSIA
21030 KUALA TERENGGANU**

1100034622

Lihat sebelah

HAK MILIK
PERPUSTAKAAN KUSTEM

CULTIVATION TRIALS OF *Gracilaria* sp. (Rhodophyta) IN POND IN PUSAT
 PENYELIDIKAN TERNAKAN AIR PAYAU (PPTAP), GELANG PATAH , JOHORE

By

Clenden Flant Langgangon

Research Report submitted in partial fulfilment of
The requirements for the degree of
Bachelor of Science (Marine Biology)

Department of Marine Science
Faculty of Science and Technology
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI MALAYSIA

1100034622

This project report should be cited as :

Flant, C.2005. Cultivation Trials of *Gracilaria* sp. (Rhodophyta) in Pond in Pusat Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johore Undergraduate thesis, Bachelor of Science in Marine Biology, Faculty of Science and Technology, Kolej Universiti Sains dan Teknologi Malaysia, Terengganu.41p.

No part of this project may be reproduced by any mechanical, photographic, or electronic process, or in the form of phonographic recording, nor it may be stored in a retrieval system, transmitted, or otherwise copied for public or private use, without written permission from the author and the supervisor (s) of the project.



JABATAN SAINS SAMUDERA
FAKULTI SAINS DAN TEKNOLOGI
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI MALAYSIA

PENGAKUAN DAN PENGESAHAN LAPORAN
PROJEK PENYELIDIKAN I DAN II

Adalah ini diakui dan disahkan bahawa laporan penyelidikan bertajuk:

*Cultivation Trials of Gracilaria sp. (Rhodophyta) In Pond In
Pusat Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johor
oleh. Denden Plant Langgamon....., No. Matrik.....uk. 2000.....*

telah diperiksa dan semua pembetulan yang disarankan telah dilakukan. Laporan ini dikemukakan kepada Jabatan Sains Samudera sebagai memenuhi sebahagian daripada keperluan memperolehi Ijazah...*Sarjana Muda Sains (Biologi Marine)*...

Fakulti Sains dan Teknologi, Kolej Universiti Sains dan Teknologi Malaysia.

Disahkan oleh:

Christine A. Orosco

Penyelia Utama: DR. SITI AISHAH ABDULLAH @
Nama: CHRISTINE A. OROSCO
Cop Rasmi: Pensyarah
Jabatan Sains Samudera
Fakulti Sains dan Teknologi Malaysia
(KUSTEM)
21030 Kuala Terengganu

Tarikh: 29/3/2005

Penyelia Kedua (jika ada)

Nama: Tarikh:

Cop Rasmi:

[Signature]
Ketua Jabatan Sains Samudera

Nama: DR. AHMAD SHAMSUDDIN B. AHMAD
Cop Rasmi: Ketua
Jabatan Sains Samudera
Fakulti Sains dan Teknologi
Kolej Universiti Sains dan Teknologi Malaysia
21030 Kuala Terengganu

Tarikh: 29/3/05

ACKNOWLEDGEMENT

First of all, I would like to express my deepest appreciation and sincere gratitude to my supervisor Dr. Siti Aishah Abdullah for her undivided attention and supervision in completing this thesis.

My appreciation also goes to Mr. Gan and Miss Fong, for their precious time and guidance throughout my final year project. To all my friends in KUSTEM, thank you for all advice and criticism towards making my project a success. It really means a lot to me.

Last but not least, I would like to dedicate this project to my family back home in Sabah who have been very supportive throughout my studies in KUSTEM. Their endless encouragement and love gave me strength to carry on towards achieving my goals. Not forgetting Sharannah Isabella who was always there to give me moral support and for getting involved with my project. I would also like to thank my friends in Taman Armon for struggling through the ups and downs with me for the past three years. Thanks a lot everyone.

TABLE OF CONTENTS

ACKNOWLEDGEMENT	ii
TABLE OF CONTENT	iii
LIST OF TABLES	vi
LIST OF FIGURES	vii
LIST OF SYMBOLS	viii
LIST OF APPENDICES	ix
ABSTRAK	x
ABSTRACT	xii
CHAPTER 1.0 INTRODUCTION	1
CHAPTER 2.0 LITERATURE REVIEW	3
2.1 Taxonomy	3
2.1 Agar	3
2.3 <i>Gracilaria</i> sp. Cultivation	4
2.4 Nutrient Contents	6
2.5 Uses of <i>Gracilaria</i>	7
CHAPTER 3.0 METHODOLOGY	9
3.1 Sampling and Culture Site	9
3.2 Methods	10
3.2.1 Line Culture Method	10
3.2.2 Cage Culture Method	11

3.2.3	Environmental Parameters	12
3.2.4	Harvest and Drying	12
3.2.5	Growth Rate	13
3.2.6	Alkaline Treatment and Agar Extraction	13
3.2.7	Determination of Agar Quality	15
	3.2.7.1 Determination of Gel Strength	15
	3.2.7.2 Determination of Gelling Temperature	16
	3.2.7.3 Determination of Melting Temperature	16
3.2.8	Determination of Nutrient Contents	17
	3.2.8.1 Determination of Ammonium Concentration	17
	3.2.8.2 Determination of Nitrate Concentration	17
	3.2.8.3 Determination of Phosphate Concentration	18
3.2.8	Statistical Analysis	19
3.3	Materials Needed	19
CHAPTER 4.0	RESULTS	20
4.1	Water Quality	20
4.2	Harvest and Drying	23
4.3	Yield and Quality of Agar from <i>Gracilaria</i> sp.	23
CHAPTER 5.0	DISCUSSION	27
5.1	Growth Rate	27
5.2	Agar Yield and Quality	28

CHAPTER 6.0	CONCLUSION	31
	RECOMMENDATION	32
LITERATURE CITED		33
APPENDIX		36
CURRICULUM VITAE		41

LIST OF TABLES

No.	Title	Page
4.1	Summary of water quality throughout the culture period from 18 September 2004 to 14 October 2004.	20
4.2	The light intensity in different depths (top, mid, bot) for line culture method.	20
4.3.	Average Relative Growth Rate (RGR) for both line and cage culture method	20
4.4	Fresh weight and dry weight of harvested <i>Gracilaria</i> sp.	23
4.5	Agar yield and quality of both native and alkali treated (60° C, 80° C, 90° C) agar from <i>Gracilaria</i> sp. and other commercial agars	23
4.6	Agar Gel Strength of both native and alkali treated (60° C, 80° C, 90° C) agar from <i>Gracilaria</i> sp. and other commercial agars using 3 different tests.	25

LIST OF FIGURES

No.	Title	Page
3.1.	Picture of <i>Gracilaria</i> sp. used in cultivation in trial pond.	9
3.2.	The line culture method used for the cultivation of <i>Gracilaria</i> sp.	10
3.3.	Net cage used in <i>Gracilaria</i> sp. cultivation.	11
3.4.	The Gel Tester used in determining the gel strength	15
3.5.	The Texture Analyzer (TA. XT plus) used in determining the gel strength	16
4.1.	Relative Growth Rate (RGR) of <i>Gracilaria</i> sp. in cage culture method	21
4.2.	Relative Growth Rate (RGR) of <i>Gracilaria</i> sp. in line culture method	21
4.3.	Relative Growth Rate (RGR) of <i>Gracilaria</i> sp. in 3 different depths.	22
4.4.	Relative Growth Rate (RGR) of <i>Gracilaria</i> sp. in both line and cage culture method.	22
4.5.	Agar gel from agar strip (China) and agar powder (Japan).	25
4.6.	Agar gel from <i>Gracilaria</i> sp. without alkali treatment (native) and with alkali treatment (60° C, 80° C, 90° C) through cage culture method.	25
4.7.	Agar gel from <i>Gracilaria</i> sp. without alkali treatment (native) and with alkali treatment (60° C, 80° C, 90° C) through line culture method.	26

LIST OF SYMBOLS

cm	-	Centimeter
mL	-	Millilitre
mg L ⁻¹	-	Milligram per Litre
%	-	Percent
‰	-	part per thousand
L	-	Litre
°C	-	Degree Celsius
mg	-	milligram
kg	-	Kilogram
DO	-	Dissolved Oxygen
ppm	-	parts per million
RGR	-	Relative Growth Rate
Sec	-	Second
NaOH	-	Sodium Hydroxide
H ₂ SO ₄	-	Sulphuric Acid

LIST OF APPENDICES

No.	Title	Page
1.	Fresh weight (g) of <i>Gracilaria</i> sp. in line (1) culture method.	36
2.	Fresh weight (g) of <i>Gracilaria</i> sp. in line (2) culture method.	36
3.	Fresh weight (g) of <i>Gracilaria</i> sp. in line (3) culture method.	36
4.	Fresh weight(g) of <i>Gracilaria</i> sp. in cage culture method.	37
5.	Average Relative Growth Rate (% , day ⁻¹) of line culture method.	37
6.	Average Relative Growth Rate (% , day ⁻¹) of cage culture method.	37
7.	Physical parameters in the pond at 9 different stations for week 0.	38
8.	Physical parameters in the pond at 9 different stations for week 4.	38
9.	Physical parameters in the pond at 9 different stations for week 8.	38
10.	Statistical analysis for RGR of different culture methods.	39
11.	Statistical analysis for RGR of different depths for line culture method.	40

Tajuk: Percubaan Pengkulturan *Gracilaria* sp. (Rhodophyta) Dalam Kolam Di

Pusat Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johor

ABSTRAK

Pengkulturan rumpai laut merah *Gracilaria* sp. telah dijalankan selama 8 minggu di sebuah kolam yang berada di Pusat Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johore. Pengkulturan *Gracilaria* sp. ini melibatkan 2 kaedah iaitu kaedah ikatan dan kandang berjaring. Kadar tumbesaran relatif (RGR, % day⁻¹) bagi kedua-dua kaedah telah diukur sepanjang tempoh pengkulturan. Begitu juga dengan kadar tumbesaran bagi 3 kedalaman yang berbeza (atas, tengah, bawah). Pada akhir tempoh pengkulturan, *Gracilaria* sp. tersebut telah dituai dan agarnya diekstrak bagi menentukan kualiti agarnya.

Secara keseluruhan, purata kadar tumbesaran relatif (RGR, % day⁻¹) bagi kaedah ikatan lebih tinggi daripada kaedah kandang berjaring dengan kadar 1.3 % day⁻¹. Di bawah keadaan yang menggalakkan, *Gracilaria* sp. mencapai kadar relative tumbesaran tertinggi iaitu 3.5 % day⁻¹. Manakala kedalaman atas mencatat kadar tumbesaran relatif yang tertinggi iaitu 2.3 %, day⁻¹.

Peratusan hasil agar daripada kedua-dua kaedah mempunyai nilai julat dari 18.4% ke 42.5% di mana kedua-dua kaedah (melalui atau tanpa melalui rawatan alkali) adalah bersilang kaitan dengan ketegangan gel. Agar yang melalui rawatan alkali mempunyai ketegangan gel yang tertinggi berbanding agar tanpa melalui rawatan alkali. Ketegangan gel untuk agar daripada *Gracilaria* sp. yang dikultur menggunakan kaedah kandang berjaring dengan 335.622 g.1.2cm⁻² merupakan yang tertinggi bagi suhu 90° C yang

melalui rawatan alkali. Suhu gel agar dan pencairan gel agar boleh dibandingkan dengan agar tepung komersil dari Jepun iaitu 42°C dan 84°C .

ABSTRACT

The cultivation of the red seaweed *Gracilaria* sp. was conducted throughout a culture period of 8 weeks in a pond in Pusat Penyelidikan Ternakan Air Payau (PPTAP), Gelang Patah, Johore. This cultivation was carried out using 2 different methods which were the line culture method and cage culture method. During the culture period, the relative growth rate (RGR, % day⁻¹) of *Gracilaria* sp. for both methods was monitored weekly. Growth rates at 3 different depths (top, middle, bottom) for line culture method were also monitored. At the end of the culture period the *Gracilaria* sp. was harvested and extracted to determine the agar quality of *Gracilaria* sp.

The overall average relative growth rate (RGR, % day⁻¹) for line culture method was higher than for cage culture method at a rate of 1.3 % day⁻¹. Under favourable conditions, the highest RGR was also recorded for line culture method at 3.5 % day⁻¹. Meanwhile the top line displayed the highest RGR compared to the other depths at 2.3 % day⁻¹.

The agar yield of samples from both culture methods ranged from 18.4% to 42.5% (extracted with or without alkali treatment) were inversely correlated with gel strength. Agar extracted after alkaline treatment had the highest gel strength. The gel strength of agar extracted from *Gracilaria* sp. in cage culture method (335.622 g.l.2cm⁻²) was the highest in 90° C alkali treatment. The gelling and melting temperatures were also comparable to the commercial agar powder from Japan at 42 ° C and 84 ° C respectively.