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The glucosidase inhibitor activity of ethanol-extract of  
aglaonema simplex culture / by Nurul Ain Zainon.

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Lihat Sebelah

THE GLUCOSIDASE INHIBITOR ACTIVITY OF ETHANOL-EXTRACT OF  
*AGLAONEMA SIMPLEX* CULTURE

By  
Nurul Ain Binti Zainon

A PITA report submitted in partial fulfillment of  
the requirements for the award of the degree of  
Bachelor of Science (Biological Sciences)

DEPARTMENT OF BIOLOGICAL SCIENCES  
FACULTY OF SCIENCE AND TECHNOLOGY  
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JABATAN SAINS BIOLOGI  
FAKULTI SAINS DAN TEKNOLOGI  
UNIVERSITI MALAYSIA TERENGGANU

**BDV/BIO 4999**  
**PENGAKUAN DAN PENGESAHAN LAPORAN PITA**

Adalah ini diakui dan disahkan bahawa laporan penyelidikan bertajuk: **The Glucosidase Inhibitor Activity of Ethanol-Extract of *Aglaonema simplex* Culture** oleh **Nurul Ain Binti Zainon**, no. matrik: **UK20388** telah diperiksa dan semua pembetulan yang disarankan telah dilakukan. Laporan ini dikemukakan kepada Jabatan Sains Biologi sebagai memenuhi sebahagian daripada keperluan memperolehi Ijazah **Sarjana Muda Sains (Sains Biologi)**, Fakulti Sains dan Teknologi, Universiti Malaysia Terengganu.

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## **DECLARATION**

I hereby declare that this Final Year Project research report entitled The Glucosidase Inhibitor Activity of Ethanol-Extract of *Aglaonema simplex* Culture is my own research except as cited in the references.

Signature :   
Name : Nurul Ain Binti Zainon  
Matric No. : UK20388  
Date : 8 July 2012

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## The Glucosidase Inhibitor Activity of Ethanol-Extract of *Aglaonema simplex* Culture

### ABSTRACT

The incidences of metabolic disorders such as diabetes mellitus are increasing. One of the therapeutic approaches for this disorder is by reducing the  $\alpha$ -glucosidase activity. In the study, the potential of *Aglaonema simplex* as therapeutic agent especially as glucosidase inhibitor due its polyhydroxylalkaloids was investigated. The in vitro plantlets of *A. simplex* were cultured in Murashige and Skoog medium for three and four months. The ethanolic extracts obtained were assayed for  $\alpha$ -glucosidase inhibition activity. Results showed that stem and root extracts exhibited the high inhibition activity with  $IC_{50}$  4.5 and 3.9 mg/ml respectively at three months culture, 4.1 and 3.2 mg/ml respectively at four months culture. Roots at four months age of culture showed the most effective inhibitor with lowest  $IC_{50}$  value, 3.2 mg/ml. The Lineweaver-Burk plots revealed that the inhibition activity of leave extracts was uncompetitive type, stems and roots were non-competitive.  $K_m$  values for stem and root extracts at three and four months were remaining constant, 5.0 mM while  $V_{max}$  for these parts were changes.  $V_{max}$  of stem and root extracts were 0.5 and 0.4  $mM \cdot min^{-1}$  respectively at three months, 0.4 and 0.3  $mM \cdot min^{-1}$  respectively at four months culture.

## **Perencatan Oleh Kultur *Aglaonema simplex* Yang Diekstrak Menggunakan Etanol Terhadap Aktiviti Glikosida**

### **ABSTRAK**

Kejadian gangguan metabolismik seperti diabetes melitus semakin meningkat. Salah satu pendekatan terapeutik yang dilakukan untuk menangani masalah adalah dengan mengurangkan aktiviti  $\alpha$ -glukosida. Dalam kajian, potensi *Aglaonema simplex* sebagai agen terapeutik terutama sebagai perencat glukosida disebabkan kandungan polihidrosilalkaloids telah dikaji. *A.simplex* dikultur di dalam Murashige dan Skoog media selama tiga dan empat bulan. Aktiviti perencatan diuji menggunakan sampel yang diekstrak menggunakan etanol. Hasil kajian menunjukkan bahawa ekstrak batang dan akar pokok menunjukkan aktiviti perencatan yang tinggi dengan  $IC_{50}$  4.5 dan 3.9 mg/ml bagi kultur tiga bulan serta 4.1 dan 3.2 mg/ml bagi kultur empat bulan. Bahagian akar yang dikultur selama empat bulan menunjukkan aktiviti perencatan yang paling efektif berdasarkan nilai  $IC_{50}$  bahagian tersebut yang paling rendah iaitu 3.2 mg/ml. Plot Lineweaver-Burk menunjukkan bahawa ekstrak daun adalah perencat tidak kompetitif manakala ekstrak batang dan akar adalah non-kompetitif. Nilai  $K_m$  bagi ekstrak batang dan akar yang dikultur selama tiga dan empat bulan adalah tidak berubah, 5.0 mM manakala  $V_{max}$  bagi ekstrak bahagian-bahagian tersebut adalah berubah.  $V_{max}$  bagi ekstrak batang dan akar ialah 0.5 dan 0.4 mM. $min^{-1}$  untuk kultur tiga bulan, 0.4 dan 0.3 mM. $min^{-1}$  untuk kultur empat bulan.