UNIVERSITI TEKNOLOGI MARA

SELF NANOEMULSIFYING DRUG DELIVERY SYSTEM OF GRISEOFULVIN

TAZQIRAH BINTI MUHAMAD

Dissertation submitted in partial fulfilment of the requirements for the Bachelor of Pharmacy (Hons.)

Faculty of Pharmacy

ADKNOWLEDGEMENT

All the praise and glory to Allah S.W.T who blessed me with good health and strength to complete my research and thesis writing accordingly. The writing of this thesis has been one of the most significant academic challenges I have ever had to face. Without the guidance and the help of several individuals, this thesis would not have been possible.

First and foremost, I would like to acknowledge the advice and guidance of Mr. Ahmad Mustafa Masoud Eid, my helpful supervisor. All his advice, encouragement, knowledge and assistance truly help the progression and smoothness of the research and thesis writing. The co-operation is much indeed appreciated.

Not forgotten is my partner, Nurul Jannah binti Sofwan, I would like to thank her, for her valuable ideas and advices. A big contribution and hard worked during the two semesters is very great indeed. My grateful thank also goes to lab assistants of Pharmaceutical Nano-Technology Laboratory, Faculty of Pharmacy, UiTM Puncak Alam for much guidance and support they gave.

I would also like to thank my faculty, Faculty of Pharmacy, which has given me the opportunity to do and complete this research. Finally, thanks to my family and friends for their unconditional love and support.

Tazqirah binti Muhamad

TABLE OF CONTENTS

		Page
TITL	E PAGE	
APPROVAL SHEET		ii
ADKNOWLEDGEMENT		iii
TABLE OF CONTENTS		iv
LIST OF TABLES		viii
LIST OF FIGURES		ix
LIST OF ABBREVIATIONS		X
ABST	TRACT	xi
CHAPTER 1 (INTRODUCTION)		1
1.1	Research Background	1
1.2	Statement of Problem	4
1.3	Significance of Study	5
1.4	Objectives	7
1.5	Hypothesis	7
CHAPTER 2 (LITERATURE REVIEW)		8
2.1	Emulsion	8
2.2	Nanoemulsion	11
2.3	Challenges in oral drug delivery	13
2.4	The Biopharmaceutical Classification Scheme	15
2.5	Lipid as Carrier	18

ABSTRACT

The oral drug delivery is fraught with low and erratic bioavailability owing to

enzymatic degradation of drug, poor permeability, poor solubility and dissolution.

The objective of the study was to develop and characterize Self Nanoemulsifying

Drug Delivery System (SNEDDS) of Griseofulvin, a Biopharmaceutical

Classification Scheme class II for improved oral delivery. The components for

nanoemulsion were identified by solubility studies and tendency for self-

emulsification in various excipients. Castor oil, Cremophor RH40 and Labrafil were

selected as oil, surfactant and co-surfactant respectively. The effects of varying ratios

of surfactants:co-surfactants:oil ratios of SNEDDS were studied. Droplet size and

zeta potential of the formulations were investigated. The SNEDDS exhibited good

efficacy for self-emulsification.

Keywords: Self emulsifying systems, Nanoemulsion, Cremophor RH40, Labrafil,

Castor oil, SNEDDS

xi

CHAPTER 1

INTRODUCTION

1.1 Research Background

An emulsion is a system consisting of two immiscible liquid phases, one of which is dispersed throughout the other in the form of fine droplets(Rodríguez-Abreu & Lazzari, 2008). The phase that is present as fine droplets is called the dispersed phase and the phase in which the droplets are suspended is the continuous phase. Most emulsions have droplets with diameters of 0.1-100µm and are inherently unstable systems. Its smaller globules exhibit colloidal behavior and the stability of a hydrophobic colloidal dispersion(Leal-Calderon, Thivilliers, & Schmitt, 2007).

Pharmaceutical emulsions usually consist of water and oil. Emulsions can be divided into two groups which are simple emulsion and multiple emulsions. For the simple emulsion, the dispersed phase is distributed throughout the continuous phase. Simple emulsions can later be divided into two types which are oil in water (o/w) emulsions and water in oil (w/o) emulsions. Oil in water in oil (o/w/o) emulsion and water-in-oil in water (w/o/w) emulsion are the subdivision of multiple emulsions(Rodríguez-Abreu & Lazzari, 2008).