

UNIVERSITI TEKNOLOGI MARA

**PREPARATION AND CHARACTERIZATION OF 3-D
PLGA SCAFFOLD FOR TISSUE ENGINEERING**

NUR KHAIRUNNISA BINTI ABDUL MANAF

Dissertation submitted in partial fulfillment of the requirements for the

Faculty of Pharmacy

October 2009

TABLE OF CONTENTS

	Page
TITLE PAGE	
APPROVAL	
ACKNOWLEDGEMENTS	ii
TABLE OF CONTENTS	iii
LIST OF TABLES	vi
LIST OF FIGURES	vii
ABSTRACT	ix
CHAPTER ONE (INTRODUCTION)	1
1.1 Background of Study	
1.2 Statement of the Problems	
1.3 Objectives	
1.4 Significance of the Study	
CHAPTER TWO (LITERATURE REVIEW)	5
2.1 Scaffold as a room for tissue growth	
2.2 Materials for Developing Scaffold	
2.2.1 Naturally Occurring Degradable Polymers	
2.2.2 Synthetic Degradable Polymers	
2.2.3 Ceramics and Composites	
2.3 Poly [L (-) lactide-co-glycolide] (PLGA)	
2.3.1 PLGA as main materials in fabrication of 3-D scaffold	
2.3.2 Degradation of different types of PLGA	
2.3.3 Fabrication methods of 3-D PLGA scaffolds.	

ACKNOWLEDGEMENT

I have a great deal of help in my completion of this study. This study cannot be completed without the help of many individuals. With the full of responsibilities and co-operation, I worked hard in order to accomplish this research.

Besides that, I extremely thank to my research supervisor, Dr. Javad Sameni Khadeem for the continuous support and encouragement. Without his critiques in his area and dedication this thesis would not be possible to accomplish.

At the same time, I would like to show my gratitude to our beloved Dean, Prof Aishah for her supports and commitment that allowed me to have this opportunity to obtained experiences as a researcher with the establishment of this subject.

Additionally, special thanks to Dr. Angela from Orthopedic Research Department, HUKM, for her willingness to share her knowledge and experiences, without her helps and dedication to the field of tissue engineering, this thesis would not be possible to be completed. It has been a privilege getting to know you.

I would like to thank Dr.Javad's lab assistant Intan as well as the science officer in the Faculty of Applied Sciences, Mrs. Masni which have help me a lot in finding handling the sorptometer to perform the porosity test.

My biggest thanks and love go out to my parents, whose love and direction have always been there for me. I am so blessed to have parents like them. To my friend Wan Nurul Huda W.Hasan, thank you, for supporting and giving me lots of help through this process.

Last but not least, I would like to give big thanks to those who has contributed directly or indirectly in completing my thesis.

ABSTRACT

Tissue engineering has emerged as a promising approach in the treatment of malfunctioning or lost organs. In this approach, a temporary scaffold is needed to serve as an adhesive substrate for the implanted cell and a physical support to guide the formation of new organs. The objective is to prepare and characterize variety PLGA scaffolds with different morphology, mechanical properties and porosity using different ratio of lactic to glycolic acid in the PLGA and different size of salt (NaCl) which range from $\geq 200\mu\text{m}$, $100\mu\text{m}$ to $<200\mu\text{m}$ and $<100\mu\text{m}$. PLGA is used in this study due to its non-toxic, biodegradable and biocompatibility effect in the human body. The method use in this study is solvent casting / particulate leaching method which involve the casting of a mixture of polymer solution and porogen in a mold, drying the mixture, followed by a leaching out of porogen with water to generate the pores. The results of the analyses showed that the size of the salt and ratio of lactic to glycolic acid content in the PLGA do give effect to the mechanical properties and porosity of the PLGA scaffold.

CHAPTER ONE

INTRODUCTION

1.1 Background of Study

Tissue engineering offers a new paradigm for the treatment of trauma and disease or aging-related loss of tissue function, combining engineering approaches with principles from the life sciences to develop biological substitutes that restore, maintain, or improve tissue function. From its very beginnings, tissue engineering followed one of three major approaches by which artificial tissue can be created within the body of a patient. The first, exemplified by the work of Vacanti and Langer (1999) combined isolated cells and biomaterial to create the artificial tissue in vitro. In this approach, cells are seeded onto a biomaterials prior to use or implantation of the device. The second approach, illustrated by the pioneering work of Yannas and Bruke (1981), they use the implantation of an acellular (e.g., cell-free) scaffold to guide the generation of new tissue in vivo. In this approach, the attachment of cells onto the biomaterial occurs only after the implantation of the device. Finally, some researchers attempt to generate functional replacement tissues in vitro and apply directly without the need of any biomaterial components. This approach is exemplified by the work of Auger, who developed a tissue culture system for the formation of functional skin tissue that can be applied directly to the patient to cover various wounds.