الآيـــة

بسم الله الرحمن الرحيم

قال تعالى:

خرر أَهُ خَيْرٌ لَكَ مِنَ الأَ وُلَى وَ لَسَو فَ فَ يَعْطِيكَ رَبُّكَ فَتَر ْضَدَى) يُعْطِيكَ رَبُّكَ فَتَر ْضَدَى)

صدق الله العظيم

سورة الضحى الأية (4-5)

Dedication

This work is dedicaTed To:

My lovely mother (AyAt el mAnsour).

Y

Dear uncle (Omer el mansOur).

Y

Lovely brothers (MohaMed & el sedig &isrra).

¥

All my family.

V

Acknowledgment

Thanks and praise first and foremost to Allah almighty forgives me strength to complete this work.

The researcher would like to express her sincere gratitude to his supervisor Dr: Ali Abdel Rahman Saeed Marouf for his help and his good guidance .My thanks are extended to Dr: Abd al-Sakhi Suleiman at the University of Neelain and Dr. Mohamed Ibrahim Medical services. For their supports, also my thanks extended to my friend awdia for her supports; I respect and thank dr Isam Ahmed attia, who giving me all support and guidance which made me complete the project duly. I am extremely thankful to him for providing such a nice support and guidance, although he had annual vacation.

I owe my deep gratitude to my examiner project Dr Abd ellatif Abass who took keen interest onmy project work and guided us all along, till the completion of my project work by providing all the necessary information for developing a good research.

I would not forget to remember dr Abd elfatah Mohamed for his encouragement more over and his support and guidance till the completion of the project work.

For all of them respect and gratitude

Abstract

There is an urgent need to develop technology for continuous in vivo glucose monitoring in subjects for diabetes mellitus. Problems with existing devices based on electrochemistry have encouraged alternative approaches to glucose sensing in recent years, and those based on fluorescence intensity and lifetime have special advantages, including sensitivity and the potential for non-invasive measurement when near infrared light is used. In this research twelve sample of blood with different concentrations of glucose have been employed to detect the fluorescence that induced using nitrogen laser. Plasma was separated from the twelve blood samples then it was exposed to nitrogen laser with wavelength 337.3 nm, power 0.04 mW, and periodic time is 100 msec. The fluorescence was collected using the fluorescence spectrometer.

The results showed that when the concentration of glucose increased, the levels or intensity of albumin and globulin exponentially decreased. At the low concentrations of glucose less than 150 Mg|Dl, the levels or intensity of protein (albumin and globulin) was very high; more than 3000 and when a concentration of glucose greater than 150, the level of protein (albumin and globulin) decreases .until reach low levels or intensity 200 At high concentrations of glucose 545 Mg/DL.

Decreasing the level of protein (albumin and globulin) indicated to attribute to a decrease in the body's insulin.

المستخلص

هناك حاجة ماسة لتطوير تكنولوجيا للرصد المستمر للجلوكوز في الجسم لمرضى السكري. وقد شجعت مشاكل الأجهزة الموجودة القائمة على الكيمياء الكهربائية طرق بديلة لإستشعار الجلوكوز في السنوات الأخيرة، وتلك التي تعتمد على شدة الفلورة والعمرالزمني لها مزايا خاصة، بما في ذلك الحساسية وإمكانية القياس غير الغازي عند إستخدام الأشعة تحت الحمراء القريبة. أستخدمت اثنتي عشر عينة من الدم بتراكيز مختلفة من الجلوكوز للكشف عن الفلورة المنتجة بإستخدام ليزر النيتروجين. فصلت البلازما من عينات الدم الاثني عشر ثم عرضت لليزر النيتروجين ذو الطول الموجي 337.3 نانومتر ، والطاقة 0.04 ميجاوات ، وزمن تكرار 100 ميللي ثانية. جمعت أشعة الفلورة باستخدام مطياف الفلورسنت .

أظهرت النتائج أنه عند زيادة تركيز الجلكوز يقل مستوى بروتيني الألبيومين والغلوبلين ؛ عند التراكيز الأقل للجلكوز يكون مستوى البروتينين (الألبيومين والغلوبلين)عالي جدا؛ أكبر من 3000 وعند زيادة تركيز الجلكوز أكثر من 150 يقل مستوى البروتينين (الألبيومين والغلوبلين) حتى يصل إلى أقل قيمة له 200 عند التراكيز العالية 545 جم/ديسلتر .

قد يشير نقصان مستوي مستوى البروتينين (الألبيومين والغلوبلين) إلى نقصان الأنسلين في الجسم .

Table of Contents

الآية	I
Dedication	II
Acknowledgement	III
Abstract (English)	IV
Abstract (Arabic)	V
Table of Contents	VI
List of figures	IX
List of tables	X

Chapter One			
Introduction and Previous Studies			
1.1	Overview	1	
1.2	Research Problem	2	
1.3	Previous Studies	2	
1.4	The Objectives of this Dissertation	4	
1.5	Research Methodology	5	
1.6	Dissertation Layout	5	

Chapter Two		
Theoretical Background		
2.1	Spectroscopy	6
2.1.1	Rotational Spectroscopy	6
2.1.2	Vibration spectroscopy	7
2.1.3	Electronic Spectroscopy	7
2.1.4	Laser Spectroscopy	8
2.1.4.1	Laser Emission Spectroscopy	8
2.1.4.2	Laser Induced Fluorescence (LIF)	9
2.1.4.3	Laser Absorption Spectroscopy	12
2.2	Blood	13
2.2.1	Blood plasma	14
2.2.1.1	Albumin	15
2.2.1.2	Globulins	16
2.2.1.3	Fibrinogen	18
2.2.2	Glucose	18
2.2.3	Blood Glucose	19

Chapter Three The Experimental Part		
3.1	The Materials	21
3.2	The Devices	22
3.2.1	Centerifuge	22
3.2.2	The Nitrogen (N ₂) Laser	22
3.3	The Method	23

Chapter Four		
The Results and Discussion		
4.1	The Results	24
4.1.1	The emission Spectra of the healthy and diabetes plasma	24
4.2	Analysis and discussions	25
4.2.1	The relation between glucose concentration and albumin	25
4.2.2	The relation between glucose concentration and globulin	27
4.3	Conclusions	29
4.4	Recommendations	29
References		31

List of Figures

No of figures	Title of Figures	Page Number
2.1	Spontaneous emission and stimulated emission	9
2.2	Jablonski Diagram	10
2.3	Laser-induce fluorescence	12
2.4	Absorption	13
2.5	Blood compound	14
2.6	Plasma compound	15
2.7	Albumins structure	15
2.8	Globulins structure	16
2.9	Fibrinogen structure	18
2.10	Glucose complex	19
3.1	Blood plasma	21
3.2	Centerifuge	22
3.3	The Nitrogen laser(N ₂) at Alneelain university	23
3.4	Experiment set up	23
4.1	Emission spectra of the twelfth samples	24
4.2	Glucose concentration and albumin	25
4.3	Glucose concentration and globulins	27

List of Tables

Number of table	Title of table	Number of page
3.1	Samples with their Concentration	21
4.1	Glucose concentration and intensity of albumin and globulin	25