

# Comparative Study of the Effect of Insulin and Green Tea on the Parotid Gland of Streptozotocin Induced Diabetic Rats

## (Histological and Electron Microscopic Study)

**Thesis** 

submitted to Faculty of Dentistry Ain Shams University in partial fulfillment of the requirements for Master Degree in Oral Biology

#### By

### **Hend Mostafa Mohamed El-Messiry**

B.D.S (Faculty of Dentistry-Ain Shams University- 2009)
Demonstrator of Oral Biology,
Faculty of Dentistry-Ain Shams University,

### **Supervisors**

## Prof. Dr. Medhat A. El-Zainy

Professor of Oral Biology Faculty of Dentistry-Ain Shams University

### **Dr. Dahlia Ghazy Mohamed**

Lecturer of Oral Biology. Faculty of Dentistry-Ain Shams University

> Faculty of Dentistry Ain Shams University 2015

# بسم الله الرحمن الرحيم قَالُوا سُبْحَانَكَ لَا عِلْمَ لَنَا إِلَّا مَا عَلَّمْتَنَا إِنَّكَ أَنْت الْمُعِيمُ الْمُكِيمُ الْمُكِيمُ

صدق الله العظيم

سورة البقرة- الآية 32

### Acknowledgement

All praise is to Allah and all thanks. He has guided and enabled me by his mercy to fulfill this thesis which I hope to be beneficial for people.

I would like to express my deepest gratitude to **Prof. Dr.**Medhat A. El-Zainy, professor of Oral Biology department, faculty of dentistry, Ain Shams University for continuous encouragement, his kind support and constructive advice that guided me to accomplish this work.

Special thanks are extended to **Dr. Dahlia Ghazy Mohamed,** lecturer of Oral Biology department, faculty of dentistry, Ain Shams University who freely gave me her time, effort and experience with continuous guidance throughout this work.

Special thanks are also extended to **Prof. Dr. Souzi Farid Shinaishin**, Professor and Head of Oral Biology Department, Faculty of Dentistry Ain Shams University, for her constant encouragement and advice whenever needed.

Finally, I am thankful to all members of Oral Biology department, faculty of dentistry, Ain Shams University for giving me the time to finish this work. Their help and support are greatly appreciated.

### **Dedication**

I dedicate this thesis work to my husband, who has been a constant source of support and encouragement throughout this project. I am truly thankful for having you in my life.

To my son, who is the best gift I've ever had.

To my parents, who have always loved me unconditionally and whose good examples have taught me to work hard for the things that I aspire to achieve.

To my brothers, who have never left my side and are very special.

# **CONTENTS**

	Page
Introduction	1
Review of Literature	3
Diabetes Mellitus	3
Streptozotocin drug	4
Effect of Diabetes Mellitus	5
Effect of Diabetes Mellitus on salivary glands	7
• Insulin	8
Green Tea	9
Effects of Green tea on Diabetes Mellitus	13
Parotid gland	13
Aim of the study	16
Materials and methods	17
Results	21
Light microscopic results	21
Electron microscopic results	34
Discussion	53

	Page
Summary	61
Conclusions and Recommendations	65
References	66
Arabic summary	-

# LIST OF ABBREVIATIONS

Abbreviation	Meaning
β	Beta
СТ	Connective tissue
DM	Diabetes Mellitus
EGCG	Epigallocatechin gallate
GT	Green tea
H&E	Haematoxylin and Eosin
NOD	Nonobese diabetic mice
PG	Parotid gland
STZ	Streptozotocin

	LIST OF FIGURES	
Fig.	Title	Page
1	A photomicrograph of the PG of the control group	22
_	(H&E original magnification x400)	
2	A photomicrograph of the PG of the control group (H&E original magnification x400)	22
3	A photomicrograph of the PG of the control group (H&E original magnification x400)	23
4	A photomicrograph of PG from the diabetic untreated group (H&E original magnification x400)	25
5	A photomicrograph of PG from the diabetic untreated group (H&E original magnification x400)	25
6	A photomicrograph of PG from the diabetic untreated group (H&E original magnification x400)	26
7	A photomicrograph of PG from the diabetic untreated group (H&E original magnification x400)	26
8	A photomicrograph of PG from the diabetic untreated group (H&E original magnification x400)	27
9	A photomicrograph of PG from the insulin treated group (H&E original magnification x400)	29
10	A photomicrograph of PG from the insulin treated group (H&E original magnification x400)	29
11	A photomicrograph of PG from the insulin treated group (H&E original magnification x400)	30

Fig.	Title	Page
No. 12	A photomicrograph of PG from green tea treated group (H&E original magnification x400)	32
13	A photomicrograph of PG from green tea treated group (H&E original magnification x400)	32
14	A photomicrograph of PG from green tea treated group (H&E original magnification x400)	33
15	An electron micrograph of PG from the control group (original magnification x1500)	35
16	An electron micrograph of PG from the control group (original magnification x6000)	35
17	An electron micrograph of PG from the control group (original magnification x8000)	36
18	An electron micrograph of PG from the control group (original magnificationx3000)	36
19	An electron micrograph of PG from the control group (original magnification x6000)	37
20	An electron micrograph of PG from diabetic untreated group (original magnification x2000)	39
21	An electron micrograph of PG from diabetic untreated group (original magnification x3000)	39
22	An electron micrograph of PG from diabetic untreated group (original magnification x1500)	40
23	An electron micrograph of PG from diabetic untreated group (original magnification x3000)	40

Fig.	Title	Page
24	An electron micrograph of PG from diabetic	41
	untreated group (original magnification x3000)	
25	An electron micrograph of PG from diabetic	41
	untreated group (original magnification x6000)	
26	An electron micrograph of PG from diabetic	42
	untreated group (original magnification x3000)	
27	An electron micrograph of PG from diabetic	42
	untreated group (original magnification x3000)	
28	An electron micrograph of PG from diabetic	43
	untreated group (original magnification x3000)	
29	An electron micrograph of PG from diabetic	43
	untreated group (original magnification x10000)	
30	An electron micrograph of PG from diabetic	44
	untreated group (original magnification x3000)	
31	An electron micrograph of PG from insulin treated	46
	group (original magnification x1500)	
32	An electron micrograph of PG from insulin treated	46
	group (original magnification x6000)	
33	An electron micrograph of PG from insulin treated	47
	group (original magnification x3000)	
34	An electron micrograph of PG from insulin treated	47
	group (original magnification x3000)	
35	An electron micrograph of PG from insulin treated	48
	group (original magnification x8000)	
37	An electron micrograph of PG from green tea treated	50
	group (original magnification x2000)	

Fig.	Title	Page
38	An electron micrograph of PG from green tea treated group (x original magnification 10000)	50
39	An electron micrograph of PG from green tea treated group (original magnification x3000)	51
40	An electron micrograph of PG from green tea treated group (original magnification x6000)	51
41	An electron micrograph of PG from green tea treated group (original magnification x3000)	52
42	An electron micrograph of PG from green tea treated group (original magnification x10000)	52

### **Abstract**

*Background:* Diabetes results in a variety of oral health complications. Green tea is a natural antioxidant that had proven to have powerful effects against diabetes.

Aim: To compare between the effect of green tea and insulin on Parotid gland of streptozotocin induced diabetic Albino rats usinglight microscopy and transmission electron microscopy.

Materials and Methods: Forty adult male Albino rats were divided into a control group and diabetic groups. The diabetic group received a single injection of 40 mg/kg of streptozotocin through the tail vein under anaesthesia and was further subdivided into three subgroups: The diabetic untreated subgroup which was left untreated for two weeks, insulin subgroup which received the treated insulin subcutaneously in a daily dose of 5 IU/kg body weight/day for two weeks. Lastly green tea treated subgroup received a ofm1/daily dose 100 body weight gm intragastricallyfortwo weeks. Rats were terminated parotid glands were dissected and processed for light and transmission electron microscopic examination.

Results: Histological examination of the diabetic untreated subgroup showed that acinar cells demonstrated

pyknotic hyperchromatic and nuclei with many vacuolations. Ultrastructurally, cytoplasmic acinar cells pleomorphism showed nuclear and dilated rough endoplasmic reticulum and mitochondria. Inflammatory cell. infiltration detected both histologically was ultrastructurally and ducts showed signs of some degeneration with of their loss normal outline stagnated secretion. However, insulin and green tea treated subgroups showed minimal degenerative changes and were almost similar to the control with minimal changes.

Conclusions: Treatment of the parotid gland of the streptozotocin induced diabetic rats with GT was closely comparable to the traditional insulin in reducing signs of histological and ultrastructural damage.

## **Introduction**

Diabetes Mellitus (DM) is one of the most prevalent, chronic diseases affecting mankind. It is a multistage, complex, chronic metabolic disorder characterized by dysregulation of carbohydrate, protein and lipid metabolism (*Artion et al.*, 1998).

The primary feature of this disorder is elevation in blood glucose levels which has been shown to affect almost all tissues in the body and associated with significant complications of multiple organs including eyes, nerves, kidneys and blood vessels (*Mealey*, 2006).

Insulin is a peptide hormone, produced by  $beta(\beta)$  cells of the pancreas, and is central to regulating carbohydrate and fat metabolism in the  $body(\textbf{\textit{Day and Bailey}, 1988})$ .

In addition to treatment of DM by insulin treatment, dietary interventions were shown to represent an effective tool to prevent and/or treat DM (*Willett et al.*, 2002).

Green tea (GT), from the plant *Camellia sinensis*, is one of the most popular beverages consumed worldwide. It is rich in antihypertensive, antiarteriosclerotic, hypoglycemic and hypocholesterolemic activity (*Wolfram*, 2007).

The present study will compare the effect of insulin and GT on Streptozotocin (STZ) induced diabetes on the parotid gland (PG) of Albino rats.