HISTOPATHOLOGIC AND IMMUNOHISTOCHEMICAL EVALUATION OF THE EFFECT OF GRAPE SEEDS EXTRACTS ON ALVEOLAR BONE AND PERIODONTAL LIGAMENT IN STREPTOZOTOCIN-INDUCED DIABETIC RATS

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ABSTRACT

Background: This study aims at clarifying if grape seed extracts (GSE) are candidates' therapeutic agents against the damaging effects of diabetes mellitus on the alveolar bone and periodontal ligaments.

Materials and methods: By randomly allocating 50 adult albino rats into five groups, each group included 10 rats as follows: Group I (negative control): served as controls; Group II: (positive control) diabetic rats receive no treatment; Group III: diabetic rats were treated with 50 mg/kg GSE for 6 weeks; Group IV: diabetic rats treated with 100 mg/kg GSE for 6 weeks; Group V: diabetic rats treated with 10 mg/kg gliclazides for 6 weeks. At the end of the experiment, extraction and preparation of rats jaws to examine their alveolar bone and periodontal ligaments histologically and immunohistochemically.

Results: Diabetic rats treated with GSE 100 mg/kg showed marked improvement of periodontal ligaments and alveolar bone with new bone formation compared to diabetic rats treated with GSE 50mg/kg and Gliclazides 10mg/kg. Immunohistochemical analysis of Bcl2 in untreated diabetic rats has statistically significantly (P≤0.001) caused an increase in Bcl2 expression in comparison with diabetic rats that receive treatments. GSE 100mg/kg has statistically significantly (P≤0.001) led the Bcl2 expression to decrease in comparison with GSE 50 mg/kg and diabetic rats treated with gliclazide.

Conclusion: GSE at dose 100mg/kg improved the periodontal ligament and alveolar bone in diabetic rats it could be used as a supplement for treating diabetic patients with diabetes Mellitus.

KEYWORDS: Streptozotocin, Diabetes Mellitus, grape seeds extract, gliclazide, Bcl2

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INTRODUCTION

Described as chronic diseases, Diabetes mellitus and periodontal disease have an effect on a large number of populations all over the world. An important long-term complication related to diabetes mellitus is the changed bone metabolism. The loss of Alveolar bone is a key outcome of periodontitis and diabetes is one of the primary factors of risk for periodontal disease. (1)

Diabetes mellitus (DM) represents a metabolic disorder that is diagnosed by the increase in plasma glucose levels, it alters the inflammatory response, impairs tissue healing, disturbs the collagen production, and reduces the angiogenesis process. (2) DM has been considered to be the major risk factor in developing serious and gradual damage to periodontal tissue. Multiple biological variables are including microbial impacts, hyperglycemia, and immune cytokines have been attributed to the increased periodontal tissue damage in DM. (3)

The advanced glycation end-products (AGES) are built up by the persistent increase in the blood glucose level, which in collagens, AGEs produce non-enzymatic cross-links thereby decreasing the flexibility of collagen matrix, which in turn leads to bone deformations. (4)

As bone density and diabetes have an intricate interplay, Diabetic patients are often led to osteopenia and extreme cases are associated with osteoporosis. In turn, this results in facilitating the loss of bone in these patients and leads them to an increased level of risk for negative outcomes in terms of bone health. (5) Several additional factors are related to hyperglycemia and have an effect on bone micro-architecture in DM. As an example, glycosuria can proportionally can lead calcium excretion to increase in urine. In addition, the interaction between hyperglycemia and the parathyroid hormone (PTH) and vitamin D system has an effect on bone turnover in the population of patients living with DM. (6)

The key reason for the development of oxidative stress and the increase in producing reactive oxygen species (ROS) that enhance cellular oxidation is hyperglycemia. (7,8,9) ROS activates nuclear factor-kappa B (NF-κB) is activated by ROS, following the increase in the expression of proinflammatory cytokine and chemokine. (10) These mediators have a significant effect on bone by stimulating the osteoclast differentiation and activity. (11) Moreover, ROS causes mitochondrial damage, which promotes the apoptotic pathway, starting with Bax pro-apoptotic proteins activation on the mitochondrial surface. (12) Consequently, the increase in the ROS stimulates the expression of Bax and downregulation of Bcl-2, which can inhibit the apoptotic cell death. (13,14)

Many active substances derived from plant-derived groups of different chemical compounds display a potent antioxidant activity, (15) such as proanthocyanidins which is a major component of grape seeds extract. They have 20 times more powerful antioxidant activity than vitamin C and 50 times more effective than vitamin E. (16,17) proanthocyanidins have a wide range of pharmacological and therapeutic effects, it has vasodilator, anticarcinogenic, anti-allergic, anti-inflammatory, antifungal, anti-arthritis, antibacterial, cardioprotective, immunostimulant, and antiviral properties which are remarkably significant. (18)

Grape seeds extracts (GSE) also efficiently reduce inflammatory cytokines’ production including interleukin (IL)-17, tumor necrosis factor (TNF)-α, nitrotyrosine, matrix metalloproteinase (MMP)-13, and osteoclastogenesis inhibition. (19,20) and acts as a safeguard against alteration induced by stress in p53 production and the antiapoptotic protein (Bcl-2) in human oral epithelial. As a natural product, the GSE can be used in managing diabetes and its complications. (21)

GSE may strongly lead to the inhibition of osteoclast differentiation, the reduction of osteoclast activity, and the stimulation of bone formation through the positive action it has on osteoblast dif-
ferentiation. As a result, it may be useful for treating the inflammation related to bone destruction.\(^{(22)}\) The anti-inflammatory effect of GSE is generated by the calibration of the delicate balance between pro-inflammatory and anti-inflammatory cytokines through the regulation of their release and gene expression.\(^{(23)}\) Nevertheless, the data on GSE effects on healthy and diseased periodontal tissues is limited. It was shown that GSE can protect against collagen breakdown,\(^{(24)}\) and have a bacteriostatic influence on the anaerobes, which can result in a significant decrease in dental biofilm maturation. Consequently, it can be employed in preventing periodontal disease.\(^{(25)}\)

This present study was designed for evaluating the influence of grape seeds extract on the alveolar bone and periodontal ligaments in diabetic rats.

**MATERIALS AND METHODS:**

This current study was approved by the Research Ethics Committee number 2021/385, Faculty of Dentistry, Suez Canal University, and all experiments were conducted according to the Guidelines for Experimental Animal Studies.

**Sample size calculation:**

By utilizing G*Power version 3.1.9.4, Faul et al., (2007), the sample size was estimated and the effect size was 0.65 using alpha (\(\alpha\)) level of 0.05 and Beta (\(\beta\)) level of 0.05, i.e., power = 95%; the minimum sample size (n) calculation was a total of 50 samples for 5 groups.\(^{(26)}\)

**Animals and experimental model**

In total, 50 male albino rats (140-180 g) were used in the experiment. Animals were kept in groups, 3 rats in each cage during the experiment. They were provided with unlimited access to rat food and tap water. In addition, they were kept in a 12 h light/12 h dark cycle at the animal house of Suez Canal University and stored in an air-conditioned room at 21-23°C and 60-65 % relative humidity.

**Grouping**

The rats were distributed to three groups (10 rats per each group) as follows:

- **Group I (\(-ve\) control):** served as a negative control
- **Group II (Diabetic Group):** a single intraperitoneal injection of streptozotocin (STZ) to given to rats.\(^{(27)}\)
- **Group III:** a single intraperitoneal injection of STZ was given to animals, the same as group II, then diabetic rats were treated with 50 mg/kg grape seed extract for 6 weeks.\(^{(28)}\)
- **Group IV:** a single intraperitoneal injection of STZ was given to animals, the same as group II, then using 100 mg/kg grape seed extract, the diabetic rats were treated for 6 weeks.\(^{(28)}\)
- **Group V:** a single intraperitoneal injection of STZ was given to animals, the same as group II, then using 10 mg/kg gliclazides, diabetic rats were treated for 6 weeks.\(^{(29)}\)

**Induction of diabetes**

Before inducing diabetes, animals were kept fasting for 24 hours. Then, 60 mg/kg body weight of STZ, which were freshly dissolved in citrate buffer (0.1 M, pH 4.5), were intraperitoneally injected as a single dose to induce diabetes, while vehicle buffer was injected for negative control rats. Following STZ injection fasting blood glucose (FBG) level was examined at 72h by obtaining blood sample from the tail vein of animals. On a glucose strip test in a glucometer (EasyGluco Blood Glucose Monitoring system, Infopia, Korea), a blood drop was placed and it was confirmed that rats are diabetic when the FBG level was above 250 mg/dl.\(^{(27)}\)

At the end of the experimental study, all rats were sacrificed using cervical dislocation. The samples of jaws of rats were collected for histological and immunohistochemical assessment.
Samples Preparation

After the sacrifice, soft tissues are dissected in each specimen and the whole mandibles are removed and bisected. For bone decalcification, the specimens were assembled and immersed in 10% formalin, then after 24-48 hours they were inserted in 10% EDTA (pH 7.4) and the solution was replaced every week for 3-5 weeks. The specimens were flushed with phosphate buffered saline PBS and then immersed in 70%, 80%, 96% ethanol (90 minutes for each one), three absolute ethanol immersions (60 minutes each), two xylol immersions (90 minutes each) and two liquid paraffin immersions (120 minutes each). Finally, mesiodistal sections (5 μm thickness in width) at the molar area were sliced with microtomes from the blocks that were deparaffinized in an oven at 60˚ overnight and in xylene for 1 h and then rehydrated to dye by using Masson’s Trichrome, hematoxylin and eosin. Sections were examined for all groups in terms of bone decalcification. Hematoxylin and Eosin results were assessed using the J Image analyzer computerized system.

Statistical analysis

In order to hold a comparison of the means of immunoreactivity of Bcl2 between different group, the ANOVA Test (F test) was used for analyzing data. After one a way ANOVA (F test), a Post hoc test was carried out to identify whether there is any significant difference between the individual groups. P-value was considered significant at p<0.05.

Hematoxylin and Eosin results

Group I (Negative control group): by examining group I, the periodontal ligament composed of cells, fibers, blood vessels & nerves were revealed. The collagen fibers which are the main components of the PDL were composed of the gingival group of fibers attached to the cervical part of the cementum and extended to the free and attached gingiva, where their fusion with the gingiva’s lamina propria took place. The interdental or trans-septal group of fibers were extending from the cementum of one tooth to the adjacent tooth’s cementum crossing above the alveolar bone’s crest. The alveolo-dental group of fibers attached to the cementum from a side and to the alveolar bone on the other side and were subdivided into: the fibers of alveolar crest, horizontal, apical, and oblique inter-radicular fibers. Cells of the PDL were predominantly fibroblasts, progenitor cells and some of the defensive cells were seen. Interstitial spaces between the bundles of fibers were demonstrated having blood vessels and alveolar connective tissue. The alveolar bone appeared normal, consisted of the proper alveolar bone, and reinforced alveolar bone. The alveolar bone proper that forms sockets’ inner walls and contains Zuckerkandl and Hirschfeld’s canals openings was formed of bundle bone and lamellar bone. The fibers of Sharpey were inserted into the bundle bone, which is the alveolar bone proper’s part that forms the innermost walls of the socket. Adjacent to it, the lamellar bone was found...
where the lamellae were arranged either parallel to each other or in the form of Haversian systems. The supporting alveolar bone is composed of spongiosa and cortical plates of compact bone. The alveolar bone showed normal turnover rate, as the number of the reversal lines were minimal. Besides, the PDL/ bone interface appeared to be smooth; no osteoclastic activity was observed. (Fig. 1)

**Group II (Diabetic Group):** examination of periodontal ligaments of rats injected with a single dose of STZ intraperitoneally for diabetes induction showed detachment and dissociation of the principle fibers, areas of vacuolization and dissolution of periodontal ligaments fibers and marked dilatation of blood vessels and areas of hemorrhage. The bone surface showed a lot of Howship’s lacuna and osteoclasts, widening of the marrow cavities & inflammatory cells infiltrate. Thinning of bone trabeculation and widening of Zuckerkandl and Hirschfeld canals also were noted. Hyercementosis was noted in some samples. (Fig. 2)

**Group III:** Diabetic animals treated with grape seeds 50mg/kg for 6 weeks revealed poor progress in the bone and periodontal ligament status. Periodontal ligaments showed more organized periodontal ligaments focal areas of detachment, and fiber dissociation, dilatation of blood vessels and inflammatory cell infiltration. A lot of Howship’s lacuna and osteoclastic activity on the bone surfaces, widening of the marrow cavities, thinning of bone trabeculation, widening of Zuckerkandl and Hirschfeld canals and hypercentnosis was recorded. (Fig. 3)

**Group IV:** animals treated with GSE in a 100mg/kg/day dose for six weeks displayed partial improvement of periodontal ligaments, mostly in the arrangement and association of the fibers of PDL. The fibers mostly regained their arrangement and association together in bundles. Dilatation of the blood vessels was still observed, however in lesser degree. Alveolar bone showed new bone formation, a lot of reversal line minimal and Howships lacanue were recorded. (Fig. 4)

**Group V:** the animals treated with Gliclazide in a 10mg/kg/day dose for 6 weeks showed improvement in periodontal ligaments orientation. Dissociation and detachment of the fibers from both tooth and bone sides were recorded. Dilatation of the blood vessels were observed, however in a lesser degree. Epithelial rest of malassezes were demonstrated in clusters or large proliferated clusters of cells. The bone–surface showed a lot of Howships lacunae and numbers of reversal lines. Zuckerkandl and Hirschfeld canals showed widening with blood vessels dilatation. Inflammatory cell infiltrations were recorded all over the periodontal ligaments. Cementum showed focal resorbed areas. (Fig 5)

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![Fig. (1) A photomicrograph of group I showing (A) gingival, transeptal (T) alveolar crest (A), horizontal (H) and oblique (O) periodontal ligaments principle fibers Interseptal bone (star). (B) Apical principle fibers of periodontal ligaments (star) and Zuckerkandl and Hirschfeld canals (arrow). (c) Interradicular fibers and interradicular bone (I) (Mag. X 200).](image-url)
Fig. (2) A photomicrograph of group II showing (A) dissociation and areas of destruction of periodontal ligaments in interradicular area, widening of Zuckerkandl and Hirschfeld canals (blue arrows). Howship’s lacunae in the interradicular bone surface (black arrows). (B) dissociation and detachment of apical fibers from tooth side with dilated blood vessel engorged with RBCs and inflammatory cell infiltration with dilated blood vessel engorged with RBCs. Notice the hypercementosis in the apical cementum. (C) detachment of oblique fibers from bone side (star) large hemorrhage area (arrow). (D) widening of Zuckerkandl and Hirschfeld canals (star), oblique fibers with focal area of dissolved fiber (blue arrow). Howship’s lacunae and osteoclasts on the bone surface (black arrows). (Mag. X 200, 400)
Fig. (3) A photomicrograph of group III showing (A) dissociation and detachment of apical periodontal ligaments from both bone and root surface, widening of Zuckerkandl and Hirschfeld canals (blue arrow). Howship’s lacunae in the interradicular bone surface (black arrows). Notice the hypercementosis in the apical cementum. (B) apical periodontal ligaments with the infiltration of inflammatory cell and the dilated blood vessel that is engorged with RBCs and the infiltration of inflammatory cell with dilated blood vessel that is engorged with RBCs. C) focal detachment of interradicular fibers from bone side Howship’s lacunae in the bone surface (black arrows) and widening of Zuckerkandl and Hirschfeld canals (star). Focal area of Cementum resorption (blue arrow) (D) higher magnification of (C) photomicrograph showing Howship’s lacunae and osteoclasts on the bone surface (black arrows) and widening of Zuckerkandl and Hirschfeld canals (star) . (Mag. X 200,400)

Fig. (4). A photomicrograph of group VI showing (A) almost normal apical periodontal ligaments with dilated blood vessel engorged with RBCs and inflammatory cell infiltration, Howship’s lacunae in the interradicular bone surface (black arrows). (B) interradicular PL. with focal areas of fiber dissociation. Reversal lines (arrows heads). (C) almost normal Transeptal (T) and alveolar crest (A) PL fibers. Reversal lines (arrows). (Mag. X 200)
Masson’s Trichrome Results

The histological examination of Masson’s trichrome staining to evaluate periodontal ligament and bone regeneration, the negative control samples showed no new bone and periodontal ligament fiber formation, most of fiber were blue colors (mature) minimal amount of new cementum formation. The periodontal ligaments and alveolar bone of rats treated with both STZ revealed mature dissociated fiber (pale blue color) with a few amount of new fibers formation, no new alveolar bone formation and new cementum formation. GSE group of dose 50mg/kg showed little amount of fibers formation and focal areas of bone formation. In a group treated with GSE in 100gm/kg dose new fiber of periodontal ligaments (strong red colored fibers) also new alveolar bone and new cementum formation (Red color). Group V treated with Gliclazide in 10mg/kg revealed a few amount of new fibers, bone and cementum formation. (Fig 6)

Immunohistochemical localization of Bcl2 Results

Group I (Negative control) showed mild staining reactivity for both periodontal ligaments and alveolar bone to Bcl2 while Group II (Diabetic group) and group III showed strong staining for both periodontal ligaments and bone to Bcl2. Group IV showed moderate to mild staining reactivity to Bcl2 for both periodontal ligaments and bone. Group V showed moderate reactivity to bcl2 for periodontal ligaments and bone. (Fig. 7)
Fig. (6) Photomicrograph of Masson’s Trichrome (A) control (-Ve control) no PDL, bone and minimal new cementum formation (Red color). (B) STZ group with minimal new fiber formation, minimal new bone and new cementum. (C) group III showed mostly pale blue colored dissociated fibers, minimal amount of new fibers, new bone formation. (D) Group IV showed new fibers formation (Strong red colors) areas of new bone formation and new cementum. (E) Gliclazide group showed minimal new fibers and little focal areas of new bone formation and new cementum notice the epithelial rest of Malassezes (arrows)
Fig. (7) A photomicrograph of the periodontal ligament and bone incubated with mouse monoclonal antibody of Bcl2 from (A) from Control groups (negative control) showing mild immunostaining reactivity of the periodontal ligaments and bone cells to Bcl2 (B) Group II (Diabetic group) showing severe immunostaining reactivity of the periodontal ligaments and bone cells to Bcl2 (C) Group III showing severe immunostaining reactivity of the periodontal ligaments and bone cells to Bcl2 (D) Group IV showing mild to moderate immunostaining reactivity of the periodontal ligaments and bone cells to Bcl2. Group V showed moderate immunostaining reactivity of the periodontal ligaments and bone cells to Bcl2 (Mag. X 250).

Statistical analysis:

Using SPSS version 22 (SPSS Inc., Chicago, IL, USA), the statistical analysis of results was performed. ANOVA (F test) was utilized for analyzing data. After one way ANOVA (F test), a Post hoc test was carried out to identify whether the individual groups are significantly different.
Concerning Bcl, the results in table (1) revealed that there is a significant difference among groups for PL and alveolar bone at P-value (P<0.05). The diabetic group gave the highest value followed by group III for PL and alveolar bone. Group V also showed a significant difference with the negative control but with a lesser value than group II and III. Group VI revealed no significance with negative control of value 97.5 and 93.57 for periodontal ligaments and alveolar bone respectively. (Table1, Fig 8)

TABLE (1). Descriptive statistics and results of one way ANOVA test for comparison between means of positive cells in the five groups

<table>
<thead>
<tr>
<th>Group</th>
<th>PL</th>
<th>Alveolar bone</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I</td>
<td>73.94±3.42</td>
<td>75.61±8.99</td>
</tr>
<tr>
<td>Group II</td>
<td>107.92±2.04</td>
<td>106.37±2.89</td>
</tr>
<tr>
<td>Group III</td>
<td>101.85±2.25</td>
<td>100.89±2.24</td>
</tr>
<tr>
<td>Group IV</td>
<td>76.49±6.46</td>
<td>77.78±4.86</td>
</tr>
<tr>
<td>Group V</td>
<td>97.29±5.09</td>
<td>93.57±2.32</td>
</tr>
<tr>
<td>F test</td>
<td>119.83</td>
<td>68.71</td>
</tr>
</tbody>
</table>

P value<0.05 <0.001** <0.001**

**;a,b,c:means significant difference between groups at same column using F-Test

DISCUSSION

It is estimated that more than 400 million adults diabetes on a worldwide basis, (30) most of them (> 90%) have type-2 diabetes mellitus that is resulting from combining insulin resistance and failing to compensate by the increase in the production of insulin. (31) Increased gingival inflammation and Periodontitis are observed in subjects with diabetes mellitus. (32, 33) Diabetes has a detrimental influence on bone, resulting in an increase in risks of fracture and osteolytic lesions’ formation such as those found in periodontitis. Many diabetic difficulties result from diabetes-enhanced inflammation. (34) In periodontal tissues, diabetes results in the enhancement of inflammation, RANKL expression, the loss of periodontal bone, and osteoclastogenesis. (35) Inducing experimental periodontal disease results in the stimulation of NF-κB activation in the fibroblasts of periodontal ligament, osteoblasts, and osteocytes, which is related to the loss of bone and the reduction of bone coupling, which represent two key factors in periodontitis. (36) Diabetes leads to the reduction of bone-lining cells’ numbers, osteoblasts, and the fibroblasts of periodontal ligament, which may lead to limiting bone coupling. (37) In diabetic animals, Osseous coupling can significantly reinforced by a TNF specific inhibitor, which can rescue bone formation and expressing bone-producing factors including FGF-2, TGFβ-1, BMP-2, and BMP-6. (38)

Oxidative stress is increased by Diabetes increases, which in turn can result in worse insulin action and secretion, consequently resulting in the acceleration of the progression to overt the disease. (39) Studies argued that oxidative stress markers Nox1, Nox2, Nox4, and p47 increase diabetes model rats. (40) Reactive Oxygen Species’ elevation (ROS) might have the most critical role to play in the establishment and progression of periodontitis. (41) The increase in ROS results in generating an impaired formation of bone through the activation of bone resorption and forkhead box transcription factors (FoxO) by decreasing Wnt signaling. (42)
Kolluru et al., reported that the endothelial dysfunction resulting in defective angiogenesis in diabetes has multiple factors that include the decreased growth factors and cytokines, the increased ROS and AGEs, and altered immune cell responses.\(^{(43)}\)

The previous studies explained the alveolar bone loss, inflammatory cell infiltration, periodontal ligament dissociation, destruction, and blood vessels dilatation which were noted in diabetic group of the present study. The H&E results were confirmed by the Masson’s trichrome stain examination which revealed pale, blue-stained dissociated fibers, with the absence of new bone formation. New cementum formation was noted might compensate for the bone loss in teeth sockets.

Proanthocyanidins are described as the natural antioxidants and complicated polymers of flavonoids.\(^{(44)}\) A higher level of proanthocyanidins is contained in legume seeds, grains, various vegetables, and particularly fruits contain. Recently, several studies have revealed their different pharmacological activities including free radical scavenging, anti-inflammatory activities, and antioxidant. Proanthocyanidins are characterized by having application prospects and possible clinical value as a therapeutic agent for diabetic comorbidities.\(^{(45)}\)

Oligomeric proanthocyanidins (PACs) are contained in grape seed extract (GSE) (\textit{Vitis vinifera} L.) and display strong antioxidant activity. It has been proven that PACs are capable of controlling malondialdehyde (MDA) and carboxymethyllysine’s higher levels, reducing superoxide dismutase, and reducing the activity glutathione (GSH) in mice suffering from induced diabetes mellitus.\(^{(46)}\)

A histopathological study,\(^{(46)}\) which was carried out in 2018, examined grape seed proanthocyanidin extract. The authors reported that GSPE is characterized by having a polyphenolic structure and having a broad spectrum of biological activity. In addition, they measured the loss of alveolar bone morphometrically by utilizing a stereomicroscope. For purposes of histopathological analyses, they performed vascular endothelial growth factor immunohistochemistry, alizarin red staining and matrix metalloproteinase, and inducible factor hypoxia. In addition they determined the relative total inflammatory cells in tetract acid-positive osteoclasts. This animal study revealed that the administration of GSPE in (100mg/kg, 200mg/kg) can cause a decrease in periodontal inflammation and the loss of alveolar bone through the reduction of levels of alpha matrix metalloproteinase and hypoxia-inducible factor, and the increase in osteoblastic activity in diabetic rats having periodontitis.\(^{(46)}\) The previous study come inconsistent with the results of group IV animals treated with grape seeds of 100mg/kg in the present investigation.

In the present investigation 2 different concentrations of grape seeds were administrated, in group III, the diabetic animals treated with grape seeds of 50mg/kg for 6 weeks showed no noticeable effect on alveolar bone resorption or periodontal ligaments degeneration. These results were confirmed by Masson’s trichrome which revealed a minimal amount of new fibers and bone formation. While in group IV, diabetic rats were treated with a concentration of 100/ kg for 6 weeks; there was marked improvement indicated through both histological examinations of H&E and Masson’s Trichrome stain, in the alveolar bone through new bone formation and periodontal ligaments showed less dissociation and detachment with new fibers formation. New cementum also demonstrated with more organized and oriental shape. This current study suggested that the grape-seeds dose in group III may be not sufficient to treat the complication of diabetes in alveolar bone and PL or may be the administrated dose need more time to induce the effect that occurred in group IV. The grape seeds dose of 100/kg group might decrease the oxidative stress on cells and increase their biological activity to regenerate.

Gliclazide represents an anti-diabetic medication, which is a second-generation sulfonylurea. Insulin is released by Sulfonylureas from pancre-
atic cells and Sulfonylureas act on insulin-sensitive tissues for the enhancement of glucose uptake. (47) Whereas these agents can directly lead to stimulating insulin secretion by the β-cell, it has been also shown that Sulfonylureas have anti-inflammatory influences. (48) The stimulation of insulin leads to activating distinct pathways that are involved in metabolic regulation such as the phosphatidylinositol-3-kinase (PI3K) cascade. Further, Gliclazide can directly influences PI3K insulin-resistant skeletal muscle for enhancing insulin signalling. (49) The pathway of PI3K signaling has an effect on the inflammatory process, which contributes to increasing neutrophil survival, (50) and the osteoclast differentiation pattern. (51) In tissue from patients with periodontitis, the expression of PI3K is higher than that in healthy gingival tissue. (52)

In the present investigation, group V treated with gliclazide 10 mg/kg for 6 weeks showed infiltration of inflammatory cells, the periodontal ligament has no noticeable improvement in their association and attachments comparing to the diabetic group, unless the orientation of the PL were better than diabetic group. Resorption of alveolar bone and a lot of Howship’s lacunae on the bone surface, numbers of reversal line revealed less new bone formation, the H&E results were confirmed by results of Masson’s trichrome stains as minimal new fibers and bone. Xing et al provided an explanation of these group V results from the current examination, where they indicated that the higher doses of gliclazides (for example 10 mg/kg) can lead to activating the pathway of PI3K/AKT and increasing periodontal bone loss. In addition, they argued that some cytokines could also lead to activating the PI3K/AKT pathway, resulting in osteoclastogenesis. (53)

Clusters and proliferation of epithelial rests of Malassezses were marked and repeated results in the samples of group V animals as they noticed concentrated in interradicular areas and apical region. Epithelial cell rests’ proliferation is frequently found in inflamed periapical lesions. The proliferation of epithelial cells rests takes place in three dimensions, resulting in forming strands or islands of epithelium. In turn, these strands of islands of epithelium are invaded by vascular fibrous connective tissue with various degrees of inflammatory infiltrates. (54) During periapical inflammation, several inflammatory mediators, growth factors, and proinflammatory cytokines are released by host cells in the periapical tissues through the innate and adaptive immune responses. (55) Brunette (56) revealed that when the intracellular level of cyclic adenosine monophosphate (cAMP) is elevated by prostaglandins (PGE2), it leads to stimulating epithelial cell rests’ growth. In the 10 mg/kg gliclazide group, there was an increase in the inflammatory cytokine IL-1. It was revealed that IL-1 stimulates osteoclastogenesis through two paralleled events: enhancing RANKL expression directly and suppressing OPG expression. (53) At low concentrations, IL-1 had a proliferative influence on the epithelial cells, which could play a role in the evocation of an inflammatory reaction and the stimulation of Malassez’s epithelial cell rests in order to proliferate to form radicular cyst. (57) Accordingly, the present investigation suggested that administration of gliclazide in a dose of 10mg/ kg could increase the release of some cytokines, particularly IL-1 that stimulate the proliferation of epithelial rest of Malassezes, Thus, the present study recommended that dental care and follow up of the patient treated with gliclazide of 10mg/kg to detect any pathosis could be developed due to activation of epithelial rest of Malassezes. The present study suggested that gliclazide of 10mg/kg could mediate an inflammatory process which causes resorption in the bone surfaces and cementum on the tooth surface.

The histological results were indicated through immunohistochemical analysis of Bcl2. Concerning Bcl2, the results showed a significant difference between groups for PDL and alveolar bone at P-value (P≤0.05). The diabetic group and Grape seeds of 50mg/kg gave the highest value followed by gliclazide group for P.L and alveolar bone comparing to negative control group. The grape seeds group of 100mg/kg showed no significant difference with the untreated control group.
Apoptosis, which is also known as the programmed death of cell, represents a form of physiological cell death. When there is infection, inflammation or tissue remodeling, it increases or decreases. Apoptosis is considered a very important process that has an effect on the normal development and tissue homeostasis. Apoptotic proteins such as bcl-2 and Bax proteins can contribute to regulating the apoptosis. In addition, the Bcl-2 is utilized as anti-apoptotic protein, whereas the Bax is utilized as a proapoptotic protein, and the balance between such proteins helps to determine the apoptotic process. 

The Bcl-2 family members has an integral role to play in apoptosis. Nevertheless, they can also contribute to several other cellular functions. Among all the Bcl-2 family members, Isoforms are recognized and some of them are well described. The therapeutics which target BCL-2 indicates a great promise of treating cancer and degenerative diseases. There is a well-recognized fact that prolonged hyperglycemia can cause the dysfunction of several organ systems. Furthermore, type 2 DM is related to the difficulties in the kidneys, eyes, and arteries, and impairing these organ systems is involved in inflammatory processes. Many studies revealed the overexpression of Bcl-2 and Bax proteins in various tissues with hyperglycemia. 

On the other hand, former studies have argued that apoptosis is involved in inflammatory periodontal disease’ pathogenesis. They also revealed that the higher frequency of Bcl-2 expression leads to progressive periodontal destruction. To the best of our knowledge, this current study is one of few trials to demonstrate the effect of GSE on the expression of Bcl2 in the alveolar bone and periodontal ligament in diabetic conditions.

A study of Güçlü et al indicated that using grape seed as treatment might be useful against apoptosis and oxidative stress in diabetic rats. The exposure to GSEs resulted in the activation of cleavage of caspase-2, caspase-3 and caspase-9 in HCT-116 cells, the induction of p53-mediated mitochondrial apoptosis signaling pathway with a concentration-dependent decrease in the level of expression of the survival protein Bcl-2, and the increase in the level of expression of the pro-apoptotic proteins, Bax and Bak. The previous study comes inconsistent with the investigation results of group seeds of dose 100mg/kg as there was no significant difference with negative control. While in group III grape seeds of dose 50mg/kg the bcl2 expression was high as mentioned, this indicates the suggestion of insufficient dose of grape seeds or insufficient time of treatment.

In group V the bcl2 expression showed significant difference from untreated controls this comes in the agreement with a study of Kanazawa et al, the authors revealed animals that are treated with gliclazide at a 10 mg/kg dose, suggesting a hypoglycemic impact at this dose and an increase in the loss of bone in periodontal disease. Another study indicated that metformin has significantly resulted in the activation of AMPK in dose- and time-dependent manners, and the induction of endothelial nitric oxide synthase (eNOS) and bone morphogenetic protein-2 (BMP-2) expressions, that may suggest the persistence of the inflammatory process and cell apoptosis and in turn explain the high bcl2 expression in group V.

According to the present study, blood glucose evaluation at the beginning and end of the experiment is recommend in further researches to investigate whether GSE has a hypoglycemic effect or not in diabetic condition

CONCLUSION

Grape seeds of 100mg/kg improve the condition of periodontal ligaments and alveolar bone of diabetic animals and could represent a supplemental medication for treatment the dental complication of diabetes mellitus. It is required to carry out further comprehensive clinical and experimental studies for investigating the beneficial influences of grape seed extracts in diabetes mellitus, which indicate decrease in Bcl2 expression.
REFERENCES


24. La VD, Bergeron C, Gafner S, and Grenier D. Grape seed extract suppresses lipopolysaccharide-induced matrix metalloproteinase (MMP) secretion by macrophages and


47. Toker H, Yuce H.B, Alpan AL, Gevrek F, and Elmastas M. Morphometric and histopathological evaluation of the ef-


