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Original Article

## Superimposed effect of ovariectomy on type 2 diabetes mellitus in Wistar rats

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### ABSTRACT

**Background:** Estrogen deprivation in the postmenopausal women plays a critical role in progression of type 2 diabetes mellitus (T2DM).

**Aim:** The present study investigated the overlaid effect of ovariectomy on T2DM and the possible underlying mechanism.

**Materials:** Forty female Wistar rats were divided into four groups (10 rats each): sham control, ovariectomized control, sham diabetic and diabetic ovariectomized groups. At the end of experiment, estimation of body weight gain percentage, food intake, fasting blood glucose concentration, and insulin tolerance test were done. Then, rats were euthanized and blood samples were taken for measurement of serum concentration of insulin, HOMA-IR, lipid parameters, tumor necrosis factor- $\alpha$ , interleukin 1 beta, interleukin 4, interleukin 10, malondialdehyde and total thiol. Also, histopathological and immunohistochemical examination of the pancreas were done.

**Results:** The present study revealed that ovariectomy aggravated the diabetic induced glucose metabolic disturbance as implied by impaired insulin tolerance test, increased insulin resistance alongside lipid dyshomeostasis. These metabolic disturbances might claim to exacerbation of oxidative stress and inflammatory response along with apparent histopathological and immunohistochemical changes on the pancreas.

**Conclusion:** We concluded that metabolic disturbances induced by diabetes might be getting worse after ovariectomy via intensification of oxidative stress and inflammatory state.

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### 1. Introduction

Type 2 diabetes mellitus (T2DM) is a progressive chronic metabolic disease characterized by elevated blood glucose concentration and insulin resistance.<sup>1</sup>

Estrogen deficiency after menopause correlates with metabolic disorders as T2DM, and cardiovascular disease;<sup>2</sup> this designates its role in the pathogenesis of these conditions.<sup>3</sup> Beside, the role of estrogens in female physiology and reproduction, now they are considering important regulators of multiple physiological and pathological processes.<sup>4</sup> Consequential to menopause, the relation between estrogen concentrations and the risk of developing diabetes is complex. Margolis et al.<sup>5</sup> showed lower risk of diabetes

in women taking hormone replacement therapy. Meanwhile, Ding et al.<sup>6</sup> reported insulin resistance and diabetes in women with elevated endogenous estrogen levels after menopause.

Low-grade inflammation plays a critical role in the pathophysiology of various chronic age related conditions such as T2DM,<sup>7</sup> alongside increased level of pro-inflammatory cytokines observed in T2DM.<sup>8</sup> Cytokines are small soluble proteins or membrane-bound proteins. Almost all nucleated cells generate these proteins and respond to them. Cytokines include “interleukins, growth factors, chemokines, hematopoietins and colony stimulating factors”.<sup>9</sup>

Estrogen deficiency after menopause is coupled with oxidative stress,<sup>10</sup> a state occurs when the production of reactive oxygen species (ROS) exceed the body's anti-oxidant defenses.<sup>11</sup> ROS are oxidative physiological processes end products<sup>12</sup> that bind with protein, lipid, carbohydrates and DNA within the cells, result in cellular damage include cell membranes and genetic materials.<sup>11</sup> Given that the amplified levels of ROS contribute to the pathogen-

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esis of T2DM,<sup>13</sup> this may suggest an increased risk of development of T2DM in postmenopausal females.

The present study was planned to examine the superimposed impact of surgically induced menopause (ovariectomy) on body weight gain percentage and food intake, glucose and lipid metabolism, insulin tolerance, and insulin resistance in a model of T2DM induced by high fat diet and low dose streptozotocin and to elucidate the potential mechanisms of this effect.

## 2. Materials and methods

### 2.1. Animals

Forty Wistar female rats, initially weighing from 150 to 200 grams, aged 3 months obtained from Animal house of Faculty of Medicine, Assiut University. Rats housed in clean stainless steel cages, the cage size was 65 cm × 25 cm × 15 cm, and each cage contained 5 rats. Rats maintained on natural light/dark cycle in an aerated room under controlled hygienic conditions and provided with free access to water. All procedures conducted in accordance to the *Guide of the Care and Use of Laboratory Animals*<sup>14</sup> and approved by the Ethical Committee for Scientific Research at the Faculty of Medicine.

Following acclimation to laboratory circumstances, 24 h food intake and body weight measured 3 times per week for 2 weeks between 8.00 a.m. and 10.00 a.m. These values utilized as baseline values. 24 h food intake estimated by offering a standard amount of rodent food pellets (70 g for each rat) every day. Then, food intake measured by subtracting the leftover food from what added at the previous day.

### 2.2. Experimental design

The rats randomly divided into four groups (10 rats per group):

1. Group 1 (Sham control group): rats subjected to simulated surgery.
2. Group 2 (Ovariectomized control group): rats subjected to bilateral ovariectomy.
3. Group 3 (Sham diabetic group): T2DM was induced then; rats underwent sham surgical procedure.
4. Group 4 (Diabetic ovariectomized group): diabetic rats were subjected to bilateral ovariectomy.

### 2.3. Induction of T2DM

T2DM was induced as mentioned by Wang et al.<sup>15</sup> by feeding rats with a high-fat diet (HFD), which prepared by adding 20% sucrose (w/w) and 10% ram tail fat (w/w) into basal diet, for 4 weeks, then rats injected with a single intraperitoneal (i.p.) administration of freshly prepared streptozotocin (STZ) dissolved in ice-cold sterile saline 9% (CAT #18883664; Sigma, Germany) in a dose of 30 mg/kg body weight, the combination of HFD and low-dose STZ surrogates as an alternative animal model resembling the human type 2 diabetes. This STZ low dose exhibited a slight trauma to pancreatic beta cells to simulate the condition of chronic hyperinsulinemic insulin resistance.<sup>16</sup> To prevent the STZ induced hypoglycemia, rats injected with 10% dextrose solution 6 h after STZ administration and for the next 24 h.<sup>17</sup> After that rats continued to feed HFD for another 4 weeks.

### 2.4. Surgical procedure: ovariectomy

After establishment of diabetes, rats sedated with pentobarbital sodium anesthesia (60 mg/kg, i.p.; Sigma–Aldrich Chemie GmbH, Steinheim, Germany). Bilateral ovariectomy performed according to the method of Khajuria et al.<sup>18</sup> Sham-operated rats anaes-

thetized, the skin and muscle layers opened; ovaries manipulated but not removed.

Rats allowed to recover in individual cages for one week, during which they received a commercial pelleted diet and water ad libitum, anti-inflammatory drug ketoprofen (2 mg/kg) for three days, and antibiotic ampicillin sodium (30 mg/kg) for five days as described by Tavares et al.<sup>19</sup>

### 2.5. Blood glucose Determination

Occurrence of diabetes was confirmed by measuring fasting blood glucose concentration (FBG) with strips using glucometer (Accu-Chek Active, Roche Diagnostic Corporation, Mannheim, Germany). Rats with FBG concentrations more than 200 mg/dl considered diabetic.<sup>20</sup>

### 2.6. Insulin tolerance test

Rats were fasted for 6 hours (from 7 a.m. to 1 p.m.) and each rat received a single intraperitoneal injection of insulin (1 IU/kg body weight) and tail blood samples used for determining concentration of blood glucose at 0, 30, 60, and 120 min.

### 2.7. Animal euthanize

Fasting rats (18 h) anesthetized and euthanized by lethal dose of pentobarbital (100 mg/kg body weight).<sup>21</sup> 4 ml of blood collected from the retro-orbital venous plexus before euthanize. Serum separated by centrifugation at 3000g for 20 min. The separated serum aliquotted and stored frozen at –20 °C until use.

### 2.8. Determination of homeostasis model of insulin resistance (HOMA-IR)

Serum insulin concentration measured using a rat insulin enzyme-linked immunosorbent assay (ELISA) kit (CAT # RAB0904; Sigma, Germany) according to the manufacturer's instructions. The insulin resistance index (IRI) assessed by homeostasis model assessment estimate of insulin resistance (HOMA-IR):

$$\text{HOMA-IR} = \frac{\text{Fasting insulin (IU/ml)} \times \text{Fasting glucose (mmol/L)}}{22.5}$$

### 2.9. Determination of serum lipids

Total cholesterol (TC), triglycerides (TG) and high density lipoprotein-cholesterol (HDL-C) determined by enzymatic methods using an automated analyzer (Dimension RxL Max, DADE Behring, Marburg, Germany). Low density lipoprotein-cholesterol (LDL-C) concentrations estimated according to the formula specified by Friedwald et al.<sup>22</sup> as follows:

$$\text{LDL-cholesterol} = \text{Total cholesterol} - [\text{HDL-cholesterol} + \text{TG}/5].$$

### 2.10. Measurement of cytokines

ELISA performed to measure serum concentrations of TNF- $\alpha$  (CAT # K0331196), IL1 $\beta$  (CAT # K0331212), IL4 (CAT # K0332133) and IL10 (CAT # K0332134) using KOMA Biotech commercial kits and following the instructions supplied with each kit.

### 2.11. Measurement of serum concentrations of malondialdehyde and total thiol

Serum concentration of malondialdehyde (MDA) estimated by the method of Uchiyama and Mihara.<sup>23</sup> Calculation of concentrations = A × 3.84 where A = absorbance.

Total thiol (TT) serum concentration measured according to the method of Ellman.<sup>24</sup> TT concentrations in mmoles/L calculated according to the following  $Co = A/E \times D$ , where Co = Original concentration, A = absorbance at 420 nm, E = extinction coefficient = 13.1000 M/cm and D = diluted factor = 36.8 M/cm. The concentrations of TT in serum =  $36.8/5 \times A$ .

2.12. Histopathological examination

Specimens took from tail area of the pancreas, fixed in buffered formalin solution, dehydrated conventionally and embedded in paraffin. 4 μm sections were cut with a Leica sliding microtome (SM 2000R, Nussbach, Germany), then the slides were stained with haematoxylin and eosin as described by the method of Durry and Wallington.<sup>25</sup>

2.13. Immunohistochemistry of pancreas

Pancreatic tissue excised, fixed, and stained for insulin as previously described by Kluth et al.<sup>26</sup> The changes in the immunohistochemical reaction of the pancreatic tissues were detected by dad chromogen and counter-staining was done using Hematoxylin.

2.14. Statistical analysis

Data expressed as mean ± SD (standard deviation) using Graph-Pad Prism software version 6 (Graph Pad Software, San Diego, CA). Differences among the groups compared by one-way analysis of variance (ANOVA) test followed by Bonferroni's multiple comparison tests. P value of <0.05 considered to be significant.

3. Results

3.1. Body weight gain percentage and food intake

Ovariectomy following diabetes exhibited no impact on the body weight gain percentage and food intake (Fig. 1A,B).

3.2. FBG, insulin concentrations and insulin resistance

FBG concentration elevated significantly in the sham diabetic and OVX diabetic rats when compared to the sham control rats (P < 0.001 for each) and OVX control rats (P < 0.001 for each) (Fig. 2A).

OVX diabetic rats demonstrated a significant increase in insulin concentrations when compared to the sham control rats

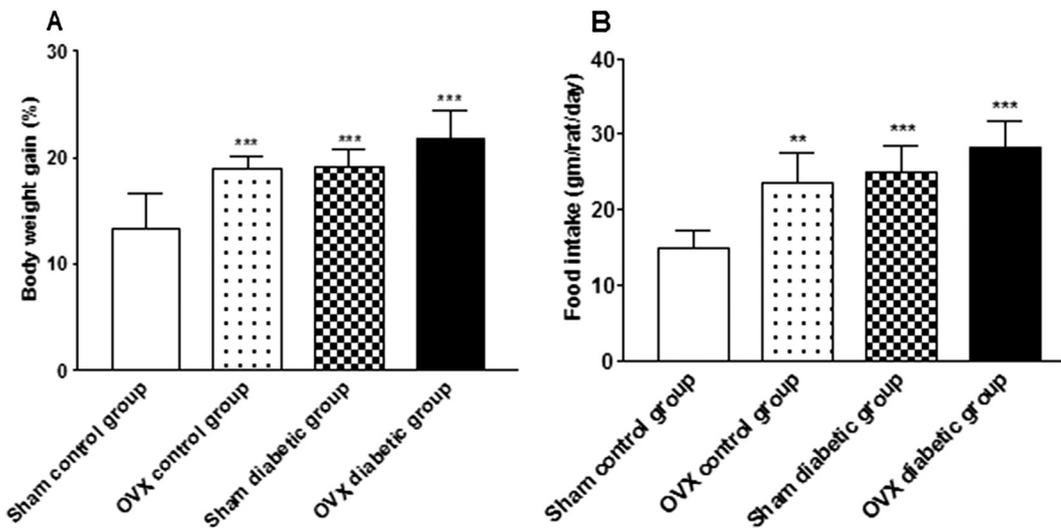


Fig. 1. Body weight gain percentage (A) and food intake (B) in the studied groups. Data were presented as mean ± standard deviation, n = 10 rats per group. OVX: ovariectomized rats. Significant difference from the sham control group \*\*: P < 0.01 and \*\*\*: P < 0.001, using one-way analysis of variance (ANOVA) followed by Bonferroni test.

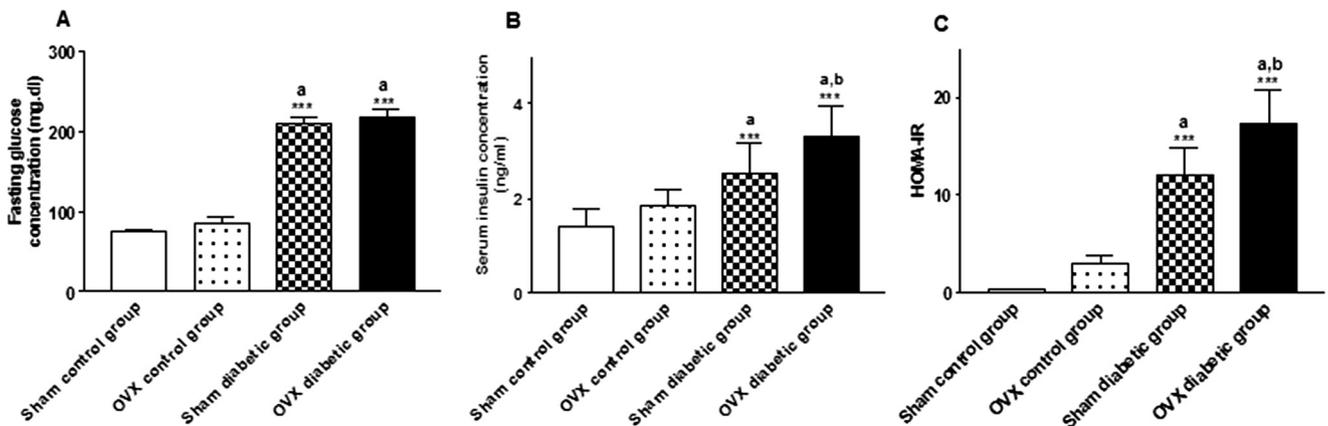
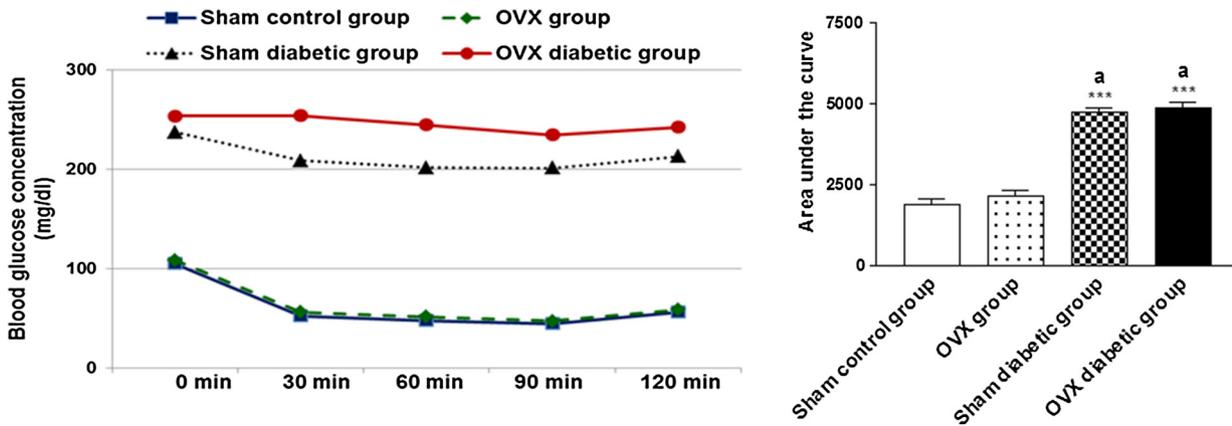


Fig. 2. Fasting blood glucose concentration (A), serum insulin concentration (B) and HOMA-IR (C) in the studied groups. Data were presented as mean ± standard deviation, n = 10 rats per group. OVX: ovariectomized rats. Significant difference from the sham control group \*\*\*: P < 0.001, a: significantly different from the OVX group and b: significantly different from the sham diabetic group, using one-way analysis of variance (ANOVA) followed by Bonferroni test.



**Fig. 3.** Insulin tolerance test in the studied groups. Data were presented as mean ± standard deviation, n = 10 rats per group. OVX: ovariectomized rats. Significant difference from the sham control group \*\*\*:  $P < 0.001$ , a: significantly different from the OVX group and b: significantly different from the sham diabetic group, using one-way analysis of variance (ANOVA) followed by Bonferroni test.

( $P < 0.001$ ), OVX control rats ( $P < 0.001$ ) and sham diabetic rats ( $P < 0.01$ ) (Fig. 2B).

OVX diabetic rats revealed a significant increase in HOMA-IR value as compared with the sham control ( $P < 0.001$ ), OVX control ( $P < 0.001$ ) and sham diabetic group ( $P < 0.001$  for each) (Fig. 2C). Thus, ovariectomy aggravated the diabetes-associated insulin resistance.

### 3.3. Insulin tolerance test

Insulin tolerance test was displayed in Fig. 3. OVX diabetic and diabetic rats exhibited a lower effect of insulin on fasting blood glucose concentrations than sham control and OVX control rats.

### 3.4. Lipid profile

The OVX diabetic rats displayed significant elevations in serum concentrations of TC, LDL-C and TG as compared to the sham control rats ( $P < 0.001$  for each parameter), the OVX control rats ( $P < 0.001$ ,  $P < 0.01$  and  $P < 0.001$ , respectively), and the sham diabetic rats ( $P < 0.01$ ,  $P < 0.01$  and  $P < 0.001$ , respectively). On the other hand, serum concentration of HDL-C exhibited a significant decline in OVX control, sham diabetic and OVX diabetic rats when

compared with,  $P < 0.001$  and  $P < 0.001$ , respectively). Additionally, HDL-C concentration decreased significantly in the OVX diabetic rats when compared with the sham control rats ( $P < 0.001$ ) and the OVX control rats ( $P < 0.05$ ) (Fig. 4).

### 3.5. Serum concentrations of TNF- $\alpha$ , IL1 $\beta$ , IL4 and IL10

As shown in Fig. 5, the OVX diabetic rats revealed a significant increase in TNF- $\alpha$  and IL1 $\beta$  when compared with the sham control rats ( $P < 0.001$  for each), OVX control ( $P < 0.01$  and  $P < 0.001$ , respectively) and sham diabetic groups ( $P < 0.01$  and  $P < 0.001$ , respectively).

Regarding serum concentration of IL4, it decreased significantly in OVX diabetic group in comparison to sham control, OVX control and sham diabetic groups ( $P < 0.001$  for each). Furthermore, serum concentration of IL10 exhibited a significant decrease in the OVX control, sham diabetic and OVX diabetic group in comparison with ( $P < 0.05$ ,  $P < 0.05$  and, respectively). the OVX diabetic group showed a significant decrease in serum concentration of IL10 as compared to the sham control group ( $P < 0.001$ ), the OVX control group ( $P < 0.01$ ) and the sham diabetic group ( $P < 0.001$ ).

#### 3.5.1. Serum concentrations of MDA and TT

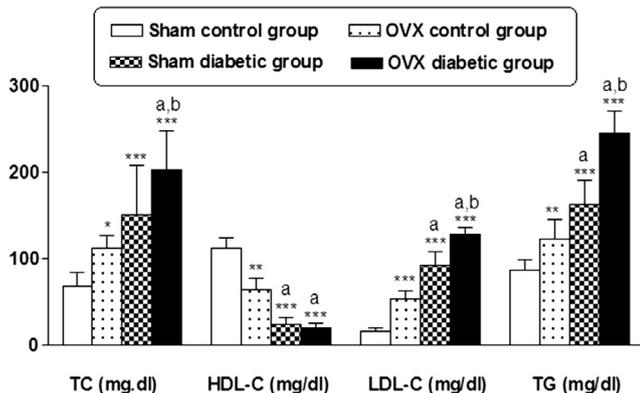
Serum concentrations of MDA and TT in studied groups displayed in Fig. 6. Concerning serum concentration of MDA, the OVX diabetic group showed a significant increase in serum concentration of MDA in comparison to the sham control group ( $P < 0.001$ ), the OVX control group ( $P < 0.001$ ) and the sham diabetic group ( $P < 0.001$ ).

With regard serum concentration of TT, it decreased significantly in the OVX diabetic group as compared to the sham control group ( $P < 0.001$ ), the OVX control group ( $P < 0.001$ ) and the sham diabetic group ( $P < 0.01$ ).

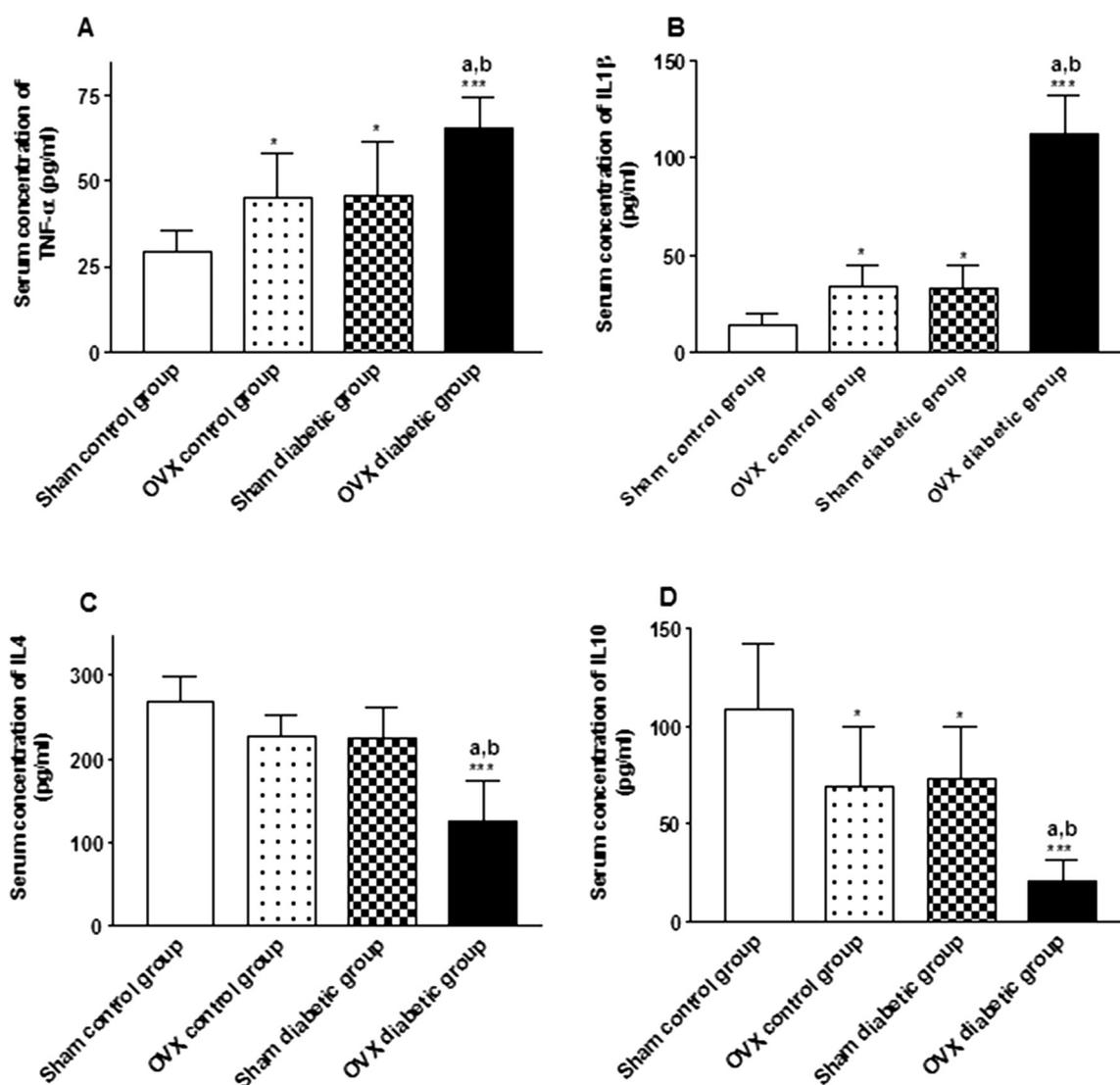
### 3.6. Histopathological results

Histopathology changes of pancreatic islets including immunohistochemistry of insulin in all experimental groups were displayed in Figs. 7 and 8. Pancreas of the sham control group showed regularly rounded islets and interlobular ducts (Fig. 7a) that showed strong immune positivity “dark brown, completely positive islets” (Fig. 8a).

Pancreas of the OVX control group showed regularly rounded islets and proliferating cells around interlobular ducts (Fig. 7b). By immune stain some islet cells were golden brown and some



**Fig. 4.** Serum concentrations of total cholesterol (TC), HDL-C (high density lipoprotein-cholesterol), LDL-C (low density lipoprotein-cholesterol) and triglycerides (TG) in the studied groups. Data were presented as mean ± standard deviation, n = 10 rats per group. OVX: ovariectomized rats. Significant difference from the sham control group \*:  $P < 0.05$ , \*\*:  $P < 0.01$ , \*\*\*:  $P < 0.001$ , a: significantly different from the OVX group and b: significantly different from the sham diabetic group, using one-way analysis of variance (ANOVA) followed by Bonferroni test.



**Fig. 5.** Serum concentrations of TNF- $\alpha$  (A), IL-1 $\beta$  (B), IL4 (C) and IL10 (D) in the studied groups. Data were presented as mean  $\pm$  standard deviation, n = 10 rats per group. OVX: ovariectomized rats; TNF- $\alpha$ : tumor necrosis factor-alpha; IL1 $\beta$ : interleukin 1 beta; IL4: interleukin 4 and IL10: interleukin 10. Significant difference from the sham control group \*:  $P < 0.05$  and \*\*\*:  $P < 0.001$ , a: significantly different from the OVX group and b: significantly different from the sham diabetic group, using one-way analysis of variance (ANOVA) followed by Bonferroni test.

were immune negative. Some of these proliferating cells around some interlobular ducts are immune positive, others related to the interlobular ducts were immune negative (Fig. 8b).

Pancreas of the sham diabetic group revealed that most islets were hyperplastic and irregular shaped islets with groups of endocrine-like proliferating cells of variable sizes observed in relation to both interlobular and interlobular ducts (Fig. 7c). By immune stain, most islets were strong positive completely immune stained (Figs. 8c). Some islets are strongly immune stained with some immune negative cells (Fig. 8d). Other islets contained only few immune positive cells (Fig. 8e). Also we observed groups of endocrine-like proliferating cells of variable sizes in relation to both interlobular and interlobular ducts (Figs. 8c, 8f); some of these cells were immune positive. Also, groups of immune positive cells of variable sizes were frequently observed within the acini of the exocrine pancreas (Figs. 8f).

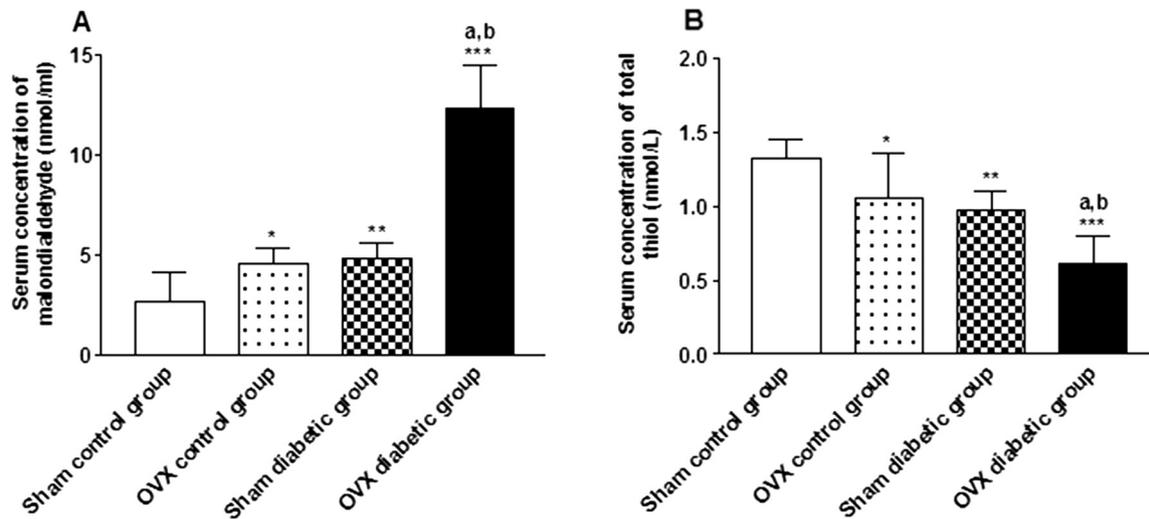
Examination of the OVX diabetic group showed islets that were mostly surrounded by a zone of cells with dark flat nuclei (Fig. 7d). Small sized islets that contain cells with vacuolated cytoplasm and dark flat nuclei were also observed (Fig. 7e). Cellular proliferation

around the interlobar and interlobular ducts was also detected (Figs. 7d). Some of these proliferating cells showed small dark nuclei (Fig. 7e). By immune stain some islets were immune positive with golden brown color and showed some peripheral immune negative cells (Fig. 8g). Occasionally, groups of proliferating immune positive cells within the acini of the exocrine pancreas together with other groups of proliferating immune negative cells were observed, most of the observed proliferating cells around the pancreatic ducts were immune negative (Fig. 8h).

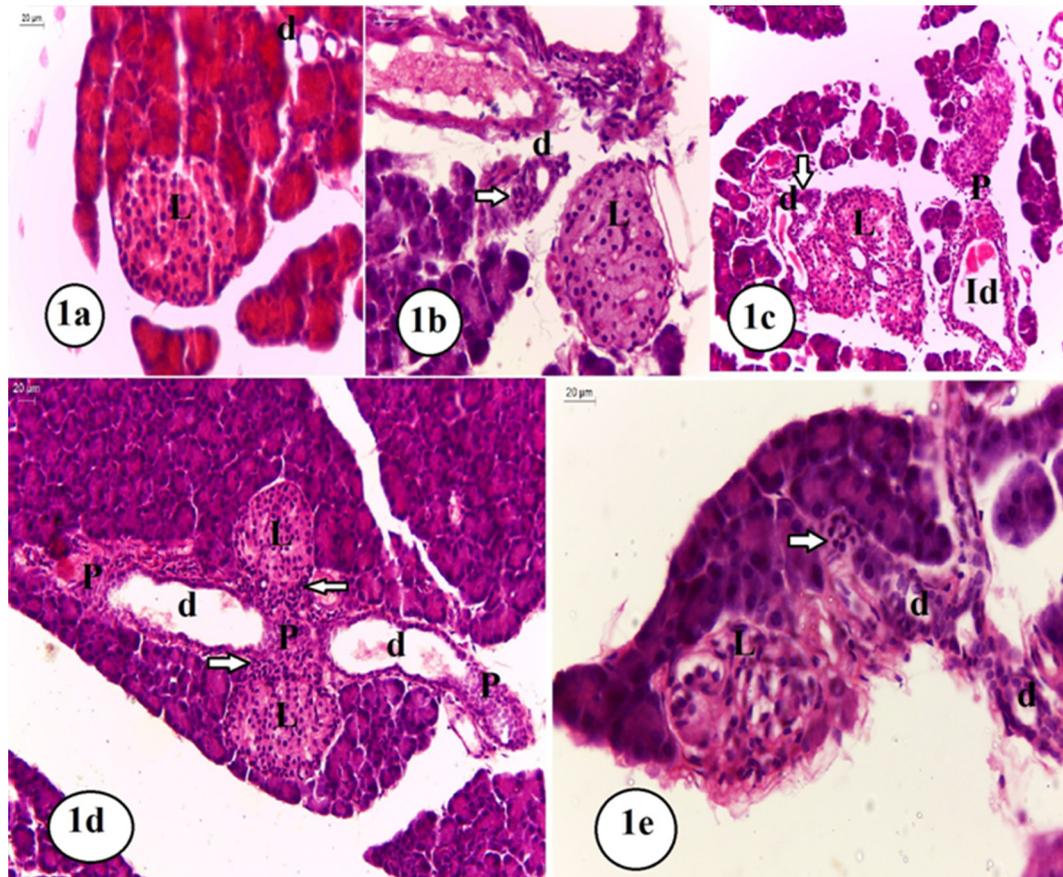
#### 4. Discussion

Menopause is an important period in women's life, characterized by loss of ovarian hormones and altered internal milieu.<sup>27</sup>

The present study revealed that ovariectomy following HFD/STZ diabetes neither affect body weight gain nor food intake. Consistent with previous studies<sup>28,29</sup> ovariectomy increased body weight gain percentage and food intake. This finding could be explained by increasing anorexigenic and decreasing orexigenic gene expression



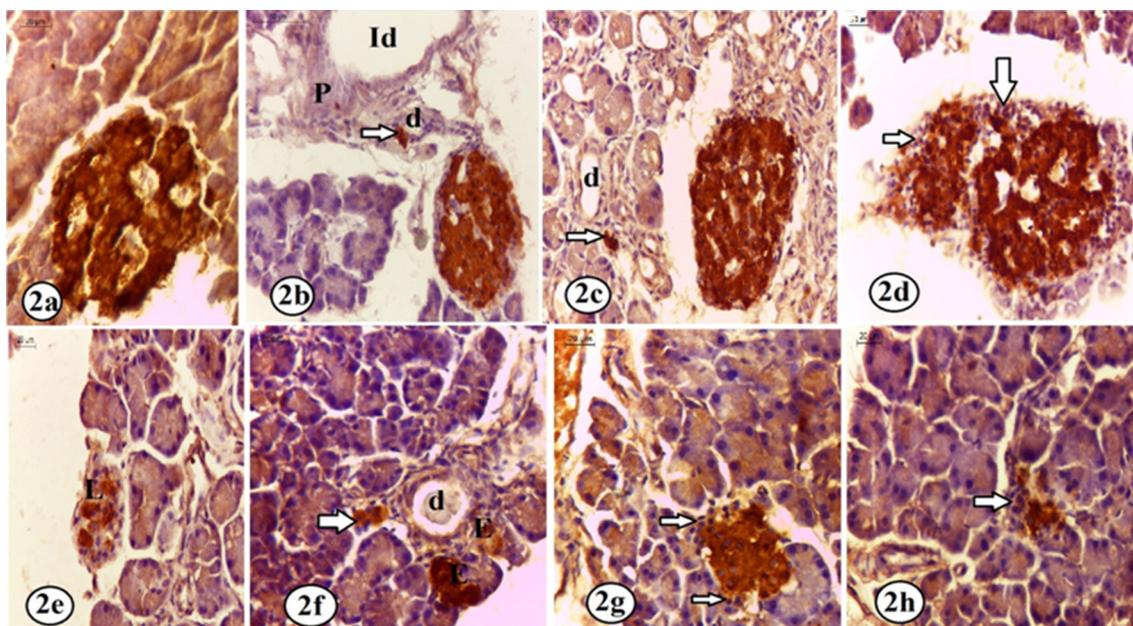
**Fig. 6.** Serum concentrations of malondialdehyde (A) and total thiol (B) in the studied groups. Data were presented as mean  $\pm$  standard deviation,  $n = 10$  rats per group. OVX: ovariectomized rats. Significant difference from the sham control group \*:  $P < 0.05$ , \*\*:  $P < 0.01$  and \*\*\*:  $P < 0.001$ , a: significantly different from the OVX group and b: significantly different from the sham diabetic group, using one-way analysis of variance (ANOVA) followed by Bonferroni test.



**Fig. 7.** Photomicrographs of paraffin sections of the pancreas in the studied groups. 1a: Pancreas of the sham control group showed regularly round islets (L) and interlobular ducts (d) (H&E, X400). 1b: Pancreas of the OVX control group showed a more or less rounded islet (L) and proliferating cells around a small rounded interlobular duct (d) (H&E, X400). 1c: Pancreas of the sham diabetic group showed a irregular shaped hyperplastic islet (L), proliferating cells (arrow) related to interlobular ducts (d), and an interlobular duct (H&E, X200). 1d & 1e: Pancreas of the OVX diabetic group, 1d: Showed islets (L) surrounded by a zone of dark cells (arrows) and interlobular ducts (d) that are related on both sides by masses of proliferating cells (P). (H&E, X200). 1e: Showed an islets (L) contained many cells with vacuolated cytoplasm and flat dark nucleus and small rounded interlobular ducts (d) surrounded by proliferating cells, some of them are small with dark nucleus (H&E, X400).

subsequent to abolish the protective effect of estrogen as proposed by Eckel.<sup>30</sup> Additionally, reduction of energy disbursement and physical activity after ovariectomy might be the underlying mech-

anism as suggested by Ludgero-Correia et al.<sup>31</sup> Moreover, increased body weight gain and food intake detected in HFD/STZ diabetic rats confirmed the finding of Liu et al.<sup>32</sup> This finding might be attribu-



**Fig. 8.** Photomicrographs of the immune stained pancreas in the studied groups. 2a: Pancreas of the sham control group showed a strong positive dark brown completely immune stained positive islet (anti-insulin antibody, X400). 2b: Pancreas of the OVX control group showed a golden brown immune positive islet with a thin rim of immune negative cells, an interlobular duct (d) surrounded by proliferating cells some of them are immune positive (arrow), and an interlobar duct (ld) with a related group of immune negative proliferating cells (P) (anti-insulin antibody, X400). 2c-2f: Pancreas of the sham diabetic group, 2c: Showed a strong positive completely immune stained islet and a group of immune positive proliferating cells (arrow) related to an interlobular duct (d) (anti-insulin antibody, X400). 2d: Showed a hyperplastic strong immune stained islet that showed some immune negative cells (arrows) (anti-insulin antibody, X400). 2e: Showed an islet that showed few immune positive cells (anti-insulin antibody, X400). 2f: Showed some immune positive proliferating cells (arrow) related to an interlobular duct (d), and other groups of immune positive cells within the acini of the exocrine pancreas (anti-insulin antibody, X400). 2g and 2h: Pancreas of the OVX diabetic group, 2g: Showed a golden brown immune stained islet with peripheral groups of immune positive cells (arrows) (anti-insulin antibody, X400). 2h: Showed proliferating cells related to the acini of the exocrine pancreas, some of them were immune positive while others were immune negative (arrow) (anti-insulin antibody, X400).

ted to utilization of an energy rich diet and fat accumulation in diverse body fat pads and decreased energy disbursement as explained by Srinivasan et al.<sup>33</sup> On the contrary, Wang et al.<sup>15</sup> showed no difference in body weight gain between HFD/STZ diabetic and control rats and they claimed this to balanced lipogenesis and lipolysis with augmentation of metabolic rate to a new value.

The present study revealed that when ovariectomy was following HFD/STZ diabetes had no impact on FBG concentration, but it worsened glucose metabolism disturbance (as indicated by impaired insulin tolerance test) and increased insulin resistance (as indicated by increased insulin concentration and elevated HOMA-IR index). This suggested that loss of ovarian hormones might be involved in insulin resistance independent of their effects on body weight. The present study confirmed the finding of previous studies demonstrating that ovariectomy disturbed glucose metabolism and impaired insulin tolerance.<sup>10,29</sup>

The result of the present study confirmed the finding of Ben-Shmuel et al.<sup>29</sup> who showed disturbance of lipid metabolism (as indicated by higher serum concentrations of TC, HDL-C and TG) in OVX diabetic rats more than either ovariectomy or HFD/STZ diabetes. It has been known that diabetes is considering as a disease of lipid metabolism disturbance more than carbohydrate.<sup>13</sup> The results of the present study were consistent with the findings of Saad et al.<sup>16</sup> and Liu et al.<sup>32</sup> that showed elevation of serum concentrations of TG, TC and free fatty acid in rat model of T2DM. This proposed to enhance lipolysis in insulin-resistant adipocytes.<sup>34</sup> Furthermore; our results were consistent with the studies whose reported impairment of lipid metabolism in OVX animals<sup>35</sup> and human.<sup>36</sup>

Total thiols, the most predominant non-protein thiol, are involved in many biological activities including neutralization of reactive oxygen species (ROS), detoxification of xenobiotics and

maintenance of -SH level in proteins.<sup>37</sup> Depletion of total thiols pool severely compromised antioxidant capacity. Thus, a compromised antioxidant defense system might be the key factor in contributing towards oxidative stress, reflected in higher levels of oxidatively damaged membrane lipids and proteins ultimately leading to tissue injury and dysfunction.<sup>38</sup>

The present study demonstrated that when OVX was following HFD/STZ diabetes, oxidative stress state exacerbated. Oxidative stress manifested by high MDA (an indicator of oxidative stress) and low TT (a major antioxidant component) concentrations. MDA and TT concentrations revealed the oxidant/antioxidant balance in the body.<sup>39</sup> This exacerbation of oxidative stress could be explained by loss of estrogens those were efficient antioxidants and diminished pro-oxidant enzymes expression concurrently with increased antioxidant enzymes expression as proposed by Sankar et al.<sup>10</sup> Oxidative stress has a critical role in the development of diabetic complications.<sup>40</sup> Matsuda and Shimomura<sup>41</sup> suggested that both oxidative stress and decreased antioxidant defense mechanisms had a role in increasing insulin resistance in diabetic rats.

The present study revealed an aggravation of inflammatory state when OVX was following HFD/STZ diabetes as indicated by high pro-inflammatory cytokines (IL1 $\beta$  and TNF- $\alpha$ ) and low anti-inflammatory cytokines (IL4 and IL10). This inflammatory state might have a role in insulin resistance. Minihane et al.<sup>7</sup> proposed that pro-inflammatory mediators released from adipose tissue impaired the insulin signaling and induced insulin resistance.

Histological findings of the pancreas revealed that pancreatic islets were clearly influenced by both ovariectomy and diabetes. Pancreas of OVX rats exhibited immune negative cells in some islets that fail to secrete insulin. This might be a result of loss of the direct hypertrophic effect of estrogen on  $\beta$  cells<sup>42</sup> or due to

cytotoxic effect of IL-1 $\beta$  on  $\beta$  cells.<sup>43</sup> In addition, OVX rats showed proliferating cells around both interlobular and interlobar ducts (nesidioblastosis) and some of them were immune positive. These changes reflected a compensating mechanism for  $\beta$  cell destruction or functional insufficiency.<sup>44</sup> HFD/STZ diabetic pancreas showed islets hyperplasia and proliferating cells around interlobular and interlobar ducts, indicating higher degree of  $\beta$  cells compensation. On the other hand, some islets showed signs of decompensation (as reflected by the presence of immune negative cells in some islets). Islet  $\beta$  cell failure in T2DM might be result from enhanced oxidative stress,<sup>45</sup> augmented endoplasmic reticulum stress, exacerbated glucolipotoxicity,<sup>46</sup> advanced glycation end product stress, inflammatory pathways activation and islet amyloid polypeptide accumulation.<sup>47</sup>

When ovariectomy was following HFD/STZ diabetes, the pancreatic changes became more apparent than the effect of any of them alone. The pancreatic architecture showed more decompensatory changes in  $\beta$  cells with evidence of degenerative changes (indicated by vacuolated cytoplasm and dark flat nuclei). Our findings were in agreement with those of Mojtaba et al.<sup>43</sup>

Thus, the present study theorized that estrogen might have a protective role against glucose and lipid disturbance induced by diabetes in rat as ovariectomy reduced this defense. Future studies using estrogen agonists will be needed to establish this protection.

In conclusion, ovariectomy-induced estrogen deficiency exacerbated glucose and lipid metabolic disturbance of diabetic female rats. This might be contributed to amplification of oxidative stress and inflammatory states. These findings suggested that estrogen deficiency in post-menopausal females could aggravate glucose and lipid dyshomeostasis. Consequently, for efficiency of the treatment, the menopausal status of diabetic female should be addressed.

## Conflict of interest

The authors declare that there is no duality of interest associated with this manuscript.

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