Short-term Effects of Date Palm Extract (Phoenix dactylifera) on Ischemia/Reperfusion Injury Induced by Testicular Torsion/Detorsion in Rats

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INTRODUCTION

Testicular torsion is a genitourinary emergency that requires an immediate surgical correction. This condition causes ischemic reperfusion which damages the testicular tissue and affects the quality of sperm.[1] It usually happens following the rotation of testis around the axis of the spermatic cord, and it is frequently observed in newborns, children, and adolescents.[2] Long-time torsion results in necrosis of the germinal cells. This may lead to subfertility or infertility.[3] Testicular tissue blood flow subsequent to the reperfusion exposes the tissue to the produced reactive oxygen species (ROS) and consequently an ischemia/reperfusion (I/R) injury.[4,5] The mechanism involved in testicular injury is not completely understood; however, oxidative stress is considered to be the main nocent in I/R injury.[6-9] Oxidative stress is the result of an imbalance between excessive ROS production and antioxidative mechanisms.[10,11]

It is demonstrated that ipsilateral testicular torsion could affect the contralateral testis too.[12-14] Under normal conditions, free radicals are produced and their effects are counterbalanced by the endogenous antioxidant system.[14] When ROS generation exceeds the defense mechanism's capacity to control, oxidative stress makes many reversible or irreversible cell injuries.[15] Antioxidants either synthetic or natural are potent scavengers of free radicals and have beneficial effects on human health and disease prevention.[16] Previous studies suggest that pretreatment with antioxidants can protect the testis against ROS insult.[16,17] Date palm (DP) is an excellent antioxidant agent due to the high concentrations of phenolic compounds, flavonoids (luteolin, methyl luteolin, quercetin, and epicatechin), and anthocyanins as well as the presence of selenoproteins, coumaric acid, and ferulic acid.[17] It is also scientifically proved to possess anti-inflammatory, immunomodulatory, vascular protective, and sex

ABSTRACT

Background: Antioxidants are potent scavengers of free radicals and have beneficial effects on human health. Objective: The aim of this study was to investigate the potential protective antioxidative activity of the edible portion of date fruit extract in an experimental testicular torsion/detorsion (T/D) model in rats. Materials and Methods: To investigate the potential protective effects of date palm (DP), 30 male Spraque-Dawley rats were divided into three groups: sham-operated, T/D, and T/D + DP-treated (500 mg/kg, PO) groups. Testicular ischemia was induced via keeping the left testis under 720° clockwise torsion for 2 h (h), afterward, detorsion was performed. All rats were sacrificed 4 h after detorsion. Serum malondialdehyde (MDA) concentration, total oxidative status (TOS), total antioxidant status (TAS), oxidative stress index (OSI), and histopathological damage score were evaluated. Results: Serum MDA, TOS, and OSI levels rose significantly in the T/D group. These values were lower in the T/D + DP group. TAS values decreased significantly in T/D group and rose in T/D + DP group. Severe injury was seen in the twisted testes of T/D group. In contrast, ipsilateral-twisted testicular tissue in the DP-treated group showed moderate-to-mild changes. Contralateral testicular tissue in the T/D group had a mild-to-moderate tissue injury; meanwhile, treated group revealed normal-to-mild changes. Spermatogenesis was significantly improved in DP-treated group when compared with the T/D group. Conclusion: The findings suggest a possible protective effect of DP against testicular oxidative damage induced by T/D; however, more detailed studies are warranted. Key words: Antioxidant markers, histopathology, ischemia/reperfusion, Phoenix dactylifera, tests

SUMMARY

Given the presence of several phenolic compounds possessing high antioxidative activity in DP, it could potentially be used to reduce testis ischemia/reperfusion-induced damage.

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hormone modulator effects. In vitro antioxidant activity of the aqueous extract of date fruit is demonstrated in many studies based on its phenolic compounds with potent free radical scavenging activity. The aim of this study was to investigate the potential protective antioxidant activity of the edible portion of date fruit extract in vivo, in an experimental testicular torsion/detorsion (T/D) model in rats.

MATERIALS AND METHODS

Preparation of extracts
Fresh date fruits (Phoenix dactylifera L., Palmae) were provided from a local market in Bam (Kerman Province, Iran), dried at room temperature, and were manually separated from the pits. The flesh of the dried P. dactylifera fruits was grinded and pulverized into powder. About 650 g of the powder was soaked in 2 L of cold distilled water. After 24 h, the solution was filtered and evaporated under vacuum and dried to a constant weight using a freeze-drier. The dry extract of the fruit was ground, and its weight was calculated in grams. About 650 g of the powder was soaked in 2 L of cold distilled water. After 24 h, the solution was filtered and evaporated under vacuum and dried to a constant weight using a freeze-drier. The dry extract of the fruit was ground, and its weight was calculated in grams.

Animals
This experiment was accomplished under the approval of the State Committee on Animal Ethics, Shiraz University, Shiraz, Iran. In addition, the recommendations of the European Council Directive (86/609/EC) of November 24, 1986, were used regarding the protection of animals used for experimental purposes. Thirty male Sprague-Dawley rats weighing 240–270 g were housed two per cage; maintained on a balanced diet and water ad libitum with a 12/12 h light-dark cycle. Animals were divided into three groups and pretreated orally for 10 days as follows: Group 1 (500 mg/kg DP extract), Group 2 (1 ml saline), and Group 3 ( sham-operated with no oral administration). The selected dose of DP was based on our previous pilot study. All groups received their treatment by oral force fed with a special gavage needle. After 10 days, rats were anesthetized using intraperitoneal administration of a combination of ketamine (80 mg/kg BW) and xylazine (5 mg/kg BW). Torsion, detorsion, and sham operation were performed through the intra-peritoneal route. Animals were anesthetized using intraperitoneal administration of a combination of ketamine (80 mg/kg BW) and xylazine (5 mg/kg BW). Torsion, detorsion, and sham operation were performed through the intra-peritoneal route. Animals were anesthetized using intraperitoneal administration of a combination of ketamine (80 mg/kg BW) and xylazine (5 mg/kg BW). Torsion, detorsion, and sham operation were performed through the intra-peritoneal route.

Biochemical analysis
Cardiac puncture was performed, and blood samples were collected in chilled nonheparinized tubes, kept in room temperature for 2 h, and centrifuged at 1500 × g for 15 min at 4°C. Separated sera evaluated for biochemical indicators included serum malondialdehyde (MDA), total antioxidant status (TAS), and total oxidant status (TOS).

Measurement of total antioxidant status
TAS level was determined using the method developed by Erel. Serum TAS levels were calculated in mmol Trolox equivalent/L.

Measurement of total oxidant status
TOS levels were determined using a novel automated and colorimetric measurement method as previously described by Erel. Serum TOS levels were calculated in μmol H2O2 equivalent/L.

Calculation of oxidative stress index
TOS: TAS ratio was used as the oxidative stress index (OSI). To perform the calculation, the unit of TAS, mmol Trolox equivalent/L, was converted to μmol Trolox equivalent/L, and OSI was calculated as follows: OSI = [(TOS, μmol H2O2 equivalent/L)/TAS, μmol Trolox equivalent/L] ×100.

Serum malondialdehyde activity assay
Lipid peroxidation in rat serum samples was determined as MDA concentration using the method described by Yagi. Tetramethoxypropane was used as a standard, and MDA levels were given as nmol/mL.

Histopathological analysis
Bilateral orchietomy was performed. The testes were immersed in 10% buffered formalin. Testes tissues were processed by routine histological methods and embedded in paraffin blocks that were placed in a rotary microtome. Sections of 5 μm thickness were obtained with disposable metal microtome blades. After deparaffinization and rehydration, all sections were stained with hematoxylin and eosin. Testicular injury was determined based on certain characteristics including hemorrhage/congestion, edema, degeneration, sloughing of germ cell layers, spermatogenesis arrest, seminiferous tubule disorganization, seminiferous tubule atrophy, presence of giant cells, and polymorphonuclear cells infiltration.

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Histopathology and testicular injury scoring
Severe injuries were seen in the twisted testis of T/D group [Figure 1b-f]. In contrast, ipsilateral twisted testicular tissue in DP-treated group showed mild-to-moderate changes [Figure 1g]. Contralateral testicular tissue had mild injury in T/D group, and also slight changes were seen in the treated group. Detailed histopathological results are shown in Table 1.

RESULTS

Histopathology and testicular injury scoring
Severe injuries were seen in the twisted testis of T/D group [Figure 1b-f]. In contrast, ipsilateral twisted testicular tissue in DP-treated group showed mild-to-moderate changes [Figure 1g]. Contralateral testicular tissue had mild injury in T/D group, and also slight changes were seen in the treated group. Detailed histopathological results are shown in Table 1.

The testes epithelial maturation score based on the Johnsen's score is shown in Table 2. The ipsilateral testes of T/D group displayed a lower Johnsen's score (P < 0.001) when compared with the sham operation group. The mean Johnsen's scores of treatment group in ipsilateral testes showed higher scores when compared with T/D group. In contralateral
Effects of Date Palm Extract on Testis I/R Injury

1.1±0.32
1.1±0.32
1.2±0.42

T/D + DP

[30]

1±0.00
1±0.00
9.1±0.57
3.7±0.48

Experimental

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1.7±0.48
1.3±0.48

G

c

Groups

Lesions

SA

STD

STA

G

c

Table 1: Means±standard deviation of histological scoring of the testicular tissue in sham and experimental groups

| Testes direction | Groups     | C            | H            | Different E | Lesions | SG |          |
|------------------|------------|--------------|--------------|-------------|---------|----|----------|
| Left             | Sham       | 1.1±0.32     | 1.1±0.32     | 1.3±0.48    | 1±0.00  | 1±0.00| 1±0.00   |
|                  | T/D        | 3.7±0.48     | 3.7±0.48     | 3.7±0.48    | 3.7±0.48 | 3.7±0.48| 3.5±0.53 |
|                  | T/D + DP   | 1.6±0.70     | 2.0±0.67     | 2.4±0.84    | 2.1±0.74 | 2±0.47| 2.2±0.42 |
| Right            | Sham       | 1.0±0.00     | 1±0.00       | 1±0.00      | 1±0.00  | 1±0.00| 1±0.00   |
|                  | T/D        | 1.4±0.52     | 1.3±0.48     | 1.7±0.48    | 1.2±0.42 | 1.1±0.32| 1.3±0.48 |
|                  | T/D + DP   | 1.2±0.42     | 1.1±0.32     | 1.4±0.52    | 1±0.00  | 1±0.00| 1±0.00   |

a,b,cAt each column, different superscript alphabets show a significant difference (P<0.05). C: Congestion; H: Hemorrhage; E: Edema; SG: Sloughed germinal cells; SA: Spermatogenesis arrest; STD: Seminiferous tubules disorganization; STA: Seminiferous tubules atrophy; G: Giant cells; T/D: Torsion/detorsion; DP: Date palm
testes, spermatogenesis in treated group significantly improved compared with T/D group.

Biochemical results

Biochemical results of the T/D, sham, and treated groups are shown in Table 2. TAS values decreased significantly in T/D group. Although it increased in T/D + DP-treated group compared to the T/D group, it was not significant. OSI and TOS values were significantly higher and lower in T/D and T/D + DP-treated groups, respectively (P≤0.05). In addition, MDA levels decreased significantly in DP pretreated group.

DISCUSSION

Oxidative stress is known as the most important guilt in testicular I/R injury,[7,15] and antioxidant agents are proved to be able to control it. Obviously, oxidative stress manifests with an increase in oxidants as well as a decrease in antioxidants. Sporadic assessment of antioxidant enzymes is time-consuming and expensive. Therefore, measuring all antioxidant levels is proposed to oxidant status evaluation. TOS and TAS as a combination of oxidant and antioxidant parameters, i.e. MDA, glutathione peroxidase, and catalase, are, therefore, valuable in terms of definition the body’s net stress. In the present study, TAS and TOS, the novel techniques developed by Erel in full determination of oxidant and antioxidant status, were used. The TOS to TAS ratio represents the OSI.[29] The OSI reflects an increased oxidant status or a decreased antioxidant status.[7]

In our study, TOS and OSI values were significantly higher in T/D group, but TAS values decreased. Lowered TAS values in the treated group suggest a protective effect of palm date against testicular T/D. Oxygen free radicals, formed during the reperfusion period, cause the oxidation of genetic materials, proteins, and membrane lipids.[30] MDA, another marker of oxidative stress, can impair a few physiological mechanisms in human body because of its ability to react with molecules such as deoxyribonucleic acid and proteins. These molecules can thus be considered to be something more than merely a lipid peroxidation product.[31] As a result, MDA level in T/D group elevated significantly compared to the sham group. In the DP-treated group, plasma MDA level reduced compared to the T/D group. Previous studies have emphasized that DP fruit is an excellent source of phenolics and therefore, possesses an extremely high antioxidant capacity.[32] Our findings support this, which would be translated to the alimentation of serum antioxidant power in treated groups.
Table 3: Profile of plasma antioxidant in sham and experimental groups

| Biochemical parameters | T/D | T/D + DP | Sham |
|------------------------|-----|---------|------|
| TAS (mmol Trolox equivalent/L) | 1.01±0.19 | 1.12±0.06 | 1.31±0.18 |
| TOS (μmol H₂O₂ equivalent/L) | 12.99±0.58 | 10.79±0.36 | 11±0.21 |
| OSI H₂O₂/Trolox | 1286.13 | 963.39 | 839.69 |
| Plasma MDA (nmol/mL) | 0.29±0.04 | 0.19±0.04 | 0.11±0.07 |

TAS: Total antioxidant status; TOS: Total oxidative status; OSI: Oxidative stress index; MDA: Malondialdehyde; T/D: Torsion/detorsion; DP: Date palm

Consistent with the serological findings, histological analysis in ipsilateral-twisted testicular tissues revealed mild-to-severe injury in pretreated rats and severe injury in T/D group. The difference between the severity of lesions at the microscopic level may seem rather arbitrary and not reproducible. However, the scoring system from a semi-quantitative perspective of histological feature showed a trend toward the protective effect by palm date [Table 1 and Figure 1].

Spermatogenesis and fertility can be significantly affected by testicular I/R injury. Based on the Johnsen tubular score, spermatogenesis significantly damaged in the I/R group. However, the Johnsen tubular score was significantly higher in DP-treated group than in the I/R group. This finding indicates that DP administration rescues spermatogenesis damage and may also prevent testicular I/R injury-induced infertility. The degree and duration of torsion have been shown to be important in testicular injuries development. In our study, after 2 h torsion and 4 h detorsion, the seminiferous tubule germinat epithelium was detached and spermatogenesis stopped. These findings were in accordance to ‘Turkmen et al.’s study[32] that reported following 2 h torsion and 4 h detorsion, germinat epithelial cells were present in the lumen of the seminiferous tubule in the T/D group. As already demonstrated, the testis injury prospers as the detorsion period extends after the torsion. Furthermore, in accordance with other assertions, contralateral testis was affected when ipsilateral testis was subjected to T/D. Decreasing testicular blood flow and subsequent tissue hypoxia are the reasons for contralateral testicular injury after T/D[33] however, the exact pathogenesis of contralateral abnormalities in patients with unilateral testicular torsion is not understood. Furthermore, it has been hypothesized that increased apoptosis in contralateral testes cells after ipsilateral testicular torsion is related to the breakdown of the blood-tissue barrier of the twisted testis, which may increase the risk of infertility[34] whereas the extent of apoptosis and necrotic changes within the twisted testis is directly correlated with the duration of torsion.

As mentioned above, the antioxidant activity of DP has been attributed to phenolic compounds, flavonoids, and anthocyanins as well as the presence of selenoproteins.[34] The effect of DP on brain I/R injury is proved too. The results have revealed that this extract has a substantial neuroprotective impact against cerebral ischemia and consequent reperfusion.[35] In another study, it was demonstrated that DP aqueous extract improves the plasma lipid profiles of the diet induced hypercholesterolemic rabbits through the augmentation of specific biosynthesis of antioxidant enzymes.[35] Consistent with these studies, our study showed that P. dactylifera is able to mitigate I/R injury induced by T/D in rat testis.

CONCLUSION

Inhibitors of oxidative stress provide a significant beneficial effect on testicular I/R injury. DP is a powerful radical scavenger, and its free radical scavenging ability should be considered to a greater extent.

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Conflficts of interest

There are no conflicts of interest.

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