AMELIORATIVE EFFECTS OF PUNICA GRANATUMON RENAL TUBULAR NECROSIS INDUCED BY NANDROLONE DECANOATE IN MICE MODEL

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ABSTRACT

Objective: To observe the effects of nandrolone decanoate on renal tubules and its protection by punicagranatum (pomegranate) in mice.

Study Design: Lab based experimental study.

Place and Duration of Study: One year in Anatomy department, Army Medical College, Rawalpindi, in association with National Institution of Health, Islamabad.

Methodology: Forty healthy BALB/c mice of both sexes with weight range of 2.5-30gms were equally disseminated into four groups, A as control while B, C and D as trial groups. Three trial groups were inoculated Nandrolone Decanoate 1mg per 100gm of body weight, through intramuscular injections in the hind limb, once weekly for eight weeks. Pomegranate nectar was administered in animals of trial group C (3ml per kg of body weight) en route oral gavage tube daily, whereas animals in trial group D was administered pomegranate peel extract (200mg per kg body weight) via oral gavage tube daily for eight weeks. Evaluation of the outcomes of trial groups B, C & D was done amongst them and with control group A.

Results: In nandrolone decanoate injected experimental group B, tubular necrosis were appreciated in comparison with control group A (p<0.001), and showed statistical improvement when evaluation was done with pomegranate nectar and pomegranate peel extract treated trial groups C (0.001) and D (p=0.001), correspondingly.

Conclusion: Punicagranatum in two forms, as nectar and peel extract, has almost identical curative effects on steroid administered renal tubular mutilation.

Keywords: Nandrolone decanoate, Necrosis, Punicagranatum, Renal tubule.

INTRODUCTION

Abuse of androgenic anabolic steroids (AAS’s) is becoming a civic health problem. AAS’s are the analogues of male sex hormone, testosterone. Nandrolone decanoate (ND), is the utmost frequently molested AAS formula in the world. Nandrolone is commonly used to handle many pathologies, including osteoporosis, HIV-associated muscle wasting, renal inadequacy, male hypogonadism and tardy puberty. However, in spite of such beneficial healing abilities, prolonged and unfettered usage of nandrolone results in adverse consequences, comprising hepatic toxicity, fluctuation of thyroid function, nephrotoxicity and cardiovascular toxicities. Even though, the AAS’s are prohibited stuff by certified sport societies and considered controlled medicines by the drug regulatory authority, they are misused widely, particularly by young athletes and teens or non-sports person as for beautifying or entertaining drives and this misuse bring about quite a lot of adversative effects.

Oxidative pressure is the primary issue that promotes organ impairment, chiefly superoxide dismutase and catalase. The humoral system can stimulate the oxidative stress that reins blood pressure, in particular through renin angiotensin coordination, which is the onemain structure compromised by means of prolonged inoculation of nandrolone decanoate, leading to damaged kidney architecture and extended collagen deposition in this tissue. Among the world’s earliest fruits, the pomegranate has had a long and interesting history. It was called the “Chinese apple,” the alternate name due to its nutritive, curative, ornamental and industrial value. The health aiding properties of pomegranate are mainly attributed to the plenty of carotenoids, vitamins, ellagitannins and other polyphenolic compounds.

Antioxidants enriched pomegranate enhanced the reduction of free radicals inside the cells, so have the ability to protect the tissues from oxidative stress damage. Thus, the present study was designed to conclude the effect of nandrolone decanoate administration on mice kidneys and protection of these effects by two forms of pomegranate.

METHODOLOGY

This lab based experimental study was done at
the department of Anatomy, Army Medical College (AMC), collaborated with National Institute of Health (NIH) Islamabad and Pathology Unit, Army Medical College, Rawalpindi. All procedural codes of behavior were agreed by ethics committee of centre for research in experimental and applied medicine (02/CREAM-A-Humaira Ali), AMC, Rawalpindi. Forty fit and well animals, including both sexes of BALB/c mice with weightrange of 25-30 grams were randomly distributed into four equal groups, each group having five male and five female. Twenty male and twenty female mice were kept in separate cages, count was made from 1-4. Animal 1 was put in group A, 2 in group B, 3 in group C and 4 in group D. This count was repeated five times for male and five times for female mice and then cages were labelled accordingly. They were provided standard laboratory diet pellets and water ad libitum for eight weeks. Group A functioned as control group and groups B, C and D were allocated as trial groups. Animals in three trial groups B, C and D were injected a shot of ND at the dose of 1mg per 100gm body weight, I/Min the hind limb once in a week for a period of two months. In addition, mice in trial group C, were administered pomegranate nectar (PN) in quantity of 3ml per kg body weight enroute oral gavage tube, every day for 2 months, and mice in trial group D were given pomegranate peel extract (PPE), in quantity of 200mg per kg body weight by oral gavage tube every day for two months.

At completion of experimental period, the mice were sacrificed. After dissection, the coronal stained sections of kidneys were studied for assessment of necrosis in kidney tubules. Necrotic changes were categorized according to the scale used by Hadjipour et al. Level 0 (none): absence of necrosis in randomly selected three views of observed slide by using light microscope at 40X. Level I (light necrosis): approximately one-two tubules displaying necrosis in three observation fields of slide by using light microscope at 40X. Level II (medium necrosis): approximately three-five tubules exhibiting necrosis in three fields of slide by using light microscope at 40X. Level III (severe necrosis): approximately six-ten tubules showing necrosis in three observational fields of slide by using light microscope at 40X. Level IV (extra severe necrosis): Ten plus tubules displaying necrosis in three observational fields of slide by using microscope at 40X. One slide of each specimen was observed.

SPSS-21 was used for data analysis. Qualitative variables were presented by frequency and percent-ages. Chi square test was applied for comparison of qualitative variables. A p-value < 0.05 was considered to be indicative of statistical significant.

RESULTS

This lab based experimental study was conducted to evaluate the “Ameliorative effect of Punica Granatum on the steroid induced histomorphological changes in the mice kidney”. For this purpose, 40 BALB/c, healthy male and female, mice were equally divided into four groups, five male and five female in each group with 25 ± 10gm of weight. Severity of necrotic changes was graded according to the scale used by Hadjipour et al., according to which changes were graded from Level 0 to Level IV. One slide of each specimen was observed.

Microscopic interpretations were elicited by analyzing the slides using light microscope at power of 40X and changes were observed in proximal as well as distal convoluted tubules of kidney. In this study, group B with H&E stained sections demonstrated individual cell damage as well as lesions of whole tubules was seen at high magnification. In 4 (40%) animals, the tubules exhibited inflamed and enlarged cells with disintegration of the nuclei. Blebbing were seen and apical domain of cells displayed blunting due to reduced microvilli in early cases of cell damage (figure), while 1 (10%) animal displayed advanced phases in which the whole cells were shrunken and shed off along with the constricting lumens of the tubules. In another 5 (50%) animals, severely necrosed tubules appeared dilated in conjunction with epithelial desquamation as well as fragmented tubular epithelial cells in lumens appeared as eosinophilic aggregates lacking identifiable cellular details and peculiarities in 50% of animals.

In all 10 (100%) specimens of control group A, no necrosis was seen in three views of any slide. In trial group B, 4 (40%) of animals had medium necrosis, 5 (50%) had severe necrosis while remaining 1 (10%) revealed extra severe necrosis. Comparison with other groups demonstrated that it was statistically significant with control group A (p<0.001), trial group C (p= 0.001) and trial group D (p<0.001) (table, figure).

Necrotic changes in group C were documented at level 0 in 5 (50%) of animal models while 5 (50%) showed light necrosis. On evaluating the outcomes of group C with group A (p-value=0.033) and B (p<0.001), the difference was statistically significant. Whereas in experimental group D, necrosis was light in 5 (50%) of the cases and rest of the 5 (50%) had no necrosis. Intergroup comparison was noticed to be statistically signi-
significant with control group A (p-value=0.03), experimental group B (p<0.001). No statistical impact was seen between groups C and D (p-value=1.000) (table, figure). Necrotic changes were categorized by tubular epithelial cells exhibiting no nuclei, intense eosinophilic homogeneous cytoplasm, but preserved shape. Necrotic cells fall into the tubular lumen, obstructing it, and determining acute kidney failure. The cells undergoing necrosis showed decreased or no basophilia and increased eosinophilia on H&E staining since cytoplasmic proteins and RNA are denatured. Basement membrane is intact, so that tubular epithelial regeneration is possible.

Pro-inflammatory cytokines are released from the infiltrate accumulated around degenerating tubules. Major source of ROS-driven oxidative kidney injury initiate from cellular reaction to pro-inflammatory cytokines. Salem et al, demonstrated cortical thickness, growth in kidney size and disturbed renal profile in AAS treated group whilst compared with the groups without AAS inoculation. This statistic remained validated by Kaufman et al, in a criticism that the use of AAS leads to renal function impairment and unexpected dying in humans. Above mentioned adverse effects were attributed to AAS-driven molecular modifications, which includes adaptations in the countenance of cardiac ion alternate proteins and heart as well hepatic and kidney enzymes after acute and chronic treatment.

Improvement was noticed in trial groups C and D in current study, but statistical significance was still present when compared with control group A. This improvement was credited to high levels of antioxidants in pomegranate which can effectually decrease not only lipids from peroxidation but also proteins from oxidation by minimizing the free radicals and preventing the proliferation. It can also inhibit DNA oxidation, particularly mitochondrial DNA.

Results of present study were comparable with the results of Pan et al who reported that ND exposure.
could also lead to DNA damage through increase in ROS generation. Trans-membrane proteins, NOX enzymes are present on nuclear membrane near DNA, increasing the risk of its damage\(^\text{a}\). The relationship among oxidative stress and DNA damage has also been proved by Gorgoulis et al. Oxidative stress is responsible for many of the chronic ailments and it imitates an imbalance between the systemic index of ROS and a genetic system's aptitude to purify the reactive intermediates or to mend the consequential damage\(^\text{a}\). High levels of phytochemicals such as polyphenol, flavonoid and tannin contents in pomegranate could enhance the reduction of free radicals inside the cells, to protect the kidney tissue from oxidative injury.

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CONCLUSION

Punicagranatum in two forms, as nector and peels extract, has almost the same preemptive effects on steroid induced changes in kidney tubules.

CONFLICT OF INTEREST

This study has no conflict of interest to be declared by any author.

REFERENCES