Letter to Editor

Sex-related Difference in Protective Role of Aerobic Exercise against Cisplatin-induced Hepatotoxicity

Farzaneh Zeynali¹, Jalaledin Noroozi¹, Zahra Pezeshki¹, Mehdi Nematbakhsh^{1,2,3}

¹Water and Electrolytes Research Center, Isfahan University of Medical Sciences, Isfahan, Iran, ²Department of Physiology, Isfahan University of Medical Sciences, Isfahan, Iran, ³Isfahan^{MN} Institute of Basic and Applied Sciences Research, Isfahan, Iran

Correspondence to:

Prof. Mehdi Nematbakhsh, Department of Physiology, Water and Electrolytes Research Center, Isfahan University of Medical Sciences, Isfahan, Iran. E-mail: nematbakhsh@med.mui.ac.ir

How to cite this article: Zeynali F, Noroozi J, Pezeshki Z, Nematbakhsh M. Sex-related difference in protective role of aerobic exercise against cisplatin-induced hepatotoxicity. Int J Prev Med 2016;7:84.

DEAR EDITOR,

Cisplatin (CP) as a potential chemotherapeutic drug is accompanied with nephrotoxicity^[1] and hepatotoxicity.^[2] Aerobic exercise could attenuate CP-induced nephrotoxicity gender dependently.^[3,4] The experimental design was described before^[3,4] briefly, sixty Wistar rats were divided into eight groups. The male animals in Group I (named EX + CP + EX) had aerobic exercise on a treadmill 1 h/day and five days/week for 8 weeks. Then, the exercise protocol was continued for another week which was accompanied with reducing the intensity of training, and during this week, the animals also received CP (2.5 mg/kg/day). Groups II (named EX + CP) had the same protocol as Group I without exercise in the last week during CP therapy. Groups III (named CP) and IV (named sham) received CP and saline respectively during the last week of the study without exercise. The female rats in Groups V-VIII had the same protocol as the male rats in Groups' I-IV. The animals were exposed to moderate exercise with the 65% oxygen consumption.^[5] Blood samples were obtained 1 week after CP administration.

CP increased the serum levels of aspartate aminotransferase (AST) in male (P < 0.05) not in female rats. The serum levels of AST decreased significantly in EX + CP + EX group compared to CP group (P < 0.05). The serum levels of alkaline phosphatase (ALP) decreased in all CP treated male and female groups compared to sham group (P < 0.05). The serum levels of alanine aminotransferase decreased in all male groups and EX + CP + EX and CP in female groups when compared with sham group [P < 0.05, Figure 1].

AST increased in male positive control (Group III) compared to sham group as reported before.^[6] CP increases toxicity via reduces antioxidant enzymes



Figure 1: The serum levels of alkaline phosphatase, alanine aminotransferase, and aspartate aminotransferase. *Indicates significant difference from negative control group (sham) and #indicates significant difference from positive control group (cisplatin) (P < 0.05) one-way ANOVA followed by the least-squares deconvolution posttest

International Journal of Preventive Medicine 2016, 7:84

and increases malondialdehyde level,^[6,7] and appropriate exercise increases antioxidant enzymes.^[8,9] Administration of CP induces magnesium deficiency^[10] whereas ALP activity was reduced due to magnesium depletion.^[11] Therefore, aerobic exercise may reduce CP-induced hepatotoxicity by increasing activation of antioxidant system in male rats.

Financial support and sponsorship Nil.

Conflicts of interest

There are no conflicts of interest.

Received: 17 Apr 16 Accepted: 20 May 16 Published: 20 Jun 16

REFERENCES

- Nematbakhsh M, Ebrahimian S, Tooyserkani M, Eshraghi-Jazi F, Talebi A, Ashrafi F. Gender difference in cisplatin-induced nephrotoxicity in a rat model: Greater intensity of damage in male than female. Nephrourol Mon 2013;5:818-21.
- Pratibha R, Sameer R, Rataboli PV, Bhiwgade DA, Dhume CY. Enzymatic studies of cisplatin induced oxidative stress in hepatic tissue of rats. Eur J Pharmacol 2006;532:290-3.
- Noroozi J, Zeynali F, Nematbakhsh M, Pezeshki Z, Talebi A. Nonpreventive role of aerobic exercise against cisplatin-induced nephrotoxicity in female rats. Int J Prev Med 2015;6:5
- Zeynali F, Nematbakhsh M, Mojtahedi H, Poorshahnazari A, Talebi A, Pezeshki Z, et al. Protective role of aerobic exercise against cisplatin-induced nephrotoxicity in rats. Asian J Sports Med 2015;6:e24901.
- 5. Powers SK, Criswell D, Lawler J, Martin D, Lieu FK, Ji LL, et al. Rigorous exercise

training increases superoxide dismutase activity in ventricular myocardium. Am J Physiol 1993;265(6 Pt 2):H2094-8.

- Mansour HH, Hafez HF, Fahmy NM. Silymarin modulates cisplatininduced oxidative stress and hepatotoxicity in rats. J Biochem Mol Biol 2006;39:656-61.
- Koc A, Duru M, Ciralik H, Akcan R, Sogut S. Protective agent, erdosteine, against cisplatin-induced hepatic oxidant injury in rats. Mol Cell Biochem 2005;278:79-84.
- Shin YA, Lee JH, Song W, Jun TW. Exercise training improves the antioxidant enzyme activity with no changes of telomere length. Mech Ageing Dev 2008;129:254-60.
- Moien-Afshari F, Ghosh S, Elmi S, Rahman MM, Sallam N, Khazaei M, et al. Exercise restores endothelial function independently of weight loss or hyperglycaemic status in db/db mice. Diabetologia 2008;51:1327-37.
- 10. Lajer H, Daugaard G. Cisplatin and hypomagnesemia. Cancer Treat Rev 1999;25:47-58.
- Pimstone B, Eisenberg E, Stallone W. Decrease in serum alkaline phosphatase activity produced by magnesium depletion in rats. Proc Soc Exp Biol Med 1966;123:201-3.

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

Access this article online	
Quick Response Code:	
	Website: www.ijpvmjournal.net/www.ijpm.ir
	DOI: 10.4103/2008-7802.184312