

## LETTER to the EDITOR

# New Insight of OCT2 Regulation as Mediator for Cisplatin- Induced Nephrotoxicity

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### Dear Editor

We read with interest the recently published review article in *Asian Pac J Cancer Prev*, by Nematbakhsh et al., entitled 'Cisplatin-Induced Nephrotoxicity; Protective Supplements and Gender Differences' (Nematbakhsh et al., 2017). The authors concluded several suggestions that may provide to reduce Cisplatin (CDDP) induced nephrotoxicity (Nematbakhsh et al., 2017). Here, we would like to add one more substantial suggestion to improve the current efforts to overcome the main dosing limiting nephrotoxic effect of CDDP. Cisplatin nephrotoxicity is sex related greater intensity of damage in male than female and these differences may be related to CDDP uptake by OCT2 due to the markedly higher renal expression of Organic cation transporter 2 (OCT2) in male than female rats. The main determinant for sex differences in OCT2 gene expression is testosterone (El-Arabey, 2015a, 2015b). Furthermore, Study demonstrated that OCT2 level was significantly reduced in mice after castration (Meetam et al., 2009). Hence, regulation of OCT2 expression act as critical mediator for CDDP induced nephrotoxicity. There are some efforts introduced toward study of protective supplement on the regulation of renal organic cation transporters (Ulu et al., 2012; El-Arabey and Salama, 2015; El-Arabey, 2016). Recently, study demonstrated that OCT2 gene variation in the South African bantu-speaking population and functional promoter variants in different populations for drug safety, response and global pharmacogenomics (Wilson et al., 2017). Therefore, we should take in our consideration when explore novel compounds against CDDP induced nephrotoxicity to examine their effects on the regulation of renal OCT2. In addition, research is needed to conduct randomized clinical trials to examine the relevance of sex differences and contribution of the renal expression of OCT2 transporter to the situation of CDDP induced nephrotoxicity in humans.

### References

- El-Arabey AA (2015a). Gender difference in Cisplatin-induced nephrotoxicity in a rat model. *Nephrourol Mon*, **7**, e23595.
- El-Arabey AA (2015b). Sex and age differences related to renal OCT2 gene expression in cisplatin-induced nephrotoxicity. *Iran J Kidney*, **9**, 335-6.
- El-Arabey AA, Salama AS (2015). Are Mice and Rats good experimental models to explore novel compounds against Cisplatin Induced Nephrotoxicity?. *Int J Nephrol Kidney, Failure*, **1**.
- El-Arabey AA (2016). Dual function of OCT2 and MATE1 in cisplatin induced nephrotoxicity. *Pharmacol Res*, **10**, 89-95.
- Kim HJ, Park DJ, Kim JH, et al (2015). Glutamine protects against cisplatin-induced nephrotoxicity by decreasing cisplatin accumulation. *J Pharmacol Sci*, **127**, 117-26.
- Meetam P, Srimaroeng C, Soodvilai S, Chatsudthipong V (2009). Regulatory role of testosterone in organic cation transport: in vivo and in vitro studies. *Biol Pharm Bull*, **32**, 982-7.
- Nematbakhsh M, Pezeshki Z, Eshraghi JF, et al (2017): Cisplatin-induced Nephrotoxicity; protective supplements and gender differences. *Asian Pac J Cancer Prev*, **18**, 295-314.
- Ulu R, Dogukan A, Tuzcu M, et al (2012). Regulation of renal organic anion and cation transporters by thymoquinone in cisplatin induced kidney injury. *Food Chem Toxicol*, **50**, 1675-9.
- Wilson NC, Choudhury A, Carstens N, Mavri-Damelin D (2017): Organic Cation Transporter 2 (OCT2/SLC22A2) gene variation in the south African Bantu-speaking population and functional promoter variants. *OMICS*, **21**, 169-76.

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