#122 - BENEFICIAL EFFECTS OF QUERCETIN ON RAT URINARY BLADDER AFTER CISPLATINUM TREATMENT

Tinay I1, Sener TE1, Ercan F2, Koroglu MK2, Tarcan T1, Sener G3

- 1. Marmara University School of Medicine Department of Urology,
- 2. Marmara University School of Medicine Department of Histology,
- 3. Marmara University School of Pharmacy Department of Pharmacology

HYPOTHESIS / AIMS OF STUDY

Cisplatinum treatment leads to cytotoxic events and generates oxidative stress, which has deleterious effects on the function of several organ systems, including the urinary bladder. The present study was designed to investigate the putative beneficial effect of quercetin (QT) against cisplatinum-induced bladder damage.

STUDY DESIGN, MATERIALS AND METHODS

Twenty-eight adult male Sprague Dawley rats were included in the study. Six-teen rats were administered intraperitoneally (i.p.) single dose cisplatin 7 mg/kg and divided in to 2 groups:

The first group (n=8) received saline i.p., whereas

the second group (n=8) fed orally with 20 mg/kg QT, respectively,

The remaining 12 rats served as the control group after i.p. saline, where 6 treated with QT.

After decapitation, bladder strips were placed in organ bath and isometric contractions to carbachol (10-8 to 10-4 M) were recorded. In order to examine oxidative tissue injury, malondialdehyde (MDA), 8-hydroxydeoxyguanosin (8-OHdG) and glutathione (GSH) levels and superoxide dismutase (SOD) and caspase-3 activities and caspase-3 protein expression in bladder tissues were measured along with histological evaluations.

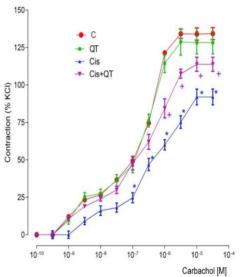


Figure 1. Concentration-response curves

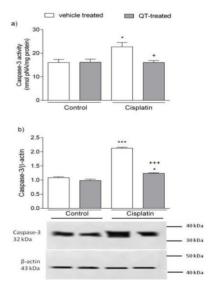


Figure 3. (A) Caspase 3 activity and (B) caspase-3 protein expression by Western blotting

RESULTS

In the cisplatinum-treated bladders, the contractile responses were lower than those of the control group and were reversed by treatment with QT (figure 1). On the other hand, increase in MDA and 8-OHdG levels (figure 2), and caspase-3 protein expression and caspase-3 activities of tissues in the cisplatinum group were significantly reversed by QT treatment (figure 3). Furthermore, treatment with QT also prevented the depletion of tissue GSH. levels and SOD activity seen in the cisplatinum group (figure 4).

INTERPRETATION OF RESULTS

According to the results, cisplatinum treatment leads to cytotoxic damage in urinary bladder, which has been reversed with QT treatment through its anti-inflammatory and antioxidant effects.

CONCLUDING MESSAGE

Quercetin exerts beneficial effects against cisplatinum-induced damage on bladder through its anti-inflammatory and antioxidant effects.

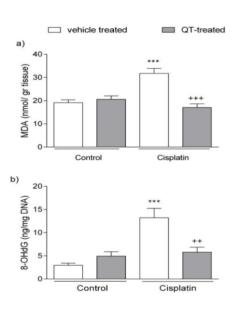


Figure 2. a) Malondialdehyde (MDA) and b) 8-hydroxydeoxyguanosin (8-OHdG) levels

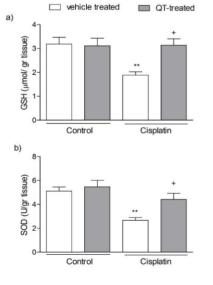


Figure 4. a) Glutathione (GSH) levels and b) superoxide dismutase (SOD) activity