

Globaqua-Cytothreat-Endetech-Scarce Workshop

PHARMACEUTICALS IN WASTEWATERS AND SURFACE WATERS UNDER MULTISTRESSORS SITUATION: Fate, Adverse effects, Risks and Removal Technologies

2-3 December 2014, Barcelona, Spain

Scientific Committee

- Damià Barceló, IDAEA-CSIC, Barcelona and ICRA, Girona, Spain
- Metka Filipič, NIB, Ljubljana, Slovenia
- Pierre-Alain Bandinelli, Da Volterra, Paris, France
- Sara Rodríguez-Mozaz, ICRA, Girona, Spain
- Miren López de Alda, IDAEA-CSIC, Barcelona

ORGANIZERS





SUPPORTING ORGANIZATIONS







Impact of in vivo and in vitro exposure to 5-fluorouracil, cisplatin, etoposide and vincristine sulphate on dna damage in haemocytes of freshwater mussels Unio pictorum AND Unio tumidus

Stoimir Kolarević¹, Zoran Gačić², Margareta Kračun-Kolarević³, Jovana Kostić^{1,2}, Karolina Sunjog^{1,2}, Momir Paunović³, Jelena Knežević-Vukčević¹ and <u>Branka Vuković-Gačić</u>¹

¹University of Belgrade, Faculty of Biology, Chair of Microbiology, Center for Genotoxicology and Ecogenotoxicology, Belgrade, Serbia,

²University of Belgrade, Institute for Multidisciplinary Research, Belgrade, Serbia

³University of Belgrade, Institute for Biological Research "SinišaStanković, Belgrade, Serbia

The impact of 5-Fluorouracil (5-FU), Cisplatin (CP), Etoposide (Eto) and Vincristine sulphate (Vin) on DNA damage level was studied *in vivo* and *in vitro* on haemocytes of freshwater mussels *Unio pictorum* and *U. tumidus* using alkaline comet assay. For *in vitro* experiments, two different approaches were applied: i) isolated haemocytes were treated for 30 min in physiological solution, ii) primary culture of haemocytes was treated for 22h. For *in vivo* experiments, mussels were exposed in static system for 72h. CdCl₂ was used as positive control. The level of DNA damage was analyzed by Comet Assay IV software. CdCl₂ induced significant increase in DNA damage *in vitro* and *in vivo*.

Treatments in vivo with 5-FU, Eto and Vin resulted in significant increase of DNA damage. Genotoxic potential was also detected in vitro for Vin after 30 min of exposure, and Eto after 22h. CP did not induced increase of DNA damage in vivo or in vitro, but post treatment with hydrogen peroxide indicated existence of DNA crosslinks.

Lack of genotoxic effects of cytostatics in isolated haemocytes can be attributed to short period of exposure (30 min), while the lack of the effects in primary cultures can be assigned to the mechanisms of action of these drugs.

This research has received funding from the European Community's FP7/2007-2013 agreement n° 265264