

## Oral communications: Drug Design Theme (Wednesday).

### Ruthenium Complexes for the Treatment of Protozoan Diseases

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Recently, some of us have shown that dinuclear thiolato-bridged arene ruthenium complexes of the type  $[(\eta^6\text{-}p\text{-cymene})_2\text{Ru}_2(\mu_2\text{-SR})_3]^+$  were among the most cytotoxic ruthenium complexes reported so far, with nanomolar  $\text{IC}_{50}$  values against both A2780 human ovarian cancer cells and their cisplatin-resistant mutant variant A2780cisR.[1] Interestingly, *in vivo* studies of *diruthenium-1*,  $[(\eta^6\text{-}p\text{-cymene})_2\text{Ru}_2(\mu_2\text{-SC}_6\text{H}_4\text{-}p\text{-Bu}^t)_3]^+$ , demonstrated a significant increase in survival of the treated mouse group.[2-3]

Motivated by the impressive results obtained in cancer therapy, we decided to test some of those compounds against *Neospora caninum* and *Toxoplasma gondii*. Two complexes, namely  $[(\eta^6\text{-}p\text{-cymene})_2\text{Ru}_2(\mu_2\text{-SC}_6\text{H}_4\text{-}p\text{-CH}_3)_3]^+$  and *diruthenium-1*, displayed  $\text{IC}_{50}$  values on *T. gondii* of 34 and 62 nM, respectively, with a high selectivity on *T. gondii* over human foreskin fibroblasts (HFF) as expressed by a selectivity index of around 20'000. Transmission electron microscopy (TEM) showed that treatment with these compounds has a dramatic impact on the parasite mitochondrion [4]. Moreover, upon treatment of human ovarian carcinoma A2780 cells with these drugs, between 71 and 97% of the ruthenium was found to accumulate in the mitochondria. This strongly suggests that these compounds are targeting the mitochondria in both parasites and cancer cells. *Giardia lamblia* that is devoid of functional mitochondria is not affected by treatments with these complexes up to a concentration of 5  $\mu\text{M}$ , thus further supporting this statement.

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