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Highly cytotoxic trithiolato-bridged dinuclear Rh(III) and Ir(III) complexesGajendra Gupta^a, Benjamin S. Murray^b, Paul J. Dyson^b, Bruno Therrien^{a,*}^a Institute of Chemistry, University of Neuchâtel, Avenue de Bellevaux 51, CH-2000 Neuchâtel, Switzerland^b Institut des Sciences et Ingénierie Chimiques, Ecole Polytechnique Fédérale de Lausanne (EPFL), CH-1015 Lausanne, Switzerland**ABSTRACT**

Water soluble trithiolato-bridged cationic complexes of the type $[(\eta^5\text{-C}_5\text{Me}_5)_2\text{M}_2(\mu\text{-SC}_6\text{H}_4\text{-}p\text{-X})_3]^+$ (M = Rh, X = H, **1**; CH₃, **3**; OCH₃, **5**; Pr^{*i*}, **7**; Bu^{*t*}, **9**; M = Ir, X = H, **2**; CH₃, **4**; OCH₃, **6**; Pr^{*i*}, **8**; Bu^{*t*}, **10**) were synthesized and isolated as their chloride salts by reacting pentamethylcyclopentadienyl rhodium and iridium dimers $[(\eta^5\text{-C}_5\text{Me}_5)_2\text{M}_2(\mu\text{-Cl})_2\text{Cl}_2]$ in ethanol with the corresponding thiophenol. All complexes were isolated in good yields and were fully characterized including single-crystal X-ray structure analysis on representative complexes. The complexes were found to have IC₅₀ values in the nanomolar range in human ovarian A2780 cancer cells, but did not display selectivity with respect to noncancerous human HEK293 embryonic kidney cells.

Keywords: Half-sandwich complexes; Dinuclear complexes; Thiolato-bridging ligands; Lipophilicity; Anticancer activity.

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