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(Hlorofenil)terpiridin Ru(II) kompleksi: Sinteza, kinetika supsticionih reakcija sa biomolekulima i interakcije sa DNA i BSA

Ana Rilak Simović, Ioannis Bratsos,* Nicola Dimitri**

*Prirodno-matematički fakultet, Univerzitet u Kragujevcu, Radoja Domanovića 12, 34000, Kragujevac, *I.N.N., Dept. of Physical Chemistry, NCSR "Demokritos", 15310 Ag. Paraskevi, Athens, Greece, **Elettra – Sincrotrone Trieste, S.S. 14 Km163.5 in Area Science Park, 34149 Basovizza, Trieste, Italy*

Sa ciljem da ispitamo uticaj aromatičnosti helatnih liganada na reaktivnost Ru(II) polipiridil kompleksa, sintetisali smo dva nova Ru(II) kompleksa [Ru(Cl-Ph-tpy)(phen)Cl]Cl (**1**) i [Ru(Cl-Ph-tpy)(*o*-bqdi)Cl]Cl (**2**) (gde je phen = 1,10-fenantrolin i *o*-bqdi = *o*-benzohinondiimin).¹ Kompleksi su okarakterisani pomoću elementalne analize i različitih spektroskopskih metoda, kao što su IR, UV-Vis i (1D i 2D) NMR, dok je njihova molekulska struktura u čvrstom stanju utvrđena pomoću rendgenske strukturne analize. Kinetika i mehanizam reakcije kompleksa **1** i **2** sa biološki relevantnim ligandom 5'-GMP ispitivani su pomoću UV-Vis spektroskopije. Proučavane su interakcije kompleksa sa CT DNA, kao i kompetitivne reakcije interkalirajućeg agensa etidijum bromida (EB). Pored toga, ispitivan je afinitet kompleksa prema govedem serum albuminu (BSA), i određene su konstante vezivanja.

(Chlorophenyl)terpyridine Ru(II) complexes: Synthesis, kinetics of the substitution reactions with small biomolecules and DNA/BSA binding studies

Ana Rilak Simović, Ioannis Bratsos,* Nicola Dimitri**

*University of Kragujevac, Faculty of Science, Department of Chemistry, R. Domanovića 12, Kragujevac, Serbia, *I.N.N., Dept. of Physical Chemistry, NCSR "Demokritos", 15310 Ag. Paraskevi, Athens, Greece, **Elettra – Sincrotrone Trieste, S.S. 14 Km163.5 in Area Science Park, 34149 Basovizza, Trieste, Italy*

In order to gain more insight into the influence of chelating ligand's aromaticity on the reactivity of Ru(II) polypyridyl complexes, we synthesized two new Ru(II) complexes [Ru(Cl-Ph-tpy)(phen)Cl]Cl (**1**) and [Ru(Cl-Ph-tpy)(*o*-bqdi)Cl]Cl (**2**) (where phen = 1,10-phenanthroline and *o*-bqdi = *o*-benzoquinonediimine).¹ The new complexes were fully characterized by elemental analysis and various spectroscopic techniques, such as IR, UV-Vis and (1D and 2D) NMR, whereas their molecular structure in solid state was determined by single crystal X-ray diffraction analysis. The kinetics and the mechanism of the reaction of complexes **1** and **2** with the biologically more relevant 5'-GMP ligand were studied by UV-Vis spectroscopy. The interaction of complexes **1** and **2** with CT DNA was studied, and a competitive study of the intercalative agent ethidium bromide (EB) was performed. Furthermore, the affinity toward bovine serum albumin (BSA) was investigated, and their binding constants were determined.

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