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Cobalt(III) complexes with 2-acetylpyridine-derived Schiff bases: studies investigating ligand release upon reduction

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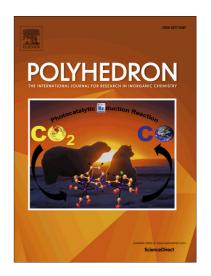
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ACCEPTED MANUSCRIPT

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Abstract

Cobalt(III) complexes $[Co(L1)_2]Cl\cdot CH_3OH\cdot 0.5H_2O$ (1), $[Co(L2)_2]Cl\cdot 1.5H_2O$ (2), 10 11 $[Co(L3)_2]Cl \cdot 2H_2O$ (3) and $[Co(L4)_2]Cl \cdot 3.5H_2O$ (4) were obtained with 2-acetylpyridine-N(4)phenylthiosemicarbazone (HL1), 2-acetylpyridine-N(4)-para-chlorophenylthiosemicarbazone 12 2-acetylpyridine-phenylhydrazone 13 (HL2),(HL3)and 2-acetylpyridine-*para*chlorophenylhydrazone (HL4). The complexes were characterized by means of 14 microanalyses, molar conductivities and their infrared and ¹H and ¹³C NMR spectra. 15 Electrochemical studies showed that the Co^{III}/Co^{II} reduction potential of complexes (3) and 16 17 (4) but not of complexes (1) and (2) are suitable for the compounds to be reduced by cellular 18 reductants. Reactivity assays showed that complex (3) undergoes reduction by sodium 19 dithionite with subsequent ligand release. The results suggested that coordination of 2-20 acetylpyridine-derived hydrazones with cytotoxic activity to cobalt(III) results in compounds which are able to release the bioactive ligand upon reduction. In addition, the cobalt(III) 21 complexes under study interacted with human serum albumin (HSA), indicating that they 22 23 could be transported by this protein.

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Keywords: hydrazones; thiosemicarbazones; cobalt(III) complexes; reduction; ligand release

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