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Reduction of rhenium(V)-oxo schiff base complexes with triphenyl phosphine ligands

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The purpose of this research is to produce radiopharmaceutical drugs for the possible use in therapy and diagnosis of cancer. A radiopharmaceutical drug is composed of a radioactive element contained within a chelating agent and linked to a biological targeting molecule. This can be achieved by complexing the radioisotope and conjugating the complex to a biologically active targeting molecule, such as a peptide antibody or antibody fragment. The biological targeting molecule directs radiation to specific peptides, antibodies and antibodies' fragments. The amount of dosage is limited to non-targeted tissue that occurs when free radionuclide is released from a decomposing molecule. Higher kinetic stability will maximize the localization of the radioisotope to the cancer sites and minimize the radiation dose to non-target tissues. This research was a continuation to seek the best pathway that would easily enable us to synthesize a kinetically inert metal complex $[\text{Re}(\text{Sal}_2\text{Phen})]$ attached to a ligand. The rigid Rhenium(V)-oxo Schiff base complex is found in the following form: $[\text{ReO}(\text{Sal}_2\text{Phen})\text{Cl}]$. Initially salicylic aldehyde reacted with bis-2-phenylenediamine to get sal_2phen . This tetradentate Schiff base ligand was reacted with TBA $[\text{ReOCl}_4]$ yielding $[\text{ReOCl}(\text{Sal}_2\text{Phen})]$. The reduction of Rhenium(V)-oxo core to Rhenium is obtained by reacting $[\text{ReO}(\text{Sal}_2\text{Phen})\text{Cl}]$ with a triphenyl phosphine (PPh_3) in dichloromethane and ethanol. Products obtained from this reaction were $^1[\text{Re}(\text{Sal}_2\text{Phen})(\text{PPh}_3)_2]$, $^2\text{ReCl}(\text{Sal}_2\text{Phen})\text{PPh}_3$, and $^3[\text{ReO}(\text{Sal}_2\text{Phen})\text{PPh}_3]$. Analysis of the major product were performed through crystals through mass spectrometry, hydrogen and phosphorus NMR, infrared spectra and x-ray crystallography.

