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Triphospha-Ferrocenes as Ligands. Crystal Structures of $[Fe(\eta^5-C_5Me_5)(\eta^5-C_2^TBu_2P_3)M(CO)_5)]$, (M= Cr, MO, W) and the Novel ruthenium and Nickel Complexes $[Fe(\eta^5-C_5Me_5)(\eta^5-C_2^TBu_2P_3)Ru_3(CO)_9]$ and $[Fe(\eta^5-C_5Me_5)(\eta^5-C_2^tBu_2P_3)Ni(Co)_9]$,

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TRIPHOSPHA-FERROCENES AS LIGANDS. CRYSTAL STRUCTURES OF [Fe(η^5 -C₂tBu₂P₃)M(CO)₅)], (M= Cr, Mo, W) AND THE NOVEL RUTHENIUM AND NICKEL COMPLEXES [Fe(η^5 -C₅Me₅) (η^5 -C₂tBu₂P₃)Ru₃(CO)₉] AND [Fe(η^5 -C₅Me₅)(η^5 -C₂tBu₂P₃)Ni(CO)₂]₂

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Abstract Syntheses and structures of penta- and hexaphosphorus analogues of ferrocene have been described recently 1 . Unlike their simple ferrocene analogues, these complexes have further ligating potential towards other transition metal centres by virtue of the availability of the ring phosphorus lone-pair electrons that are not involved in the η^5 -coordination. We now describe the first examples of coordination compounds of the triphospha-ferrocene [Fe(η^5 -C₅Me₅) (η^5 -C₂tBu₂P₃]. In the ruthenium complex [Fe(η^5 -C₅Me₅)(η^5 -C₂tBu₂P₃) Ru₃(CO)₉] 2 two adjacent phosphorus atoms of the η^5 -C₂tBu₂P₃ ring are interlinked by a ruthenium carbonyl cluster in which all three ruthenium atoms interact with the phosphorus atoms. The tetrametallic nickel complex [Fe(η^5 -C₅Me₅)(η^5 -C₂tBu₂P₃)Ni(CO)₂]₂ 3 represents the first example of intermolecular interlinkage of two phospha-ferrocene systems by two metal centres.

Key Words: Metal complexes, phospha-ferrocenes, 31P NMR spectroscopy, X-ray crystal structure analysis

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