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DIPHOSPHIRENIUM SALT: A NEW VERSATILE LIGAND

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Strained aromatic cyclopropenylium cations are versatile ligands for transition metal complexes: η^{1} -, η^{2} - and η^{3} -coordination modes have been observed. Here we report on the ligand properties of the related diphosphirenium salt A[1]. Treatment of a dichloromethane solution of diphosphirenium salt A with an equimolar amount of tetrakis(triphenylphosphine) palladium afforded 1,3-diphospha-2-bis(triphenylphosphine)pallada(II)cyclobutene B in 70% yield. Exchange of the triphenylphosphine ligands occurs with various phosphines, and the structures of these new diphosphametallacyclobutenes have been elucidated by NMR and in one case by a single X-ray diffraction study[2].

Surprisingly, when a dichloromethane solution of complex B was refluxed overnight in the presence of diphosphirenium salt A a new complex C was formed. This cationic 1,3-diphospha-2,4-dipallada(II)tricyclo[1.1.1]pentane is the first compound featuring pyramidal µ²-phosphinidene units.

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