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Synthesis and structure of palladium complexes: $[Ph_4Sb]^+_2[Pd_2I_6]^{2-}$, $[Ph_4Sb(DMSO)]^+_2[Pd_2I_6]^{2-}$, $[Ph_3AmP]^+_2[Pd_2I_6]^{2-}$ and $[Ph_3EtP]^+_2[Pd_2I_6]^{2-}$

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Abstract

By the reaction of palladium diiodide with tetraphenylantimony iodide in aqueous solution of HBr obtained complex $[Ph_4Sb]^+_2[Pd_2I_6]^{2-}$ (I), whose recrystallization from dimethylsulfoxide led to the complex $[Ph_4Sb(DMSO)]^+_2[Pd_2I_6]^{2-}$ (II). Similarly synthesized complexes $[Ph_3AmP]^+_2[Pd_2I_6]^{2-}$ (III) and $[Ph_3EtP]^+_2[Pd_2I_6]^{2-}$ (IV), whose composition is not changed by recrystallization. According to the X-Ray date, the complex II consists of trigonal-bipyramidal cation [Ph₄Sb(DMSO)]⁺ with the oxygen atom of the ligand dimethylsulfoxide in the axial position (Sb···O 2.481(4) Å) and planar centrosymmetric binuclear anions $[Pd_2I_6]^{2-}$ $(Pd-I_b 2.5875(5), 2.5965(7) \text{ Å}, Pd-I_t 2.5836(8), 2.6053(8) \text{ Å}; <math>\mu\nu$ -IPdI 85.35(2)°-93.35(3)°; $\mu\nu$ -IPdI 175.75(3)°, 175.82(3)°). The crystals III, IV contain tetrahedral cations tetraorganylphosphonium (III P-C 1.785(4)-1.803(4) Å, CPC 108.6(2)°-110.6(2)°; IV P-C 1.792(3)-1.802(3) Å, CPC 106.55(13)°-112.10(14)°) and planar centrosymmetric binuclear anions (III Pd-I_b 2.6061(4), 2.6093(4) Å, Pd-I_t 2.6002(4), 2.6093(4) Å; цис-IPdI 84.224(14)°-95.775(15)°, транс-IPdI 175.778(16)°, 175.958(17)°; IV Pd-I_b 2.6079(3), 2.6129(3) Å, Pd-I_t 2.5913(3), 2.6065(3) Å; uuc-IPdI 85.874(9)°-94.126(9)°, транс-IPdI 172.808(12)°, 173.125(12)°).

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