

Erratum

Erratum to “*Ortho*-metallated transition metal complexes derived from tertiary phosphine and arsine ligands” [Coord. Chem. Rev. 250 (2006) 1851–1888]

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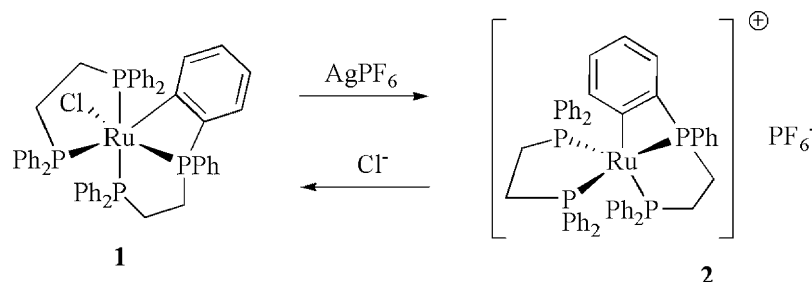
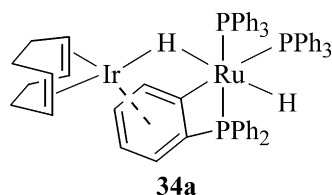
Available online 20 January 2007

The following paragraph was inadvertently omitted from Section 4.3 of the original manuscript:

Ortho-metallation has also been observed in Ru complexes of dppe. The sole product of the reaction of *trans*-[RuCl₂(dppe)₂] with neat AlMe₃ at 90 °C is the *ortho*-metallated complex [RuCl{2-C₆H₄P(Ph)CH₂CH₂PPh₂}(dppe)] (**1**), which has a distorted octahedral structure. Abstraction of Cl[−] from **1** with AgPF₆ generates the 16-electron Ru(II) cation [Ru{2-C₆H₄P(Ph)CH₂CH₂PPh₂}(dppe)]⁺ (**2**) (Scheme 1) in which the *ortho*-metallated carbon atom occupies the apical site of the square pyramidal geometry about ruthenium. This reaction is reversed by treatment of **2** with dodecyltrimethylammonium chloride. The initial product of the reaction of *trans*-[RuCl₂(dppe)₂] with AlMe₃ is *trans*-[RuClMe(dppe)₂], which can be isolated. The proposed sequence leading to **1** is: (1) AlMe₃-promoted loss of Cl[−] from *trans*-[RuCl₂(dppe)₂] giving the 16-electron cation [RuMe(dppe)₂]⁺; (2) loss of methane from this species, generating cation **2**; (3) reaction of **2** with Cl[−] [1].

[1] K. Umezawa-Vizzini, T.R. Lee, Organometallics 16 (1997) 5613.

On page 1859 Structure **34a** should be:



Scheme 1.

DOI of original article: 10.1016/j.ccr.2005.10.003.

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