

Mixed-Valence Properties in L-type Complexes

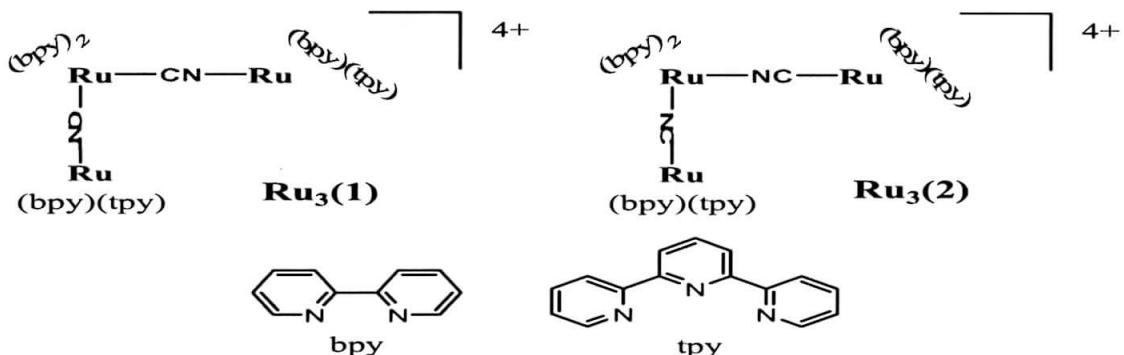
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Abstract

Mixed-Valence chemistry in ruthenium is the transition metal choice to study electron transfer or exchange because low spin d^5 - d^6 centers form relatively stable coordinate complexes.¹⁻³ Hush published the well-known basis of mixed-valence chemistry which focused on the two-state model.⁴ Creutz, Newton and Sutin reported a new aspect of the superexchange coupling model with the three-state system.^{5,6} Few reports provided significant points of real systems for superexchange coupling.⁷⁻⁹

The focus of this report is the synthesis and study of L-type tri-ruthenium complexes that exhibit interesting intervalence absorption and strong comproportionation between terminal metal centers. The electrochemistry of $\Delta E_{1/2}$ between terminal centers for Ru₃(1) and Ru₃(2) ranged between 0.12 and 0.13 V.



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