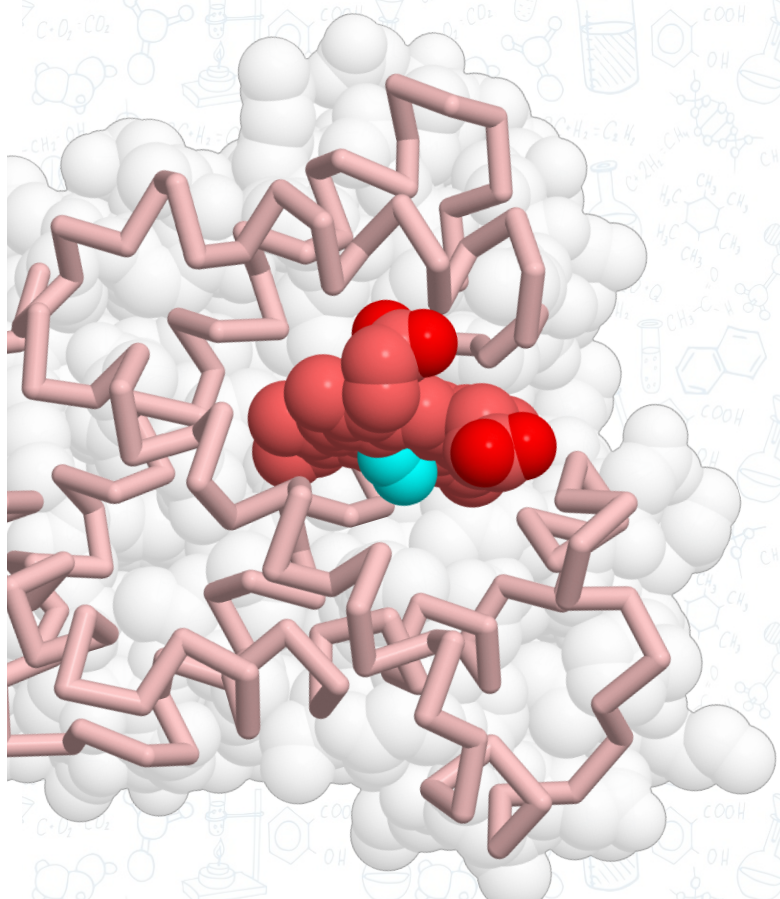


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Osmium Diazoalkane Derivatives with Pentamethylcyclopentadienyl Ligand: Synthesis and Reactivity

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Diazoalkane complexes $[\text{Os}(\eta^5\text{-C}_5\text{Me}_5)(\text{N}_2\text{CAr}_1\text{Ar}_2)(\text{PPh}_3)\{\text{P}(\text{OR})_3\}]\text{BPh}_4$ (**1**, **2**) were prepared by reacting bromo-compounds $\text{OsBr}(\eta^5\text{-C}_5\text{Me}_5)(\text{PPh}_3)\{\text{P}(\text{OR})_3\}$ with diazoalkane $\text{Ar}_1\text{Ar}_2\text{CN}_2$ in the presence of NaBPh_4 .

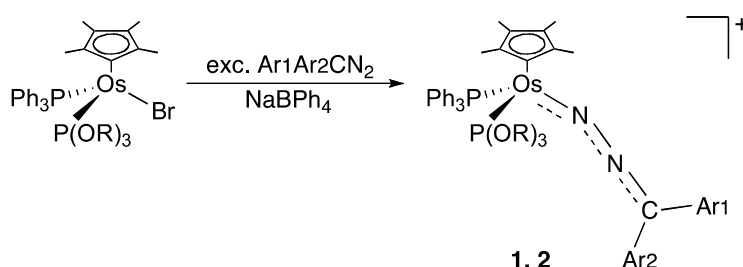


Figure 1: $R = \text{Me}$ (**1**), Et (**2**); $\text{Ar}_1 = \text{Ar}_2 = \text{Ph}$ (**a**); $\text{Ar}_1 = \text{Ph}$, $\text{Ar}_2 = p\text{-tolyl}$ (**b**); $\text{Ar}_1\text{Ar}_2 = \text{C}_{12}\text{H}_8$ (**c**)

Treatment of complexes **1** and **2** with air afforded dioxygen derivatives $[\text{Os}(\eta^5\text{-C}_5\text{Me}_5)(\eta^2\text{-O}_2)(\text{PPh}_3)\{\text{P}(\text{OR})_3\}]\text{BPh}_4$ (**3**, **4**).

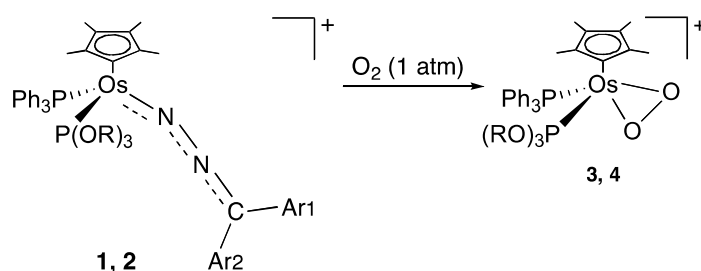


Figure 2: $R = \text{Me}$ (**3**), Et (**4**); $\text{Ar}_1 = \text{Ar}_2 = \text{Ph}$ (**a**); $\text{Ar}_1 = \text{Ph}$, $\text{Ar}_2 = p\text{-tolyl}$ (**b**); $\text{Ar}_1\text{Ar}_2 = \text{C}_{12}\text{H}_8$ (**c**)

Vinylidene $[\text{Os}(\eta^5\text{-C}_5\text{Me}_5)\{\text{C}=\text{C}(\text{H})\text{R}_1\}(\text{PPh}_3)\{\text{P}(\text{OR})_3\}]\text{BPh}_4$ and propadienylidene derivatives $[\text{Os}(\eta^5\text{-C}_5\text{Me}_5)(\text{C}=\text{C}=\text{CR}_1\text{R}_2)(\text{PPh}_3)\{\text{P}(\text{OR})_3\}]\text{BPh}_4$ were also prepared by allowing complexes **1** and **2** to react with terminal alkynes $\text{HC}\equiv\text{CR}_1$ and $\text{HC}\equiv\text{CCR}_1\text{R}_2(\text{OH})$, respectively.

All the complexes were characterised spectroscopically (IR and NMR) and by X-ray crystal structure determinations.