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Coordinating Properties of Dipod and Tripod Phosphorus Ligands with Separated Donor and Acceptor Sites

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COORDINATING PROPERTIES OF DIPOD AND TRIPOD PHOSPHORUS LIGANDS WITH SEPARATED DONOR AND ACCEPTOR SITES

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The coordinating properties of dipod and tripod phosphorus ligands $L_{\rm I} = R_2 M' ({\rm OCH_2PMe_2})_n ({\rm CH_2CH_2PMe_2})_{2-n}$ and $L_{\rm II} = {\rm RM'} ({\rm OCH_2PMe_2})_n ({\rm CH_2CH_2PMe_2})_{3-n}$ (M' = Si, Ge) with separated donor and acceptor centres have been investigated using electron rich metal complex fragments, e. g. ${\rm M(CO)}_m$ (M = Cr, Mo, W), π -C₅H₅Co, RhCl(CO) or Ni(CO), as bonding partners.

The poster will present information about a) the preparation of the ligands $L_{\rm I}$ and $L_{\rm II}$, b) the synthesis of complexes of these ligands, c) the effects of coordination on the spectroscopic data [ν (CO), $\Delta \delta$, ΔJ], allowing conclusions about metal/ligand interactions, and d) the molecular structures of selected examples, two of which are shown below:

(A) MeSi(OCH₂PMe₂)₃Ni(CO); (B) [RhCl(CO)(PMe₂CH₂CH₂)₂SiMe₂]₂.

So far, interaction between the basic centres M and the acceptor sites M' could not be established. Mechanisms for reducing the enhanced electron density on M will be discussed.

