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## PH-Functional Methylenebisphosphanes RR'P-CH<sub>2</sub>-PRH (R'= R, H) - Reactive Bridging Ligands with A Small Bite Angle

G. Hasselku $\beta^a$ ; S. Hietkamp $^a$ ; P. C. Knüppel $^a$ ; O. Stelzer $^a$ 

<sup>a</sup> Anorganische Chemie, Bergische Universität-GH Gauβstr. 20, Wuppertal

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## PH-Functional Methylenebisphosphanes RR'P-CH<sub>2</sub>-PRH (R'= R, H) - Reactive Bridging Ligands with A Small Bite Angle

G. Hasselkuß, S. Hietkamp, P. C. Knüppel and O. Stelzer\*
Fachbereich 9, Anorganische Chemie, Bergische Universität-GH
Gaußstr. 20, D-5600 Wuppertal 1

PH-functional methylenbisphosphanes RR'P-CH<sub>2</sub>-PRH (R' = R, H) are versatile ligands in coordination chemistry. They may oxidatively add with their PH-bonds to transition metals in low oxidation numbers and form cluster compounds with P-C-P-bridges. P-C-P-bond cleavage reactions afford phosphinidenes  $\langle$  PR? (through a formal 1,2-hydrogen shift from phosphorus to the CH<sub>2</sub>-group, RR'P-CH<sub>2</sub>-PRH  $\rightarrow$  RR'P-CH<sub>3</sub> +  $\langle$  PR>) which are incorporated into the oligometallic cluster framework formed. As reactive bidentate donors with a small bite angle and low steric demand they also may bridge isolated and directly bonded transition metals in various manners (I - III), e.g.

They add to M=M-multiple bonds to give binuclear complexes with a  $\rm M_2P_2C$  five membered PH-functional framework, e.g.

$$(C_5H_5)_4M_0_2(CO)_4 + RR'P-CH_2-PRH \rightarrow (C_5H_5)(CO)_2M_0-M_0(CO)_2(C_5H_5)$$

The structure of compounds of type I - IV will be discussed using  $^{1}\text{H-}$ ,  $^{13}\text{C-}$ ,  $^{31}\text{P-nmr}$  data and x-ray structures. Reactions at the PH-bonds of the ligands RR'P-CH<sub>2</sub>-PRH in complexes of type I - IV will be reported.