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First Example for Competitive Formation of 1H-Phosphirene and η^1 -1-Phosphaallene Complexes

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2*H*-azaphosphirene complex $1^{[1]}$ serves as an efficient precursor for terminal phosphanediyl complex 2 ($R = CH(SiMe_3)_2$), which was trapped with acetylene derivatives 3a-e ($R_1E = Me_3Si$ (3a), Ph_3Si (3b), Me_3Sn (3c), Ph_3Sn (3d), Bu_3Sn (3e)) to yield 2,3-bifunctionalized 1*H*-phosphirene 4 and/or η^1 -1-phosphaallene complexes 4a-d were formed in the case of 3a-d, whereas η^1 -1-phosphaallene complexes 5a,b were generated only, if stannyl-substituted acteylenes 3c,d were used. In these cases, 4c,d were obtained as byproducts. If 3e was employed, having tert-butyl

$$\begin{array}{c} R \\ W(CO)_5 \\ \hline \\ R_3EC = COEt (3) \\ \hline \\ PhCN \\ \hline \\ PhCN \\ \hline \\ (OC)_5W|RP=C=C(OEt)SnR_3 (5a,b) \\ \hline \\ + 4cd \\ \hline \end{array}$$

groups at tin, then no η^1 -1-phosphallene complexes were formed, but, instead, <u>diastereoisomeric</u> complexes **6a,b** obtained, exclusively. These complexes are related to 1*H*-phosphirenes **4a-d**, but have zwitterionic ring systems, owing, most probably, to a strong π -donor electron interaction of the ethoxy group. [3] NMR spectroscopic data of **4a-d**, **5a,b** and **6a,b** and single crystal X-ray structures of **4a**, **4d** and **6a** will be presented.

References

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