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# Phosphorus, Sulfur, and Silicon and the Related Elements

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# P-Metalated Phosphaalkenes

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### P-METALATED PHOSPHAALKENES

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Structure and reactivity of metalated species resulted from insertion to reactive element-halogen bond is of great interest since their reliability to Grignard reagents, one of the most powerful tools in the arsenal of synthetic chemistry. Here we present the study of reactivity of group 14 carbenoids (germane- and stananediyls) toward P—Cl bond in iminophosphines and methylenephosphines. Treatment of Mes\*N=PCl with M(NR<sub>2</sub>)<sub>2</sub> gives just common substitution products<sup>2</sup> of the type Mes\*N=P-NR<sub>2</sub>. But when (Me<sub>3</sub>Si)<sub>2</sub>C=PCl is used, either substitution proceeds or metal insertion products are isolated in high yields depending on nucleophilic properties of amide substituent:

$$(Me_3Si)_2C=P-NRR'$$

$$1a,b: M=Ge, Sn; R=H, R'=Ttb$$

$$(Me_3Si)_2C=P-M(NRR')_2CI$$

$$2a-c: M=Ge; R=SiMe_3, R'=SiMe_3, Ttb, Tip$$

$$2d: M=Sn; R=R'=SiMe_3$$

All the compounds have been characterized by <sup>1</sup>H, <sup>31</sup>P, <sup>13</sup>C spectral data, and x-ray structure for **2a** has been determined. Long P—Ge bond implies possible rich scope of reactivity of germetalated phosphaalkenes.

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