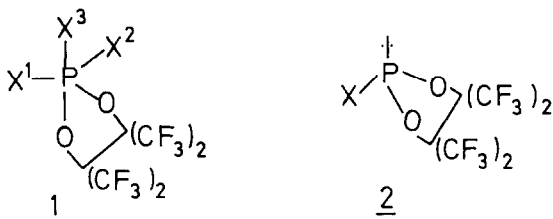


THE PERFLUOROPINACOLYL GROUP, A STABILIZING SUBSTITUENT FOR UNUSUAL PHOSPHINES AND PHOSPHORANES

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The perfluoropinacolyl group (=PFP) stabilises halogenophosphoranes of the type 1 e.g.



$\text{X}^1=\text{X}^2=\text{X}^3=\text{F}, \text{Cl}, \text{Br}$ or $\text{X}^1=\text{X}^2=\text{Cl}, \text{Br}$, $\text{X}^3=\text{F}$, covalent species which are able to react with Li_2PFP to form spirocyclic systems [1]. Precursors of compounds 1 are the phosphites 2 ($\text{X}=\text{F}, \text{Cl}, \text{Br}$). The aminophosphite with $\text{X}=\text{NH}_2$ shows interesting behaviour towards hexafluoroacetone forming a spiro system containing a four and a five membered ring. Phosphites with $\text{X}=\text{NR}^1\text{R}^2$ add readily chlorine to give stable chlorophosphoranes. Products of the ammonolysis of the trifluorophosphoranes ($\text{X}^1=\text{X}^2=\text{X}^3=\text{F}$) were the aminophosphoranes $(\text{H}_2\text{N})_n\text{PF}_{3-n}$ (PFP) ($n=1, 2, 3$) in good yields. Dynamic processes of the latter compounds were discussed on the bases of low temperature ^{19}F n.m.r. spectroscopy.

1 G.-V. Rösenthaller, J.A. Gibson, K. Sauerbrey, and R. Schmutzler, Z. anorg. allg. Chem. 450 (1979)

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