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The Regiochemistry of the Reaction of Dihalotris(hexafluoroisopropoxy)-phosphoranes with 1,2-Epoxy-3-trimethylsiloxypropane

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THE REGIOCHEMISTRY OF THE REACTION OF DIHALOTRIS(HEXAFLUOROISOPROPOXY)-PHOSPHORANES WITH 1,2-EPOXY-3-TRIMETHYLSILOXYPROPANE

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The interaction of trimethylsilylepoxyde **1** with fluorinated halogenophosphoranes is investigated. The reaction of silylepoxyde **1** with phosphorane **2** has been found to give the phospholane 3 (70%, δ_P -60.1 ppm, X=Cl; 27%, δ_P -60.3 ppm, X=Br) or phosphate **4** (15%, X=Cl; 71%, X=Br) as the major product depending on the nature of substituent X at low temperature. The initial opening of epoxyde occurs

OSiMe3 + (RFO)3PX2
$$\xrightarrow{-Me3SiX}$$
 (RFO)3P O X

 $\xrightarrow{1}$ $\xrightarrow{2}$ (RFO)3PX2 \xrightarrow{X} OSiMe3 (RFO)4PCl $\xrightarrow{1}$ $\xrightarrow{\Delta}$ 3 + (RFO)4P-O OSiMe3 + $\xrightarrow{6}$ + (RFO)4P-O OSiMe3 RF = $\xrightarrow{CF3}$, X = Cl, Br $\xrightarrow{7}$

SCHEME 1

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in reaction between compound 5 and silylepoxyde 1 under heating, phosphorane 6 being the main product. The partial cyclization with eliminating of R_FOSiMe_3 and formation f phosphorane 3 (X = Cl) takes place in this reaction. The structure of compounds 3, 6, 7 was confirmed by $^{13}C,\,^{31}P$ NMR.