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Synthesis of the First Stable Pentacoordinate Selenaphosphirane

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SYNTHESIS OF THE FIRST STABLE PENTACOORDINATE SELENAPHOSPHIRANE

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Although heteracyclopropanes containing a highly coordinate phosphorus atom are expected to have interesting structures and reactivities, only few examples have been reported so far. Here we report the synthesis of the first stable pentacoordinate selenaphosphirane 4 by taking advantage of the stabilizing effects of the Martin ligand. Alkylidenephosphorane 3, which was prepared by the reaction of fluorophosphorane 1 with trimethylsilyl trifluoromethanesulfonate followed by

F₃C CF₃

Me

TMSOSO₂CF₃

$$Et_2O$$

THF

THF

Ph

Me

 Cp_2TiSe_5

THF

r.t.

 F_3C
 CF_3
 O
 OSO_2CF_3
 OS

SCHEME 1

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deprotonation of the resulting phosphonium salt 2 with 1.2 equiv of 2,4,6-tirmethylphenyllithium (MesLi), was allowed to react with Cp_2TiSe_5 in THF at room temperature to give the pentacoordinate selenaphosphirane 4. The pentacoordinate structure of 4 was determined by X-ray crystallographic analysis.