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Preliminary communication

HYDRIDE MIGRATION IN THE MASS SPECTRA OF SOME 2-PHENYL-1,3,2-DIOXASILACYCLOALKANES

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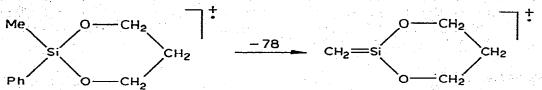
Summary

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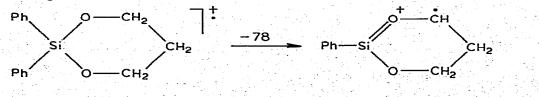
The mass spectra of a series of 2-phenyl-1,3,2-dioxasilacycloalkanes all contain a peak at m/e P - 78, which we suggest arises from hydride migration from the ring.

We have for some time been interested in the mass spectral fragmentation of heterocyclic organoboranes [1, 2] and have recently begun a detailed investigation of heterocyclic silanes. Although there have been a few reports of loss of C_6H_6 from parent ions [3, 4, 5] the importance of this as a general process for alkoxyphenylsilanes and 2-phenyl-1,3,2-dioxasilacycloalkanes does not appear to have been commented on.

Our initial investigations involved a study of a series of 2-methyl-2-phenyl-1,-3,2-dioxasilacycloalkanes, and it was observed that the mass spectrum of each compound contained a peak at m/e P - 78. It is attractive to suggest that the loss of 78 from the parent ion resulted in the formation of a species involving a carbon—silicon double bond.



However, in further investigations involving diphenyl- or chlorophenyl-silanes it has been found that all the mass spectra show the loss of 78 from the molecular ion, which strongly suggests that there is a hydrogen atom migration from the ring.



Some indication of the generality of this process can be seen from the examples given in Table 1. It is also of interest that the mass spectra of diethoxydiphenylsilane and dimethoxydiphenylsilane also contained peaks at m/e P - 78.

TABLE 1

RELATIVE INTENSITIES OF P AND P — 78 PEAKS IN THE MASS SPECTRA OF A SERIES OF 2-PHENYL-1,3,2-DIOXACYCLOSILAALKANES

Compound	P 7. base	2-78 % base	Compound	P % base	P-78 % base
0-CH2-CH2 Ph(Me)5i 0-CH2-CH2	15	21	Ph251 CMe2	15	97
Pn(Me)S1 0	8	з	Ph(Me)Si O-CHMe CHMe	5	20
Ph2Si CH2 O-CH2 O-CH2	26	50	Ph ₂ Si O-CHMe CHMe	7	48
Ph(Me)Si O-CHMe CH2 O-CH2	9	10	Ph(Me)Si CH ₂ CH ₂ CH ₂	22	2
Ph ₂ Si CH ₂ O-CH ₂ CH ₂	33	100	Ph ₂ Si CH ₂ CH ₂	34	15
Ph(Me)Si CH ₂ CH ₂	8	10	Ph (CI) SI O-CHMe CHMe CHMe	64	10
Ph2Si CH2	15	64	Ph(Me)Si	100	58
Ph (Me)Si O-CH2 CMe2	8	24	Ph ₂ Si	100	63

Experimental

The mass spectra were recorded, using an A.E.I. MS902 mass spectrometer at 70 eV. The source was maintained at 170° C and the compounds were introduced as neat liquids or solids using an unheated direct-insertion probe.

Acknowledgement

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