Preliminary communication

Cyclopentadienyltin(II) halides. A novel type of divalent organotin compounds

K.D. BOS, E.J. BULTEN and J.G. NOLTES Institute for Organic Chemistry TNO, Utrecht (The Netherlands) (Received April 27th, 1972)

So far only a few true divalent organotin compounds are known, *viz*. dicyclopentadienyltin¹, its boron trifluoride complex², bis(methylcyclopentadienyl)tin³ and bis(phenylbarenyl)tin⁴. The divalent character of these compounds has been confirmed by Mössbauer spectroscopy. Very recently the preparation of a dialkylstannylene—pentacarbonylchromium complex has been reported⁵. However, no conclusive Mössbauer data were presented.

We report here the synthesis of divalent organotin compounds of the type C_5H_5SnX (X = Cl, Br). Although such species have been occasionally suggested as intermediates^{2, 6, 7} they have not been isolated as such.

Cyclopentadienyltin halides precipitate as white, crystalline solids upon mixing concentrated THF solutions of dicyclopentadienyltin and the tin dihalide.

$$(C_5H_5)_2Sn + SnX_2 \xrightarrow{\text{THF}} 2C_5H_5SnX_2 \downarrow$$
(I)
(IIa) X = Cl
(IIb) X = Br

Analytical and molecular weight data are given in Table 1.

TABLE I

Compound C ₅ H ₅ SnX	Analysis ^a Found (calcd.)			u	Mol. wt. ^b Found (calcd.)	М.р. (°С)
	с	н	x	Sn		
X = Cl	27.10	2.24	16.27 (16.17)	53.7	217 (219)	130-133
X = Br	22.81 (22.77)	1.98 (1.91)	30.19 (30.30)	45.39 (45.01)	256 (264)	160–162

ANALYTICAL AND MOLECULAR WEIGHT DATA FOR C5H3SnX

^aThe elemental analyses were carried out at the Micro-Analytical Department of our Institute. ^bEbulliometric in THF.

J. Organometal Chem., 39 (1972)

PRELIMINARY COMMUNICATION

¹¹⁹ ^mSn MOSSBAUER DATA^a

The ^{119 m}Sn Mössbauer spectral parameters are given in Table 2.

TABLE 2

	IS ± 0.06 (mm/sec)	QS ± 0.06 (mm/sec)
C,H,SnCl	1.65	1.05
C, H, SnBr	1.64	0.99
(C,H,),Sn	1.66 (1.64) ^D	0.81 (0 . 86) ^b
SnCl.	2.00 (2.07) ^c	0.56 (0) ^C
SnBr ₂	1.88 (1.93) ^c	0 (0) ^c

^aRecorded at 77°K. The IS values are relative to α -Sn at 77°K. ^b Ref. 2. ^c Ref. 8.

The Mössbauer data clearly show the compounds to be derivatives of divalent tin. Furthermore, it can be concluded from the spectra that the products contain only one kind of tin atom. Thus, in the solid state a complex such as $(C_5H_5)_2Sn-SnX_2$ can be ruled out.

The mass spectrum of C_5H_5SnCl displays $C_5H_5Sn^+$ as the most abundant tin-containing fragment. The parent peak $[C_5H_5SnCl]^+$ has a relative intensity of 9.3 percent^{*}.

The PMR spectra of IIa and IIb in THF show a singlet at $\delta \approx 6.05$ ppm. No Sn-C-H coupling was observed. This can be explained by a rapid exchange of

the cyclopentadienyl groups $[cf.^{2} (C_{5}H_{5})_{2}Sn : \delta = 5.89 \text{ ppm}, J(^{117}, ^{119}Sn-C-^{1}H) = 15.5 \text{ Hz}].$



For these cyclopentadienyltin(II) halides a structure such as III seems probable. This structure involves stabilization by intermolecular tin—halogen bridging, which precludes polymerization with formation of tin(IV) species containing metal—metal bonds (*cf.* ref. 9). The synthesis of other compounds of the type C_5H_5SnX in which X is a monoanionic ligand with pronounced donor properties would thus be of interest.

Our observation that C_5H_5SnCl can also be prepared by the reaction of I with anhydrous hydrochloric acid in THF suggests that the reaction of I with acidic compounds is generally applicable for the synthesis of C_5H_5SnX compounds^{**}. E.g., the reaction of I

J. Organometal Chem., 39 (1972)

^{*}For the mass spectrum of $(C_5H_5)_2$ Sn, see ref. 2.

 $[\]star$ Reactions of (C₅H₅)₂Sn with proton active species leading to inorganic tin(II) derivatives have been studied recently¹⁰.

with phenol in THF gives rise to the formation of cyclopentadienyltin phenoxide, $C_5H_5SnOC_6H_5$.

ACKNOWLEDGEMENTS

Financial support by the International Tin Research Council, London, is gratefully acknowledged. We are much indebted to Ir. A.M. van der Kraan of the Interuniversitair Reactor Instituut, Delft, for performing the Mössbauer measurements.

REFERENCES

- 1 E.O. Fischer and H. Grubert, Z. Naturforsch. B, 11 (1956) 423.
- 2 P.G. Harrison and J.J. Zuckerman, J. Amer. Chem. Soc., 92 (1970) 2577.
- 3 L.D. Dave, D.F. Evans and G. Wilkinson, J. Chem. Soc., (1959) 3684.
- 4 Yu. A. Aleksandrov, V.I. Bregadse, V.I. Goldanskii, L.J. Zakhardin, O.Yu. Okhlobystin and V.V. Khrapov, Dokl. Akad. Nauk SSSR, 165 (1965) 593.
- 5 T.J. Marks, J. Amer. Chem. Soc., 93 (1971) 7090.
- 6 P.G. Harrison and J.J. Zuckerman, J. Amer. Chem. Soc., 91 (1969) 6885.
- 7 F.J.A. des Tombe, Ph. D. Thesis, University of Utrecht, 1969.
- 8 J.D. Donaldson and D.G. Nicholson, J. Chem. Soc. A, (1970) 145.
- 9 M. Massol, J. Satgé, P. Rivière and J. Barrau, J. Organometal. Chem., 22 (1970) 599.
- 10 P.G. Harrison, J. Chem. Soc., Chem. Commun., in press.

J. Organometal. Chem., 39 (1972)