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GEOMETRICAL ISOMERS OF STIBINOUS AND STIBINIC PROPENYL COMPOUNDS

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IN continuation of stereochemical investigation of organometallic alkenyl compounds syntheses of a number of geometrical isomers of stibinous and stibinic propenyl compounds have been effected.

<u>Cis-</u> and <u>trans-</u>lithiumpropenyl react with stibinous chloride to form <u>cis-</u> and <u>trans-</u>tripropenyl stibine, respectively.

The reaction of the isomers with halogens leads to a series of isomeric stibinic compounds.

$$(CH_3CH=CH)_3Sb + X_2 \longrightarrow (CH_3CH=CH)_3SbX_2 ; X=C1,Br,J$$

The same compounds, <u>cis</u>- and <u>trans</u>- respectively, have been isolated by the reaction of <u>cis</u>- and <u>trans</u>-propenyl stibine with thallic chloride.

<u>Cis</u>-propenylstibine bromide and chloride are crystallinic solids, and the <u>trans</u>-isomers are liquids.

The liquid geometrical isomers of pentapropenyl stibine have been prepared by the reaction of <u>cis-</u> and <u>trans-</u>tripropenylstibine dibromide and the corresponding isomers of lithium propenyl.

(CH₃CH=CH)₃SbBr₂ + 2CH₃CH=CHLi - (CH₃CH=CH)₃Sb + 2LiBr

These differ by the refractive index (<u>cis</u>-compound n_D^{2O} 1.5610, <u>trans</u>-

compound n_D^{20} 1.5490) and the infra-red spectra.

Two tetrapropenylstibonium bromide isomers with different melting points (cis-compound m.p. 140-143°, trans-compound m.p. 45-48°) and infrared spectra have been prepared by treatment of these isomers with the definite quantity of bromine.

NN	Compounds	Infra-red spectra in cm ⁻¹	
		cis	trans
1	(CH ₃ CH=CH) ₃ Sb	920 w, 970 w, 1182 m, 1615 m	935 m, 970 s, 1068 m, 1200 s, 1620 s
2	(CH ₃ CH=CH) ₃ SbCl ₂	923 w, 1200 w, 1600 w	955 s, 1070 m, 1188 s, 1615 s
3	(CH3CH=CH)3SbBr2	925 w, 1600 w	953 s, 1068 m, 1188 s, 1620 s
4	(CH ₃ CH=CH) ₄ SbBr	925 w, 1600-1610 m	965 s, 1068 s, 1190 s, 1620 s
5	(CH ₃ CH=CH) ₃ SbJ ₂	928 w, 1470 s, 1610 m	948 s, 1065 m, 1625 s
6	(сн ₃ сн=сн) ₅ sь	920 w, 974 w, 1200 w, 1595 s	972 s, 1068 m, 1185 m, 1600 s

s - strong, m - middle, w - weak

The configurations of the compounds prepared were assigned on the basis of infra-red analysis and are in agreement with the regularity drawn by us. 1

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