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ACCEPTED MANUSCRIPT

Synthesis and structure of chlorotriphenylantimony pentafluoro- and pentachloroaroxides

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Graphical abstract



Highlights

- A facile synthesis of fluoro- and chloro-containing antimony(V) derivatives via ligand redistribution reaction was reported
- New chloro(pentafluorophenoxy)- and chloro(pentachlorophenoxy)triphenylantimony were first obtained in high yields
- The structure of synthesized compounds was unambiguously confirmed by spectral analysis and X-ray diffraction study

ABSTRACT

Chloro(pentafluorophenoxy)triphenylantimony $Ph_3SbCl(OC_6F_5)$ (3a) and chloro(pentachlorophenoxy)triphenylantimony Ph₃SbCl(OC₆Cl₅) (**3b**) were synthesized triphenylantimony dichloride (1) by the reactions of with bis(pentafluorophenoxy)triphenylantimony (2a) and bis(pentachlorophenoxy)triphenylantimony (2b), respectively. The structure of compounds **3a,b** was confirmed by IR, ¹H, ¹³C{¹H} and ¹⁹F{¹H} NMR spectroscopy, elemental analysis and single-crystal Xray diffraction study. According to X-ray diffraction data, two types of crystallographically independent molecules exist in crystals 3a and 3b. The antimony atoms have a distorted trigonal bipyramidal coordination with the chlorine and oxygen atoms in axial positions. The axial OSbCl angles in **3a** and **3b** are 177.07(19)°, 177.09(19)° and 178.25(5)°, 178.28(5)°, respectively; the equatorial angles are 118.0(9)°-122.2(4)° (3a) and 113.49(8)°-129.23(8)° (3b). The Sb-C equatorial bond lengths are

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