

P-10

RECENT INVESTIGATION OF DIVALENT METAL HEXAFLUOROANTIMONATES

Darja Gantar*, Ivan Leban, Boris Frlec

'Jožef Stefan' Institute, 'Edvard Kardelj' University Ljubljana, 61000 Ljubljana
(Yugoslavia)

and John H. Holloway

The University, Leicester LE1 7RH (U.K.)

Reactions between metal difluorides ($M = \text{Cr, Fe, Ni, Ca, Sr, Ba}$) and excess antimony pentafluoride in anhydrous hydrogen fluoride yielded adducts with nonstoichiometric compositions. After prolonged pumping of volatiles at room temperature $\text{MF}_2 \cdot 2\text{SbF}_5$ adducts were isolated for $M = \text{Cr, Fe, Ni}$. In the case of Ca, Sr and Ba the preparation described led only to nonstoichiometric compounds. The solid adducts were characterized by chemical analysis, vibrational spectra and X-ray powder diffraction patterns.

A single-crystal X-ray study on $\text{AgF}_2 \cdot 2\text{SbF}_5$ showed that the crystals are triclinic, space group $P\bar{1}$, with unit-cell dimensions $a = 5.224$ (1), $b = 5.466$ (2), $c = 8.779$ (2) Å, $\alpha = 75.78$ (2), $\beta = 89.02$ (2), $\gamma = 65.29$ (2)°, and $Z = 1$. The Ag^{2+} ion has a square-planar coordination; two Ag-F distances are 2.09 and 2.13 Å, with two other F atoms at 2.43 Å, completing a distorted octahedron. There are also distorted SbF_6^- octahedra in the structure: four Sb-F distances range from 1.83-1.87 Å, while two distances are of 1.94 and 1.96 Å.