

## ORGANOBROMINE(III)-COMPOUNDS [1]

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Investigations of nucleophilic substitution on  $\text{BrF}_3$  enabled us to prepare  $\text{C}_6\text{F}_5\text{BrF}_2$  - the first covalent organobromine(III) compound. Its spectroscopic properties and its chemical behaviour show that the formerly published [2] compound formed by reaction of  $\text{C}_6\text{F}_5\text{Br}$  with fluorine was not  $\text{C}_6\text{F}_5\text{BrF}_2$ .

The  $^{19}\text{F}$ -NMR-spectrum of  $\text{C}_6\text{F}_5\text{BrF}_2$  shows four resonances with correct intensities and shifts in the expected regions.  $\text{C}_6\text{F}_5\text{BrF}_2$  - compared with  $\text{BrF}_3$  and  $\text{C}_6\text{F}_5\text{Br}$  - shows significant shifts: highfield for bromine bonded fluorine and lowfield for aromatic para-fluorine; both observations are in good agreement with results known from aryl iodine fluorides [3].

Reaction of  $\text{C}_6\text{F}_5\text{BrF}_2$  with  $[\text{CF}_3\text{C}(\text{O})]_2\text{O}$  yields  $\text{C}_6\text{F}_5\text{Br}[\text{O}(\text{O})\text{CCF}_3]_2$  - the first carboxylate of positive bromine stable above room temperature.

$\text{C}_6\text{F}_5\text{BrF}_2$  is a good source for the preparation of symmetric and asymmetric bromonium-salts:  $[\text{Ar}_2\text{Br}]^+\text{X}^-$  or  $[\text{ArAr}'\text{Br}]^+\text{X}^-$ .

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