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ALKYLIDENE SULFUR TETRAFLUORIDES AND ALKYL TETRAFLUOROSULFURANONIUM IONS

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The preparation of $\text{H}_2\text{C}=\text{SF}_4$, $\text{CH}_3\text{-CH}=\text{SF}_4$, $\text{CF}_3\text{-CH}=\text{SF}_4$, $\text{CF}_3(\text{CH}_3)\text{C}=\text{SF}_4$ and other alkylidene sulfur tetrafluorides is presented.

The structure of these materials are all alike: trigonal bipyramidal coordination of sulfur, the double bonded carbon occupying an equatorial position, and the carbon substituents (H, CH_3 , CF_3) being fixed in the axial plane. Furthermore, all such compounds turn out to be rigid. The CS bond is regarded as a true double bond. Reactions of these compounds are either the cleavage into SF_4 and the carbene $\text{RR}'\text{C}$, or simple cis-addition across the double bond.

The alkyl-tetrafluoro-sulfuranonium ions are only transient species, and are detected with ^1H and ^{19}F nmr spectroscopy at low temperatures. They are formed by fluoride ion abstraction from R-SF_5 compounds. Their ease of formation and stability are discussed as a function of R, the solvent, and the fluoride ion abstractor.