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Note

Chlorotrifluoroethylene 에 브롬화 수소의 부가반응. 소량 생성물들의 확인

박 달 조*·J. P. McClure. J. R. Lacher

미국 콜로라도대학교 **화학과** (1973, 1. 18 접수)

The Addition of Hydrogen Bromide to Chlorotrifluoroethylene. The Identification of the Minor Products

Joshep D. Park*, J. P. McClure and J. R. Lacher

Department of Chemistry, University of Colorado, Boulder, Colorado, U.S.A. (Received Jan. 18, 1973)

The catalytic addition of hydrogen bromide to chlorotrifluoroethylene has been previously reported in the ltierature¹⁻⁴ and the only product reported is 1-bromo-2-chloro-1, 1, 2-trifluoroethane. The present work reports the isolation and identification of the following products, 1, 2-dichloro-2-bromo-1, 1, 2-trifluoroethane, 1, 2dibromo-1, 1, 2-trifluorethane and 1, 2-dibromo 2-chloro-1, 1, 2-trifluoroethane along with the major product 1-bromo-2-chloro-1, 1, 2-trifluoroethane (I). These minor products are all explainable on the basis of the operation of a free radical brought about by pyrolysis of the major product (I),

$$BrCF_2 - CFClH \rightarrow \begin{cases} \rightarrow \mathring{C}F_2 - CFCl + \mathring{B}r \\ \rightarrow \mathring{C}FH - CF_2Br + \mathring{C}l \\ \rightarrow \mathring{C}FCl - CF_2Br + \mathring{B}r \end{cases}$$

and the pyroloysis of $CF_2Br-CFClBr$ to yield. $CF_2-CFClBr$

Experimental

The steel catalyst chamber filled with 16-mesh activated carbon is heated to 175° and a large excess of olefin (6 parts) to hydrogen bromide (1 part) is used. The flow rate of trifluorochloroethylene is kept at about 40-45 g/hr. Over 3000 g. of product was collected and subjected

I CF ₂ Br-CFClH ^{1,2}	% by weight b. p. /630mm. Hg			
	92.5		46°	·
II CF ₂ Cl-CFClBr ⁵	0. 5; n ^e n ^s	1. 3920 1. 910	66°	
III CF ₂ Br-CFBrH ³	2.4		66. 5°	
IV CF ₂ Br-CFClBr ⁶	2.7		86. 5	
V $CF_2 = CFCl$	1.9			

*Present Address; The Korea Advanced Institute of Science, Seoul, Korea

Chlorotrifluoroethylene에 브롬화 수소와 부가반응, 소량 생성물들의 확인

to fractionation. The product distribution was

References

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- 1. U.S. Patent 2, 363, 516 assigned to duPont
- 2. J. D. Park, M. L. Sharrah and J. R. Lacher, J. Amer. Chem. Soc., 71, 2340 (1949).
- 3. J. D. Park, W. R. Lycan and J. R. Lacher, J. Amer. Chem. Soc., 73, 711 (1951).
- 4. R. N. Haszeldine and B. R. Steele, J. Chem. Soc., 3747 (1954).
- 5. R. W. Lamb, Ph. D. Thesis, University of Colorado, 1961
- 6. R. N. Haszeldine, J. Chem. Soc., 4423 (1952).

as follows: Compound I, II, III and IV were identified

by comparison of their physical properties and infrared spectra with authentic known samples.

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