General Anesthetics. 1. Halogenated Methyl Ethyl Ethers as Anesthetic Agents

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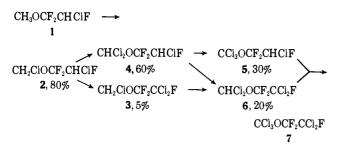
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Thirty-six halogenated Me Et ethers have been synthesized for evaluation as volatile general anesthetics. Eleven of the ethers were too unstable to test, and, of the remaining 25, 13 had promising anesthetic properties in mice and are suitable for study in larger animals. Those ethers having one H with at least 2 halogens other than F or 2 or more H with at least one Br or one Cl were the best anesthetics.

The first study of the anesthetic properties of fluorinated hydrocarbons was reported by Robbins¹ in 1946. Since that time many fluorinated compounds, both hydrocarbons and ethers, have been found to have anesthetic properties in laboratory animals, and several have progressed to clinical trials in humans.^{2,3} Three are presently in clinical use: fluroxene, $CF_3CH_2OCH=$ CH_2 ; halothane, $CF_3CHClBr$; and methoxyflurane, $CH_3OCF_2CHCl_2$.

In an attempt to find a new agent superior to these three we have synthesized 36 new halogenated Me Et ethers by photochlorination and thermal bromination; 25 of these have been evaluated as anesthetics in mice (Table I). The remaining **11** were too unstable to test.

Synthesis.—All compounds were synthesized by photochlorination or thermal bromination of 9 fluorinated Me Et ethers. Chlorination of CH_3OCF_2CHClF (1) following the published procedure^{4,5} gave mixtures of 6 chlorination products resulting from simultaneous chlorination of starting material and reaction products as outlined in the following equation. The percentage figures given represent the maximum percentages found in any chlorination mixture. The maximum amount of monochloro product was formed after reaction of about 1 mole of Cl_2 , the maximum amounts of dichloro products after reaction of about 2 moles of Cl_2 , and so forth. Yields of perhalogenated products were above 95%.



These results do not agree with those reported by Park^{4,5} who described the reaction as "stepwise" and reported that the H on the CHClF carbon was the last to be replaced.

Similar results were found in the chlorination of both CH_3OCF_2CHBrF (8) and $CH_3OCF_2CHCl_2$ (14) except that the H of the OCF_2CHCl_2 group was more reactive

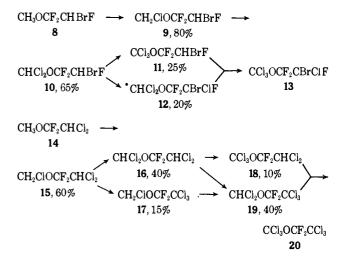
(I) B. H. Robbins, J. Pharmacol. Exp. Ther., 86, 197 (1946).

- (2) J. C. Krantz, Jr., and F. G. Rudo, in "Handbook of Experimental Pharmacology," Vol. XX/1. O. Eichler, A. Farah, H. Herken, and A. D. Welch, Ed., Springer, Berlin, 1966, pp 501-564.
- (3) E. R. Larsen, Fluorine Chem. Rev., 3, 1 (1969).

(4) J. D. Park, D. M. Griffin, and J. R. Lacher, J. Amer. Chem. Soc., 74, 2293 (1952).

(5) J. D. Park, B. Stricklin, and J. R. Lacher, ibid., 76, 1387 (1954).

and larger amounts of $CH_2ClOCF_2CCl_3$ (17) (corresponding to 3 were formed.



The reactivity of the CF_2CHCl_2 group compared to CF_2CHClF or CF_2CHBrF shows that chlorination is directed away from a C having an F substituent as well as away from a C adjacent to a F-substituted C as reported by Park.^{4,5} This directive influence of F on the same C is also clearly shown in the chlorination of $CH_3OCF_2CHF_2$ (21) where the H on the CHF_2 group reacts very slowly and good conversions into the 3 products chlorinated on Me are found.

$$\begin{array}{c} \mathrm{CH}_{3}\mathrm{OCF}_{2}\mathrm{CHF}_{2} \longrightarrow \mathrm{CH}_{2}\mathrm{CloCF}_{2}\mathrm{CHF}_{2} \longrightarrow \\ \mathbf{21} \qquad \mathbf{22}, \ 80\% \\ \mathrm{CHCl}_{2}\mathrm{OCF}_{2}\mathrm{CHF}_{2} \longrightarrow \mathrm{CCl}_{3}\mathrm{OCF}_{2}\mathrm{CHF}_{2} \longrightarrow \\ \mathbf{23}, \ 80\% \qquad \mathbf{24}, \ 80\% \\ \mathrm{CCl}_{3}\mathrm{OCF}_{2}\mathrm{CClF}_{2} \\ \mathbf{25} \end{array}$$

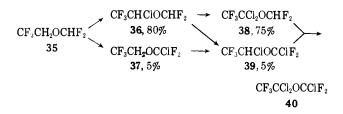
Additional examples of the directive influence of F substitution on the same C and on an adjacent C are found in the chlorination of $CF_3CH_2OCH_3$ (29) where CH_3O is more reactive than OCH_2CF_3 and good yields of $CHCl_2OCH_2CF_3$ (31) are formed and in the chlorination of $CF_3CH_2OHCF_2$ (35) where the $OCHF_2$ group is much less reactive than the CH_2 group adjacent to CF_3 and good yields of $CF_3CHClOCHF_2$ (36) and CF_3 - CCl_2OCHF_2 (38) are formed.

$$\begin{array}{cccc} \mathrm{CF_3CH_2OCH_3} & \longrightarrow & \mathrm{CF_3CH_2OCH_2Cl} & \longrightarrow \\ & & & & & & \\ \mathbf{29} & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & &$$

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$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	1	$CH_3OCF_2CHClF^a$	70	1.3338		Light anesthesia at 2.5%	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	2	CH ₂ ClOCF ₂ CHClF ^c	110	1.3740		Too unstable to test	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		$CH_2ClOCF_2CCl_2F$	120	1.3885	C, H; umr	Deep anesthesia at 1.5%	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		CHCl ₂ OCF ₂ CHClF ^c	118	1.3872			
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		$\mathrm{CCl}_3\mathrm{OCF}_2\mathrm{CHClF}^c$	62 (50)			Anesthetic and convulsant at	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	6	$CHCl_2OCF_2CCl_2F$	53(50)	1.3982	C, H, F; umr		
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		CHF ₂ CH ₂ OCHF ₂	55	1.2966		Very weak anesthetic at 5%	
		CHF ₂ CHClOCHF ₂	74	1.3351	С, Н, F	Anesthetic at 1.5%	
	50	$\rm CF_3CHBrCl^{e}$	50.2			Good anesthetic at 2.5%	
51 $CF_3CH_4OCH=CH_2^e$ 43.7 Good anesthetic at 7.5%	51	CF ₃ CH ₂ OCH=CH ₂ ^e	43.7			Good anesthetic at 7.5%	

TABLE I PROPERTIES OF HALOGENATED METHYL ETHYL ETHERS

^a J. D. Park, D. K. Vail, K. R. Lea, and J. R. Lacher, J. Amer. Chem. Soc., **70**, 1550 (1948). ^b Microanalyses were done for the elements indicated and all results were within $\pm 0.4\%$ of the theoretical values. Nmr spectra were recorded where indicated. All spectra were routine and were consistent with the assigned structures. ^c J. D. Park, D. M. Griffin, and J. R. Lacher, J. Amer. Chem. Soc., **74**, 2293 (1952); J. D. Park, B. Stricklin, and J. R. Lacher, *ibid.*, **76**, 1387 (1954). ^d A. Demiel, J. Org. Chem., **25**, 993 (1960). ^e Reference standard for comparison. ^d W. T. Miller, E. W. Fager, and P. H. Griswold, J. Amer. Chem. Soc., **70**, 431 (1948). ^e A. Van Poznak and J. F. Artusio, Jr., *Toxicol. Appl. Pharmacol.*, **2**, 374 (1960). ^b A. L. Henne and M. A. Smook, J. Amer. Chem. Soc., **72**, 4378 (1950). ^d R. C. Terrell, U. S. Patent 3,469,011 (to Air Reduction Co., Inc., New York, N. Y.), Sept 23, 1969 [Chem. Abstr., **72**, 3025 (1970)].



The OCHF₂ group is also less reactive than OCF₂-CHClF or OCF₂CHF₂ since chlorination of **42** and **45** gives good yields of the β -substitution products.

The thermal bromination of both 42 and 45 gives the corresponding bromo ethers 44 and 47, but in lower yields, and the bromination of 35 gave 41 in low yield with many degradation products resulting from cleav-

age of the C–O bonds. In none of the bromination reactions was any evidence of bromination of the $\mathrm{CHF}_2\mathrm{O}$ group found.

 $\begin{array}{ccc} \mathrm{CHF_2OCF_2CHF_2} &\longrightarrow \mathrm{CHF_2OCF_2CBrF_2} \\ & \mathbf{42} & \mathbf{44} \ (25\% \ \mathrm{conversion}, \ 60\% \ \mathrm{yield}) \\ \mathrm{CHF_2OCF_2CHClF} &\longrightarrow \mathrm{CHF_2OCF_2CBrClF} \\ & \mathbf{45} & \mathbf{47} \ (60\% \ \mathrm{conversion}, \ 70\% \ \mathrm{yield}) \\ \mathrm{CF_3CH_2OCHF_2} &\longrightarrow \mathrm{CF_3CHBrOCHF_2} \\ & \mathbf{35} & \mathbf{41} \ (15\% \ \mathrm{yield}) \end{array}$

Pharmacology.—All compounds which were stable were evaluated as anesthetic agents in mice. In general, those ethers having OCH₂Cl, OCHCl₂, or OCCl₃ groups were the least stable although some could be stabilized sufficiently to permit testing by the addition of K₂CO₃. Eleven new compounds (9, 10, 16, 17, 18, 19, 27, 28, 30, 32, and 33) and 4 known compounds were too unstable to test (2, 4, 15, 22).

The anesthetic properties of 9 compounds (1, 7, 8, 14, 26, 29, 35) plus CF₃CHBrCl and CF₃CH₂OCH=CH₂ have been reported previously^{1-3,6} and testing of these compounds was repeated for comparison with the new compounds. Our screening results were in good agreement with those reported.

About one-half of the compounds screened had good anesthetic activity in mice with the remainder about equally divided between very weak or inert compounds and convulsants.

In general, it was necessary to have at least one H present in order to have anesthetic activity as all the perhalogenated compounds (7, 13, 20, 25, 40) were weak anesthetics or convulsants. These results are in agreement with the observations of Krantz.²

Those compounds which had one H with at least 2 halogens other than F or 2 or more H with at least one Br or one Cl were the best anesthetics. Among these were 14 new compounds (3, 11, 12, 23, 24, 39, 46, 31, 36, 38, 41, 45, 47, 49), and all, except 31 which is irritating, are suitable for study in larger animals. Some of this work has been reported⁶⁻⁸ and other studies are in progress.

(6) R. W. Virtue, I., O. Lund, McK. Phelps, Jr., J. H. K. Vogel, H. Beckivitt, and M. Heron, Can. Anaesth. Soc. J., **13**, 233 (1966).

(7) A. B. Dobkin, R. G. Heinrich, J. S. Israel, A. A. Levy, J. F. Neville, Jr., and K. Ounkasen, Anesthesiology, 29, 275 (1968).

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Experimental Section

Pharmacology.—All compds screened were routinely checked for purity by gas chromatography and all were 99.5% pure or better.

All screening was done using mice, and 6, 11, 12, 13, 20, 29, and 34 were administered by ip injection as 0.6 M emulsions.⁹ The remainder were administered by inhalation in admixture with O_2 .⁹ All pharmacology was done by J. C. Krantz, Jr., F. G. Rudo, and H. F. Cascorbi at the Department of Pharmacology, University of Maryland School of Medicine, Baltimore, Md., and The Huntingdon Research Center, Inc., Baltimore, Md., and A. B. Dobkin and P. H. Byles at the Department of Anesthesiology, State University of New York, Upstate Medical Center, Syracuse, N. Y.

Synthesis.—Boiling points were determined by distn or by the Siwoloboff method and are uncorr. Nmr spectra were detd in CCl₄ (Me₄Si) using a Varian A-60 spectrophotometer.

Compds 1, 8, 14, 21, 29, 42, and 45 were prepared according to published procedures, referenced in Table I.

 $CH_3OCF_2CHBrCl$ (26).—Na (75 g) was dissolved in abs MeOH (700 g), and CF₃CHBrCl³ (540 g) was added slowly. After refluxing for 24 hr, the mixt was poured into H₂O (3 l.). The crude product (500 g) was fractionated to yield unreacted material (200 g), CH₃OCF₂CHBrCl (125 g), bp 68° (100 mm), and (CH₃O)₃CCHBrCl (125 g), bp 86° (10 mm), $n^{20}D$ 1.4727. Anal. (C₃H₁₀BrClO₃) C, H.

CF₃CH₂OCHF₂ (35) and CHF₂OCH₂CHF₂ (48).-Into the stainless steel liner of a 1-l. autoclave was placed a solu of 60 g (1 mole) of KOH pellets in 374 g (3.7 moles) of CF₃CH₂OH and 26 g (1.4 moles) of $H_2()$. The autoclave was then sealed and while being stirred and heated at 80-95°, 60 ml (0.8 mole) of liquefied CHClF₂ was added in increments to reach an antogenous pressure of 8.08 kg/cm². The autoclave was maintained at approx 90° for 2 hr after the addition of the reactants and was then cooled. The gases from the autoclave were vented through a Dry-Ice trap condensing 13 g of liq. The contents of the liner were distd to give 49 g of crude CF3CH2OCHF2 (bp 27-40°) plus an additional 27 g collected in a Dry-Ice trap connected to the still. Low-temp distu of the combined 40 g from the Dry-Ice traps gave 24 g of recovered diffuorochloromethane and 16 g more of CF₃CH₂OCHF₂. The combined product, 65 g, represents a 56.5% conversion of CHClF₂.

 $CHF_2OCH_2CHF_2$ was synthesized from CHF_2CH_2OH in the same way with about 25% conversion.

Chlorination of Ethers.—All chlorinations were done by the method of Park.^{4,5} The amount of Cl_2 was estimated by titration of the effluent HCl gas which was scrubbed into H₂O. Reaction mixts were analyzed using an F and M Model 202 chroniatograph with thermistor detectors and a 3-m 10% nonylphenoxy-(poly(ethylenoxy))ethanol, 15% Ucon LB-550-X on "Chromosorb" column, or a Wilkins aerograph Model A-350 with a hot wire detector and a 2-m Se 30 column. He flow was 50 cm³/min. Products were isolated by fractional distu, prep chroniatography using a Wilkins "autoprep" chromatograph, or a combination of the two.

Bromination of Ethers.—Brominations were done by passing a stream of N_2 into a mixt of Br_2 and the ether and then through a 30×2.5 cm glass tube at $450-475^\circ$. Products were collected in a Dry-Ice trap and purified by distn or gas chromatography.

(9) H. F. Cascorbi and F. G. Rudo, Anesth. Analg. (Cleveland), 43, 163 (1964).