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ELECTROCHEMICAL FLUORINATION OF SOME CYCLIC TERTIARY AMINES

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In the course of our research concerning the preparation of perfluorinated compounds suitable as blood substitutes, we have studied the electrochemical fluorination of $\underline{N}, \underline{N}$ -diethylcyclo-hexylamine (DECHA) and \underline{N} -ethyldicyclohexylamine (EDCHA) using the classical equipment of Simons. The two compounds behaved in a different way:

1) DECHA showed a process yield (based on the theoretical amount of the corresponding perfluoroamine) of about 51%, while the yield for EDCHA was about 16%;

2) in the case of EDCHA the reaction mixture was more complex and no identification of its components was possible; for DECHA the more significant products of the reaction, corresponding to about 65% of the total amount, were identified as:

$C_{6}F_{11}N(C_{2}F_{5})_{2}$ (molar yield: 13.8%);	$n-C_{6}F_{13}N(C_{2}F_{5})_{2}$	(9.5%);
$C_{6}F_{11}N(CF_{3})C_{2}F_{5}$ (5.3%);	C $_{6}$ F $_{10}$ HN (C $_{2}$ F $_{5}$) $_{2}$	(4.4%);
$n-C_{6}F_{1_{3}}N(CF_{3})C_{2}F_{5}$ (4.7%);	C ₆ F ₁₀ HN(CF ₃)C ₂ F ₅	; (0.9%).

Therefore the following remarks can be made:

- a more complex structure of the molecule undergoing the electrochemical process produced lower yields, due to a greater extent of destructive side reactions;
- fragmentation reactions producing ring opening were more important with respect to those involving linear chains;
- 3) hydrogen atoms belonging to carbon linked to a nitrogen atom were the last to be substituted, in accordance with the EC_bEC_N mechanism proposed for the electrochemical fluorination of organic compounds.