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Swarts fluorination

Swarts fluorination is a process whereby the <u>chlorine</u> <u>atoms</u> in a compound – generally an organic compound, but experiments have been performed using <u>silanes</u> – are replaced with <u>fluorine</u>, by treatment with <u>antimony trifluoride</u> in the presence of chlorine or of <u>antimony pentachloride</u>. The active species is antimony trifluorodichloride, which is produced <u>in situ</u>; this compound can also be produced in bulk, according to a patent of John Weaver.^[1]

Swarts fluorination	
Named after	Frédéric Jean Edmond Swarts
Reaction type	Substitution reaction

The process was initially described by Frédéric Jean Edmond Swarts in 1892.^[2]

References

- 1. "Preparation of antimony trifluorodichloride and fluorination of fluorinatable hydrocarbons and halocarbons therewith Patent # 4438088 PatentGenius" (http://www.patentgenius.com/patent/4438088.html). www.patentgenius.com.
- 2. Acad. Roy. Belg 3(24) p.474 (1892)

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