

K781917 MUCUS ASPIRATORDec 4, 1978
21 days to decisionK781917 · Product code: **FMF** · General HospitalSource: <https://www.510kdatabase.net/k781917/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 13, 1978
Decision date	Dec 4, 1978
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Ovutime, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k781917/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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