FDA 510(k) Premarket Notification

K781917 MUCUS ASPIRATOR

CLEARED

Dec 4, 1978
21 days to decision

K781917 · Product code: **FMF** · General Hospital
Source: 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