FDA 510(k) Premarket Notification

K833399 SYRINGE

CLEARED

Jan 10, 1984
97 days to decision

K833399 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k833399/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1983
Decision date	Jan 10, 1984
Days to decision	97 days
Third-party review	No

APPLICANT

Company	**I M, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1976-1992