FDA 510(k) Premarket Notification

K894335 MODEL #535 PACU STRETCHER/BED & #530 ED/TRAUMA STR

CLEARED

Sep 12, 1989
60 days to decision

K894335 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k894335/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 14, 1989
Decision date	Sep 12, 1989
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Midmark Corp.**
Location	Versailles, OH, US
Contact	JOHN OLDIGES
510(k) history	31 submissions · 31 cleared · 1981-2013