FDA 510(k) Premarket Notification

K801839 DISPOSABLE HYPODERMIC NEEDLES

CLEARED

Sep 16, 1980
46 days to decision

K801839 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k801839/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Aug 1, 1980
Decision date	Sep 16, 1980
Days to decision	46 days
Third-party review	No

APPLICANT

Company	**Martin'S Maine Exchange**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1981