FDA 510(k) Premarket Notification

K861081 DISPOSABLE HYPODERMIC NEEDLE SAFETY SHEATH

CLEARED

Apr 9, 1986
19 days to decision

K861081 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k861081/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 21, 1986
Decision date	Apr 9, 1986
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Perkins & Perkins, Inc.**
Location	Potomac, MD, US
Contact	JERRI BARDEN-PERKINS
510(k) history	1 submissions · 1 cleared · 1986-1986