

K861081 DISPOSABLE HYPODERMIC NEEDLE SAFETY SHEATHApr 9, 1986
19 days to decisionK861081 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k861081/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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