FDA 510(k) Premarket Notification

K874024 ACU-MICRO-NEEDLE

CLEARED

Nov 23, 1987
52 days to decision

K874024 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k874024/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 2, 1987
Decision date	Nov 23, 1987
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Acuderm, Inc.**
Location	Mchenry, IL, US
Contact	CHARLES R YEH
510(k) history	13 submissions · 13 cleared · 1983-1994