

K874024 ACU-MICRO-NEEDLENov 23, 1987
52 days to decisionK874024 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k874024/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026