

K881866 NEEDLE REMOVERNov 14, 1988
195 days to decisionK881866 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k881866/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	May 3, 1988
Decision date	Nov 14, 1988
Days to decision	195 days
Third-party review	No

APPLICANT

Company	Howard, Hills & Harsh
Location	Salt Lake City, UT, US
Contact	LINDA R HILLS
510(k) history	2 submissions · 2 cleared · 1988-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881866/>; 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 July 8, 2026