

K895769 EXEL HUBER NEEDLEFeb 16, 1990
143 days to decisionK895769 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k895769/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 26, 1989
Decision date	Feb 16, 1990
Days to decision	143 days
Third-party review	No

APPLICANT

Company	Exel Intl.
Location	Culver City, CA, US
Contact	ESHAGH HAMID
510(k) history	18 submissions · 18 cleared · 1986-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895769/>; 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 June 30, 2026