FDA 510(k) Premarket Notification

K895769 EXEL HUBER NEEDLE

CLEARED

Feb 16, 1990
143 days to decision

K895769 · 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