FDA 510(k) Premarket Notification

K900857 NON STERILE PARENTERAL SUPPLY KIT

CLEARED

May 22, 1990
88 days to decision

K900857 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k900857/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 23, 1990
Decision date	May 22, 1990
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Intermed, Inc.**
Location	Mchenry, IL, US
Contact	NORM RUEDT
510(k) history	16 submissions · 15 cleared · 1976-1990