FDA 510(k) Premarket Notification

K900282 NEEDLE-SHIELD

CLEARED

Feb 28, 1990
38 days to decision

K900282 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k900282/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 21, 1990
Decision date	Feb 28, 1990
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Macbrud Corp. Medical Div.**
Location	Miami, FL, US
Contact	GRAY LARY
510(k) history	4 submissions · 4 cleared · 1989-1991