FDA 510(k) Premarket Notification

K905305 SAFE-SITE NEEDLE COVER

CLEARED

Feb 12, 1991
76 days to decision

K905305 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k905305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Nov 28, 1990
Decision date	Feb 12, 1991
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**North American Medical Products, Inc.**
Location	Walker, MI, US
Contact	ARTHUR GIANAKOS
510(k) history	6 submissions · 6 cleared · 1984-2001