FDA 510(k) Premarket Notification

K020921 NO-POKES NEEDLE SAFETY DEVICE

CLEARED

Jul 11, 2002
112 days to decision

K020921 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k020921/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 21, 2002
Decision date	Jul 11, 2002
Days to decision	112 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bemis Mfg. Co.**
Location	Sheboygan Falls, WI, US
Contact	JOHN B HOWELL
510(k) history	12 submissions · 12 cleared · 1985-2002