FDA 510(k) Premarket Notification

K960698 NON-CORING NEEDLE

CLEARED

Apr 15, 1996
55 days to decision

K960698 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k960698/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Feb 20, 1996
Decision date	Apr 15, 1996
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Med Institute, Inc.**
Location	West Lafayette, IN, US
Contact	NEAL E FEARNOT, PH.D., E.E.
510(k) history	26 submissions · 23 cleared · 1990-2000