FDA 510(k) Premarket Notification

K954189 EASI-JECTOR CONTROL SYSTEM

CLEARED

Oct 19, 1995
43 days to decision

K954189 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k954189/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 6, 1995
Decision date	Oct 19, 1995
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**North, Inc.**
Location	Northfield, MN, US
Contact	JIM POKORNEY
510(k) history	2 submissions · 2 cleared · 1995-1997