FDA 510(k) Premarket Notification

K945275 PRO-TEC DISPOSBLE SHARPS CONTAINERS Q1

CLEARED

Dec 14, 1994
47 days to decision

K945275 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k945275/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 28, 1994
Decision date	Dec 14, 1994
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pro-Tec Containers, Inc.**
Location	Irvine, CA, US
Contact	TREESA SPENCER
510(k) history	6 submissions · 6 cleared · 1989-1994