FDA 510(k) Premarket Notification

K945273 PRO-TECH DISPOSABLE SHARPS CONTAINERS VNC2

CLEARED

Dec 14, 1994
47 days to decision

K945273 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k945273/

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