FDA 510(k) Premarket Notification

K000714 VACUFLOW+SAFE BLOOD COLLECTION SET AND VACUFLOW+SAFE WITH HOLDER BLOOD COLLECTION SET

CLEARED

Dec 21, 2000
294 days to decision

K000714 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k000714/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 2, 2000
Decision date	Dec 21, 2000
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Med-Pro Technologies, Inc.**
Location	Appollo Beach, FL, US
Contact	ART WARD
510(k) history	1 submissions · 1 cleared · 2000-2000