FDA 510(k) Premarket Notification

K960933 BIO-PLEXUS NEEDLE DISPOSAL CONTAINER

CLEARED

Aug 28, 1996
174 days to decision

K960933 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k960933/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 7, 1996
Decision date	Aug 28, 1996
Days to decision	174 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bio-Plexus, Inc.**
Location	Tolland, CT, US
Contact	CARL SAHI
510(k) history	6 submissions · 6 cleared · 1990-2001