

K960933 BIO-PLEXUS NEEDLE DISPOSAL CONTAINERAug 28, 1996
174 days to decisionK960933 · Product code: **FMI** · General HospitalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960933/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026