FDA 510(k) Premarket Notification

K925875 HORIZONTAL NEEDLE GUIDE ATTACHMENT

CLEARED

Apr 6, 1993
138 days to decision

K925875 · Product code: **GDF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k925875/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Needle, Surgical (GDF)
Date received	Nov 19, 1992
Decision date	Apr 6, 1993
Days to decision	138 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bip USA, Inc.**
Location	Niagara Falls, NY, US
Contact	GARY HORNER
510(k) history	7 submissions · 7 cleared · 1993-2003