FDA 510(k) Premarket Notification

K923840 NEEDLE-GRABBER

CLEARED

Jul 13, 1993
347 days to decision

K923840 · Product code: **GDF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k923840/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Needle, Surgical (GDF)
Date received	Jul 31, 1992
Decision date	Jul 13, 1993
Days to decision	347 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Guardian Angel Products, Inc.**
Location	Huntingdon Valley, PA, US
Contact	HOWARD M HOLSTEIN
510(k) history	5 submissions · 5 cleared · 1990-2003