

K923840 NEEDLE-GRABBERJul 13, 1993
347 days to decisionK923840 · Product code: **GDF** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923840/> 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 July 3, 2026