

**K925875 HORIZONTAL NEEDLE GUIDE ATTACHMENT**Apr 6, 1993  
138 days to decisionK925875 · Product code: **GDF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k925875/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k925875/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026