

**K900903 KOPANS HOOKWIRE LESION LOCALIZATION  
NEEDLE**May 3, 1990  
65 days to decisionK900903 · Product code: **GDF** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k900903/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Needle, Surgical (GDF)
Date received	Feb 27, 1990
Decision date	May 3, 1990
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Milex Products, Inc.</b>
Location	Mchenry, IL, US
Contact	ROBERT SHAW
510(k) history	8 submissions · 8 cleared · 1976-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900903/>; 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 29, 2026