

**K903079 ENDOGUIDE (R), MICROGUIDE (TM), FLEXIGUIDE (TM)**Oct 4, 1990  
84 days to decisionK903079 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k903079/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 12, 1990
Decision date	Oct 4, 1990
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Luxar Corp.</b>
Location	Bothell, WA, US
Contact	KATHERINE LAAKMANN
510(k) history	17 submissions · 17 cleared · 1988-1996

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903079/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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