

**K931070 CLOSED END FIBEROPTIC LASER DELIVERY SYSTEM**Oct 5, 1993  
217 days to decisionK931070 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k931070/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 2, 1993
Decision date	Oct 5, 1993
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lca, Inc.</b>
Location	Mountain View, CA, US
Contact	MICHAEL D JOHNSON
510(k) history	12 submissions · 12 cleared · 1990-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931070/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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