FDA 510(k) Premarket Notification

K931070 CLOSED END FIBEROPTIC LASER DELIVERY SYSTEM

CLEARED

Oct 5, 1993
217 days to decision

K931070 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k931070/

SUBMISSION DETAILS

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

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