

**K001211 MODIFICATION TO INTRALASE 600C LASER
KERATOME**

Dec 4, 2000
234 days to decision

K001211 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k001211/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Apr 14, 2000
Decision date	Dec 4, 2000
Days to decision	234 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Intra Lase Corp.
Location	Irvine, CA, US
Contact	CHARLINE GAUTHIER
510(k) history	9 submissions · 9 cleared · 1999-2008

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k001211/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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