

K895559 MODIFIED ENDOGUIDEOct 20, 1989
49 days to decisionK895559 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k895559/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Powered Laser Surgical Instrument (GEX) |
| Date received | Sep 1, 1989 |
| Decision date | Oct 20, 1989 |
| Days to decision | 49 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Luxar Corp. |
| Location | Bothell, WA, US |
| Contact | D LAAKMANN |
| 510(k) history | 17 submissions · 17 cleared · 1988-1996 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895559/>; Data sourced from FDA 510(k) public records (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. - Generated July 5, 2026