

K913758 KTP/532 AND KTP/YAGNov 20, 1991
90 days to decisionK913758 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k913758/>**SUBMISSION DETAILS**

| | |
|-----------------------|-----------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Powered Laser Surgical Instrument (GEX) |
| Date received | Aug 22, 1991 |
| Decision date | Nov 20, 1991 |
| Days to decision | 90 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|-----------------------------------------|
| Company | Laserscope |
| Location | Santa Clara, CA, US |
| Contact | DONNA PAGE |
| 510(k) history | 60 submissions · 60 cleared · 1985-2006 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913758/> 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 June 29, 2026