FDA 510(k) Premarket Notification

# K040289 SMARTLITE D FREQUENCY DOUBLED ND:YAG LASER

**CLEARED**

Apr 30, 2004
84 days to decision

K040289 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k040289/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Powered Laser Surgical Instrument (GEX) |
| Date received | Feb 6, 2004 |
| Decision date | Apr 30, 2004 |
| Days to decision | 84 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Cynosure, Inc.** |
| Location | Bedford, MA, US |
| Contact | GEORGE CHO |
| 510(k) history | 98 submissions · 98 cleared · 1992-2019 |