

K040289 SMARTLITE D FREQUENCY DOUBLED ND:YAG LASERApr 30, 2004
84 days to decisionK040289 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040289/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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