FDA 510(k) Premarket Notification

K080121 CYNOSURE SMARTLIPO MULTIWAVLENGTH LASER

CLEARED

Jan 31, 2008
14 days to decision

K080121 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k080121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 17, 2008
Decision date	Jan 31, 2008
Days to decision	14 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