FDA 510(k) Premarket Notification

K992765 CYNOSURE DIODERM

CLEARED

Sep 8, 1999
22 days to decision

K992765 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k992765/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 17, 1999
Decision date	Sep 8, 1999
Days to decision	22 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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