CLEARED

K021724 DORNIER DIODE LASER FAMILY, INCLUDING MEDILAS D FIBERTOM, MEDILAS D SKINPULSE & MEDILAS D SKINPULSE S LASER SYSTEMS

Aug 19, 2002
87 days to decision

K021724 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k021724/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 24, 2002
Decision date	Aug 19, 2002
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dornier Medtech America, Inc.**
Location	Marietta, GA, US
Contact	Tim Thomas
510(k) history	40 submissions · 40 cleared · 1990-2023