

**K021724 DORNIER DIODE LASER FAMILY, INCLUDING  
MEDILAS D FIBERTOM, MEDILAS D SKINPULSE & MEDILAS D  
SKINPULSE S LASER SYSTEMS**Aug 19, 2002  
87 days to decisionK021724 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k021724/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021724/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026