

**K020339 DORNIER DIODE LASER FAMILY MEDILAS D,  
MEDILAS D SKINPULSE & MEDILAS D SKINPULSE S LASER  
SYSTEM**Oct 25, 2002  
266 days to decisionK020339 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k020339/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 1, 2002
Decision date	Oct 25, 2002
Days to decision	266 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/k020339/>; 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