

K110364 DORNIER MEDILAS D LITEBEAM+ 1470Dec 7, 2011
303 days to decisionK110364 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k110364/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 7, 2011
Decision date	Dec 7, 2011
Days to decision	303 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k110364/> Data sourced from FDA 510(k) public records (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. - Generated June 15, 2026