

**K020673 WAVE FORM MANUFACTURING PROLASE GENERAL
SHAPED FIBER**May 30, 2002
90 days to decisionK020673 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k020673/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 1, 2002
Decision date	May 30, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Wave Form Mfg., Inc.
Location	Panama City Beach, FL, US
Contact	JOE D BROWN
510(k) history	1 submissions · 1 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020673/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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