FDA 510(k) Premarket Notification

CLEARED

K020673 WAVE FORM MANUFACTURING PROLASE GENERAL SHAPED FIBER

May 30, 2002
90 days to decision

K020673 · Product code: **GEX** · General & Plastic Surgery
Source: 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