FDA 510(k) Premarket Notification

K922917 ENDOPROBES

CLEARED

Sep 28, 1992
103 days to decision

K922917 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k922917/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 17, 1992
Decision date	Sep 28, 1992
Days to decision	103 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Endooptiks, Inc.**
Location	Marlboro, NJ, US
Contact	MARTIN S KNOPF
510(k) history	5 submissions · 5 cleared · 1992-2009

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k922917/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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