FDA 510(k) Premarket Notification

K914520 DIOMED 25 SURGICAL DIODE LASER, UROLOGY APPLICAT

CLEARED

Jan 22, 1993
470 days to decision

K914520 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k914520/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 10, 1991
Decision date	Jan 22, 1993
Days to decision	470 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dio-Med Corp.**
Location	Mchenry, IL, US
Contact	ILKKA MANNONEN
510(k) history	7 submissions · 7 cleared · 1981-1994

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k914520/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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