

**K914521 DIOMED 25 SURGICAL DIODE LASER, GEN SURG
APPLICAT**Jan 15, 1993
463 days to decisionK914521 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k914521/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 10, 1991
Decision date	Jan 15, 1993
Days to decision	463 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.netDevice record: <https://www.510kdatabase.net/k914521/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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