FDA 510(k) Premarket Notification

K914521 DIOMED 25 SURGICAL DIODE LASER, GEN SURG APPLICAT

CLEARED

Jan 15, 1993
463 days to decision

K914521 · 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