FDA 510(k) Premarket Notification

K914519 DIOMED 25 SURGICAL DIODE LASER, GYNECOLOGY APPLIC

CLEARED

Sep 16, 1993
707 days to decision

K914519 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k914519/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 10, 1991
Decision date	Sep 16, 1993
Days to decision	707 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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