FDA 510(k) Premarket Notification

K952340 MEGABEAM ASPIRATING ENDOCULAR PROBE

CLEARED

Jun 12, 1995
24 days to decision

K952340 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k952340/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 19, 1995
Decision date	Jun 12, 1995
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ceram Optec, Inc.**
Location	Washington, DC, US
Contact	JONATHAN S KAHAN
510(k) history	30 submissions · 30 cleared · 1992-2000