FDA 510(k) Premarket Notification

K944626 MODEL 210-AS ASPIRATING ENDOPHOTOCOAGULATOR LASER

CLEARED

Nov 3, 1994
44 days to decision

K944626 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k944626/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 20, 1994
Decision date	Nov 3, 1994
Days to decision	44 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mnlase, Inc.**
Location	Minneapolis, MN, US
Contact	BRUCE HEYMANN
510(k) history	22 submissions · 22 cleared · 1992-1994