FDA 510(k) Premarket Notification

K990725 NUVOLASE 532 LASER SYSTEM

CLEARED

Mar 22, 1999
17 days to decision

K990725 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k990725/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 5, 1999
Decision date	Mar 22, 1999
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**American Laser Corp.**
Location	Mchenry, IL, US
Contact	DANIEL HOFFER
510(k) history	5 submissions · 5 cleared · 1982-2001