FDA 510(k) Premarket Notification

K022266 AURORA SR

CLEARED

Oct 3, 2002
83 days to decision

K022266 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k022266/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 12, 2002
Decision date	Oct 3, 2002
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Syneron Medical, Ltd.**
Location	Yokneam Elite, IL
Contact	AMIR WALDMAN
510(k) history	35 submissions · 35 cleared · 2002-2017