FDA 510(k) Premarket Notification

K940173 CANDELA Q-SWITCHED ALEXANDRITE LASER

CLEARED

Dec 7, 1994
330 days to decision

K940173 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k940173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 11, 1994
Decision date	Dec 7, 1994
Days to decision	330 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Candela Laser Corp.**
Location	Wayland, MA, US
Contact	THOMAS R VERRICCHIONE
510(k) history	43 submissions · 43 cleared · 1988-1996