FDA 510(k) Premarket Notification

K013623 CLEARLIGHT PHOTOTHERAPHY SYSTEM, MODEL CL 420

CLEARED

Aug 16, 2002
284 days to decision

K013623 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k013623/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 5, 2001
Decision date	Aug 16, 2002
Days to decision	284 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Curelight, Ltd.**
Location	Washington, DC, US
Contact	JONATHAN S KAHAN
510(k) history	6 submissions · 6 cleared · 2002-2004