FDA 510(k) Premarket Notification

K020591 BCLEAR

CLEARED

May 16, 2002
83 days to decision

K020591 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k020591/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 22, 2002
Decision date	May 16, 2002
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumenis**
Location	Pleasanton, CA, US
Contact	C. ROBERT PAYNE JR.
Website	http://www.lumenis.com/
510(k) history	5 submissions · 5 cleared · 2001-2010