FDA 510(k) Premarket Notification

K170489 Skylit Phototherapy System

CLEARED

May 23, 2017
95 days to decision

K170489 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k170489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 17, 2017
Decision date	May 23, 2017
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Skylit Medical**
Location	La Jolla, CA, US
Contact	Martyn Gross
510(k) history	1 submissions · 1 cleared · 2017-2017