FDA 510(k) Premarket Notification

K181430 LENSAR Laser System - fs 3D (LLS-fs 3D)

CLEARED

Aug 9, 2018
69 days to decision

K181430 · Product code: **OOE** · Ophthalmic
Source: https://www.510kdatabase.net/k181430/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmic Femtosecond Laser (OOE)
Date received	Jun 1, 2018
Decision date	Aug 9, 2018
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lensar, Inc.**
Location	Winter Park, FL, US
Contact	Keith Peck
510(k) history	14 submissions · 14 cleared · 2010-2022