FDA 510(k) Premarket Notification

K122905 PEREGRINE 23GA CURVED LASER PROBE

CLEARED

Apr 30, 2013
221 days to decision

K122905 · Product code: **HQB** · Ophthalmic
Source: https://www.510kdatabase.net/k122905/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Photocoagulator And Accessories (HQB)
Date received	Sep 21, 2012
Decision date	Apr 30, 2013
Days to decision	221 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Peregrine Surgical , Ltd.**
Location	Doylestown, PA, US
Contact	RYAN O'LEARY
510(k) history	19 submissions · 19 cleared · 1994-2015