FDA 510(k) Premarket Notification

K090938 FLOW CONTROLLER

CLEARED

Jun 4, 2009
62 days to decision

K090938 · Product code: **LZU** · Ophthalmic
Source: https://www.510kdatabase.net/k090938/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plug, Punctum (LZU)
Date received	Apr 3, 2009
Decision date	Jun 4, 2009
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Odyssey Medical, Inc.**
Location	Memphis, TN, US
Contact	TERRY R GREEN
510(k) history	4 submissions · 4 cleared · 1997-2009