FDA 510(k) Premarket Notification

K171539 Contact Lens Case

CLEARED

Jul 14, 2017
49 days to decision

K171539 · Product code: **LRX** · Ophthalmic
Source: https://www.510kdatabase.net/k171539/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	May 26, 2017
Decision date	Jul 14, 2017
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**E-Link Plastic & Metal Industrial Co.,Ltd**
Location	New Taipei, CN
Contact	Marvis Lee
510(k) history	1 submissions · 1 cleared · 2017-2017