

K171539 Contact Lens CaseJul 14, 2017
49 days to decisionK171539 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171539/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 1, 2026