

K130753 STANDARD CONTACT LENS CASEJun 13, 2013
86 days to decisionK130753 · Product code: **LRX** · Ophthalmic
Source: <https://www.510kdatabase.net/k130753/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Mar 19, 2013
Decision date	Jun 13, 2013
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary
Other names	FLIP N SLIDE CONTACT LENS CASE

APPLICANT

Company	Watchdog Group, LC
Location	Clayton, MO, US
Contact	JOHN O'&HARA
510(k) history	1 submissions · 1 cleared · 2013-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130753/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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