FDA 510(k) Premarket Notification

K252175 LANDR Contact LENS Case

CLEARED

Feb 10, 2026
214 days to decision

K252175 · Product code: **LRX** · Ophthalmic
Source: https://www.510kdatabase.net/k252175/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Jul 11, 2025
Decision date	Feb 10, 2026
Days to decision	214 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Fourth Axis, LLC**
Location	Boise, ID, US
Contact	Christine Brown
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	**Bdra Consulting, LLC**
Contact	Srinagesh Koushik

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov