

K252175 LANDR Contact LENS CaseFeb 10, 2026
214 days to decisionK252175 · 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

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