

**K943921 RYDER CONTACT LENS CASE**Oct 11, 1994  
60 days to decisionK943921 · Product code: **LRX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k943921/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Aug 12, 1994
Decision date	Oct 11, 1994
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Clinical Research Assoc.</b>
Location	Brunswick, NJ, US
Contact	PATRICIA W BRADSTREET
510(k) history	1 submissions · 1 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k943921/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026