

**K920064 OPTI-LENS CONTACT LENS CASE**Mar 4, 1992  
57 days to decisionK920064 · Product code: **LRX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k920064/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Jan 7, 1992
Decision date	Mar 4, 1992
Days to decision	57 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ryder Intl. Corp.</b>
Location	Arab, AL, US
Contact	PATRICIA BRADSTREET
510(k) history	10 submissions · 10 cleared · 1985-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920064/> 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