

**K874657 RIGID-LENS CASE**Dec 8, 1987  
19 days to decisionK874657 · Product code: LRX · Ophthalmic  
Source: <https://www.510kdatabase.net/k874657/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Nov 19, 1987
Decision date	Dec 8, 1987
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Sola Ophthalmics</b>
Location	Phoenix, AZ, US
Contact	KRISTINE A FOSS
510(k) history	1 submissions · 1 cleared · 1987-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874657/>, 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 30, 2026