

**K900777 ITEM #273 CONTACT LENS CASES**Mar 30, 1990  
38 days to decisionK900777 · Product code: LRX · Ophthalmic  
Source: <https://www.510kdatabase.net/k900777/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Feb 20, 1990
Decision date	Mar 30, 1990
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Magnivision, Inc.</b>
Location	Miami, FL, US
Contact	ROY HIRSCHFELD
510(k) history	8 submissions · 8 cleared · 1987-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900777/>, 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