

**K973192 FLEXLENS, FLEXLENS TORIC**Nov 20, 1997  
87 days to decisionK973192 · Product code: LPL · Ophthalmic  
Source: <https://www.510kdatabase.net/k973192/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Aug 25, 1997
Decision date	Nov 20, 1997
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Specialty Ultravision, Inc.</b>
Location	Campbell, CA, US
Contact	MARTIN S KNOPF
510(k) history	12 submissions · 12 cleared · 1996-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973192/> 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