

**K896681 SOFT CORNEAL LIGHT SHIELD**Jan 11, 1990  
45 days to decisionK896681 · Product code: **HOY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k896681/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shield, Eye, Ophthalmic (including Sunlamp Protective Eyewear And Post-mydratic Eyewear) (HOY)
Date received	Nov 27, 1989
Decision date	Jan 11, 1990
Days to decision	45 days
Third-party review	No

**APPLICANT**

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Company	<b>Visitec Co.</b>
Location	Walker, MI, US
Contact	LIAQUAT ALLARAKHIA
510(k) history	49 submissions · 49 cleared · 1979-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896681/>, 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 May 23, 2026