

**K895364 LID LOC**Oct 31, 1989  
64 days to decisionK895364 · Product code: **HOY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k895364/>**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	Aug 28, 1989
Decision date	Oct 31, 1989
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Spectrum Co.</b>
Location	Mchenry, IL, US
Contact	AMY PICKEREL
510(k) history	3 submissions · 3 cleared · 1982-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895364/>; 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