

**K833259 MULTIPLE**Apr 2, 1984  
195 days to decisionK833259 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k833259/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Sep 20, 1983
Decision date	Apr 2, 1984
Days to decision	195 days
Third-party review	No

**APPLICANT**

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Company	<b>Sun House Products Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833259/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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