

**K790251 FUL-VUE ASPHERIC CATARACT LENS**Feb 26, 1979  
20 days to decisionK790251 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k790251/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Feb 6, 1979
Decision date	Feb 26, 1979
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>American Optical Corp.</b>
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995

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Device record: <https://www.510kdatabase.net/k790251/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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