

**K861684 SEIKO P-2 & SEIKO P-3 (SPECTACLE LENS)**Jun 9, 1986  
38 days to decisionK861684 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k861684/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	May 2, 1986
Decision date	Jun 9, 1986
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Seiko Optical Products of America, Inc.</b>
Location	Mahwah, NJ, US
Contact	ROBERT CURLEY
510(k) history	4 submissions · 4 cleared · 1986-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861684/>; 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