

**K882814 NEOLENS PRESCRIPTION SPECTACLE LENS**Jul 29, 1988  
21 days to decisionK882814 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k882814/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Jul 8, 1988
Decision date	Jul 29, 1988
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>Neolens, Inc.</b>
Location	Miami, FL, US
Contact	CLAYMAN, MD
510(k) history	2 submissions · 2 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882814/>; 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