

**K821398 CORNING CPF 550 LENS**Jun 25, 1982  
45 days to decisionK821398 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k821398/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	May 11, 1982
Decision date	Jun 25, 1982
Days to decision	45 days
Third-party review	No

**APPLICANT**

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Company	<b>Corning Medical &amp; Scientific</b>
Location	Mchenry, IL, US
Website	<a href="https://www.corning.com">https://www.corning.com</a>
510(k) history	111 submissions · 111 cleared · 1976-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821398/>; 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 May 14, 2026