

**K920170 SPECTACLE FRAME**Mar 11, 1992  
58 days to decisionK920170 · Product code: **HQZ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k920170/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Jan 13, 1992
Decision date	Mar 11, 1992
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sciara Intl., Inc.</b>
Location	Springfield Gardens, NY, US
Contact	MARILYN MAHLER
510(k) history	1 submissions · 1 cleared · 1992-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920170/> 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