

**K932773 FRAMES, SPECTACLES**Sep 20, 1993  
104 days to decisionK932773 · Product code: **HQZ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k932773/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Jun 8, 1993
Decision date	Sep 20, 1993
Days to decision	104 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Diaco, Inc.</b>
Location	Miami, FL, US
Contact	AGGIE GRAUPERA
510(k) history	1 submissions · 1 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932773/> 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