

**K934466 SPECTACLE FRAMES**Feb 22, 1994  
165 days to decisionK934466 · Product code: **HQZ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k934466/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Sep 10, 1993
Decision date	Feb 22, 1994
Days to decision	165 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Airess, Inc.</b>
Location	Miami, FL, US
Contact	BOBY REGE
510(k) history	2 submissions · 2 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k934466/> 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