

**K863429 MODIFIED DEVICE FOR SELF EXAMINATION OF EYES**Jan 20, 1987  
138 days to decisionK863429 · Product code: **HOQ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k863429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Grid, Amsler (HOQ)
Date received	Sep 4, 1986
Decision date	Jan 20, 1987
Days to decision	138 days
Third-party review	No

**APPLICANT**

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Company	<b>Wheel Checkers</b>
Location	Denver, CO, US
Contact	GIESKIENG
510(k) history	2 submissions · 2 cleared · 1985-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863429/>; 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