

**K970029 KEELER SPECTRA INDIRECT OPHTHALMOSCOPE**Apr 2, 1997  
89 days to decisionK970029 · Product code: **HLJ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k970029/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Battery-powered (HLJ)
Date received	Jan 3, 1997
Decision date	Apr 2, 1997
Days to decision	89 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Keeler Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	EUGENE R VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970029/> 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 20, 2026