

K893172 VISTA OPHTHALMOSCOPE (BATTERY AND AC-POWERED)Jun 5, 1989
41 days to decisionK893172 · Product code: **HLI** · Ophthalmic
Source: <https://www.510kdatabase.net/k893172/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Ac-powered (HLI)
Date received	Apr 25, 1989
Decision date	Jun 5, 1989
Days to decision	41 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893172/> Data sourced from FDA 510(k) public records (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. - Generated June 20, 2026