

**K893171 VISTA STREAK RETINOSCOPE (AC-POWERED)**Sep 21, 1989  
149 days to decisionK893171 · Product code: **HKL** · Ophthalmic  
Source: <https://www.510kdatabase.net/k893171/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retinoscope, Ac-powered (HKL)
Date received	Apr 25, 1989
Decision date	Sep 21, 1989
Days to decision	149 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893171/>, 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