

K942104 KEELER VANTAGE INDIRECT OPHTHALMOSCOPEAug 4, 1994
94 days to decisionK942104 · Product code: **HLI** · Ophthalmic
Source: <https://www.510kdatabase.net/k942104/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Ac-powered (HLI)
Date received	May 2, 1994
Decision date	Aug 4, 1994
Days to decision	94 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942104/> 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