FDA 510(k) Premarket Notification

K893173 VISTA SPECIALIST OPHTHALMOSCOPE (BATTERY & AC)

CLEARED

Jun 5, 1989
41 days to decision

K893173 · Product code: **HLI** · Ophthalmic
Source: https://www.510kdatabase.net/k893173/

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.net
Device record: https://www.510kdatabase.net/k893173/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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