FDA 510(k) Premarket Notification

CLEARED

K893171 VISTA STREAK RETINOSCOPE (AC-POWERED)

Sep 21, 1989
149 days to decision

K893171 · 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	**Keeler Instruments, Inc.**
Location	Mchenry, IL, US
Contact	VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019