FDA 510(k) Premarket Notification

K891610 HENSON CFA 3000 CENTRAL VISUAL FIELD SCREENER

CLEARED

Jun 5, 1989
77 days to decision

K891610 · Product code: **HOD** · Ophthalmic
Source: https://www.510kdatabase.net/k891610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Eyelid, Ophthalmic (HOD)
Date received	Mar 20, 1989
Decision date	Jun 5, 1989
Days to decision	77 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