

**K891610 HENSON CFA 3000 CENTRAL VISUAL FIELD
SCREENER**Jun 5, 1989
77 days to decisionK891610 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891610/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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