

K830770 CONTRAST SENSITIVITY EQUIPMENTApr 28, 1983
48 days to decisionK830770 · Product code: **HLX** · Ophthalmic
Source: <https://www.510kdatabase.net/k830770/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photostimulator, Ac-powered (HLX)
Date received	Mar 11, 1983
Decision date	Apr 28, 1983
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Cadwell Laboratories, Inc.
Location	Walker, MI, US
510(k) history	46 submissions · 46 cleared · 1979-2007

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Device record: <https://www.510kdatabase.net/k830770/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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