FDA 510(k) Premarket Notification

K830770 CONTRAST SENSITIVITY EQUIPMENT

CLEARED

Apr 28, 1983
48 days to decision

K830770 · 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