FDA 510(k) Premarket Notification

K881358 PG CONSEN CONTRAST SENSITIVITY TESTING PROGRAM

CLEARED

Jun 27, 1988
88 days to decision

K881358 · Product code: **HOX** · Ophthalmic
Source: https://www.510kdatabase.net/k881358/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chart, Visual Acuity (HOX)
Date received	Mar 31, 1988
Decision date	Jun 27, 1988
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Neuroscientific Corp.**
Location	Farmingdale, NY, US
Contact	ALAN P SCHWARTZ
510(k) history	2 submissions · 2 cleared · 1988-1988