

**K881358 PG CONSEN CONTRAST SENSITIVITY TESTING PROGRAM**Jun 27, 1988  
88 days to decisionK881358 · 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	<b>Neuroscientific Corp.</b>
Location	Farmingdale, NY, US
Contact	ALAN P SCHWARTZ
510(k) history	2 submissions · 2 cleared · 1988-1988

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881358/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026