

K914393 RETINA TESTERMar 2, 1992
153 days to decisionK914393 · Product code: **HLX** · Ophthalmic
Source: <https://www.510kdatabase.net/k914393/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photostimulator, Ac-powered (HLX)
Date received	Oct 1, 1991
Decision date	Mar 2, 1992
Days to decision	153 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Retina Tester
Location	Memphis, TN, US
Contact	IRBY SEAY
510(k) history	1 submissions · 1 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914393/> 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