FDA 510(k) Premarket Notification

K914393 RETINA TESTER

CLEARED

Mar 2, 1992
153 days to decision

K914393 · 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