

**K950793 SPOT RETINOSCOPE RX-3SP**Mar 28, 1995  
35 days to decisionK950793 · Product code: **HKM** · Ophthalmic  
Source: <https://www.510kdatabase.net/k950793/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retinoscope, Battery-powered (HKM)
Date received	Feb 21, 1995
Decision date	Mar 28, 1995
Days to decision	35 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Neitz Instruments Company, Ltd.</b>
Location	Washington, DC, US
Contact	YASUO KAWANO
510(k) history	24 submissions · 24 cleared · 1994-1997

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950793/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026