

K942242 AUTO REF-KERATOMETER, RK-3Sep 2, 1994
116 days to decisionK942242 · Product code: **HKO** · Ophthalmic
Source: <https://www.510kdatabase.net/k942242/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	May 9, 1994
Decision date	Sep 2, 1994
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cannon U.S.A., Inc.
Location	Lake Success, NY, US
Contact	HIROYUKI TAKAHASHI
510(k) history	2 submissions · 2 cleared · 1994-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942242/> 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 May 20, 2026