FDA 510(k) Premarket Notification

K942242 AUTO REF-KERATOMETER, RK-3

CLEARED

Sep 2, 1994
116 days to decision

K942242 · 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