FDA 510(k) Premarket Notification

K841861 CANON AUTOREF R-10

CLEARED

Oct 5, 1984
154 days to decision

K841861 · Product code: **HKO** · Ophthalmic
Source: https://www.510kdatabase.net/k841861/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	May 4, 1984
Decision date	Oct 5, 1984
Days to decision	154 days
Third-party review	No

APPLICANT

Company	**Canon USA, Inc.**
Location	Mchenry, IL, US
510(k) history	48 submissions · 48 cleared · 1984-2009