FDA 510(k) Premarket Notification

K861066 AR-1000 AUTOMATIC REFRACTOR

CLEARED

Apr 4, 1986
15 days to decision

K861066 · Product code: **HKO** · Ophthalmic
Source: https://www.510kdatabase.net/k861066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Mar 20, 1986
Decision date	Apr 4, 1986
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Nidek, Inc.**
Location	Mchenry, IL, US
Contact	JOHN BRATKOWSKY
510(k) history	77 submissions · 77 cleared · 1983-2005