FDA 510(k) Premarket Notification

CLEARED

K802183 SR-IV PROGRAMMED SUBJECTIVE REFRACTOR

Apr 23, 1981
226 days to decision

K802183 · Product code: **HKO** · Ophthalmic
Source: https://www.510kdatabase.net/k802183/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Sep 9, 1980
Decision date	Apr 23, 1981
Days to decision	226 days
Third-party review	No

APPLICANT

Company	**American Optical Corp.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995