FDA 510(k) Premarket Notification

K801906 DIOPTRON III AUTOREFRACTOR

CLEARED

Aug 27, 1980
16 days to decision

K801906 · Product code: **HKO** · Ophthalmic
Source: https://www.510kdatabase.net/k801906/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Aug 11, 1980
Decision date	Aug 27, 1980
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Coherent Medical Division**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1990