FDA 510(k) Premarket Notification

K790251 FUL-VUE ASPHERIC CATARACT LENS

CLEARED

Feb 26, 1979
20 days to decision

K790251 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k790251/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Feb 6, 1979
Decision date	Feb 26, 1979
Days to decision	20 days
Third-party review	No

APPLICANT

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