FDA 510(k) Premarket Notification

K833259 MULTIPLE

CLEARED

Apr 2, 1984
195 days to decision

K833259 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k833259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Sep 20, 1983
Decision date	Apr 2, 1984
Days to decision	195 days
Third-party review	No

APPLICANT

Company	**Sun House Products Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984