FDA 510(k) Premarket Notification

K812441 DI-CROWN

CLEARED

Oct 6, 1981
41 days to decision

K812441 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k812441/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Aug 26, 1981
Decision date	Oct 6, 1981
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Coburn Optical Ind., Inc.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1978-1989