FDA 510(k) Premarket Notification

K821398 CORNING CPF 550 LENS

CLEARED

Jun 25, 1982
45 days to decision

K821398 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k821398/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	May 11, 1982
Decision date	Jun 25, 1982
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Corning Medical & Scientific**
Location	Mchenry, IL, US
Website	https://www.corning.com
510(k) history	111 submissions · 111 cleared · 1976-1988