FDA 510(k) Premarket Notification

K772024 RX LENS SERIES

CLEARED

Nov 9, 1977
14 days to decision

K772024 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k772024/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Oct 26, 1977
Decision date	Nov 9, 1977
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Gentex Corporation**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1977-2009