FDA 510(k) Premarket Notification

K813156 ULTRA-WIDE SAFETY SPECTACLE FRAME

CLEARED

Dec 14, 1981
38 days to decision

K813156 · Product code: **HQZ** · Ophthalmic
Source: https://www.510kdatabase.net/k813156/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Nov 6, 1981
Decision date	Dec 14, 1981
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Zee Medical Products Co., Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-1983