FDA 510(k) Premarket Notification

K895364 LID LOC

CLEARED

Oct 31, 1989
64 days to decision

K895364 · Product code: **HOY** · Ophthalmic
Source: https://www.510kdatabase.net/k895364/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shield, Eye, Ophthalmic (including Sunlamp Protective Eyewear And Post-mydriatic Eyewear) (HOY)
Date received	Aug 28, 1989
Decision date	Oct 31, 1989
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Spectrum Co.**
Location	Mchenry, IL, US
Contact	AMY PICKEREL
510(k) history	3 submissions · 3 cleared · 1982-1989