FDA 510(k) Premarket Notification

K896681 SOFT CORNEAL LIGHT SHIELD

CLEARED

Jan 11, 1990
45 days to decision

K896681 · Product code: **HOY** · Ophthalmic
Source: https://www.510kdatabase.net/k896681/

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	Nov 27, 1989
Decision date	Jan 11, 1990
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Visitec Co.**
Location	Walker, MI, US
Contact	LIAQUAT ALLARAKHIA
510(k) history	49 submissions · 49 cleared · 1979-1995