FDA 510(k) Premarket Notification

K861684 SEIKO P-2 & SEIKO P-3 (SPECTACLE LENS)

CLEARED

Jun 9, 1986
38 days to decision

K861684 · Product code: **HQG** · Ophthalmic
Source: https://www.510kdatabase.net/k861684/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	May 2, 1986
Decision date	Jun 9, 1986
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Seiko Optical Products of America, Inc.**
Location	Mahwah, NJ, US
Contact	ROBERT CURLEY
510(k) history	4 submissions · 4 cleared · 1986-1994