FDA 510(k) Premarket Notification

K842835 BRILLETTE MODEL

CLEARED

Nov 21, 1984
124 days to decision

K842835 · Product code: **HOI** · Ophthalmic
Source: https://www.510kdatabase.net/k842835/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spectacle, Magnifying (HOI)
Date received	Jul 20, 1984
Decision date	Nov 21, 1984
Days to decision	124 days
Third-party review	No

APPLICANT

Company	**Cillus Company, Inc.**
Location	Novato, CA, US
Contact	JORGEN L KREJSBOL
510(k) history	1 submissions · 1 cleared · 1984-1984