FDA 510(k) Premarket Notification

K920170 SPECTACLE FRAME

CLEARED

Mar 11, 1992
58 days to decision

K920170 · Product code: **HQZ** · Ophthalmic
Source: https://www.510kdatabase.net/k920170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Jan 13, 1992
Decision date	Mar 11, 1992
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sciara Intl., Inc.**
Location	Springfield Gardens, NY, US
Contact	MARILYN MAHLER
510(k) history	1 submissions · 1 cleared · 1992-1992