FDA 510(k) Premarket Notification

K933891 SPECTACLE FRAMES

CLEARED

Oct 18, 1993
69 days to decision

K933891 · Product code: **HQZ** · Ophthalmic
Source: https://www.510kdatabase.net/k933891/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Aug 10, 1993
Decision date	Oct 18, 1993
Days to decision	69 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Euro Frame Design, Inc.**
Location	Wayne, NJ, US
Contact	THOMAS BONIFACE
510(k) history	1 submissions · 1 cleared · 1993-1993