FDA 510(k) Premarket Notification

K932773 FRAMES, SPECTACLES

CLEARED

Sep 20, 1993
104 days to decision

K932773 · Product code: **HQZ** · Ophthalmic
Source: https://www.510kdatabase.net/k932773/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Jun 8, 1993
Decision date	Sep 20, 1993
Days to decision	104 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Diaco, Inc.**
Location	Miami, FL, US
Contact	AGGIE GRAUPERA
510(k) history	1 submissions · 1 cleared · 1993-1993