

K933891 SPECTACLE FRAMESOct 18, 1993
69 days to decisionK933891 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933891/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026