FDA 510(k) Premarket Notification

K813548 UVEX MINI-SPECS

CLEARED

Mar 2, 1982
71 days to decision

K813548 · Product code: **HOI** · Ophthalmic
Source: https://www.510kdatabase.net/k813548/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spectacle, Magnifying (HOI)
Date received	Dec 21, 1981
Decision date	Mar 2, 1982
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**Bacou USA, Inc. and Uvex Safety, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1992