

K813548 UVEX MINI-SPECSMar 2, 1982
71 days to decisionK813548 · 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

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Device record: <https://www.510kdatabase.net/k813548/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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