

K813156 ULTRA-WIDE SAFETY SPECTACLE FRAMEDec 14, 1981
38 days to decisionK813156 · Product code: **HQZ** · Ophthalmic
Source: <https://www.510kdatabase.net/k813156/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Frame, Spectacle (HQZ)
Date received	Nov 6, 1981
Decision date	Dec 14, 1981
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Zee Medical Products Co., Inc.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k813156/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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