

K801733 FENZL RETICLESep 16, 1980
53 days to decisionK801733 · Product code: **HOI** · Ophthalmic
Source: <https://www.510kdatabase.net/k801733/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spectacle, Magnifying (HOI)
Date received	Jul 25, 1980
Decision date	Sep 16, 1980
Days to decision	53 days
Third-party review	No

APPLICANT

Company	Robert E. Fenzi, M.D.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k801733/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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