FDA 510(k) Premarket Notification

K801733 FENZL RETICLE

CLEARED

Sep 16, 1980
53 days to decision

K801733 · 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. Fenzl, M.D.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980