

K830468 MODEL 200Apr 12, 1983
56 days to decisionK830468 · Product code: **FTZ** · Ophthalmic
Source: <https://www.510kdatabase.net/k830468/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Locator, Magnetic (FTZ)
Date received	Feb 15, 1983
Decision date	Apr 12, 1983
Days to decision	56 days
Third-party review	No

APPLICANT

Company	Western Laboratories Corp.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1978-1984

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Device record: <https://www.510kdatabase.net/k830468/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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