FDA 510(k) Premarket Notification

K830468 MODEL 200

CLEARED

Apr 12, 1983
56 days to decision

K830468 · 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