FDA 510(k) Premarket Notification

K780125 FLUORESOFT

K780125 · Product code: **KYC** · Ophthalmic
Source: https://www.510kdatabase.net/k780125/

CLEARED

Mar 15, 1978
50 days to decision

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jan 24, 1978
Decision date	Mar 15, 1978
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Holles Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k780125/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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