

K830624 FLUOROMETER, GENERAL USEApr 5, 1983
36 days to decisionK830624 · Product code: **KHO** · Toxicology
Source: <https://www.510kdatabase.net/k830624/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fluorometer, For Clinical Use (KHO)
Date received	Feb 28, 1983
Decision date	Apr 5, 1983
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Abbott Diagnostics
Location	Mountain View, CA, US
510(k) history	28 submissions · 28 cleared · 1981-1997

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Device record: <https://www.510kdatabase.net/k830624/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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