

K813540 A FLUOROMETERDec 31, 1981
10 days to decisionK813540 · Product code: **KHO** · Chemistry
Source: <https://www.510kdatabase.net/k813540/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fluorometer, For Clinical Use (KHO)
Date received	Dec 21, 1981
Decision date	Dec 31, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

Company	American Diagnostic Corp.
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1980-2017

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Device record: <https://www.510kdatabase.net/k813540/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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