FDA 510(k) Premarket Notification

K813540 A FLUOROMETER

CLEARED

Dec 31, 1981
10 days to decision

K813540 · 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