FDA 510(k) Premarket Notification

K802468 T3 UPTAKE TEST SYSTEM

CLEARED

Oct 31, 1980
22 days to decision

K802468 · Product code: **KHQ** · Chemistry
Source: https://www.510kdatabase.net/k802468/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Oct 9, 1980
Decision date	Oct 31, 1980
Days to decision	22 days
Third-party review	No

APPLICANT

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