FDA 510(k) Premarket Notification

K821722 T3 UPTAKE DIAGNOSTIC KIT

CLEARED

Jun 24, 1982
14 days to decision

K821722 · Product code: **KHQ** · Chemistry
Source: https://www.510kdatabase.net/k821722/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Jun 10, 1982
Decision date	Jun 24, 1982
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Diagnostic Reagents, Inc.**
Location	Walker, MI, US
510(k) history	66 submissions · 66 cleared · 1982-1998