FDA 510(k) Premarket Notification

K820340 T3 UPTAKE-SQUIBB DIAGNOSTIC KIT

CLEARED

Mar 4, 1982
24 days to decision

K820340 · Product code: **KHQ** · Toxicology
Source: https://www.510kdatabase.net/k820340/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Feb 8, 1982
Decision date	Mar 4, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**E. R. Squibb & Sons, Inc.**
Location	New York, NY, US
510(k) history	32 submissions · 32 cleared · 1977-1982