FDA 510(k) Premarket Notification

K885259 ACA THYRONINE UPTAKE (TU E) METHOD

CLEARED

Feb 17, 1989
56 days to decision

K885259 · Product code: **KHQ** · Chemistry
Source: https://www.510kdatabase.net/k885259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Dec 23, 1988
Decision date	Feb 17, 1989
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**The DU Pont Co.**
Location	Wilmington, DE, US
Contact	RICHARD M VAUGHT
510(k) history	13 submissions · 13 cleared · 1988-1995

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k885259/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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