FDA 510(k) Premarket Notification

K884170 ANALYST THEOPHYLLINE CALIBRATOR

CLEARED

Nov 16, 1988
56 days to decision

K884170 · Product code: **DLJ** · Toxicology
Source: https://www.510kdatabase.net/k884170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	Sep 21, 1988
Decision date	Nov 16, 1988
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/k884170/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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