FDA 510(k) Premarket Notification

K884745 DU PONT ACA METHOTREXATE CALIBRATOR

CLEARED

Jan 30, 1989
77 days to decision

K884745 · Product code: **DLJ** · Toxicology
Source: https://www.510kdatabase.net/k884745/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	Nov 14, 1988
Decision date	Jan 30, 1989
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**E.I. Dupont DE Nemours & Co., Inc.**
Location	Mchenry, IL, US
Contact	CHRISTOPHER BENTSEN
510(k) history	253 submissions · 252 cleared · 1976-1996