FDA 510(k) Premarket Notification

K833380 LIDOAINE CALIBRATOR

CLEARED

Dec 27, 1983
89 days to decision

K833380 · Product code: **DLJ** · Chemistry
Source: https://www.510kdatabase.net/k833380/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	Sep 29, 1983
Decision date	Dec 27, 1983
Days to decision	89 days
Third-party review	No

APPLICANT

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