FDA 510(k) Premarket Notification

K771805 ACA SALICYLATE CALIBRATOR

CLEARED

Oct 4, 1977
11 days to decision

K771805 · Product code: **DLJ** · Toxicology
Source: https://www.510kdatabase.net/k771805/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	Sep 23, 1977
Decision date	Oct 4, 1977
Days to decision	11 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