

K771805 ACA SALICYLATE CALIBRATOROct 4, 1977
11 days to decisionK771805 · 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

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Device record: <https://www.510kdatabase.net/k771805/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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