

**K833380 LIDOINE CALIBRATOR**Dec 27, 1983  
89 days to decisionK833380 · Product code: **DLJ** · Chemistry  
Source: <https://www.510kdatabase.net/k833380/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k833380/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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