

K780731 LIDOCAINE CALIBRATOR KITJan 17, 1979
260 days to decisionK780731 · Product code: **DLJ** · Toxicology
Source: <https://www.510kdatabase.net/k780731/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	May 2, 1978
Decision date	Jan 17, 1979
Days to decision	260 days
Third-party review	No

APPLICANT

Company	Vitek Systems, Inc.
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1978-1993

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Device record: <https://www.510kdatabase.net/k780731/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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