

**K780732 PROCAINAMIDE CALIBRATOR KIT**Nov 22, 1978  
204 days to decisionK780732 · Product code: **DLJ** · Toxicology  
Source: <https://www.510kdatabase.net/k780732/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	May 2, 1978
Decision date	Nov 22, 1978
Days to decision	204 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k780732/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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