

**K780734 ANTICONVULSANTS II CALIBRATOR KIT**Jun 22, 1978  
51 days to decisionK780734 · Product code: **DIY** · Toxicology  
Source: <https://www.510kdatabase.net/k780734/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gas Chromatography, Ethosuximide (DIY)
Date received	May 2, 1978
Decision date	Jun 22, 1978
Days to decision	51 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/k780734/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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