FDA 510(k) Premarket Notification

K780734 ANTICONVULSANTS II CALIBRATOR KIT

CLEARED

Jun 22, 1978
51 days to decision

K780734 · Product code: **DIY** · Toxicology
Source: https://www.510kdatabase.net/k780734/

SUBMISSION DETAILS

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

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