FDA 510(k) Premarket Notification

K780731 LIDOCAINE CALIBRATOR KIT

CLEARED

Jan 17, 1979
260 days to decision

K780731 · 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