FDA 510(k) Premarket Notification

K780732 PROCAINAMIDE CALIBRATOR KIT

CLEARED

Nov 22, 1978
204 days to decision

K780732 · Product code: **DLJ** · Toxicology
Source: https://www.510kdatabase.net/k780732/

SUBMISSION DETAILS

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

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