FDA 510(k) Premarket Notification

K780960 SERUM, CONTROL, DRUGS, ANTIARRYTHMIC

CLEARED

Dec 7, 1978
178 days to decision

K780960 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k780960/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jun 12, 1978
Decision date	Dec 7, 1978
Days to decision	178 days
Third-party review	No

APPLICANT

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