

K780960 SERUM, CONTROL, DRUGS, ANTIARRYTHMICDec 7, 1978
178 days to decisionK780960 · 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

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Device record: <https://www.510kdatabase.net/k780960/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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