FDA 510(k) Premarket Notification

CLEARED

K971111 TRI-TECH INC., EVACUATED BLOOD COLLECTION TUBE, GRAY STOPPER

Jun 16, 1997
82 days to decision

K971111 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k971111/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Mar 26, 1997
Decision date	Jun 16, 1997
Days to decision	82 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tri-Tech, Inc.**
Location	Mchenry, IL, US
Contact	WENDIE WALKER
510(k) history	11 submissions · 9 cleared · 1981-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k971111/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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