FDA 510(k) Premarket Notification

K011699 VACUETTE EVACUATED BLOOD COLLECTION TUBES

CLEARED

Jun 28, 2001
27 days to decision

K011699 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k011699/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Jun 1, 2001
Decision date	Jun 28, 2001
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Greiner Bio-One Vacuette North America**
Location	Baldwin, MD, US
Contact	Judi Smith
510(k) history	5 submissions · 5 cleared · 2001-2008

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k011699/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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