FDA 510(k) Premarket Notification

K971236 GREINER VACUETTE BLOOD COLLECTION TUBE

CLEARED

May 12, 1997
40 days to decision

K971236 · Product code: **JKA** · Toxicology
Source: https://www.510kdatabase.net/k971236/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Apr 2, 1997
Decision date	May 12, 1997
Days to decision	40 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Greiner America, Inc.**
Location	Wilmington, DE, US
Contact	ED MAIER
510(k) history	7 submissions · 7 cleared · 1996-1997