FDA 510(k) Premarket Notification

K960860 GREINER VACUETTE BLOOD COLLECTION TUBE E/EDTA K3

CLEARED

Sep 5, 1996
188 days to decision

K960860 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k960860/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Mar 1, 1996
Decision date	Sep 5, 1996
Days to decision	188 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