FDA 510(k) Premarket Notification

K002777 VACUETTE WITH PPACK

CLEARED

Oct 5, 2000
29 days to decision

K002777 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k002777/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Sep 6, 2000
Decision date	Oct 5, 2000
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Greiner Vacuette North America, Inc.**
Location	Ellicott City, MD, US
Contact	Judi Smith
510(k) history	5 submissions · 5 cleared · 2000-2002