FDA 510(k) Premarket Notification

K933939 BACT/ALERT BLOOD COLLECTION ADAPTER CAP AND INSERT

CLEARED

Feb 7, 1994
179 days to decision

K933939 · Product code: **GIM** · Microbiology
Source: https://www.510kdatabase.net/k933939/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vacuum Sample, With Anticoagulant (GIM)
Date received	Aug 12, 1993
Decision date	Feb 7, 1994
Days to decision	179 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Organon Teknika Corp.**
Location	Mchenry, IL, US
Contact	ANN M QUINN
510(k) history	130 submissions · 129 cleared · 1980-2000