

**K933939 BACT/ALERT BLOOD COLLECTION ADAPTER CAP
AND INSERT**Feb 7, 1994
179 days to decisionK933939 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933939/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026