

K233979 VACUETTE® QUICKSHIELD Complete (Plus)Jul 19, 2024
214 days to decisionK233979 · Product code: **JKA** · General Hospital
Source: <https://www.510kdatabase.net/k233979/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Dec 18, 2023
Decision date	Jul 19, 2024
Days to decision	214 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Greiner Bio-One North America, Inc.
Location	Baldwin, MD, US
Contact	Manfred Abel
510(k) history	10 submissions · 10 cleared · 2006-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233979/> 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 13, 2026