

K222299 Steripath Micro Blood Collection SystemNov 30, 2022
121 days to decisionK222299 · Product code: **JKA** · General Hospital
Source: <https://www.510kdatabase.net/k222299/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Aug 1, 2022
Decision date	Nov 30, 2022
Days to decision	121 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Magnolia Medical Technologies
Location	Seattle, WA, US
Contact	Greg Bullington
510(k) history	2 submissions · 2 cleared · 2022-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k222299/> 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 25, 2026