

K072320 GREINER VACUETTE QUICKSHIELD COMPLETESep 14, 2007
25 days to decisionK072320 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k072320/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Aug 20, 2007
Decision date	Sep 14, 2007
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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