FDA 510(k) Premarket Notification

K072320 GREINER VACUETTE QUICKSHIELD COMPLETE

CLEARED

Sep 14, 2007
25 days to decision

K072320 · Product code: **FMI** · General Hospital
Source: 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