

**K061483 VACUETTE VISIO PLUS BLOOD COLLECTION
NEEDLES**Aug 14, 2006
76 days to decisionK061483 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k061483/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 30, 2006
Decision date	Aug 14, 2006
Days to decision	76 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061483/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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