

**K111424 VACUETTE PREMIUM SAFETY NEEDLE SYSTEM
TUBE-TOUCH**Jul 29, 2011
67 days to decisionK111424 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k111424/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 23, 2011
Decision date	Jul 29, 2011
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medpro Safety Products, Inc.
Location	Lexington, KY, US
Contact	Judi Smith
510(k) history	3 submissions · 3 cleared · 2008-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k111424/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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