FDA 510(k) Premarket Notification

K111424 VACUETTE PREMIUM SAFETY NEEDLE SYSTEM TUBE-TOUCH

CLEARED

Jul 29, 2011
67 days to decision

K111424 · 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