FDA 510(k) Premarket Notification

K141749 CAREFINE PEN NEEDLE FAMILY INCLUDING QUINTAPOINT AND SUPERPOINT

CLEARED

Jul 29, 2014
29 days to decision

K141749 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k141749/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 30, 2014
Decision date	Jul 29, 2014
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Facet Technologies, LLC**
Location	Kennesaw, GA, US
Contact	JENNIFER REGISTER
510(k) history	7 submissions · 7 cleared · 2014-2024