

**K141749 CAREFINE PEN NEEDLE FAMILY INCLUDING
QUINTAPOINT AND SUPERPOINT**Jul 29, 2014
29 days to decisionK141749 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k141749/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 25, 2026