

K122373 NEOMED ORAL/ENTERAL SYRINGESep 20, 2012
45 days to decisionK122373 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k122373/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Syringe, Piston (FMF)
Date received	Aug 6, 2012
Decision date	Sep 20, 2012
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Neomed, Inc.
Location	Swanee, GA, US
Contact	MELINDA HARRISON
510(k) history	13 submissions · 13 cleared · 2007-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122373/> 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 June 20, 2026