FDA 510(k) Premarket Notification

K122373 NEOMED ORAL/ENTERAL SYRINGE

CLEARED

Sep 20, 2012
45 days to decision

K122373 · 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