

K161812 Freedom SyringeMar 27, 2017
269 days to decisionK161812 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k161812/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Syringe, Piston (FMF)
Date received	Jul 1, 2016
Decision date	Mar 27, 2017
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Procedure Products, Inc.
Location	Mchenry, IL, US
Contact	Doug Rowley
510(k) history	16 submissions · 16 cleared · 1981-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161812/> 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 1, 2026