

K023752 BD INTEGRA SYRINGEFeb 7, 2003
91 days to decisionK023752 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k023752/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 8, 2002
Decision date	Feb 7, 2003
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bd
Location	Vernon Hills, IL, US
Contact	PASQUALE AMATO
510(k) history	15 submissions · 15 cleared · 2001-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k023752/> 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 24, 2026