

**K021475 BD SINGLE LUMEN NEEDLE, SYRINGE AND BLOOD COLLECTION SET**Jul 19, 2002  
72 days to decisionK021475 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k021475/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 8, 2002
Decision date	Jul 19, 2002
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021475/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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