

**K001690 PROTECTIV SAFETY BLOOD COLLECTION NEEDLE**Jul 31, 2000  
59 days to decisionK001690 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k001690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 2, 2000
Decision date	Jul 31, 2000
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Medical, Inc.</b>
Location	Arlington, TX, US
Contact	DORENE MARKWIESE
510(k) history	53 submissions · 50 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001690/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026