

**K110105 BD PEN NEEDLE**Apr 28, 2011  
105 days to decisionK110105 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k110105/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 13, 2011
Decision date	Apr 28, 2011
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Becton, Dickinson and Company (BD)</b>
Location	Vernon Hills, IL, US
Contact	PASQUALE AMATO
510(k) history	5 submissions · 5 cleared · 2009-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110105/> 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 May 14, 2026