

**K020921 NO-POKES NEEDLE SAFETY DEVICE**Jul 11, 2002  
112 days to decisionK020921 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k020921/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 21, 2002
Decision date	Jul 11, 2002
Days to decision	112 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Bemis Mfg. Co.</b>
Location	Sheboygan Falls, WI, US
Contact	JOHN B HOWELL
510(k) history	12 submissions · 12 cleared · 1985-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020921/> 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 July 2, 2026