

**K960982 MONOJECT BLUNT I.V. ACCESS  
CANNULA/MONOJECT HYPODERMIC SYRINGE WITH BLUNT  
I.V. ACCESS CANNULA**Dec 23, 1996  
287 days to decisionK960982 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k960982/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 11, 1996
Decision date	Dec 23, 1996
Days to decision	287 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sherwood Medical Co.</b>
Location	Mchenry, IL, US
Contact	CHERLY WINTERS-HEARD
510(k) history	191 submissions · 177 cleared · 1976-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960982/> 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