

**K913317 MEGA ARTHROGRAM KIT**Oct 31, 1991  
98 days to decisionK913317 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k913317/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 25, 1991
Decision date	Oct 31, 1991
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mega Medical, Inc.</b>
Location	Tampa, FL, US
Contact	RAY VAHAB
510(k) history	7 submissions · 4 cleared · 1991-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913317/> 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 26, 2026