

**K103200 PROPEP DELIVERY DEVICE**Apr 12, 2011  
162 days to decisionK103200 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k103200/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 1, 2010
Decision date	Apr 12, 2011
Days to decision	162 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rhythmink International, LLC</b>
Location	Cayce, SC, US
Contact	JAMES M MEWBORNE
510(k) history	18 submissions · 18 cleared · 2002-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103200/> 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 27, 2026