

**K091386 MICROFUSE DUAL RATE INFUSER**Oct 2, 2009  
144 days to decisionK091386 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k091386/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Infusion (FRN)
Date received	May 11, 2009
Decision date	Oct 2, 2009
Days to decision	144 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Numia Medical Technology, LLC</b>
Location	Lyndonville, VT, US
Contact	ERIC FLACHBART
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091386/> 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 June 22, 2026