

**K091315 COVIDIEN EXTENDED TIP APPLICATOR**Jun 5, 2009  
32 days to decisionK091315 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k091315/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Syringe, Piston (FMF)
Date received	May 4, 2009
Decision date	Jun 5, 2009
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Confluent Surgical,Inc</b>
Location	Waltham, MA, US
Contact	SEAN REYNOLDS
510(k) history	6 submissions · 6 cleared · 2004-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091315/> 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 23, 2026