

**K042588 CONFLUENT SURGICAL DUAL LIQUID APPLICATOR,  
MODEL 70-2005093**

Dec 9, 2004  
78 days to decision

K042588 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k042588/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 22, 2004
Decision date	Dec 9, 2004
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Confluent Surgical, Inc</b>
Location	Waltham, MA, US
Contact	Eric Ankerud
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/k042588/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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