

**K070548 C-PORT FLEXA DISTAL ANASTOMOSIS SYSTEM**Mar 29, 2007  
31 days to decisionK070548 · Product code: **FZP** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k070548/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clip, Implantable (FZP)
Date received	Feb 26, 2007
Decision date	Mar 29, 2007
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardica, Inc.</b>
Location	Redwood City, CA, US
Contact	TIFFINI LALUDE
510(k) history	15 submissions · 15 cleared · 2005-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070548/> 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 2, 2026