

**K121168 APTUS 18FR HELIFX GUIDE-22 DEFLECTABLE LENGTH**Aug 8, 2012  
113 days to decisionK121168 · Product code: **OTD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k121168/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endovascular Suturing System (OTD)
Date received	Apr 17, 2012
Decision date	Aug 8, 2012
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aptus Endosystems, Inc.</b>
Location	Sunnyvale, CA, US
Contact	BURT GOODSON
510(k) history	4 submissions · 3 cleared · 2011-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121168/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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