

**K122585 CRUX VENA CAVA FILTER SYSTEM**Jan 17, 2013  
146 days to decisionK122585 · Product code: **DTK** · Cardiovascular  
Source: <https://www.510kdatabase.net/k122585/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filter, Intravascular, Cardiovascular (DTK)
Date received	Aug 24, 2012
Decision date	Jan 17, 2013
Days to decision	146 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Crux Biomedical</b>
Location	Menlo Park, CA, US
Contact	ELISA HEBB
510(k) history	2 submissions · 2 cleared · 2012-2013

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122585/> 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 9, 2026