

**K092175 THE CROSSER S6 CATHETER, THE CROSSER SYSTEM ELECTRONICS, MODEL CRUS6, GEN200**Feb 17, 2010  
211 days to decisionK092175 · Product code: **PDU** · Cardiovascular  
Source: <https://www.510kdatabase.net/k092175/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter For Crossing Total Occlusions (PDU)
Date received	Jul 21, 2009
Decision date	Feb 17, 2010
Days to decision	211 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Flowcardia, Inc.</b>
Location	Sunnyvale, CA, US
Contact	Dustin Michaels
510(k) history	11 submissions · 11 cleared · 2005-2010

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092175/> 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 21, 2026