

**K102725 CROSSBOSS CATHETER MODEL M-2000, STINGRAY  
ORIENTING BALLOON CATHETER MODEL M-1000, STINGRAY  
GUIDEWIRE MODEL M-3004 AND M**May 10, 2011  
231 days to decisionK102725 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k102725/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 21, 2010
Decision date	May 10, 2011
Days to decision	231 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Bridge Medical, Inc.</b>
Location	Plymouth, MN, US
Contact	JILL MUNSINGER
510(k) history	2 submissions · 2 cleared · 1998-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102725/> 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 19, 2026