

**K102721 LIVEWIRE ELECTROPHYSIOLOGY CATHETER**Oct 18, 2010  
27 days to decisionK102721 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k102721/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Sep 21, 2010
Decision date	Oct 18, 2010
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>St Jude Medical</b>
Location	Minnetonka, MN, US
Contact	NICOLE BOWDEN
Website	<a href="http://www.sjm.com/">http://www.sjm.com/</a>
510(k) history	105 submissions · 105 cleared · 2000-2018

St Jude Medical was a global medical device company headquartered in Little Canada, Minnesota. The company operated more than 20 principal facilities worldwide and sold products in over 100 countries. St Jude Medical received FDA 510(k) clearances from total submissions between 2000 and 2018. The company's regulatory focus centered on Cardiovascular devices, which represented 91% of all submissions. Notable cleared products include cardiac mapping systems, pacing catheters, and mobile cardiac applications. Now part of Abbott Laboratories following its acquisition in Janua...

---