

K131592 ENLIGHTN RENAL GUIDE CATHETERJan 31, 2014
242 days to decisionK131592 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k131592/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 3, 2013
Decision date	Jan 31, 2014
Days to decision	242 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	St. Jude Medical, Inc.
Location	Salt Lake City, UT, US
Contact	BLAIR SCHWARTZ
Website	http://www.sjm.com/
510(k) history	23 submissions · 22 cleared · 1989-2018

St. Jude Medical, Inc. 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 1989 and 2018. The company specialized exclusively in Cardiovascular devices, establishing a focused portfolio in cardiac monitoring, catheter systems, and related interventional technologies. Founded in 1976 and publicly listed in 1977, St. Jude Medical achieved Fortune 500 status annually...

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