FDA 510(k) Premarket Notification

CLEARED

K083402 AGILIS NXT STEERABLE INTRODUCER, MODEL G408322 AND G408323

Dec 15, 2008
27 days to decision

K083402 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k083402/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Introducer, Catheter (DYB)
Date received	Nov 18, 2008
Decision date	Dec 15, 2008
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**St Jude Medical**
Location	Minnetonka, MN, US
Contact	MAC MCKEEN
Website	http://www.sjm.com/
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...