

**K014037 SJM SEGUIN ANNULOPLASTY RING (SEGUIN RING),  
MODEL SARP-(SIZE)**Jan 4, 2002  
28 days to decisionK014037 · Product code: KRH · Cardiovascular  
Source: <https://www.510kdatabase.net/k014037/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Ring, Annuloplasty (KRH)
Date received	Dec 7, 2001
Decision date	Jan 4, 2002
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>St. Jude Medical, Inc.</b>
Location	Salt Lake City, UT, US
Contact	WILLIAM MCKELVEY
Website	<a href="http://www.sjm.com/">http://www.sjm.com/</a>
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...