

**K131816 SORIN CONNECT DATA MANAGEMENT SYSTEM**Jul 23, 2013  
33 days to decisionK131816 · Product code: **DXJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k131816/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Jun 20, 2013
Decision date	Jul 23, 2013
Days to decision	33 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sorin Group Deutschland GmbH</b>
Location	North Attleboro, MA, US
Contact	RENATE GOEBERT
510(k) history	20 submissions · 20 cleared · 2006-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131816/> 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 May 24, 2026