

**K200950 Ventricular Catheter**May 8, 2020  
29 days to decisionK200950 · Product code: **JXG** · Neurology  
Source: <https://www.510kdatabase.net/k200950/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Shunt, Central Nervous System And Components (JXG)
Date received	Apr 9, 2020
Decision date	May 8, 2020
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	Kathy A. Racosky
510(k) history	207 submissions · 201 cleared · 1991-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200950/> 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 14, 2026