

**K992318 MEDTRONIC AVE BRIDGE STENT**Aug 2, 1999  
21 days to decisionK992318 · Product code: **FGE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k992318/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Jul 12, 1999
Decision date	Aug 2, 1999
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Ave, Inc.</b>
Location	Santa Rosa, CA, US
Contact	SUSAN WALTON
510(k) history	13 submissions · 2 cleared · 1999-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992318/> 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