

K992569 MEDTRONIC AVE BRIDGE STENTAug 31, 1999
29 days to decisionK992569 · Product code: **FGE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k992569/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Aug 2, 1999
Decision date	Aug 31, 1999
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992569/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026