FDA 510(k) Premarket Notification

K992569 MEDTRONIC AVE BRIDGE STENT

CLEARED

Aug 31, 1999
29 days to decision

K992569 · Product code: **FGE** · Gastroenterology & Urology
Source: 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