FDA 510(k) Premarket Notification

K992318 MEDTRONIC AVE BRIDGE STENT

CLEARED

Aug 2, 1999
21 days to decision

K992318 · Product code: **FGE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k992318/

SUBMISSION DETAILS

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

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