

K032929 MODIFICATION TO:XCEED NITINOL SELF-EXPANDING TRANSHEPATIC BILIARY STENT SYSTEM, MODELS 14819, 14820, 14821, 14822, 14822Oct 22, 2003
30 days to decisionK032929 · Product code: **FGE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k032929/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Sep 22, 2003
Decision date	Oct 22, 2003
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Abbott Vascular, Inc.
Location	Redwood, CA, US
Contact	JOANNA KUSKOWSKI
510(k) history	20 submissions · 17 cleared · 2000-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032929/> 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