

**K122546 SUPERA VERITAS INTERWOVEN SELF-EXPANDING
NITINOL STENT TRANSHEPATIC BILIARY SYSTEM**Dec 7, 2012
108 days to decisionK122546 · Product code: **FGE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k122546/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Aug 21, 2012
Decision date	Dec 7, 2012
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Idev Technologies, Inc.
Location	Houston, TX, US
Contact	DARLENE GARNER
510(k) history	11 submissions · 4 cleared · 2003-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122546/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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