

**K130792 LIFESTENT BILIARY STENT SYSTEM, LIFESTENT XL
BILIARY STENT SYSTEM**Aug 30, 2013
161 days to decisionK130792 · Product code: **FGE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k130792/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Mar 22, 2013
Decision date	Aug 30, 2013
Days to decision	161 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
Contact	Christoph Wagner von Hoff
Website	https://www.bd.com
510(k) history	645 submissions · 609 cleared · 1976-2026

C.R. Bard, Inc. is a developer, manufacturer, and marketer of medical technologies headquartered in Covington, US. The company specializes in vascular medicine, urology, oncology, and surgical specialty devices. C.R. Bard maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions spanning 1976 to 2026. The company's portfolio encompasses cardiovascular devices, gastroenterology and urology products, and general surgical technologies. Recent clearances include temporary pacing electrode catheters, thrombectomy systems, and ...

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Device record: <https://www.510kdatabase.net/k130792/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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