CLEARED

K993588 MODIFICATION TO RX HERCULINK 14 BILIARY STENT SYSTEM, MODELS 1005126-13, 1005128-13, 1005130-13, 1005132-13, 1005134-13,

Nov 17, 1999
26 days to decision

K993588 · Product code: **FGE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k993588/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Oct 22, 1999
Decision date	Nov 17, 1999
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Guidant Corp.**
Location	Santa Clara, CA, US
Contact	SANTA SUNDELL
510(k) history	71 submissions · 56 cleared · 1997-2006

Guidant Corp. is a medical device manufacturer specializing in cardiovascular devices and surgical products. Headquartered in Indianapolis, Indiana, the company designs and manufactures artificial cardiac pacemakers, implantable cardioverter-defibrillators, stents, and related cardiovascular medical products. Guidant received FDA 510(k) clearances from total submissions between 1997 and 2006. The company's regulatory portfolio is dominated by cardiovascular devices, including guide wires, embolic protection systems, stents, and hemostasis valves. The company also cleared ...