FDA 510(k) Premarket Notification

CLEARED

K211410 GlidePath 13F Long-Term Hemodialysis Catheter

Jun 4, 2021
29 days to decision

K211410 · Product code: **MSD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k211410/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Special
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	May 6, 2021
Decision date	Jun 4, 2021
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Bard Peripheral Vascular, Inc.**
Location	Tempe, AZ, US
Contact	Joan Bergstrom
Website	https://www.bd.com
510(k) history	34 submissions · 30 cleared · 2004-2026

Bard Peripheral Vascular, Inc. is a medical device manufacturer based in Tempe, Arizona. The company specializes in cardiovascular and surgical devices for minimally invasive procedures. FDA 510(k) regulatory activity spans from 2004 to 2026. The company has received FDA 510(k) clearances from total submissions. Cardiovascular devices represent a dominant category, including PTA balloons, atherectomy systems, and vascular access solutions. The company remains actively engaged in device development, with the latest clearance in 2026. Recent cleared devices reflect expertis...