

K961841 BARD V-MAX ROLLER BAR (3551XX)Oct 9, 1996
149 days to decisionK961841 · Product code: **FAS** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k961841/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrosurgical, Active, Urological (FAS)
Date received	May 13, 1996
Decision date	Oct 9, 1996
Days to decision	149 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	XX=01,02,03,04,05, OR 06)

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
Contact	EVANS YI-WEN WUU
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 ...