FDA 510(k) Premarket Notification

K810314 WILLIAM HARVEY FLOWMETER SYSTEM

CLEARED

Feb 19, 1981
14 days to decision

K810314 · Product code: **BYM** · Anesthesiology
Source: https://www.510kdatabase.net/k810314/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Thorpe, Uncompensated (BYM)
Date received	Feb 5, 1981
Decision date	Feb 19, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**C.R. Bard, Inc.**
Location	Covington, GA, US
Website	https://www.bd.com
510(k) history	644 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 ...