FDA 510(k) Premarket Notification

K803037 WILLIAM HARVEY VENTED MEDIASTINAL DRAIN

CLEARED

Feb 4, 1981
65 days to decision

K803037 · Product code: **DWM** · Anesthesiology
Source: https://www.510kdatabase.net/k803037/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Patient Care (DWM)
Date received	Dec 1, 1980
Decision date	Feb 4, 1981
Days to decision	65 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 ...