FDA 510(k) Premarket Notification

CLEARED

K960176 DAVOL NASOGASTRIC SUMP TUBE W/PREVENT FILTER

Jul 24, 1996
190 days to decision

K960176 · Product code: **FEG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k960176/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Double Lumen For Intestinal Decompression And/or Intubation (FEG)
Date received	Jan 16, 1996
Decision date	Jul 24, 1996
Days to decision	190 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**C.R. Bard, Inc.**
Location	Covington, GA, US
Contact	ROBIN DRAGO
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 ...