FDA 510(k) Premarket Notification

K971137 BARD RAPIDFIRE MULTIPLE BAND LIGATOR

CLEARED

Oct 16, 1997
202 days to decision

K971137 · Product code: **MND** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k971137/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ligator, Esophageal (MND)
Date received	Mar 28, 1997
Decision date	Oct 16, 1997
Days to decision	202 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	BETH A ROCHETTE
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 ...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k971137/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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