

**K904779 BUTTON REPLACEMENT GASTROSTOMY DEVICE**Jan 18, 1991  
88 days to decisionK904779 · Product code: **KGC** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k904779/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Gastro-enterostomy (KGC)
Date received	Oct 22, 1990
Decision date	Jan 18, 1991
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No

**APPLICANT**

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Company	<b>C.R. Bard, Inc.</b>
Location	Covington, GA, US
Contact	ERNEST MANFREDO
Website	<a href="https://www.bd.com">https://www.bd.com</a>
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 ...

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