

K890152 USCI PROBING CATHETERAug 17, 1989
216 days to decisionK890152 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k890152/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 13, 1989
Decision date	Aug 17, 1989
Days to decision	216 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
Contact	JANICE T PIASECKI
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 ...
