

K771394 ELECTRONIC STETHESCOPENov 9, 1977
104 days to decisionK771394 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k771394/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jul 28, 1977
Decision date	Nov 9, 1977
Days to decision	104 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
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 ...
