

K772295 INSPIRON MANIFOLD (SINGLE)Jan 17, 1978
35 days to decisionK772295 · Product code: **CAF** · AnesthesiologySource: <https://www.510kdatabase.net/k772295/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Dec 13, 1977
Decision date	Jan 17, 1978
Days to decision	35 days
Third-party review	No

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
Website	https://www.bd.com
510(k) history	644 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 ...
