FDA 510(k) Premarket Notification

CLEARED

K884063 BARD ABG 100 INTRAVASCULAR OXYGEN MONITOR SYSTEM

Dec 16, 1988
81 days to decision

K884063 · Product code: **CCE** · Anesthesiology
Source: https://www.510kdatabase.net/k884063/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Partial Pressure, Blood-phase, Indwelling (CCE)
Date received	Sep 26, 1988
Decision date	Dec 16, 1988
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**C.R. Bard, Inc.**
Location	Covington, GA, US
Contact	KEVIN E DALY
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 ...