

**K872125 BARD ABG 100 INTRAVASCULAR OXYGEN MONITOR.
SYSTEM**Dec 11, 1987
192 days to decisionK872125 · Product code: **CCE** · Anesthesiology
Source: <https://www.510kdatabase.net/k872125/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Partial Pressure, Blood-phase, Indwelling (CCE)
Date received	Jun 2, 1987
Decision date	Dec 11, 1987
Days to decision	192 days
Third-party review	No

APPLICANT

Company	C.R. Bard, Inc.
Location	Covington, GA, US
Contact	JOSEPH KICEINA
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 ...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k872125/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026