FDA 510(k) Premarket Notification

K141936 MICROBLAB, MICROLOOP

CLEARED

Dec 29, 2014
165 days to decision

K141936 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k141936/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jul 17, 2014
Decision date	Dec 29, 2014
Days to decision	165 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Carefusion Germany 234 GmbH**
Location	Hoechberg, DE
Contact	ELMAR NIEDERMEYER
510(k) history	12 submissions · 12 cleared · 2010-2016

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k141936/ 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 24, 2026