

K925459 GS-MODULAR (GOLD STANDARD - MODULAR)Feb 4, 1994
463 days to decisionK925459 · Product code: **BZG** · Anesthesiology
Source: <https://www.510kdatabase.net/k925459/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Oct 29, 1992
Decision date	Feb 4, 1994
Days to decision	463 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Warren E. Collins, Inc.
Location	Mchenry, IL, US
Contact	JAMES J TULLY
510(k) history	14 submissions · 14 cleared · 1978-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925459/> 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 26, 2026