

K914272 KOKO SPIROMETERJul 2, 1992
282 days to decisionK914272 · Product code: **BZG** · Anesthesiology
Source: <https://www.510kdatabase.net/k914272/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Sep 24, 1991
Decision date	Jul 2, 1992
Days to decision	282 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ferraris Respiratory, Inc.
Location	Louisville, CO, US
Contact	ARLIN LEHMAN
510(k) history	2 submissions · 2 cleared · 1992-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914272/> 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 25, 2026