FDA 510(k) Premarket Notification

K903674 MODEL 295 ELECTRONIC RESPIROMETER

CLEARED

Jan 4, 1991
144 days to decision

K903674 · Product code: **BZK** · Anesthesiology
Source: https://www.510kdatabase.net/k903674/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Monitoring (w/wo Alarm) (BZK)
Date received	Aug 13, 1990
Decision date	Jan 4, 1991
Days to decision	144 days
Third-party review	No

APPLICANT

Company	**National Medical Sales, Inc.**
Location	Mission Viejo, CA, US
Contact	WILLIAM D BURCH
510(k) history	2 submissions · 2 cleared · 1986-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k903674/ 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