

K931542 NEWPORT COMPASS EXHALED TIDAL VOLUME MONITOROct 25, 1993
210 days to decisionK931542 · Product code: **BZK** · Anesthesiology
Source: <https://www.510kdatabase.net/k931542/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Monitoring (w/wo Alarm) (BZK)
Date received	Mar 29, 1993
Decision date	Oct 25, 1993
Days to decision	210 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Newport Medical Instruments, Inc.
Location	Mchenry, IL, US
Contact	CYNDY MILLER
510(k) history	22 submissions · 19 cleared · 1982-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931542/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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