FDA 510(k) Premarket Notification

K931542 NEWPORT COMPASS EXHALED TIDAL VOLUME MONITOR

CLEARED

Oct 25, 1993
210 days to decision

K931542 · 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