FDA 510(k) Premarket Notification

CLEARED

K142303 NeumoFilt Ergo/Neumo Filt BiteOn

Nov 20, 2014
94 days to decision

K142303 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k142303/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Aug 18, 2014
Decision date	Nov 20, 2014
Days to decision	94 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medical Respiratory Devices S.L.**
Location	Coslada(Madrid), ES
Contact	Jose Alberto Hernandez
510(k) history	1 submissions · 1 cleared · 2014-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k142303/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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