FDA 510(k) Premarket Notification

K141152 ADVANTAGE

CLEARED

Oct 17, 2014
165 days to decision

K141152 · Product code: **BZD** · Anesthesiology
Source: https://www.510kdatabase.net/k141152/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	May 5, 2014
Decision date	Oct 17, 2014
Days to decision	165 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stevenson Industries, Inc.**
Location	Simi Valley, CA, US
Contact	SERRAH NAMINI
510(k) history	2 submissions · 2 cleared · 1999-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k141152/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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