FDA 510(k) Premarket Notification

K030843 VPAP III

CLEARED

Aug 15, 2003
151 days to decision

K030843 · Product code: **BZD** · Anesthesiology
Source: https://www.510kdatabase.net/k030843/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	Mar 17, 2003
Decision date	Aug 15, 2003
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Resmed, Ltd.**
Location	Poway, CA, US
Contact	ROGER KOTTER
Website	http://www.resmed.com/
510(k) history	103 submissions · 103 cleared · 1996-2019