FDA 510(k) Premarket Notification

K090888 NEWPORT HT70 FAMILY OF VENTILATORS

CLEARED

Nov 18, 2009
232 days to decision

K090888 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k090888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Mar 31, 2009
Decision date	Nov 18, 2009
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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