FDA 510(k) Premarket Notification

K093094 VELA VENTILATOR

CLEARED

May 11, 2010
222 days to decision

K093094 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k093094/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Oct 1, 2009
Decision date	May 11, 2010
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardinal Health 207, Inc.**
Location	Yorba Linda, CA, US
Contact	FAROKH ETEMADIEH
510(k) history	5 submissions · 5 cleared · 2008-2010

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k093094/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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