

K090888 NEWPORT HT70 FAMILY OF VENTILATORSNov 18, 2009
232 days to decisionK090888 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 24, 2026