

K031863 CARESTATION 126S VIDEOPHONEOct 10, 2003
116 days to decisionK031863 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k031863/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jun 16, 2003
Decision date	Oct 10, 2003
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Motion Media Technology, Inc.
Location	Wilmington, NC, US
Contact	STUART R ROSS
510(k) history	1 submissions · 1 cleared · 2003-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k031863/> 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 July 3, 2026