

K032142 DATACTORAug 8, 2003
25 days to decisionK032142 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k032142/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jul 14, 2003
Decision date	Aug 8, 2003
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Capsule Technologie
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	5 submissions · 5 cleared · 2001-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032142/> 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 June 28, 2026