

K161517 Mortara Surveyor Patient MonitorJan 11, 2017
223 days to decisionK161517 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k161517/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 2, 2016
Decision date	Jan 11, 2017
Days to decision	223 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mortara Instrument, Inc.
Location	Walker, MI, US
Contact	SARAH WEBER
510(k) history	51 submissions · 51 cleared · 1983-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161517/> 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 19, 2026