

K173765 Surveyor Patient MonitorSep 4, 2018
267 days to decisionK173765 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k173765/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Dec 11, 2017
Decision date	Sep 4, 2018
Days to decision	267 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173765/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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