

K141020 SURVEYOR S4 MOBILE MONITORDec 3, 2014
226 days to decisionK141020 · Product code: **DRG** · Cardiovascular
Source: <https://www.510kdatabase.net/k141020/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Apr 21, 2014
Decision date	Dec 3, 2014
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k141020/> 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