FDA 510(k) Premarket Notification

CLEARED

K161517 Mortara Surveyor Patient Monitor

Jan 11, 2017
223 days to decision

K161517 · Product code: **MHX** · Cardiovascular
Source: 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