

K060135 MORTARA SURVEYOR TELEMETRY CENTRAL STATION

May 15, 2006
116 days to decision

K060135 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k060135/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jan 19, 2006
Decision date	May 15, 2006
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mortara Instrument, Inc.
Location	Walker, MI, US
Contact	HARLAN L VAN MATRE
510(k) history	51 submissions · 51 cleared · 1983-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k060135/> 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