

K001359 DASH 3000/4000 PATIENT MONITORJul 18, 2000
81 days to decisionK001359 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k001359/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 28, 2000
Decision date	Jul 18, 2000
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	General Electric Medical Systems Information Techn
Location	Sugarland, TX, US
Contact	DAVID WAHLIG
510(k) history	33 submissions · 33 cleared · 1999-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001359/> 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 July 3, 2026