FDA 510(k) Premarket Notification

K001359 DASH 3000/4000 PATIENT MONITOR

CLEARED

Jul 18, 2000
81 days to decision

K001359 · 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