

K962291 SIEMENS SC9000/SC9015 BEDSIDE MONITORING SYSTEMJan 29, 1997
229 days to decisionK962291 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k962291/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jun 14, 1996
Decision date	Jan 29, 1997
Days to decision	229 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	Jacqueline Emery
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962291/> 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 May 15, 2026