

K952054 SIEMENS SIRECUST MONITORING (1200, 1200 NEO, 960)Jun 7, 1996
403 days to decisionK952054 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k952054/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	May 1, 1995
Decision date	Jun 7, 1996
Days to decision	403 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k952054/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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