FDA 510(k) Premarket Notification

K962291 SIEMENS SC9000/SC9015 BEDSIDE MONITORING SYSTEM

CLEARED

Jan 29, 1997
229 days to decision

K962291 · 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