FDA 510(k) Premarket Notification

K980625 SIEMENS MULTIVIEW WORKSTATION ENHANCED WITH DIAGNOSTIC STATEMENTS (REST ECG)

CLEARED

May 19, 1998
90 days to decision

K980625 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k980625/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 18, 1998
Decision date	May 19, 1998
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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