FDA 510(k) Premarket Notification

K023569 SIEMENS MULTIVIEW WORKSTATION MODIFICATIONS

CLEARED

Nov 4, 2002
11 days to decision

K023569 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k023569/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Oct 24, 2002
Decision date	Nov 4, 2002
Days to decision	11 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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