

**K023569 SIEMENS MULTIVIEW WORKSTATION  
MODIFICATIONS**Nov 4, 2002  
11 days to decisionK023569 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k023569/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023569/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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