

K022453 MODIFICATION TO ACUITY CENTRAL MONITORING SYSTEMAug 21, 2002
26 days to decisionK022453 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k022453/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 26, 2002
Decision date	Aug 21, 2002
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Welch Allyn Protocol, Inc.
Location	Beaverton, OR, US
Contact	DON M ABBEY
510(k) history	7 submissions · 7 cleared · 2001-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022453/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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