

K841083 QUIKPAGE ARRHYTHMIA REPORT GENERATORFeb 7, 1985
331 days to decisionK841083 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k841083/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 13, 1984
Decision date	Feb 7, 1985
Days to decision	331 days
Third-party review	No

APPLICANT

Company	Agilent Technologies
Location	Walker, MI, US
Contact	WILLIAM B RICH
Website	http://www.agilent.com
510(k) history	13 submissions · 13 cleared · 1982-2001

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