

K812149 CARDIO-DATA SYSTEM, #MARK IIIAug 31, 1981
34 days to decisionK812149 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k812149/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 28, 1981
Decision date	Aug 31, 1981
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Cardio Data
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1977-1985

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Device record: <https://www.510kdatabase.net/k812149/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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