

K820856 EVENTMASTER 2Apr 29, 1982
31 days to decisionK820856 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k820856/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 29, 1982
Decision date	Apr 29, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	Instruments For Cardiac Research
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1977-1984

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Device record: <https://www.510kdatabase.net/k820856/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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