

K821173 EVENTMASTER 3 HOLTER EVENT SCANNERMay 18, 1982
22 days to decisionK821173 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k821173/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Apr 26, 1982
Decision date	May 18, 1982
Days to decision	22 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/k821173/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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