

**K834397 HOLTER CARDIOGRAPHY SYS 6201**Apr 17, 1984  
125 days to decisionK834397 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k834397/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 14, 1983
Decision date	Apr 17, 1984
Days to decision	125 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k834397/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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