

**K811527 CC-103**Jun 16, 1981  
15 days to decisionK811527 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k811527/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jun 1, 1981
Decision date	Jun 16, 1981
Days to decision	15 days
Third-party review	No

**APPLICANT**

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Company	<b>Anthro-Metrics Corp.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1981

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

Device record: <https://www.510kdatabase.net/k811527/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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