

**K802312 CC101 ECG MONITORING SYSTEM**Jan 7, 1981  
106 days to decisionK802312 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k802312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 23, 1980
Decision date	Jan 7, 1981
Days to decision	106 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/k802312/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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