

**K801762 MICRO-PROCESSOR UNIT #A3600AC**Aug 27, 1980  
30 days to decisionK801762 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801762/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 28, 1980
Decision date	Aug 27, 1980
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>General Electric Co.</b>
Location	Mchenry, IL, US
510(k) history	254 submissions · 254 cleared · 1976-2011

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Device record: <https://www.510kdatabase.net/k801762/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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