

**K791632 COMPU-SCAN**Nov 13, 1979  
84 days to decisionK791632 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k791632/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 21, 1979
Decision date	Nov 13, 1979
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Hittman Medical Systems</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1983

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

Device record: <https://www.510kdatabase.net/k791632/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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