

**K900560 ACUSON HEARTSOUND/PULSE/RESPIRATION  
MODULE**Dec 3, 1990  
299 days to decisionK900560 · Product code: **DQC** · Cardiovascular  
Source: <https://www.510kdatabase.net/k900560/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Phonocardiograph (DQC)
Date received	Feb 7, 1990
Decision date	Dec 3, 1990
Days to decision	299 days
Third-party review	No

**APPLICANT**

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Company	<b>Acuson Corp.</b>
Location	Mountain View, CA, US
Contact	T JOHNSON
510(k) history	37 submissions · 37 cleared · 1988-2002

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900560/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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