

**K812608 VASCULAB #R12B**Oct 2, 1981  
17 days to decisionK812608 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k812608/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Sep 15, 1981
Decision date	Oct 2, 1981
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Medasonics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1978-1999

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

Device record: <https://www.510kdatabase.net/k812608/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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