

K820820 CARDIO REVUE CENTERApr 29, 1982
36 days to decisionK820820 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k820820/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Mar 24, 1982
Decision date	Apr 29, 1982
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Diasonics, Inc.
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1978-1997

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Device record: <https://www.510kdatabase.net/k820820/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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