

K771924 HEARTBEATERNov 28, 1977
46 days to decisionK771924 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k771924/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Oct 13, 1977
Decision date	Nov 28, 1977
Days to decision	46 days
Third-party review	No

APPLICANT

Company	Dunn Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1977-1985

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Device record: <https://www.510kdatabase.net/k771924/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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