

K800519 HEARTWATCHSep 26, 1980
204 days to decisionK800519 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k800519/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 6, 1980
Decision date	Sep 26, 1980
Days to decision	204 days
Third-party review	No

APPLICANT

Company	D. Jenkins
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800519/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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