

K781407 ERM-101Sep 27, 1978
44 days to decisionK781407 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k781407/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 14, 1978
Decision date	Sep 27, 1978
Days to decision	44 days
Third-party review	No

APPLICANT

Company	Cynergy
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1981

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Device record: <https://www.510kdatabase.net/k781407/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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