

K780117 CARDIOMETERFeb 3, 1978
11 days to decisionK780117 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k780117/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 23, 1978
Decision date	Feb 3, 1978
Days to decision	11 days
Third-party review	No

APPLICANT

Company	Med General
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1977-1979

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k780117/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 28, 2026