

K820671 CM10 CENTRAL MONITORING SYSTEMApr 1, 1982
21 days to decisionK820671 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k820671/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Mar 11, 1982
Decision date	Apr 1, 1982
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Medtel Pty. , Ltd.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1979-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820671/> 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 27, 2026