

K852887 CR5 CARDIAC MONITORNov 14, 1985
128 days to decisionK852887 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k852887/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Jul 9, 1985
Decision date	Nov 14, 1985
Days to decision	128 days
Third-party review	No

APPLICANT

Company	Cardiac Recorders, Ltd.
Location	London, GB
Contact	KEITH I BEVIS
510(k) history	8 submissions · 8 cleared · 1985-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k852887/>; 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 July 7, 2026