FDA 510(k) Premarket Notification

K852887 CR5 CARDIAC MONITOR

CLEARED

Nov 14, 1985
128 days to decision

K852887 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k852887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & 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