FDA 510(k) Premarket Notification

K853673 CARDIORATER CR 51

CLEARED

Dec 11, 1985
98 days to decision

K853673 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k853673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Sep 4, 1985
Decision date	Dec 11, 1985
Days to decision	98 days
Third-party review	No

APPLICANT

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