FDA 510(k) Premarket Notification

K852808 CR26, CR26A, CR26B, CR26C(CARDIAC DC DEFIBRILLATO)

CLEARED

Sep 16, 1985
76 days to decision

K852808 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k852808/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 2, 1985
Decision date	Sep 16, 1985
Days to decision	76 days
Third-party review	No

APPLICANT

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