FDA 510(k) Premarket Notification

K871251 CIRCAMED FULL DISCLOSURE SYSTEM

CLEARED

May 22, 1987
53 days to decision

K871251 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k871251/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 30, 1987
Decision date	May 22, 1987
Days to decision	53 days
Third-party review	No

APPLICANT

Company	**Circadian, Inc.**
Location	Mchenry, IL, US
Contact	MATT S MAGOUN
510(k) history	13 submissions · 13 cleared · 1980-1992