FDA 510(k) Premarket Notification

K885085 ERGOMED 840/ERGOMED 840L

CLEARED

Oct 3, 1989
298 days to decision

K885085 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k885085/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 9, 1988
Decision date	Oct 3, 1989
Days to decision	298 days
Third-party review	No

APPLICANT

Company	**Burdick Corp.**
Location	Mchenry, IL, US
Contact	FRANCIS I DOMINY
Website	https://www.cardinalhealth.com
510(k) history	38 submissions · 38 cleared · 1976-1994