FDA 510(k) Premarket Notification

K840717 ECS 502

CLEARED

Aug 19, 1985
549 days to decision

K840717 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k840717/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Feb 17, 1984
Decision date	Aug 19, 1985
Days to decision	549 days
Third-party review	No

APPLICANT

Company	**Robert Bosch Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 18 cleared · 1979-1986