FDA 510(k) Premarket Notification

K792312 HEART RATE MONITOR MD-1

CLEARED

Nov 21, 1979
5 days to decision

K792312 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k792312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 16, 1979
Decision date	Nov 21, 1979
Days to decision	5 days
Third-party review	No

APPLICANT

Company	**Medical Devices, Inc.**
Location	Mchenry, IL, US
510(k) history	49 submissions · 47 cleared · 1977-2001