FDA 510(k) Premarket Notification

K772332 MONITOR, CARDIAC, BEDSIDE

CLEARED

Feb 13, 1978
53 days to decision

K772332 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k772332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 22, 1977
Decision date	Feb 13, 1978
Days to decision	53 days
Third-party review	No

APPLICANT

Company	**Telectronics, Inc.**
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990