FDA 510(k) Premarket Notification

K820671 CM10 CENTRAL MONITORING SYSTEM

CLEARED

Apr 1, 1982
21 days to decision

K820671 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k820671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 11, 1982
Decision date	Apr 1, 1982
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Medtel Pty. , Ltd.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1979-1983