FDA 510(k) Premarket Notification

K812271 CENTRAL STATION MONITOR

CLEARED

Aug 31, 1981
18 days to decision

K812271 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k812271/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 13, 1981
Decision date	Aug 31, 1981
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019