FDA 510(k) Premarket Notification

CLEARED

K772056 PHYSIOLOGICAL MONITOR MODEL 871

Nov 22, 1977
22 days to decision

K772056 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k772056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Oct 31, 1977
Decision date	Nov 22, 1977
Days to decision	22 days
Third-party review	No

APPLICANT

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