

K812271 CENTRAL STATION MONITORAug 31, 1981
18 days to decisionK812271 · Product code: **DRT** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k812271/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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