

K772056 PHYSIOLOGICAL MONITOR MODEL 871Nov 22, 1977
22 days to decisionK772056 · Product code: **DRT** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k772056/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 20, 2026