

K832041 SERVOMED STANDARD SMS 104Oct 26, 1983
124 days to decisionK832041 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k832041/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 24, 1983
Decision date	Oct 26, 1983
Days to decision	124 days
Third-party review	No

APPLICANT

Company	Litton Medical Electronics
Location	Walker, MI, US
510(k) history	38 submissions · 38 cleared · 1982-1985

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k832041/> 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 May 27, 2026