FDA 510(k) Premarket Notification

K832041 SERVOMED STANDARD SMS 104

CLEARED

Oct 26, 1983
124 days to decision

K832041 · 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