FDA 510(k) Premarket Notification

CLEARED

K832250 SERVOMED BASIC BEDSIDE UNIT SMA 102-

Jun 1, 1984
326 days to decision

K832250 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k832250/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 11, 1983
Decision date	Jun 1, 1984
Days to decision	326 days
Third-party review	No

APPLICANT

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