FDA 510(k) Premarket Notification

K840523 SERVOMED PULSE AMPLIFIER MODULE-206-

CLEARED

Oct 29, 1984
265 days to decision

K840523 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k840523/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Feb 7, 1984
Decision date	Oct 29, 1984
Days to decision	265 days
Third-party review	No

APPLICANT

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