

K834600 RECOMB ECG PREAMPLIFIER/206 069 03Jun 1, 1984
154 days to decisionK834600 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k834600/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 30, 1983
Decision date	Jun 1, 1984
Days to decision	154 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k834600/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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