FDA 510(k) Premarket Notification

K833000 I-CHANNEL RECORDER

CLEARED

Jun 1, 1984
288 days to decision

K833000 · Product code: **DSF** · Cardiovascular
Source: https://www.510kdatabase.net/k833000/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Aug 18, 1983
Decision date	Jun 1, 1984
Days to decision	288 days
Third-party review	No

APPLICANT

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