FDA 510(k) Premarket Notification

K820881 MEMOPORT C

CLEARED

May 18, 1982
49 days to decision

K820881 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k820881/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 30, 1982
Decision date	May 18, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820881/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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