FDA 510(k) Premarket Notification

K822432 CENTURION CHART RECORDER

CLEARED

Nov 1, 1982
81 days to decision

K822432 · Product code: **DSF** · Cardiovascular
Source: https://www.510kdatabase.net/k822432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Aug 12, 1982
Decision date	Nov 1, 1982
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**Clinical Data, Inc.**
Location	Mchenry, IL, US
510(k) history	36 submissions · 36 cleared · 1982-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k822432/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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