

K801485 SERIES 602 CKG RECORDER (MODIFIED)Jul 8, 1980
14 days to decisionK801485 · Product code: **DSF** · Cardiovascular
Source: <https://www.510kdatabase.net/k801485/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Jun 24, 1980
Decision date	Jul 8, 1980
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Cardio Kinetics, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1979-1982

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Device record: <https://www.510kdatabase.net/k801485/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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