FDA 510(k) Premarket Notification

K801485 SERIES 602 CKG RECORDER (MODIFIED)

CLEARED

Jul 8, 1980
14 days to decision

K801485 · 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