

K792051 CKG CARDIOKYMOGRAPHNov 13, 1979
29 days to decisionK792051 · Product code: **DSB** · CardiovascularSource: <https://www.510kdatabase.net/k792051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Impedance (DSB)
Date received	Oct 15, 1979
Decision date	Nov 13, 1979
Days to decision	29 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k792051/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 9, 2026