

K812586 KANTRON 504 HEARTSTATIONDec 29, 1981
109 days to decisionK812586 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k812586/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 11, 1981
Decision date	Dec 29, 1981
Days to decision	109 days
Third-party review	No

APPLICANT

Company	Kantron Cardiovascular, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812586/> 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 May 28, 2026