

K854307 KONTRON DEFIBRILLATOR 7501Dec 4, 1985
40 days to decisionK854307 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k854307/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 25, 1985
Decision date	Dec 4, 1985
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Kontron Instruments, Inc.
Location	Mchenry, IL, US
Contact	GEORGE CHO
510(k) history	57 submissions · 57 cleared · 1981-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k854307/>; 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 21, 2026