FDA 510(k) Premarket Notification

K854307 KONTRON DEFIBRILLATOR 7501

CLEARED

Dec 4, 1985
40 days to decision

K854307 · Product code: **LDD** · Cardiovascular
Source: 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