FDA 510(k) Premarket Notification

K874204 450SLL DEFIBRILLATOR

CLEARED

Jan 28, 1988
105 days to decision

K874204 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k874204/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 15, 1987
Decision date	Jan 28, 1988
Days to decision	105 days
Third-party review	No

APPLICANT

Company	**Medical Research Laboratories, Inc.**
Location	Mchenry, IL, US
Contact	DILIP MEHTA
510(k) history	19 submissions · 15 cleared · 1981-2002