FDA 510(k) Premarket Notification

K902429 MODIFIED HEARTSTART DEFIBRILLATION ELECTRODE

CLEARED

Sep 27, 1990
128 days to decision

K902429 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k902429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	May 22, 1990
Decision date	Sep 27, 1990
Days to decision	128 days
Third-party review	No

APPLICANT

Company	**Laerdal Mfg. Corp.**
Location	Portland, OR, US
Contact	JOHN L KARPOWICZ
510(k) history	5 submissions · 5 cleared · 1988-1990