FDA 510(k) Premarket Notification

K900169 MODIFIED HEARTSTART 2000 DEFIBRILLATION ELECTRODE

CLEARED

Jan 26, 1990
14 days to decision

K900169 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k900169/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jan 12, 1990
Decision date	Jan 26, 1990
Days to decision	14 days
Third-party review	No

APPLICANT

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