FDA 510(k) Premarket Notification

K812519 2000 SERIES PORTABLE RESUSCITATION SYST.

CLEARED

Sep 25, 1981
23 days to decision

K812519 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k812519/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 2, 1981
Decision date	Sep 25, 1981
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Life Science Instrumentation, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985