FDA 510(k) Premarket Notification

K822294 VPD-261 DEFIBRILLATOR MONITOR

CLEARED

Aug 31, 1982
29 days to decision

K822294 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k822294/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 2, 1982
Decision date	Aug 31, 1982
Days to decision	29 days
Third-party review	No

APPLICANT

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