FDA 510(k) Premarket Notification

K895533 LIFE DEFENSE PLUS(TM), DEFIBRILLATOR/MONITOR/PACER

CLEARED

Jan 26, 1990
135 days to decision

K895533 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k895533/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 13, 1989
Decision date	Jan 26, 1990
Days to decision	135 days
Third-party review	No

APPLICANT

Company	**Matrix Medica, Inc.**
Location	Minden, NV, US
Contact	WYNNE, P.E.
510(k) history	14 submissions · 13 cleared · 1985-1996