

**K895533 LIFE DEFENSE PLUS(TM),
DEFIBRILLATOR/MONITOR/PACER**Jan 26, 1990
135 days to decisionK895533 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895533/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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