FDA 510(k) Premarket Notification

K991241 IF8000

CLEARED

Nov 2, 1999
204 days to decision

K991241 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k991241/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Apr 12, 1999
Decision date	Nov 2, 1999
Days to decision	204 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dan Med, Inc.**
Location	Denver, CO, US
Contact	THOMAS SANDGAARD
510(k) history	6 submissions · 6 cleared · 1997-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k991241/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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