FDA 510(k) Premarket Notification

K790033 HVG-100

CLEARED

Feb 9, 1979
37 days to decision

K790033 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k790033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jan 3, 1979
Decision date	Feb 9, 1979
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Elmed, Inc.**
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-2001