

K790033 HVG-100Feb 9, 1979
37 days to decisionK790033 · Product code: **IPF** · Physical MedicineSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k790033/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026