FDA 510(k) Premarket Notification

# K852963 MYO-ACTUATOR

**CLEARED**

Jan 21, 1986
193 days to decision

K852963 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k852963/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Device, Biofeedback (HCC) |
| Date received | Jul 12, 1985 |
| Decision date | Jan 21, 1986 |
| Days to decision | 193 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Verimed Holdings, Inc.** |
| Location | Mchenry, IL, US |
| Contact | WILLIAM W MEE |
| 510(k) history | 12 submissions · 12 cleared · 1983-1998 |