

**K852963 MYO-ACTUATOR**Jan 21, 1986  
193 days to decisionK852963 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k852963/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852963/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026