

K832714 VERIMED MYOEXORCISEROct 28, 1983
77 days to decisionK832714 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k832714/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Aug 12, 1983
Decision date	Oct 28, 1983
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Verimed Holdings, Inc.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1983-1998

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Device record: <https://www.510kdatabase.net/k832714/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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