FDA 510(k) Premarket Notification

K832714 VERIMED MYOEXORCISER

CLEARED

Oct 28, 1983
77 days to decision

K832714 · 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