FDA 510(k) Premarket Notification

K823511 EMG BIOFEEDBACK DEVICE

CLEARED

Mar 31, 1983
122 days to decision

K823511 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k823511/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Nov 29, 1982
Decision date	Mar 31, 1983
Days to decision	122 days
Third-party review	No

APPLICANT

Company	**Nemectron Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 12 cleared · 1980-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k823511/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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