FDA 510(k) Premarket Notification

# K842354 EMG MODEL 133

**CLEARED**

Jul 17, 1984
71 days to decision

K842354 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k842354/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Device, Biofeedback (HCC) |
| Date received | May 7, 1984 |
| Decision date | Jul 17, 1984 |
| Days to decision | 71 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Calnor of EL Paseo, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 6 submissions · 6 cleared · 1977-1984 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k842354/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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