

K841895 GSR, 126, EEG 130-TEMP. 132, EMG 133Jul 17, 1984
71 days to decisionK841895 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k841895/>**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/k841895/> Data sourced from FDA 510(k) public records (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. - Generated July 1, 2026