FDA 510(k) Premarket Notification

K841895 GSR, 126, EEG 130-TEMP. 132, EMG 133

CLEARED

Jul 17, 1984
71 days to decision

K841895 · 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