FDA 510(k) Premarket Notification

K960170 ELECTROENCEPHALOGRAPH (EEG)

CLEARED

Mar 1, 1996
45 days to decision

K960170 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k960170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jan 16, 1996
Decision date	Mar 1, 1996
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Moberg Medical, Inc.**
Location	Ambler, PA, US
Contact	LARRY ENGLE
510(k) history	2 submissions · 2 cleared · 1992-1996