FDA 510(k) Premarket Notification

K122879 EEGER4 MODEL 4.3

CLEARED

Feb 6, 2013
140 days to decision

K122879 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k122879/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Sep 19, 2012
Decision date	Feb 6, 2013
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Eeg Software, LLC**
Location	Northridge, CA, US
Contact	HOWARD LIGHTSTONE
510(k) history	1 submissions · 1 cleared · 2013-2013