FDA 510(k) Premarket Notification

K990538 BRAINMASTER 2E MODULE AND SOFTWARE BIOFEEDBACK DEVICE

CLEARED

May 19, 1999
89 days to decision

K990538 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k990538/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 19, 1999
Decision date	May 19, 1999
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Brainmaster**
Location	Great Neck, NY, US
Contact	SUSAN D GOLDSTEIN-FALK
510(k) history	1 submissions · 1 cleared · 1999-1999