

**K990538 BRAINMASTER 2E MODULE AND SOFTWARE
BIOFEEDBACK DEVICE**May 19, 1999
89 days to decisionK990538 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990538/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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