

K811848 MODEL SIX BEAM SYSTEMSep 8, 1981
69 days to decisionK811848 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k811848/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jul 1, 1981
Decision date	Sep 8, 1981
Days to decision	69 days
Third-party review	No

APPLICANT

Company	Braintech, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k811848/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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