

K832333 FEEDBACK EMG SYSTEM PE-020Sep 1, 1983
45 days to decisionK832333 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k832333/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Jul 18, 1983
Decision date	Sep 1, 1983
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Bio-Feedback Systems, Inc.
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1977-1991

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Device record: <https://www.510kdatabase.net/k832333/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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