

K802390 EMG-UT 30Jan 9, 1981
99 days to decisionK802390 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k802390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Oct 2, 1980
Decision date	Jan 9, 1981
Days to decision	99 days
Third-party review	No

APPLICANT

Company	American Scientific Corp.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k802390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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