

K780539 STIMULATOR MODEL 8021Jun 22, 1978
80 days to decisionK780539 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k780539/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Apr 3, 1978
Decision date	Jun 22, 1978
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Life-Tech Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1977-1981

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Device record: <https://www.510kdatabase.net/k780539/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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