

K810182 PROGRAMMABLE SPINAL CORAL STIM. SYSTEMApr 3, 1981
71 days to decisionK810182 · Product code: **GZB** · Neurology
Source: <https://www.510kdatabase.net/k810182/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Jan 22, 1981
Decision date	Apr 3, 1981
Days to decision	71 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k810182/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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