

K862915 MULTISTIM(4-CHANNEL), MULTISTIM-OCTRODE(8 CHANNEL)Oct 6, 1986
66 days to decisionK862915 · Product code: **GZB** · Neurology
Source: <https://www.510kdatabase.net/k862915/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Aug 1, 1986
Decision date	Oct 6, 1986
Days to decision	66 days
Third-party review	No

APPLICANT

Company	Neuromed, Inc.
Location	Ft. Lauderdale, FL, US
Contact	FRANK LISKA
510(k) history	15 submissions · 15 cleared · 1985-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862915/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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