

**K833809 NEURAL STIMULATOR MONITOR**Dec 1, 1983  
29 days to decisionK833809 · Product code: **GZB** · Neurology  
Source: <https://www.510kdatabase.net/k833809/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Nov 2, 1983
Decision date	Dec 1, 1983
Days to decision	29 days
Third-party review	No

**APPLICANT**

---

Company	<b>Instromedix, Inc.</b>
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1977-1997

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833809/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026