

**K843822 MEA110**Oct 22, 1984  
24 days to decisionK843822 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k843822/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 28, 1984
Decision date	Oct 22, 1984
Days to decision	24 days
Third-party review	No

**APPLICANT**

---

Company	<b>Canadian Medical Products</b>
Location	Mchenry, IL, US
Contact	ALAN L RUSSELL
510(k) history	2 submissions · 2 cleared · 1982-1984

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843822/>, 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 July 1, 2026