

**K823299 MODEL 402P SEQUENCER FOR EMPI TEMS DEV**Dec 15, 1982  
40 days to decisionK823299 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k823299/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 5, 1982
Decision date	Dec 15, 1982
Days to decision	40 days
Third-party review	No

**APPLICANT**

---

Company	<b>Epmi, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823299/> 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 21, 2026