

**K833037 POINTER F-3**Nov 29, 1983  
83 days to decisionK833037 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k833037/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 7, 1983
Decision date	Nov 29, 1983
Days to decision	83 days
Third-party review	No

**APPLICANT**

---

Company	<b>Petra Intl. Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

---

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

Device record: <https://www.510kdatabase.net/k833037/> 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