

**K955470 ACTION POTENTIAL SIMULATION THERAPY DEVICE**Oct 22, 1996  
327 days to decisionK955470 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k955470/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 30, 1995
Decision date	Oct 22, 1996
Days to decision	327 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Neurotech, Inc.</b>
Location	Washington, DC, US
Contact	J J M WILKINS
510(k) history	3 submissions · 3 cleared · 1986-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955470/> 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 30, 2026