

**K903017 MEGA-TENS DUAL CHANNEL T.E.N.S.**Oct 26, 1990  
108 days to decisionK903017 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k903017/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 10, 1990
Decision date	Oct 26, 1990
Days to decision	108 days
Third-party review	No

**APPLICANT**

---

Company	<b>Futuremed Div. of Future Impex Corp.</b>
Location	Deerpark, NY, US
Contact	MIKE DAVIDSON
510(k) history	17 submissions · 17 cleared · 1985-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903017/>; 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 May 26, 2026