

**K931424 MAXIMA I**Jun 4, 1993  
74 days to decisionK931424 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k931424/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 22, 1993
Decision date	Jun 4, 1993
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Stadyne, Inc.</b>
Location	Longmont, CO, US
Contact	THOMAS W ROBIRDS
510(k) history	9 submissions · 9 cleared · 1992-1995

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931424/> 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 29, 2026