

**K871666 MODEL HP (OR MODEL 1001) TRANSCUTANEOUS STIMULATOR**

Jun 25, 1987  
58 days to decision

K871666 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k871666/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 28, 1987
Decision date	Jun 25, 1987
Days to decision	58 days
Third-party review	No

**APPLICANT**

---

Company	<b>Oriental Medical Supplies, Inc.</b>
Location	Braintree, MA, US
Contact	THOMAS RIIHIMAKI
510(k) history	6 submissions · 6 cleared · 1985-1989

---

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

Device record: <https://www.510kdatabase.net/k871666/> Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026