

**K033857 T-TECH DIGITAL, MODEL WL-22XXA SERIES**Dec 19, 2003  
8 days to decisionK033857 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k033857/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 11, 2003
Decision date	Dec 19, 2003
Days to decision	8 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Well-Life Healthcare, Inc.</b>
Location	Taichung, R.O.C., TW
Contact	JENNY HSIAH
510(k) history	6 submissions · 6 cleared · 2002-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033857/> 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