

K120398 WITOUCH PROAug 16, 2012
190 days to decisionK120398 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k120398/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ) |
| Date received | Feb 8, 2012 |
| Decision date | Aug 16, 2012 |
| Days to decision | 190 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Hollywog, LLC |
| Location | Chattanooga, TN, US |
| Contact | MICHAEL W TREAS |
| 510(k) history | 4 submissions · 4 cleared · 2012-2017 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120398/> 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 May 31, 2026