

K833354 AUTO-TENS MODEL VI DUAL CHANNEL BATTNov 28, 1983
61 days to decisionK833354 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k833354/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 28, 1983
Decision date	Nov 28, 1983
Days to decision	61 days
Third-party review	No

APPLICANT

Company	Physio Tronics Corp.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k833354/>; 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 July 1, 2026