

**K051773 MODEL 3873 1* 8 AND MODEL 3874 1* 8 COMPACT
TEST STIMULATION LEADS**Jul 28, 2005
27 days to decisionK051773 · Product code: **GZB** · Neurology
Source: <https://www.510kdatabase.net/k051773/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Jul 1, 2005
Decision date	Jul 28, 2005
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Inc.,Neurological Division.
Location	Minneapolis,, MN, US
Contact	PAULA CORDERO
510(k) history	1 submissions · 1 cleared · 2005-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051773/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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