

**K013120 PERCUTANEOUS INTRODUCER SET, MODEL 042294
AND KIT MODEL 3550-18**Jan 24, 2002
128 days to decisionK013120 · Product code: **GZB** · Neurology
Source: <https://www.510kdatabase.net/k013120/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Sep 18, 2001
Decision date	Jan 24, 2002
Days to decision	128 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medamicus, Inc.
Location	Minneapolis, MN, US
Contact	DENNIS S MADISON
510(k) history	20 submissions · 20 cleared · 1989-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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