

K171366 Receiver Kit, Trial Kit, Spare Lead Kit, Sterile Revision Kit, SWAG Kit, SWAG Accessory Kit, Charger KitAug 4, 2017
87 days to decisionK171366 · Product code: **GZF** · Neurology
Source: <https://www.510kdatabase.net/k171366/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Peripheral Nerve, Implanted (pain Relief) (GZF)
Date received	May 9, 2017
Decision date	Aug 4, 2017
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stimwave Technologies Inc., Db a Stimq, LLC
Location	Fort Lauderdale, FL, US
Contact	Elizabeth Greene
510(k) history	1 submissions · 1 cleared · 2017-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171366/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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