FDA 510(k) Premarket Notification

**CLEARED**

# K171366 Receiver Kit, Trial Kit, Spare Lead Kit, Sterile Revision Kit, SWAG Kit, SWAG Accessory Kit, Charger Kit

Aug 4, 2017
87 days to decision

K171366 · 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., Dba Stimq, LLC** |
| Location | Fort Lauderdale, FL, US |
| Contact | Elizabeth Greene |
| 510(k) history | 1 submissions · 1 cleared · 2017-2017 |