FDA 510(k) Premarket Notification

# K131910 MPP

**CLEARED**

Jan 24, 2014
212 days to decision

K131910 · Product code: **NUH** · Neurology
Source: https://www.510kdatabase.net/k131910/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH) |
| Date received | Jun 26, 2013 |
| Decision date | Jan 24, 2014 |
| Days to decision | 212 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Marc Pro** |
| Location | Irvine, CA, US |
| Contact | GREG HOLLAND |
| 510(k) history | 1 submissions · 1 cleared · 2014-2014 |