

**K131910 MPP**Jan 24, 2014  
212 days to decisionK131910 · Product code: **NUH** · Neurology  
Source: <https://www.510kdatabase.net/k131910/>**SUBMISSION DETAILS**

---

|                       |   |
|-----------------------|---|
| 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        | <b>Marc Pro</b>                       |
| Location       | Irvine, CA, US                        |
| Contact        | GREG HOLLAND                          |
| 510(k) history | 1 submissions · 1 cleared · 2014-2014 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131910/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 7, 2026