

K151035 PulseReliefJul 21, 2015
95 days to decisionK151035 · Product code: **NUH** · Neurology
Source: <https://www.510kdatabase.net/k151035/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Apr 17, 2015
Decision date	Jul 21, 2015
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Consumer Lifestyle
Location	Stamford, CT, US
Contact	Marta Walker
510(k) history	2 submissions · 2 cleared · 2015-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k151035/> Data sourced from FDA 510(k) public records (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. - Generated June 4, 2026