FDA 510(k) Premarket Notification

K151035 PulseRelief

CLEARED

Jul 21, 2015
95 days to decision

K151035 · 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