FDA 510(k) Premarket Notification

K171599 WiTouch Pro, WiTouch, Neubac

CLEARED

Sep 22, 2017
113 days to decision

K171599 · Product code: **NUH** · Neurology
Source: https://www.510kdatabase.net/k171599/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Jun 1, 2017
Decision date	Sep 22, 2017
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Hollywog, LLC**
Location	Chattanooga, TN, US
Contact	Michael Treas
510(k) history	4 submissions · 4 cleared · 2012-2017