FDA 510(k) Premarket Notification

K120398 WITOUCH PRO

CLEARED

Aug 16, 2012
190 days to decision

K120398 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k120398/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Feb 8, 2012
Decision date	Aug 16, 2012
Days to decision	190 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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