FDA 510(k) Premarket Notification

K170659 IQ Technologies

CLEARED

May 26, 2017
84 days to decision

K170659 · Product code: **NUH** · Neurology
Source: https://www.510kdatabase.net/k170659/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Mar 3, 2017
Decision date	May 26, 2017
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Iq Technologies, Inc.**
Location	Las Vegas, NV, US
Contact	Elli Josef
510(k) history	2 submissions · 2 cleared · 2014-2017