FDA 510(k) Premarket Notification

K183692 Avid IF2

CLEARED

Jun 12, 2019
163 days to decision

K183692 · Product code: **LIH** · Neurology
Source: https://www.510kdatabase.net/k183692/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interferential Current Therapy (LIH)
Date received	Dec 31, 2018
Decision date	Jun 12, 2019
Days to decision	163 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Vision Quest Industries Inc./Dba VQ Orthocare**
Location	Vista, CA, US
Contact	Mohamed Ouerghi
510(k) history	2 submissions · 2 cleared · 2019-2020