FDA 510(k) Premarket Notification

K202490 Avid CT2

CLEARED

Nov 16, 2020
77 days to decision

K202490 · Product code: **IPF** · Neurology
Source: https://www.510kdatabase.net/k202490/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Aug 31, 2020
Decision date	Nov 16, 2020
Days to decision	77 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