

K202490 Avid CT2Nov 16, 2020
77 days to decisionK202490 · 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./Db a VQ Orthocare
Location	Vista, CA, US
Contact	Mohamed Ouerghi
510(k) history	2 submissions · 2 cleared · 2019-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k202490/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 14, 2026