

K183692 Avid IF2Jun 12, 2019
163 days to decisionK183692 · 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./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/k183692/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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