

K203331 Quick-20mMay 20, 2021
189 days to decisionK203331 · Product code: **GWL** · Neurology
Source: <https://www.510kdatabase.net/k203331/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Amplifier, Physiological Signal (GWL)
Date received	Nov 12, 2020
Decision date	May 20, 2021
Days to decision	189 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cgx, LLC
Location	San Diego, CA, US
Contact	Spencer Linton
510(k) history	1 submissions · 1 cleared · 2021-2021

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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