

**K232749 DOOPANG**Jun 11, 2024  
277 days to decisionK232749 · Product code: **PCC** · Neurology  
Source: <https://www.510kdatabase.net/k232749/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Electrical, Transcutaneous, For Migraine (PCC)
Date received	Sep 8, 2023
Decision date	Jun 11, 2024
Days to decision	277 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ybrain, Inc.</b>
Location	Daejeon, KR
Contact	Kiwon Lee
510(k) history	1 submissions · 1 cleared · 2024-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Gms Consulting</b>
Contact	Jonghyun Kim

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