

**K241512 Electrode Pad**Jan 15, 2025  
231 days to decisionK241512 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k241512/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	May 29, 2024
Decision date	Jan 15, 2025
Days to decision	231 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Deliduo Medical Technology Co.,Ltd</b>
Location	Huizhou, CN
Contact	Andy Liu
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shanghai SUNGO Management Consulting Co., Ltd.</b>
Contact	Wu Jarvis

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241512/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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