

**K190700 ELECTRODES PAD**Jun 14, 2019  
88 days to decisionK190700 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k190700/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 18, 2019
Decision date	Jun 14, 2019
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Bestpad Technology Development Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Junwei Xu
510(k) history	1 submissions · 1 cleared · 2019-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>Feiying Drug &amp; Medical Consulting Technical Service Group</b>
Contact	Rain Yip

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/k190700/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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