

**K133980 PANTHEON ELECTROSTIMULATOR**Nov 7, 2014  
315 days to decisionK133980 · Product code: **BWK** · Neurology  
Source: <https://www.510kdatabase.net/k133980/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electro-acupuncture (BWK)
Date received	Dec 27, 2013
Decision date	Nov 7, 2014
Days to decision	315 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pantheon Research</b>
Location	Crofton, MD, US
Contact	YOLANDA SMITH
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133980/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 5, 2026